Methods of treating sporadic inclusion body myositis

This disclosure claims priority to US Provisional Patent Application No. 61/865861, filed 2013-08-14 and US Provisional Patent Application No. 61/983567 filed 2014-04-24, the disclosure of which are incorporated by reference herein in their entirety.

TECHNICAL FIELD

This disclosure is in the field of myostatin antagonists, e.g., myostatin binding molecules or Activin receptor II (ActRII) binding molecules, e.g., an antagonist antibody to myostatin or to ActRII, e.g, BYM338. In particular, it relates to the treatment of sporadic inclusion body myositis (sIBM), and novel dosing regimens for treating it which employ a therapeutically effective amount of an ActRII antagonist, e.g., an Activin receptor II (ActRII) binding molecule, e.g., an anti-Activin receptor II (ActRII) antibody, such as the BYM338 antibody (which is also known as “bimagrumab”).

BACKGROUND OF THE DISCLOSURE

Sporadic inclusion-body myositis (sIBM) is a very rare disease. While there are limited epidemiological literature regarding the prevalence of the disease based on modern diagnostic criteria, it is sporadic with a prevalence estimate of 15-71 per million for all ages, and 51 per million over age 50. Men are more often affected than women approximately 2:1. There are notable regional differences in prevalence rates, which may involve both genetic and environmental factors. The primary underlying etiology of sIBM is unknown and is unfortunately refractory to any known treatments despite evidence of possibly secondary degenerative and inflammatory features (Hohlfeld 2011).

Although rare, sIBM represents the most common form for idiopathic inflammatory myopathies affecting those >50 years of age (Dimachkie and Barohn 2013, Griggs 2006, Peng et al 2000) and accounts for a total of 30% of all inflammatory myopathies. The disease is characterized clinically by the insidious and asymmetric onset of proximal and distal weakness which generally develops after the age of 50 years and progresses with time (Needham et al 2008). Diagnosis is often not made until 5–8 years after the onset of symptoms, mainly due to the slow evolution of the disease and the difficulty in recognizing the subtle early symptoms and the complex diagnostic criteria, including need for biopsy muscle pathology expertise. Lower extremity complaints come typically

in the form of difficulty arising from chairs, and walking upstairs or downstairs. As the disease progresses, lower extremity weakness leads to frequent falls with potential injuries. Falls in sIBM patients may occur as a result of proximal muscle weakness, but distal lower extremity weakness (foot drop) occurs as well and may play a contributing role to such falls. In addition there is early onset of hand and finger weakness which eventually impairs the ability to retain independence in activities of daily living (e.g. writing, feeding, bathing, dressing, brushing teeth). Of other important symptoms, dysphagia occurs in at least 40% of patients due to esophageal and pharyngeal muscle involvement. This can lead to weight loss or aspiration with consequent pneumonia (Amato & Barohn 2009, Oh et al 2008). Disease progression is relatively slow but relentless and virtually all patients with sIBM require an assistive device, such as a cane or walker within a few years and a wheelchair, by about ten years of onset (Griggs et al 1995, Dalakas 2006). The irreversible morbidity of sIBM is well-recognized by the clinical and patient community (Hohlfeld 2011).

Long-term follow up studies on a relatively larger number of sIBM patients have confirmed the seriously debilitating nature of sIBM (Benveniste et al 2011, Cox et al 2011). The highly debilitating nature of sIBM was confirmed by a national epidemiological study of 64 patients conducted in the Netherlands conducted over a period of 10–13 years focusing on the progressive decline in muscle strength, functional status and life expectancy with sIBM (Cox et al 2011). Although not a formal prospective registry, the authors reported that life expectancy was normal at 81 years, but activities of daily life were clearly restricted. At follow-up, all the 15 surviving patients who consented to further follow-up were found to be using a wheelchair, seven of them (47%) being completely wheelchair-bound. After a mean disease duration of 20 years, three patients were living in a nursing home and 12 at home with adaptations (stair lift, no thresholds, standup chairs). Nearly all patients required considerable help with daily activities from their partners or other caregivers; 40% of patients were completely or severely dependent (Barthel index <10), and 20% of patients were moderately dependent (Barthel index 10–15). In three patients (6.5%), euthanasia was requested because of “unbearable suffering and severe loss of quality of life due to extensive weakness” and in another three (6.5%), requests for continuous deep sedation due to severe disabling dysphagia, cachexia, and dehydration was granted. The fact that end-of-life care interventions were reported in six of these Dutch patients (13%) reflects the severe disability and loss of quality of life at the end stage of this disease.

There is no drug approved for the treatment of sIBM as no treatments have been found to slow or reverse the progression of muscle weakness in sIBM (Greenberg 2009, Aggarwal and Oddis 2012). Moreover, patients with sIBM have not demonstrated a

clinically meaningful response to agents used traditionally to treat inflammatory myopathies, including corticosteroids, methotrexate, azathioprine, or cyclophosphamide (Griggs 2006, Mann and Vencovský 2011, Needham and Mastaglia 2007, Solorzano and Phillips 2011). Intravenous immunoglobulin is used off-label in some centers, but there is no evidence to support its long-term effectiveness in this condition. Similar overall conclusions can be drawn on the efficacy of different immunotherapies such as the anti-T lymphocyte inhibitor, the anti-TNF medication (etanercept) and beta-inteferon 1A. Oxandrolone is still in an explorative phase and further data are required before reaching conclusions on its potential benefits. Therefore, there is currently a clear, unmet medical need in the treatment of patients with sIBM.

The mean rate of decline in muscle strength is 3.5-5.4% per year based on different methods and scores, with the potential rate of progression considerably faster for individual patients (Hohlfeld 2011, Cox et al 2011). Since sIBM causes dramatic skeletal muscle atrophy, treatments that target atrophy pathways in muscle, like bimagrumab, may be effective in this disease.

Preliminary data from Dr. Steven Greenberg's laboratory at Brigham & Women's Hospital suggest that the signaling pathway downstream of ActRIIB and other receptors may be inappropriately activated in sIBM, further supporting the hypothesis that ActRIIB inhibition could help patients with sIBM. Based on the data from 17 patients with sIBM compared to 12 patients with polymyositis or dermatomyositis and 5 normal controls, patients with sIBM had prominent phosphorylated SMAD (pSMAD) signaling (which is downstream of the ActRIIB receptor and acts as a second messenger for TGFβ signaling), with a mean 27-fold increase compared to normal controls (Greenberg et al 2013).

On this basis, the hypothesis is that inappropriate signaling through the TGFβ pathway is contributing to the pathogenesis of sIBM. While it is not known if this TGFβ up-regulation occurs via the ActRIIB receptor or another TGFβ receptor, it seemed reasonable to test whether ActRIIB inhibition by bimagrumab can intervene on muscle atrophy signaling pathways and target weakness symptoms experienced by patients with sIBM.

Moreover, the study by Wojcik et al included analyses of biopsies from 12 sIBM patients and suggests that the myostatin/myostatin precursor, either alone, or bound to amyloid-β, may play a role in the pathogenesis of sIBM (Wojcik et al 2005).

Bimagrumab (BYM338) or is a monoclonal antibody developed to bind competitively to activin receptor type II (ActRII) with greater affinity than myostatin or activin, its natural ligands. Bimagrumab is a fully human antibody (modified IgG1, 234-235-Ala-Ala, λ_2) which binds to the ligand binding domain of ActRII, thereby preventing binding and subsequent signaling of its ligands, one of which is myostatin and activin. Myostatin, a

member of the transforming growth factor beta (TGF-β) superfamily, is a secreted protein that negatively regulates skeletal muscle mass in animals and humans. Myostatin signaling occurs at ActRII and its proposed mechanism of action is through the Smad 2/3 pathway to inhibit protein synthesis and myocyte differentiation and proliferation. Myostatin inhibition or genetic ablation increases muscle mass and strength (Lee et al 2005; Lee and McPherron 2001; Whittemore et al 2003).

BYM338 is cross-reactive with human and mouse ActRIIB and effective on human, cynomolgus, mouse and rat skeletal muscle cells. BYM338 is formulated for both intravenous (i.v.) and subcutaneous (s.c.) administration.

Data from study CBYM338X2205 on 14 patients with sIBM (11 active, 3 placebo) showed statistically significant increases in BYM338 relative to placebo for both muscle volume and lean body mass after a single dose of BYM338 30 mg/kg i.v. was administered. Eight weeks after dosing, the mean change from baseline in dominant thigh muscle volume (TMV) favored BYM338 (+ 6.5%; P=0.024). Non-dominant TMV and whole body lean mass (measured by DXA) also favored BYM338 (+ 7.6% and +5.7% respectively; P=0.009 and P=0.014). Patients were followed for 24 weeks after the single dose and demonstrated a numerical increase in muscle strength in several muscle groups as measured by both Quantitative Muscle Testing (QMT) and Manual Muscle Testing. Data also suggested benefits 16 weeks after the single BYM338 administration in physical function and mobility as shown by a 14.6% statistically significant increase (p=0.008) in the 6-minute walking distance test (6MWD) as compared to placebo.

As illustrated by data from patients with sporadic inclusion body myositis, a progressive muscle degenerative disease, the rapid increases in lean body mass (>5 % from baseline) induced by a single injection of bimagrumab (30 mg/kg) are able to trigger significant increases in physical performance (**Figure 1**). Importantly, improvement of functional following muscle mass increase require a period of lag time possibly reflecting the structural/functional remodeling of skeletal muscle before becoming fully matured and ready to serve increased contractile activities.

The present disclosure is further studying these favorable preliminary single dose findings and investigate how long-term treatment with different doses of bimagrumab influence the changes of muscle mass, muscle strength, physical function, and mobility in ambulatory sIBM patients.

Myostatin, ActRIIB Receptor and ActRIIB recptor antibodies

Bimagrumab, also known as BYM338, is a human monoclonal antibody developed to bind competitively to activin receptor type II B (ActRIIB) with greater affinity than myostatin, its principal natural ligand. Bimagrumab is disclosed in WO2010/125003, which is incorporated by reference herein in its entirety. Myostatin, a member of the transforming growth factor beta (TGF-β) superfamily, is a secreted protein that negatively regulates skeletal muscle mass in animals and humans, throughout the lifecycle. Myostatin signaling occurs at ActRIIB and its proposed mechanism of action is through the Smad 2/3 pathway to inhibit protein synthesis and myocyte differentiation and proliferation. The absence of myostatin in developing animals and humans results in a hypermuscular phenotype with an increased number and size of muscle fibers. Reducing the level of myostatin postpartum results in the hypertrophy of skeletal muscle due to an increase in the size of existing myofibers. In the adult, myostatin is produced in skeletal muscle and circulated in the blood in part as a latent inactive complex.

Consistent with the role of myostatin as an endogenous inhibitor of skeletal muscle mass, BYM338 dramatically increased skeletal muscle mass in preclinical murine models of disuse and steroid-induced atrophy and in toxicology studies with healthy cynomolgus monkeys. In addition, the increased mass in mouse and rat resulted in a corresponding increase in muscle strength (force production). Following i.v. and s.c. administration to mice and cynomolgus monkey, bimagrumab showed a consistent IgG1 pharmacokinetic (PK) profile with target mediated drug disposition (TMDD) and was well tolerated.

An analysis of the six dose levels of the first in human, single ascending dose study, suggests that single i.v. doses of 0.1, 0.3, 1, 3, 10 and 30 mg/kg of bimagrumab are safe, well tolerated, and produce a PK profile that is predictable from modeled preclinical data. At four weeks doses of 3-30 mg/kg result in a measurable increase in thigh muscle volume of 2.7-5.2% from baseline over placebo.

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The potential clinical application of bimagrumab is in conditions with muscle wasting. Of particular note are clinical scenarios that require the recovery of muscle mass from atrophy resulting from disuse, cachexia, corticosteroid use, and sarcopenia. Sporadic inclusion-body mmyositis (sIBM) is a clinical situation in which elements of all of these are likely to play a part.

SUMMARY OF THE DISCLOSURE

Intervening in a patient population with sporadic inclusion body myositis (sIBM) would be highly innovative and would meet a high unmet medical need. Indeed, there is currently no therapeutic option to treat sIBM. This objective is achieved by the use, methods and dosing regimen provided within this disclosure.

A first subject matter of the disclosure therefore relates to methods or uses for treating sIBM of compositions comprising a myostatin antagonist, which can be a myostatin binding molecule or an ActRII binding molecule. The myostatin binding molecule can be, e.g., an antagonist antibody to myostatin. The ActRII binding molecule can be, e.g., an antagonist antibody to ActRII, e.g., bimagrumab also known as BYM338.

"Myostatin antagonist" as used herein refers to a molecule capable of antagonizing (e.g., reducing, inhibiting, decreasing, delaying) myostatin function, expression and/or signalling (e.g., by blocking the binding of myostatin to the myostatin receptor, i.e., ActRIIB). Non-limiting examples of antagonists include myostatin binding molecules and ActRIIB receptor binding molecules. In some embodiments of the disclosed methods, regimens, kits, processes, uses and compositions, a myostatin antagonist is employed.

By "myostatin binding molecule" is meant any molecule capable of binding to the human myostatin antigen either alone or associated with other molecules. The binding reaction may be shown by standard methods (qualitative assays) including, for example, a binding assay, competition assay or a bioassay for determining the inhibition of myostatin binding to its receptor or any kind of binding assays, with reference to a negative control test in which an antibody of unrelated specificity, but ideally of the same isotype, e.g., an anti-CD25 antibody, is used. Non-limiting examples of myostatin binding molecules include small molecules, myostatin receptor decoys, and antibodies that bind to myostatin as produced by B-cells or hybridomas and chimeric, CDR-grafted or human antibodies or any fragment thereof, e.g., $F(ab')_2$ and Fab fragments, as well as single chain or single domain antibodies. Preferably the myostatin binding molecule antagonizes (e.g., reduces, inhibits, decreases, delays) myostatin function, expression and/or signalling. In some embodiments of the disclosed methods, regimens, kits, processes, uses and compositions, a myostatin binding molecule is employed.

By "ActRII binding molecule" is meant any molecule capable of binding to the human ActRII receptor (ActRII A and/or ActRIIB) either alone or associated with other molecules. The binding reaction may be shown by standard methods (qualitative assays) including, for example, a binding assay, competition assay or a bioassay for determining the inhibition of ActRII receptor binding to myostatin or any kind of binding assays, with reference to a negative control test in which an antibody of unrelated

specificity, but ideally of the same isotype, e.g., an anti-CD25 antibody, is used. Non-limiting examples of ActRII receptor binding molecules include small molecules, myostatin decoys, and antibodies to the ActRII receptor as produced by B-cells or hybridomas and chimeric, CDR-grafted or human antibodies or any fragment thereof, e.g., $F(ab')_2$ and Fab fragments, as well as single chain or single domain antibodies. Preferably the ActRII receptor binding molecule antagonizes (e.g., reduces, inhibits, decreases, delays) myostatin function, expression and/or signalling. In some embodiments of the disclosed methods, regimens, kits, processes, uses and compositions, an ActRIIB receptor binding molecule is employed.

In another embodiment the composition comprises an anti-ActRII antibody which binds to a binding domain consisting of amino acids 19-134 of SEQ ID NO: 181 (SEQ ID NO:182), or to an epitope comprising or consisting of (a) amino acids 78-83 of SEQ ID NO: 181 (WLDDFN – SEQ ID NO:188); (b) amino acids 76-84 of SEQ ID NO:181 (GCWLDDFNC – SEQ ID NO:186); (c) amino acids 75-85 of SEQ ID NO:181 (KGCWLDDFNCY – SEQ ID NO:190); (d) amino acids 52-56 of SEQ ID NO:181 (EQDKR – SEQ ID NO:189); (e) amino acids 49-63 of SEQ ID NO:181 (CEGEQDKRLHCYASW – SEQ ID NO:187); (f) amino acids 29-41 of SEQ ID NO:181 (CIYYNANWELERT– SEQ ID NO:191); (g) amino acids 100-110 of SEQ ID NO:181 (YFCCCEGNFCN – SEQ ID NO:192); or (h) amino acids 78-83 of SEQ ID NO:181 (WLDDFN) and amino acids 52-56 of SEQ ID NO:181 (EQDKR).

In a yet further alternative embodiment, the above mentioned compositions comprise an anti-ActRII antibody which binds ActRIIB with a 10-fold or greater affinity than it binds to ActRIIA.

Additionally, the disclosure relates to composition wherein the anti-ActRIIB antibody comprises a heavy chain variable region CDR1 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-14; a heavy chain variable region CDR2 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 15-28; a heavy chain variable region CDR3 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 29-42; a light chain variable region CDR1 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 43-56; a light chain variable region CDR2 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 57-70; and a light chain variable region CDR3 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 71-84.

In certain embodiments, the disclosure provides compositions wherein the anti-ActRII antibody comprises: (a) a heavy chain variable region CDR1 of SEQ ID NO: 1; a heavy chain variable region CDR2 of SEQ ID NO: 15; a heavy chain variable region CDR3 of SEQ ID NO: 29; a light chain variable region CDR1 of SEQ ID NO: 43; a light chain variable region CDR2 of SEQ ID NO: 57; and a light chain variable region CDR3 of SEQ ID NO: 71, (b) a heavy chain variable region CDR1 of SEQ ID NO: 2; a heavy chain variable region CDR2 of SEQ ID NO: 16; a heavy chain variable region CDR3 of SEQ ID NO: 30; a light chain variable region CDR1 of SEQ ID NO: 44; a light chain variable region CDR2 of SEQ ID NO: 58; and a light chain variable region CDR3 of SEQ ID NO: 72, (c) a heavy chain variable region CDR1 of SEQ ID NO: 3; a heavy chain variable region CDR2 of SEQ ID NO: 17; a heavy chain variable region CDR3 of SEQ ID NO: 31; a light chain variable region CDR1 of SEQ ID NO: 45; a light chain variable region CDR2 of SEQ ID NO: 59; and a light chain variable region CDR3 of SEQ ID NO: 73, (d) a heavy chain variable region CDR1 of SEQ ID NO: 4; a heavy chain variable region CDR2 of SEQ ID NO: 18; a heavy chain variable region CDR3 of SEQ ID NO: 32; a light chain variable region CDR1 of SEQ ID NO: 46; a light chain variable region CDR2 of SEQ ID NO: 60; and a light chain variable region CDR3 of SEQ ID NO: 74, (e) a heavy chain variable region CDR1 of SEQ ID NO: 5; a heavy chain variable region CDR2 of SEQ ID NO: 19; a heavy chain variable region CDR3 of SEQ ID NO: 33; a light chain variable region CDR1 of SEQ ID NO: 47; a light chain variable region CDR2 of SEQ ID NO: 61; and a light chain variable region CDR3 of SEQ ID NO: 75, (f) a heavy chain variable region CDR1 of SEQ ID NO: 6; a heavy chain variable region CDR2 of SEQ ID NO: 20; a heavy chain variable region CDR3 of SEQ ID NO: 34; a light chain variable region CDR1 of SEQ ID NO: 48; a light chain variable region CDR2 of SEQ ID NO: 62; and a light chain variable region CDR3 of SEQ ID NO: 76, (g) a heavy chain variable region CDR1 of SEQ ID NO: 7; a heavy chain variable region CDR2 of SEQ ID NO: 21; a heavy chain variable region CDR3 of SEQ ID NO: 35; a light chain variable region CDR1 of SEQ ID NO: 49; a light chain variable region CDR2 of SEQ ID NO: 63; and a light chain variable region CDR3 of SEQ ID NO: 77, (h) a heavy chain variable region CDR1 of SEQ ID NO: 8; a heavy chain variable region CDR2 of SEQ ID NO: 22; a heavy chain variable region CDR3 of SEQ ID NO: 36; a light chain variable region CDR1 of SEQ ID NO: 50 a light chain variable region CDR2 of SEQ ID NO: 64; and a light chain variable region CDR3 of SEQ ID NO: 78, (i) a heavy chain variable region CDR1 of SEQ ID NO: 9; a heavy chain variable region CDR2 of SEQ ID NO: 23; a heavy chain variable region CDR3 of SEQ ID NO: 37; a light chain variable region CDR1 of SEQ ID NO: 51; a light chain variable region CDR2 of SEQ ID NO: 65; and a light chain variable region CDR3 of SEQ ID NO: 79, (j) a heavy chain variable region CDR1 of SEQ ID NO: 10; a heavy chain variable region CDR2 of SEQ ID NO: 24; a heavy chain variable region CDR3 of SEQ ID NO: 38;

a light chain variable region CDR1 of SEQ ID NO: 52; a light chain variable region CDR2 of SEQ ID NO: 66; and a light chain variable region CDR3 of SEQ ID NO: 80, (k) a heavy chain variable region CDR1 of SEQ ID NO: 11; a heavy chain variable region CDR2 of SEQ ID NO: 25; a heavy chain variable region CDR3 of SEQ ID NO: 39; a light chain variable region CDR1 of SEQ ID NO: 53; a light chain variable region CDR2 of SEQ ID NO: 67; and a light chain variable region CDR3 of SEQ ID NO: 81, (l) a heavy chain variable region CDR1 of SEQ ID NO: 12; a heavy chain variable region CDR2 of SEQ ID NO: 26; a heavy chain variable region CDR3 of SEQ ID NO: 40; a light chain variable region CDR1 of SEQ ID NO: 54; a light chain variable region CDR2 of SEQ ID NO: 68; and a light chain variable region CDR3 of SEQ ID NO: 82, (m) a heavy chain variable region CDR1 of SEQ ID NO: 13; a heavy chain variable region CDR2 of SEQ ID NO: 27; a heavy chain variable region CDR3 of SEQ ID NO: 41; a light chain variable region CDR1 of SEQ ID NO: 55; a light chain variable region CDR2 of SEQ ID NO: 69; and a light chain variable region CDR3 of SEQ ID NO: 83, or (n) a heavy chain variable region CDR1 of SEQ ID NO: 14; a heavy chain variable region CDR2 of SEQ ID NO: 28; a heavy chain variable region CDR3 of SEQ ID NO: 42; a light chain variable region CDR1 of SEQ ID NO: 56; a light chain variable region CDR2 of SEQ ID NO: 70; and a light chain variable region CDR3 of SEQ ID NO: 84.

In yet another embodiment, the above mentioned anti-ActRII antibody comprises (i) a full length heavy chain amino acid sequence having at least 95% sequence identity to at least one sequence selected from the group consisting of SEQ ID NOs:146-150 and 156-160, (ii) a full length light chain amino acid sequence having at least 95% sequence identity to at least one sequence selected from the group consisting of SEQ ID NOs:141-145 and 151-155 or (iii) (a) the variable heavy chain sequence of SEQ ID NO: 99 and variable light chain sequence of SEQ ID NO: 85; (b) the variable heavy chain sequence of SEQ ID NO: 100 and variable light chain sequence of SEQ ID NO: 86; (c) the variable heavy chain sequence of SEQ ID NO: 101 and variable light chain sequence of SEQ ID NO: 87; (d) the variable heavy chain sequence of SEQ ID NO: 102 and variable light chain sequence of SEQ ID NO: 88; (e) the variable heavy chain sequence of SEQ ID NO: 103 and variable light chain sequence of SEQ ID NO: 89; (f) the variable heavy chain sequence of SEQ ID NO: 104 and variable light chain sequence of SEQ ID NO: 90; (g) the variable heavy chain sequence of SEQ ID NO: 105 and variable light chain sequence of SEQ ID NO: 91; (h) the variable heavy chain sequence of SEQ ID NO: 106 and variable light chain sequence of SEQ ID NO: 92; (i) the variable heavy chain sequence of SEQ ID NO: 107 and variable light chain sequence of SEQ ID NO: 93; (j) the variable heavy chain sequence of SEQ ID NO: 108 and variable light chain sequence of SEQ ID NO: 94; (k) the variable heavy chain sequence of SEQ ID NO: 109 and

variable light chain sequence of SEQ ID NO: 95; (l) the variable heavy chain sequence of SEQ ID NO: 110 and variable light chain sequence of SEQ ID NO: 96; (m) the variable heavy chain sequence of SEQ ID NO: 111 and variable light chain sequence of SEQ ID NO: 97; or (n) the variable heavy chain sequence of SEQ ID NO: 112 and variable light chain sequence of SEQ ID NO: 98.

In certain aspects the disclosure relates to the above described compositions, wherein the comprised anti-ActRII antibody comprises (a) the heavy chain sequence of SEQ ID NO: 146 and light chain sequence of SEQ ID NO: 141; (b) the heavy chain sequence of SEQ ID NO: 147 and light chain sequence of SEQ ID NO: 142; (c) the heavy chain sequence of SEQ ID NO: 148 and light chain sequence of SEQ ID NO: 143; (d) the heavy chain sequence of SEQ ID NO: 149 and light chain sequence of SEQ ID NO: 144; (e) the heavy chain sequence of SEQ ID NO: 150 and light chain sequence of SEQ ID NO: 145; (f) the heavy chain sequence of SEQ ID NO: 156 and light chain sequence of SEQ ID NO: 151; (g) the heavy chain sequence of SEQ ID NO: 157 and light chain sequence of SEQ ID NO: 152; (h) the heavy chain sequence of SEQ ID NO: 158 and light chain sequence of SEQ ID NO: 153; (i) the heavy chain sequence of SEQ ID NO: 159 and light chain sequence of SEQ ID NO: 154; or (j) the heavy chain sequence of SEQ ID NO: 160 and light chain sequence of SEQ ID NO: 155.

An additional subject matter of the disclosure relates to composition, wherein (i) the anti-ActRII antibody cross-blocks or is cross blocked by one of the above described antibodies, (ii) has altered effector function through mutation of the Fc region and/or (iii) binds to an epitope recognized by one of the above described antibodies.

In yet another embodiment, the disclosed composition comprises an anti-ActRII antibody encoded by pBW522 (DSM22873) or pBW524 (DSM22874).

BRIEF DESCRIPTION OF THE FIGURES

Figure 1: Bimagrumab-induced changes of lean body mass (LBM), quadriceps strength (QMT) and 6-minute walking distance (6MWD) from baseline in sporadic inclusion body myositis patients. Note the time-lag between increase in LBM (at Week 8 to Week 16) and significant increases in muscle strength and physical performance starting at Week 16.

Figure 2: Study design

DEFINITIONS

In order that the present disclosure may be more readily understood, certain terms are first defined. Additional definitions are set forth throughout the detailed description.

The term "comprising" means "including" *e.g.* a composition "comprising" X may consist exclusively of X or may include something additional *e.g.* X + Y.

The term "about" in relation to a numerical value *x* means, for example, *x*±10%.

The following exemplifies a possible pre-clinical treatment regimes to evaluate possible effects of a treatment with a myostatin antagonist, e.g., myostatin binding molecule or ActRII binding molecule, preferably ActRII binding molecule, more preferably an antagonist antibody to ActRII, e.g, bimagrumab.

The treatment is exemplified by using cynomolgus monkeys, but the described experiments are not limited to monkeys and the skilled person knows how to set up suitable experiments or dosing regimens for other species, in particular for humans: the anti-ActRII antibody, e.g., bimagrumab, can be administered once a week for 3 months to male and female cynomolgus monkeys by intravenous injection. 32 cynomolgus monkeys (16/sex) can be assigned to one of four treatment groups (3 to 5 animals/sex/group) and can be administered intravenous injections of either vehicle or the ActRIIB antibody, e.g., BYM338, at 10, 30, or 100 mg/kg once weekly for 13 weeks (total of 14 doses; doses shall be selected on the basis of muscle hypertrophy activity in monkey).

The terms "ActRII", "ActRIIA" and "ActRIIB" refer to Activin receptors. Activins signal through a heterodimeric complex of receptor serine kinases which include at least two type I (I and IB) and two type II "ActRII" (IIA and IIB, aka ACVR2A and ACVR2B) receptors. These receptors are all transmembrane proteins, composed of a ligand-binding extracellular domain with a cysteine-rich region, a transmembrane domain, and a cytoplasmic domain with predicted serine/threonine specificity. Type I receptors are essential for signaling while type II receptors are required for binding ligands and for expression/recruitment of type I receptors. Type I and II receptors form a stable complex after ligand binding resulting in the phosphorylation of type I receptors by type II receptors. The activin receptor II (ActRII) is a receptor for myostatin. The activin receptor

type II B (ActRIIB) is a receptor for myostatin. The activin receptor type II A (ActRIIA) is also a receptor for myostatin. The term ActRIIB or Act IIB receptor refers to human ActRIIB as defined in SEQ ID NO: 181 (AAC64515.1, GI:3769443). Research grade polyclonal and monoclonal anti-ActRIIB antibodies are known in the art, such as those made by R&D Systems®, MN, USA. Of course, antibodies could be raised against ActRIIB from other species and used to treat pathological conditions in those species.

The term "immune response" refers to the action of, for example, lymphocytes, antigen presenting cells, phagocytic cells, granulocytes, and soluble macromolecules produced by the above cells or the liver (e.g. antibodies, cytokines, and complement) that results in selective damage to, destruction of, or elimination from the human body of invading pathogens, cells or tissues infected with pathogens, cancerous cells, or, in cases of autoimmunity or pathological inflammation, normal human cells or tissues.

A “signaling activity” refers to a biochemical causal relationship generally initiated by a protein-protein interaction such as binding of a growth factor to a receptor, resulting in transmission of a signal from one portion of a cell to another portion of a cell. In general, the transmission involves specific phosphorylation of one or more tyrosine, serine, or threonine residues on one or more proteins in the series of reactions causing signal transduction. Penultimate processes typically include nuclear events, resulting in a change in gene expression.

The term "antibody" as referred to herein includes whole antibodies and any antigen binding fragment (*i.e.* "antigen-binding portion") or single chains thereof. A naturally occurring "antibody" is a glycoprotein comprising at least two heavy (H) chains and two light (L) chains inter-connected by disulfide bonds. Each heavy chain is comprised of a heavy chain variable region (abbreviated herein as V_H) and a heavy chain constant region. The heavy chain constant region is comprised of three domains, CH1, CH2 and CH3. Each light chain is comprised of a light chain variable region (abbreviated herein as V_L) and a light chain constant region. The light chain constant region is comprised of one domain, C_L. The V_H and V_L regions can be further subdivided into regions of hypervariability, termed complementarity determining regions (CDR), interspersed with regions that are more conserved, termed framework regions (FR). Each V_H and V_L is composed of three CDRs and four FRs arranged from amino-terminus to carboxy-terminus in the following order: FR1, CDR1, FR2, CDR2, FR3, CDR3, FR4. The variable regions of the heavy and light chains contain a binding domain that interacts with an antigen. The constant regions of the antibodies may mediate the binding of the

immunoglobulin to host tissues or factors, including various cells of the immune system (*e.g.* effector cells) and the first component (Clq) of the classical complement system.

The term "antigen-binding portion" of an antibody (or simply "antigen portion"), as used herein, refers to full length or one or more fragments of an antibody that retain the ability to specifically bind to an antigen (*e.g.* a portion of ActRIIB). It has been shown that the antigen-binding function of an antibody can be performed by fragments of a full-length antibody. Examples of binding fragments encompassed within the term "antigen-binding portion" of an antibody include a Fab fragment, a monovalent fragment consisting of the V_L, V_H, C_L and CH1 domains; a $F(ab)_2$ fragment, a bivalent fragment comprising two Fab fragments, each of which binds to the same antigen, linked by a disulfide bridge at the hinge region; a Fd fragment consisting of the V_H and CH1 domains; a Fv fragment consisting of the V_L and V_H domains of a single arm of an antibody; a dAb fragment (Ward *et al.*, 1989 Nature 341:544-546), which consists of a V_H domain; and an isolated complementarity determining region (CDR).

Furthermore, although the two domains of the Fv fragment, V_L and V_H, are coded for by separate genes, they can be joined, using recombinant methods, by a synthetic linker that enables them to be made as a single protein chain in which the V_L and V_H regions pair to form monovalent molecules (known as single chain Fv (scFv); see *e.g.* Bird *et al.*, 1988 Science 242:423-426; and Huston *et al.*, 1988 Proc. Natl. Acad. Sci. 85:5879-5883). Such single chain antibodies are also intended to be encompassed within the term "antigen-binding region" of an antibody. These antibody fragments are obtained using conventional techniques known to those of skill in the art, and the fragments are screened for utility in the same manner as are intact antibodies.

An "isolated antibody", as used herein, refers to an antibody that is substantially free of other antibodies having different antigenic specificities (*e.g.*, an isolated antibody that specifically binds ActRIIB is substantially free of antibodies that specifically bind antigens other than ActRIIB). An isolated antibody that specifically binds ActRIIB may, however, have cross-reactivity to other antigens, such as ActRIIB molecules from other species. Moreover, an isolated antibody may be substantially free of other cellular material and/or chemicals.

The terms "cross-block", "cross-blocked" and "cross-blocking" are used interchangeably herein to mean the ability of an antibody or other binding agent to interfere with the binding of other antibodies or binding agents to ActRIIB, particularly the ligand binding domain, in a standard competitive binding assay.

The terms "monoclonal antibody" or "monoclonal antibody composition" as used herein refer to a preparation of antibody molecules of single molecular composition. A monoclonal antibody composition displays a single binding specificity and affinity for a particular epitope.

The term "human antibody", as used herein, is intended to include antibodies having variable regions in which both the framework and CDR regions are derived from sequences of human origin. Furthermore, if the antibody contains a constant region, the constant region also is derived from such human sequences, *e.g.* human germline sequences, or mutated versions of human germline sequences or antibody containing consensus framework sequences derived from human framework sequences analysis, for example, as described in Knappik, *et al.* (2000. J Mol Biol 296, 57-86). The human antibodies of the disclosure may include amino acid residues not encoded by human sequences (*e.g.* mutations introduced by random or site-specific mutagenesis *in vitro* or by somatic mutation *in vivo*). However, the term "human antibody", as used herein, is not intended to include antibodies in which CDR sequences derived from the germline of another mammalian species, such as a mouse, have been grafted onto human framework sequences.

The term "human monoclonal antibody" refers to antibodies displaying a single binding specificity which have variable regions in which both the framework and CDR regions are derived from human sequences. In one embodiment, the human monoclonal antibodies are produced by a hybridoma which includes a B cell obtained from a transgenic nonhuman animal, *e.g.* a transgenic mouse, having a genome comprising a human heavy chain transgene and a light chain transgene fused to an immortalized cell.

The term "recombinant human antibody", as used herein, includes all human antibodies that are prepared, expressed, created or isolated by recombinant means, such as antibodies isolated from an animal (*e.g.* a mouse) that is transgenic or transchromosomal for human immunoglobulin genes or a hybridoma prepared therefrom, antibodies isolated from a host cell transformed to express the human antibody, *e.g.* from a transfectoma, antibodies isolated from a recombinant, combinatorial human antibody library, and antibodies prepared, expressed, created or isolated by any other means that involve splicing of all or a portion of a human immunoglobulin gene, sequences to other DNA sequences. Such recombinant human antibodies have variable regions in which the framework and CDR regions are derived from human germline immunoglobulin sequences. In certain embodiments, however, such recombinant human antibodies can

be subjected to *in vitro* mutagenesis (or, when an animal transgenic for human Ig sequences is used, *in vivo* somatic mutagenesis) and thus the amino acid sequences of the V_H and V_L regions of the recombinant antibodies are sequences that, while derived from and related to human germline V_H and V_L sequences, may not naturally exist within the human antibody germline repertoire *in vivo*.

As used herein, "isotype" refers to the antibody class (*e.g.* IgM, IgE, IgG such as IgG1 or IgG2) that is provided by the heavy chain constant region genes.

The phrases "an antibody recognizing an antigen" and "an antibody specific for an antigen" are used interchangeably herein with the term "an antibody which binds specifically to an antigen".

As used herein, an antibody that "specifically binds to ActRIIB polypeptide" is intended to refer to an antibody that binds to human ActRIIB polypeptide with a K_D of a about 100nM or less, about 10nM or less, about 1nM or less. An antibody that "cross-reacts with an antigen other than ActRIIB" is intended to refer to an antibody that binds that antigen with a K_D of about 10×10^{-9} M or less, about 5×10^{-9} M or less, or about 2×10^{-9} M or less. An antibody that "does not cross-react with a particular antigen" is intended to refer to an antibody that binds to that antigen, with a K_D of about 1.5×10^{-8} M or greater, or a K_D of about 5-10 $\times 10^{-8}$ M, or about 1×10^{-7} M or greater. In certain embodiments, such antibodies that do not cross-react with the antigen exhibit essentially undetectable binding against these proteins in standard binding assays. K_D may be determined using a biosensor system, such as a Biacore® system, or Solution Equilibrium Titration.

As used herein, the term "antagonist antibody" is intended to refer to an antibody that inhibits ActRIIB induced signaling activity in the presence of myostatin or of other ActRIIB ligands such as activins or GDF-11 and/or to an antibody that inhibits ActRIIA induced signaling activity in the presence of myostatin or of other ActRIIA ligands such as activins or GDF-11. Examples of an assay to detect this include inhibition of myostatin induced signalling (for instance by a Smad dependent reporter gene assay), inhibition of myostatin induced Smad phosphorylation (P-Smad ELISA) and inhibition of myostatin induced inhibition of skeletal muscle cell differentiation (for instance by a creatine kinase assay).

In some embodiments, the antibodies inhibit myostatin induced signalling as measured in a Smad dependent reporter gene assay at an IC_{50} of about 10nM or less, about 1nM or less, or about 100pM or less.

As used herein, an antibody with “no agonistic activity" is intended to refer to an antibody that does not significantly increase ActRIIB mediated signaling activity in the absence of myostatin in a cell-based assay, such as inihibition of myostatin induced signalling (for instance by a Smad dependent reporter gene assay), inhibition of myostatin induced Smad phosphorylation (P-Smad ELISA) and inhibition of myostatin induced inhibition of skeletal muscle cell differentiation (for instance by a creatine kinase assay). Such assays are described in more details in the examples below.

The term "K_{assoc}" or "K_a", as used herein, is intended to refer to the association rate of a particular antibody-antigen interaction, whereas the term "K_{dis}" or "K_d", as used herein, is intended to refer to the dissociation rate of a particular antibody-antigen interaction. The term "K_D", as used herein, is intended to refer to the dissociation constant, which is obtained from the ratio of K_d to K_a (*i.e.* K_d/K_a) and is expressed as a molar concentration (M). K_D values for antibodies can be determined using methods well established in the art. A method for determining the K_D of an antibody is by using surface plasmon resonance, such as the biosensor system of Biacore®, or Solution Equilibrium Titration (SET) (see Friguet B *et al.* (1985) J. Immunol Methods; 77(2): 305-319, and Hanel C et al. (2005) Anal Biochem; 339(1): 182-184).

As used herein, the term "Affinity" refers to the strength of interaction between antibody and antigen at single antigenic sites. Within each antigenic site, the variable region of the antibody “arm” interacts through weak non-covalent forces with antigen at numerous sites; the more interactions, the stronger the affinity.

As used herein, the term "Avidity" refers to an informative measure of the overall stability or strength of the antibody-antigen complex. It is controlled by three major factors: antibody epitope affinity; the valency of both the antigen and antibody; and the structural arrangement of the interacting parts. Ultimately these factors define the specificity of the antibody, that is, the likelihood that the particular antibody is binding to a precise antigen epitope.

As used herein, the term “ADCC” or “antibody dependent cellular cytotoxicity” activity refers to human B cell depleting activity. ADCC activity can be measured by the human B cell depleting assays known in the art.

In order to get a higher avidity probe, a dimeric conjugate (two molecules of an antibody protein coupled to a FACS marker) can be constructed, thus making low affinity

interactions (such as with the germline antibody) more readily detected by FACS. In addition, another means to increase the avidity of antigen binding involves generating dimers, trimers or multimers of any of the constructs described herein of the anti-ActRIIB antibodies. Such multimers may be generated through covalent binding between individual modules, for example, by imitating the natural C-to-N-terminus binding or by imitating antibody dimers that are held together through their constant regions. The bonds engineered into the Fc/Fc interface may be covalent or non-covalent. In addition, dimerizing or multimerizing partners other than Fc can be used in ActRIIB hybrids to create such higher order structures. For example, it is possible to use multimerizing domains such as the trimerizing domain described in WO2004/039841 or pentamerizing domain described in WO98/18943.

As used herein, the term “selectivity” for an antibody refers to an antibody that binds to a certain target polypeptide but not to closely related polypeptides.

As used herein, the term "high affinity" for an antibody refers to an antibody having a K_D of 1nM or less for a target antigen. As used herein, the term "subject" includes any human or nonhuman animal.

The term "nonhuman animal" includes all vertebrates, *e.g.* mammals and non-mammals, such as nonhuman primates, sheep, dogs, cats, horses, cows, chickens, amphibians, reptiles, *etc.*

As used herein, the term, "optimized" means that a nucleotide sequence has been altered to encode an amino acid sequence using codons that are preferred in the production cell or organism, generally a eukaryotic cell, for example, a cell of *Pichia*, a cell of *Trichoderma*, a Chinese Hamster Ovary cell (CHO) or a human cell. The optimized nucleotide sequence is engineered to retain completely or as much as possible the amino acid sequence originally encoded by the starting nucleotide sequence, which is also known as the "parental" sequence. The optimized sequences herein have been engineered to have codons that are preferred in CHO mammalian cells, however optimized expression of these sequences in other eukaryotic cells is also envisioned herein. The amino acid sequences encoded by optimized nucleotide sequences are also referred to as optimized.

DETAILED DESCRIPTION OF THE DISCLOSURE

It has been discovered that antibodies directed to the ActRII receptors, e.g, bimagrumab, can prevent myostatin from binding to the receptor, thus treating sIBM patients.

Therefore, in one aspect, the disclosure provides a composition comprising a myostatin antagonist, e.g., myostatin binding molecule or ActRII binding molecule, preferably ActRII binding molecule, more preferably an anti-ActRII antibody, e.g, bimagrumab or a functional protein comprising an antigen-binding portion of said antibody for use. In one embodiment, the ActRIIB is human ActRIIB. The polypeptide sequence of human ActRIIB is recited in SEQ ID NO: 181 (AAC64515.1, GI:3769443). In one embodiment, the antibody or functional protein is from a mammal, having an origin such as human or camelid. Thus the antibody comprised in the disclosed composition may be a chimeric, human or a humanized antibody. In a particular embodiment, the anti-ActRIIB antibody comprised in the disclosed composition is characterized as having an antigen-binding region that is specific for the target protein ActRIIB and binds to ActRIIB or a fragment of ActRIIB.

The disclosed composition and regimen are also suitable for use in treating age related mobility disability, cancer cachexia, chronic obstructive pulmonary disease (COPD) and joint replacement, e.g, knee arthroplasty or hip arthroplasty, or hip fracture.

In one embodiment, the antibodies comprised in the disclosed composition are ActRII antagonists with no or low agonistic activity. In another embodiment, the antibody or functional fragment comprised in the disclosed composition binds the target protein ActRII and decreases the binding of myostatin to ActRII to a basal level. In a further aspect of this embodiment, the antibody or functional fragment comprised in the disclosed composition completely prevents myostatin from binding to ActRII. In a further embodiment, the antibody or functional fragment comprised in the disclosed composition inhibits Smad activation. In a further embodiment, the antibody or functional fragment comprised in the disclosed composition inhibits activin receptor type IIB mediated myostatin-induced inhibition of skeletal differentiation via the Smad-dependent pathway.

The binding may be determined by one or more assays that can be used to measure an activity which is either antagonism or agonism by the antibody. Preferably, the assays measure at least one of the effects of the antibody on ActRIIB that include: inhibition of myostatin binding to ActRIIB by ELISA, inihibition of myostatin induced signalling (for instance by a Smad dependent reporter gene assay), inhibition of myostatin induced

Smad phosphorylation (P-Smad ELISA) and inhibition of myostatin induced inhibition of skeletal muscle cell differentiation (for instance by a creatine kinase assay).

In one embodiment, the disclosure provides compositions comprising antibodies that specifically bind to the myostatin binding region (*i.e.* ligand binding domain) of ActRIIB. This ligand binding domain consists of amino acids 19-134 of SEQ ID NO: 181 and has been assigned SEQ ID NO: 182 herein. The ligand biding domain comprises several below described epitopes.

In one embodiment, the antibodies comprised in the disclosed composition bind to ActRIIB with a K_D of about 100nM or less, about 10nM or less, about 1nM or less. Preferably, the antibodies comprised in the disclosed composition bind to ActRIIB with an affinity of 100pM or less (*i.e.* about 100pM, about 50pM, about 10pM, about 2 pM, about 1pM or less). In one embodiment, the antibodies comprised in the disclosed composition bind to ActRIIB with an affinity of between about 1 and about 10pM.

In one embodiment, the antibodies comprised in the disclosed composition do not cross-react with an ActRIIB related protein, particularly do not cross-react with human ActRIIA (NP_001607.1, GI:4501897). In another embodiment, the antibodies comprised in the disclosed composition cross-react with Act RIIA and bind to ActRIIB with equivalent affinity, or about 1, 2, 3 , 4 or 5-fold greater affinity than they bind to ActRIIA, more preferably about 10-fold, still more preferably about 20-, 30- ,40- or 50-fold, still more preferably about 100-fold.

In one embodiment, the antibodies comprised in the disclosed composition bind to ActRIIA with an affinity of 100pM or more (*i.e.* about 250pM, about 500pM, about 1nM, about 5nM or more).

In one embodiment the antibodies comprised in the disclosed composition are of the IgG_2 isotype.

In another embodiment, the antibodies comprised in the disclosed composition are of the IgG_1 isotype. In a further embodiment, the antibodies comprised in the disclosed composition are of the IgG1 isotype and have an altered effector function through mutation of the Fc region. Said altered effector function may be a reduced ADCC and CDC activity. In one embodiment, said altered effector function is silenced ADCC and CDC activity.

In another related embodiment, the antibodies comprised in the disclosed composition are fully human or humanized IgG1 antibodies with no antibody dependent cellular cytotoxicity (ADCC) activity or CDC activity and bind to a region of ActRIIB consisting of amino acids 19-134 of SEQ ID NO:181.

In another related embodiment, the antibodies comprised in the disclosed composition are fully human or humanized IgG1 antibodies with reduced antibody dependent cellular cytotoxicity (ADCC) activity or CDC activity and bind to a region of ActRIIB consisting of amino acids 19-134 of SEQ ID NO:181.

The present disclosure relates to compositions comprising human or humanized anti-ActRIIB antibodies for use reducing time to mechanical ventilation liberation in an intensive care patient with failure to wean from the mechanical ventilation.

In certain embodiments, the antibodies comprised in the disclosed composition are derived from particular heavy and light chain sequences and/or comprise particular structural features such as CDR regions comprising particular amino acid sequences. The disclosure provides isolated ActRIIB antibodies, methods of making such antibodies, immunoconjugates and multivalent or multispecific molecules comprising such antibodies and pharmaceutical compositions containing the antibodies, immunoconjugates or bispecific molecules.

In alternative embodiments the disclosure relates to compositions comprising a myostatin antagonist for use according to the following aspects:

1. A myostatin antagonist for use in treating sporadic inclusion body myositis.

2. A myostatin antagonist for use according to aspect 1, wherein said myostatin antagonist is to be administered to a patient in need thereof at a dose of about 1-10 mg/kg.

3. A myostatin antagonist for use according to aspects 1-2, wherein said myostatin antagonist is to be administered at a dose of about 1, about 3 or about 10 mg/kg body weight.

4. A myostatin antagonist for use according to aspects 1-3, wherein said myostatin antagonist is to be administered intraveneously.

5. A myostatin antagonist for use according to anyone of aspects 1-4, wherein said myostatin anatagonis is to be administred every four weeks.

6. A myostatin antagonist for use according to anyone of aspects 1-5, wherein said patient is ambulatory.

7. A myostatin antagonist for use according to anyone of aspects 1-6, wherein treating sporadic inclusion body myositis comprises slowing down the progression of the disease or improving physical function and mobility.

8. A myostatin antagonist for use according to anyone of aspects 1-6, wherein treating sporadic inclusion body myositis comprises improving dysphagia or swallowing difficulties.

9. A myostatin antagonist for use according to anyone of aspects 1-6, wherein treating sporadic inclusion body myositis comprises improving upper extremity strength.

10. A myostatin antagonist for use according to anyone of aspects 1-6, wherein treating sporadic inclusion body myositis comprises reducing incidence of falls or preventing falls.

11. A myostatin antagonist for use according to anyone of aspects 1-10, wherein the myostatin antagonist is a myostatin receptor binding molecule.

12. A myostatin antagonist for use according to anyone of aspects 1-11, wherein the myostatin antagonist is an ActRII receptor antagonist.

13. A myostatin antagonist for use according to anyone of aspects 1-12, wherein the myostatin antagonist is an anti-ActRII receptor antibody.

14. A myostatin antagonist for use according to anyone of aspects 1-13, wherein the anti-ActRII receptor antibody is bimagrumab.

15. A myostatin antagonist for use according to any one of aspects 13-14, wherein the myostatin antagonist is an anti-ActRII antibody that binds to an epitope of ActRIIB consisting of amino acids 19-134 of SEQ ID NO: 181 (SEQ ID NO: 182).

16. A myostatin antagonist for use according to anyone of aspects 13-15, wherein the anti-ActRII antibody binds to an epitope of ActRIIB comprising or consisting of:

(a) amino acids 78-83 of SEQ ID NO: 181 (WLDDFN – SEQ ID NO:188);

(b) amino acids 76-84 of SEQ ID NO: 181 (GCWLDDFNC – SEQ ID NO:186);

(c) amino acids 75-85 of SEQ ID NO: 181 (KGCWLDDFNCY – SEQ ID NO:190);

(d) amino acids 52-56 of SEQ ID NO: 181 (EQDKR – SEQ ID NO:189);

(e) amino acids 49-63 of SEQ ID NO: 181 (CEGEQDKRLHCYASW – SEQ ID NO:187);

(f) amino acids 29-41 of SEQ ID NO: 181 (CIYYNANWELERT– SEQ ID NO:191);

(g) amino acids 100-110 of SEQ ID NO: 181 (YFCCCEGNFCN – SEQ ID NO:192); or

(h) amino acids 78-83 of SEQ ID NO: 181 (WLDDFN) and amino acids 52-56 of SEQ ID NO: 181 (EQDKR).

17. A myostatin antagonist for use according to any of aspects 13-16, wherein the anti-ActRIIB antibody is selected from the group consisting of:

a) an anti-ActRIIB antibody that binds to an epitope of ActRIIB comprising :

(a) amino acids 78-83 of SEQ ID NO: 181 (WLDDFN – SEQ ID NO:188);

(b) amino acids 76-84 of SEQ ID NO: 181 (GCWLDDFNC – SEQ ID NO:186);

(c) amino acids 75-85 of SEQ ID NO: 181 (KGCWLDDFNCY – SEQ ID NO:190);

(d) amino acids 52-56 of SEQ ID NO: 181 (EQDKR – SEQ ID NO:189);

(e) amino acids 49-63 of SEQ ID NO: 181 (CEGEQDKRLHCYASW – SEQ ID NO:187);

(f) amino acids 29-41 of SEQ ID NO: 181 (CIYYNANWELERT– SEQ ID NO:191);

(g) amino acids 100-110 of SEQ ID NO: 181 (YFCCCEGNFCN – SEQ ID NO:192); or

(h) amino acids 78-83 of SEQ ID NO: 181 (WLDDFN) and amino acids 52-56 of SEQ ID NO: 181 (EQDKR).

; and b) an antagonist antibody to ActRIIB that binds to an epitope of ActRIIB comprising amino acids 78-83 of SEQ ID NO: 181 (WLDDFN – SEQ ID NO:188);

(b) amino acids 76-84 of SEQ ID NO: 181 (GCWLDDFNC – SEQ ID NO:186);

(c) amino acids 75-85 of SEQ ID NO: 181 (KGCWLDDFNCY – SEQ ID NO:190);

(d) amino acids 52-56 of SEQ ID NO: 181 (EQDKR – SEQ ID NO:189);

(e) amino acids 49-63 of SEQ ID NO: 181 (CEGEQDKRLHCYASW – SEQ ID NO:187);

(f) amino acids 29-41 of SEQ ID NO: 181 (CIYYNANWELERT– SEQ ID NO:191);

(g) amino acids 100-110 of SEQ ID NO: 181 (YFCCCEGNFCN – SEQ ID NO:192); or

(h) amino acids 78-83 of SEQ ID NO: 181 (WLDDFN) and amino acids 52-56 of SEQ ID NO: 181 (EQDKR), wherein the antibody has a K_D of about 2 pM.

18. A myostatin antagonist for use according to any of aspects 13-17, wherein the antibody binds to ActRIIB with a 10-fold or greater affinity than it binds to ActRIIA.

19. A myostatin antagonist for use according to anyone of aspects 13-18, wherein the antibody comprises a heavy chain variable region CDR1 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-14; a heavy chain variable region CDR2 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 15-28; a heavy chain variable region CDR3 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 29-42; a light chain variable region CDR1 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 43-56; a light chain variable region CDR2 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 57-70; and a light chain variable region CDR3 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 71-84.

20. A myostatin antagonist for use according to any of aspects 13-19 wherein the antibody comprises:

(a) a heavy chain variable region CDR1 of SEQ ID NO: 1; a heavy chain variable region CDR2 of SEQ ID NO: 15; a heavy chain variable region CDR3 of SEQ ID NO: 29; a light chain variable region CDR1 of SEQ ID NO: 43; a light chain variable region CDR2 of SEQ ID NO: 57; and a light chain variable region CDR3 of SEQ ID NO: 71,

(b) a heavy chain variable region CDR1 of SEQ ID NO: 2; a heavy chain variable region CDR2 of SEQ ID NO: 16; a heavy chain variable region CDR3 of SEQ ID NO: 30; a light chain variable region CDR1 of SEQ ID NO: 44; a light chain variable region CDR2 of SEQ ID NO: 58; and a light chain variable region CDR3 of SEQ ID NO: 72,

(c) a heavy chain variable region CDR1 of SEQ ID NO: 3; a heavy chain variable region CDR2 of SEQ ID NO: 17; a heavy chain variable region CDR3 of SEQ ID NO: 31; a light chain variable region CDR1 of SEQ ID NO: 45; a light chain variable region CDR2 of SEQ ID NO: 59; and a light chain variable region CDR3 of SEQ ID NO: 73,

(d) a heavy chain variable region CDR1 of SEQ ID NO: 4; a heavy chain variable region CDR2 of SEQ ID NO: 18; a heavy chain variable region CDR3 of SEQ ID NO: 32; a light chain variable region CDR1 of SEQ ID NO: 46; a light chain variable region CDR2 of SEQ ID NO: 60; and a light chain variable region CDR3 of SEQ ID NO: 74,

(e) a heavy chain variable region CDR1 of SEQ ID NO: 5; a heavy chain variable region CDR2 of SEQ ID NO: 19; a heavy chain variable region CDR3 of SEQ ID NO: 33; a light chain variable region CDR1 of SEQ ID NO: 47; a light chain variable region CDR2 of SEQ ID NO: 61; and a light chain variable region CDR3 of SEQ ID NO: 75,

(f) a heavy chain variable region CDR1 of SEQ ID NO: 6; a heavy chain variable region CDR2 of SEQ ID NO: 20; a heavy chain variable region CDR3 of SEQ ID NO: 34; a light chain variable region CDR1 of SEQ ID NO: 48; a light chain variable region CDR2 of SEQ ID NO: 62; and a light chain variable region CDR3 of SEQ ID NO: 76,

(g) a heavy chain variable region CDR1 of SEQ ID NO: 7; a heavy chain variable region CDR2 of SEQ ID NO: 21; a heavy chain variable region CDR3 of SEQ ID NO: 35; a light chain variable region CDR1 of SEQ ID NO: 49; a light chain variable region CDR2 of SEQ ID NO: 63; and a light chain variable region CDR3 of SEQ ID NO: 77,

(h) a heavy chain variable region CDR1 of SEQ ID NO: 8; a heavy chain variable region CDR2 of SEQ ID NO: 22; a heavy chain variable region CDR3 of SEQ ID NO: 36; a light chain variable region CDR1 of SEQ ID NO: 50 a light chain variable region CDR2 of SEQ ID NO: 64; and a light chain variable region CDR3 of SEQ ID NO: 78,

(i) a heavy chain variable region CDR1 of SEQ ID NO: 9; a heavy chain variable region CDR2 of SEQ ID NO: 23; a heavy chain variable region CDR3 of SEQ ID NO: 37; a light chain variable region CDR1 of SEQ ID NO: 51; a light chain variable region CDR2 of SEQ ID NO: 65; and a light chain variable region CDR3 of SEQ ID NO: 79,

(j) a heavy chain variable region CDR1 of SEQ ID NO: 10; a heavy chain variable region CDR2 of SEQ ID NO: 24; a heavy chain variable region CDR3 of SEQ ID NO: 38; a light chain variable region CDR1 of SEQ ID NO: 52; a light chain variable region CDR2 of SEQ ID NO: 66; and a light chain variable region CDR3 of SEQ ID NO: 80,

(k) a heavy chain variable region CDR1 of SEQ ID NO: 11; a heavy chain variable region CDR2 of SEQ ID NO: 25; a heavy chain variable region CDR3 of SEQ ID NO: 39; a light chain variable region CDR1 of SEQ ID NO: 53; a light chain variable region CDR2 of SEQ ID NO: 67; and a light chain variable region CDR3 of SEQ ID NO: 81,

(l) a heavy chain variable region CDR1 of SEQ ID NO: 12; a heavy chain variable region CDR2 of SEQ ID NO: 26; a heavy chain variable region CDR3 of SEQ ID NO: 40; a light chain variable region CDR1 of SEQ ID NO: 54; a light chain variable region CDR2 of SEQ ID NO: 68; and a light chain variable region CDR3 of SEQ ID NO: 82,

(m) a heavy chain variable region CDR1 of SEQ ID NO: 13; a heavy chain variable region CDR2 of SEQ ID NO: 27; a heavy chain variable region CDR3 of SEQ ID NO: 41; a light chain variable region CDR1 of SEQ ID NO: 55; a light chain variable region CDR2 of SEQ ID NO: 69; and a light chain variable region CDR3 of SEQ ID NO: 83, or

(n) a heavy chain variable region CDR1 of SEQ ID NO: 14; a heavy chain variable region CDR2 of SEQ ID NO: 28; a heavy chain variable region CDR3 of SEQ ID NO: 42; a light chain variable region CDR1 of SEQ ID NO: 56; a light chain variable region CDR2 of SEQ ID NO: 70; and a light chain variable region CDR3 of SEQ ID NO: 84.

21. A myostatin antagonist for use according to according to any of aspects 13-20, wherein the antibody comprises a full length heavy chain amino acid sequence having at least 95% sequence identity to at least one sequence selected from the group consisting of SEQ ID NOs:146-150 and 156-160.

22. A myostatin antagonist for use according to according to any of aspects 13-21, wherein the antibody comprises a full length light chain amino acid sequence having at least 95% sequence identity to at least one sequence selected from the group consisting of SEQ ID NOs:141-145 and 151-155.

23. A myostatin antagonist for use according to according to any of aspects 13-22, wherein the antibody comprises:

(a) the variable heavy chain sequence of SEQ ID NO: 99 and variable light chain sequence of SEQ ID NO: 85;

(b) the variable heavy chain sequence of SEQ ID NO: 100 and variable light chain sequence of SEQ ID NO: 86;

(c) the variable heavy chain sequence of SEQ ID NO: 101 and variable light chain sequence of SEQ ID NO: 87;
(d) the variable heavy chain sequence of SEQ ID NO: 102 and variable light chain sequence of SEQ ID NO: 88;
(e) the variable heavy chain sequence of SEQ ID NO: 103 and variable light chain sequence of SEQ ID NO: 89;
(f) the variable heavy chain sequence of SEQ ID NO: 104 and variable light chain sequence of SEQ ID NO: 90;
(g) the variable heavy chain sequence of SEQ ID NO: 105 and variable light chain sequence of SEQ ID NO: 91;
(h) the variable heavy chain sequence of SEQ ID NO: 106 and variable light chain sequence of SEQ ID NO: 92;
(i) the variable heavy chain sequence of SEQ ID NO: 107 and variable light chain sequence of SEQ ID NO: 93;
(j) the variable heavy chain sequence of SEQ ID NO: 108 and variable light chain sequence of SEQ ID NO: 94;
(k) the variable heavy chain sequence of SEQ ID NO: 109 and variable light chain sequence of SEQ ID NO: 95;
(l) the variable heavy chain sequence of SEQ ID NO: 110 and variable light chain sequence of SEQ ID NO: 96;
(m) the variable heavy chain sequence of SEQ ID NO: 111 and variable light chain sequence of SEQ ID NO: 97; or
(n) the variable heavy chain sequence of SEQ ID NO: 112 and variable light chain sequence of SEQ ID NO: 98.

24. A myostatin antagonist for use according to according to any of aspects 13-23, wherein the antibody comprises:

(a) the heavy chain sequence of SEQ ID NO: 146 and light chain sequence of SEQ ID NO: 141;
(b) the heavy chain sequence of SEQ ID NO: 147 and light chain sequence of SEQ ID NO: 142;
(c) the heavy chain sequence of SEQ ID NO: 148 and light chain sequence of SEQ ID NO: 143;
(d) the heavy chain sequence of SEQ ID NO: 149 and light chain sequence of SEQ ID NO: 144;
(e) the heavy chain sequence of SEQ ID NO: 150 and light chain sequence of SEQ ID NO: 145;

(f) the heavy chain sequence of SEQ ID NO: 156 and light chain sequence of SEQ ID NO: 151;
(g) the heavy chain sequence of SEQ ID NO: 157 and light chain sequence of SEQ ID NO: 152;
(h) the heavy chain sequence of SEQ ID NO: 158 and light chain sequence of SEQ ID NO: 153;
(i) the heavy chain sequence of SEQ ID NO: 159 and light chain sequence of SEQ ID NO: 154; or
(j) the heavy chain sequence of SEQ ID NO: 160 and light chain sequence of SEQ ID NO: 155.

25. A myostatin antagonist for use according to according to any of aspects 13-24, wherein the antibody comprised in said composition cross-blocks or is cross blocked by at least one antibody of claim 10 from binding to ActRIIB.

26. A myostatin antagonist for use according to according to any of aspects 13-25, wherein the antibody comprised in said composition has altered effector function through mutation of the Fc region.

27. A myostatin antagonist for use according to according to any of aspects 13-26, wherein the antibody comprised in said composition binds to an epitope recognised by an antibody listed in aspects 15-17.

28. A myostatin antagonist for use according to any of aspects 13-27, wherein the antibody is encoded by pBW522 (DSM22873) or pBW524 (DSM22874).

29. Bimaghumab for use in treating sporadic inclusion body myositis, wherein bimagrumab is to be administered intraveneously at a dose of about 1-10 mg/kg body weight every four weeks.

30. Bimagrumab for use in treating sporadic inclusion body myositis, wherein bimagrumab is to be administered intraveneously at a dose of about 1 mg/kg body weight every four weeks.

31. Bimagrumab for use in treating sporadic inclusion body myositis, wherein bimagrumab is to be administered intraveneously at a dose of about 3 mg/kg body weight every four weeks.

32. Bimagrumab for use in treating sporadic inclusion body myositis, wherein bimagrumab is to be administered intraveneously at a dose of about 10 mg/kg body weight every four weeks.

33. A method of treating sporadic inclusion body myositis comprising administering a therapeutically effective amount of a myostatin antagonist to a patient in need thereof.

34. A method of treating sporadic inclusion body myositis according to claim 33, comprising adminstering said myostatin antagonist at a dose of about 1-10 mg/kg.

35. A method of treating sporadic inclusion body myositis according to any of aspects 33-34, comprising administering at a dose of about 1, about 3 or about 10 mg/kg body weight.

36. A method of treating sporadic inclusion body myositis according to any of aspects 33-35, comprising administering said myostatin antagonist intraveneously.

37. A method of treating sporadic inclusion body myositis according to any of aspects 33-36, comprising administering said myostatin anatagonist every four weeks.

38. A method of treating sporadic inclusion body myositis according to any of aspects 33-37, wherein said patient is ambulatory.

39. A method of treating sporadic inclusion body myositis according to any of aspects 33-38, wherein treating sporadic inclusion body myositis comprises slowing down the progression of the disease or improving physical function and mobility.

40. A method of treating sporadic inclusion body myositis according to any of aspects 33-38, wherein treating sporadic inclusion body myositis comprises improving dysphagia or swallowing difficulties.

41. A method of treating sporadic inclusion body myositis according to any of aspects 33-38, wherein treating sporadic inclusion body myositis comprises improving upper extremity strength.

42. A method of treating sporadic inclusion body myositis according to any of aspects 33-38, wherein treating sporadic inclusion body myositis comprises reducing incidence of falls or preventing falls.

43. A method of treating sporadic inclusion body myositis according to any of aspects 33-42, wherein the myostatin antagonist is a myostatin receptor binding molecule.

44. A method of treating sporadic inclusion body myositis according to any of aspects 33-43, wherein the myostatin antagonist is an ActRII receptor antagonist.

45. A method of treating sporadic inclusion body myositis according to any of aspects 33-44, wherein the myostatin antagonist is an anti-ActRII receptor antibody.

46. A method of treating sporadic inclusion body myositis according to any of aspects 33-45, wherein the anti-ActRII receptor antibody is bimagrumab.

47. A method of treating sporadic inclusion body myositis according to any of aspects 45 or 46, wherein the myostatin antagonist is an anti-ActRII antibody that binds to an epitope of ActRIIB consisting of amino acids 19-134 of SEQ ID NO: 181 (SEQ ID NO: 182).

48. A method of treating sporadic inclusion body myositis according to any of aspects 45-47, wherein the anti-ActRII antibody binds to an epitope of ActRIIB comprising or consisting of:

(a) amino acids 78-83 of SEQ ID NO: 181 (WLDDFN – SEQ ID NO:188);

(b) amino acids 76-84 of SEQ ID NO: 181 (GCWLDDFNC – SEQ ID NO:186);

(c) amino acids 75-85 of SEQ ID NO: 181 (KGCWLDDFNCY – SEQ ID NO:190);

(d) amino acids 52-56 of SEQ ID NO: 181 (EQDKR – SEQ ID NO:189);

(e) amino acids 49-63 of SEQ ID NO: 181 (CEGEQDKRLHCYASW – SEQ ID NO:187);

(f) amino acids 29-41 of SEQ ID NO: 181 (CIYYNANWELERT– SEQ ID NO:191);

(g) amino acids 100-110 of SEQ ID NO: 181 (YFCCCEGNFCN – SEQ ID NO:192); or

(h) amino acids 78-83 of SEQ ID NO: 181 (WLDDFN) and amino acids 52-56 of SEQ ID NO: 181 (EQDKR).

49. A method of treating sporadic inclusion body myositis according to any of aspects 45-48, wherein the anti-ActRIIB antibody is selected from the group consisting of:

a) an anti-ActRIIB antibody that binds to an epitope of ActRIIB comprising :

(a) amino acids 78-83 of SEQ ID NO: 181 (WLDDFN – SEQ ID NO:188);

(b) amino acids 76-84 of SEQ ID NO: 181 (GCWLDDFNC – SEQ ID NO:186);

(c) amino acids 75-85 of SEQ ID NO: 181 (KGCWLDDFNCY – SEQ ID NO:190);

(d) amino acids 52-56 of SEQ ID NO: 181 (EQDKR – SEQ ID NO:189);

(e) amino acids 49-63 of SEQ ID NO: 181 (CEGEQDKRLHCYASW – SEQ ID NO:187);

(f) amino acids 29-41 of SEQ ID NO: 181 (CIYYNANWELERT– SEQ ID NO:191);

(g) amino acids 100-110 of SEQ ID NO: 181 (YFCCCEGNFCN – SEQ ID NO:192); or

(h) amino acids 78-83 of SEQ ID NO: 181 (WLDDFN) and amino acids 52-56 of SEQ ID NO: 181 (EQDKR).

; and b) an antagonist antibody to ActRIIB that binds to an epitope of ActRIIB comprising amino acids 78-83 of SEQ ID NO: 181 (WLDDFN – SEQ ID NO:188);

(b) amino acids 76-84 of SEQ ID NO: 181 (GCWLDDFNC – SEQ ID NO:186);

(c) amino acids 75-85 of SEQ ID NO: 181 (KGCWLDDFNCY – SEQ ID NO:190);

(d) amino acids 52-56 of SEQ ID NO: 181 (EQDKR – SEQ ID NO:189);

(e) amino acids 49-63 of SEQ ID NO: 181 (CEGEQDKRLHCYASW – SEQ ID NO:187);

(f) amino acids 29-41 of SEQ ID NO: 181 (CIYYNANWELERT– SEQ ID NO:191);

(g) amino acids 100-110 of SEQ ID NO: 181 (YFCCCEGNFCN – SEQ ID NO:192); or

(h) amino acids 78-83 of SEQ ID NO: 181 (WLDDFN) and amino acids 52-56 of SEQ ID NO: 181 (EQDKR), wherein the antibody has a K_D of about 2 pM.

50. A method of treating sporadic inclusion body myositis according to any of aspects 45-49, wherein the antibody binds to ActRIIB with a 10-fold or greater affinity than it binds to ActRIIA.

51. A method of treating sporadic inclusion body myositis to any of aspects 45-50, wherein the antibody comprises a heavy chain variable region CDR1 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-14; a heavy chain variable region CDR2 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 15-28; a heavy chain variable region CDR3 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 29-42; a light chain variable region CDR1 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 43-56; a light chain variable region CDR2 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 57-70; and a

light chain variable region CDR3 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 71-84.

52. A method of treating sporadic inclusion body myositis according to any of aspects 45-51, wherein the antibody comprises:

(a) a heavy chain variable region CDR1 of SEQ ID NO: 1; a heavy chain variable region CDR2 of SEQ ID NO: 15; a heavy chain variable region CDR3 of SEQ ID NO: 29; a light chain variable region CDR1 of SEQ ID NO: 43; a light chain variable region CDR2 of SEQ ID NO: 57; and a light chain variable region CDR3 of SEQ ID NO: 71,

(b) a heavy chain variable region CDR1 of SEQ ID NO: 2; a heavy chain variable region CDR2 of SEQ ID NO: 16; a heavy chain variable region CDR3 of SEQ ID NO: 30; a light chain variable region CDR1 of SEQ ID NO: 44; a light chain variable region CDR2 of SEQ ID NO: 58; and a light chain variable region CDR3 of SEQ ID NO: 72,

(c) a heavy chain variable region CDR1 of SEQ ID NO: 3; a heavy chain variable region CDR2 of SEQ ID NO: 17; a heavy chain variable region CDR3 of SEQ ID NO: 31; a light chain variable region CDR1 of SEQ ID NO: 45; a light chain variable region CDR2 of SEQ ID NO: 59; and a light chain variable region CDR3 of SEQ ID NO: 73,

(d) a heavy chain variable region CDR1 of SEQ ID NO: 4; a heavy chain variable region CDR2 of SEQ ID NO: 18; a heavy chain variable region CDR3 of SEQ ID NO: 32; a light chain variable region CDR1 of SEQ ID NO: 46; a light chain variable region CDR2 of SEQ ID NO: 60; and a light chain variable region CDR3 of SEQ ID NO: 74,

(e) a heavy chain variable region CDR1 of SEQ ID NO: 5; a heavy chain variable region CDR2 of SEQ ID NO: 19; a heavy chain variable region CDR3 of SEQ ID NO: 33; a light chain variable region CDR1 of SEQ ID NO: 47; a light chain variable region CDR2 of SEQ ID NO: 61; and a light chain variable region CDR3 of SEQ ID NO: 75,

(f) a heavy chain variable region CDR1 of SEQ ID NO: 6; a heavy chain variable region CDR2 of SEQ ID NO: 20; a heavy chain variable region CDR3 of SEQ ID NO: 34; a light chain variable region CDR1 of SEQ ID NO: 48; a light chain variable region CDR2 of SEQ ID NO: 62; and a light chain variable region CDR3 of SEQ ID NO: 76,

(g) a heavy chain variable region CDR1 of SEQ ID NO: 7; a heavy chain variable region CDR2 of SEQ ID NO: 21; a heavy chain variable region CDR3 of SEQ ID NO: 35; a light chain variable region CDR1 of SEQ ID NO: 49; a light chain variable region CDR2 of SEQ ID NO: 63; and a light chain variable region CDR3 of SEQ ID NO: 77,

(h) a heavy chain variable region CDR1 of SEQ ID NO: 8; a heavy chain variable region CDR2 of SEQ ID NO: 22; a heavy chain variable region CDR3 of SEQ ID NO: 36; a light chain variable region CDR1 of SEQ ID NO: 50 a light chain variable region CDR2 of SEQ ID NO: 64; and a light chain variable region CDR3 of SEQ ID NO: 78,

(i) a heavy chain variable region CDR1 of SEQ ID NO: 9; a heavy chain variable region CDR2 of SEQ ID NO: 23; a heavy chain variable region CDR3 of SEQ ID NO: 37; a light chain variable region CDR1 of SEQ ID NO: 51; a light chain variable region CDR2 of SEQ ID NO: 65; and a light chain variable region CDR3 of SEQ ID NO: 79,

(j) a heavy chain variable region CDR1 of SEQ ID NO: 10; a heavy chain variable region CDR2 of SEQ ID NO: 24; a heavy chain variable region CDR3 of SEQ ID NO: 38; a light chain variable region CDR1 of SEQ ID NO: 52; a light chain variable region CDR2 of SEQ ID NO: 66; and a light chain variable region CDR3 of SEQ ID NO: 80,

(k) a heavy chain variable region CDR1 of SEQ ID NO: 11; a heavy chain variable region CDR2 of SEQ ID NO: 25; a heavy chain variable region CDR3 of SEQ ID NO: 39; a light chain variable region CDR1 of SEQ ID NO: 53; a light chain variable region CDR2 of SEQ ID NO: 67; and a light chain variable region CDR3 of SEQ ID NO: 81,

(l) a heavy chain variable region CDR1 of SEQ ID NO: 12; a heavy chain variable region CDR2 of SEQ ID NO: 26; a heavy chain variable region CDR3 of SEQ ID NO: 40; a light chain variable region CDR1 of SEQ ID NO: 54; a light chain variable region CDR2 of SEQ ID NO: 68; and a light chain variable region CDR3 of SEQ ID NO: 82,

(m) a heavy chain variable region CDR1 of SEQ ID NO: 13; a heavy chain variable region CDR2 of SEQ ID NO: 27; a heavy chain variable region CDR3 of SEQ ID NO: 41; a light chain variable region CDR1 of SEQ ID NO: 55; a light chain variable region CDR2 of SEQ ID NO: 69; and a light chain variable region CDR3 of SEQ ID NO: 83, or

(n) a heavy chain variable region CDR1 of SEQ ID NO: 14; a heavy chain variable region CDR2 of SEQ ID NO: 28; a heavy chain variable region CDR3 of SEQ ID NO: 42; a light chain variable region CDR1 of SEQ ID NO: 56; a light chain variable region CDR2 of SEQ ID NO: 70; and a light chain variable region CDR3 of SEQ ID NO: 84.

53. A method of treating sporadic inclusion body myositis according to any of aspects 45-52, wherein the antibody comprises a full length heavy chain amino acid sequence having at least 95% sequence identity to at least one sequence selected from the group consisting of SEQ ID NOs:146-150 and 156-160.

54. A method of treating sporadic inclusion body myositis to according to any of aspects 45-53, wherein the antibody comprises a full length light chain amino acid sequence having at least 95% sequence identity to at least one sequence selected from the group consisting of SEQ ID NOs:141-145 and 151-155.

55. A method of treating sporadic inclusion body myositis according to any of aspects 45-54, wherein the antibody comprises:

(a) the variable heavy chain sequence of SEQ ID NO: 99 and variable light chain sequence of SEQ ID NO: 85;

(b) the variable heavy chain sequence of SEQ ID NO: 100 and variable light chain sequence of SEQ ID NO: 86;

(c) the variable heavy chain sequence of SEQ ID NO: 101 and variable light chain sequence of SEQ ID NO: 87;

(d) the variable heavy chain sequence of SEQ ID NO: 102 and variable light chain sequence of SEQ ID NO: 88;

(e) the variable heavy chain sequence of SEQ ID NO: 103 and variable light chain sequence of SEQ ID NO: 89;

(f) the variable heavy chain sequence of SEQ ID NO: 104 and variable light chain sequence of SEQ ID NO: 90;

(g) the variable heavy chain sequence of SEQ ID NO: 105 and variable light chain sequence of SEQ ID NO: 91;

(h) the variable heavy chain sequence of SEQ ID NO: 106 and variable light chain sequence of SEQ ID NO: 92;

(i) the variable heavy chain sequence of SEQ ID NO: 107 and variable light chain sequence of SEQ ID NO: 93;

(j) the variable heavy chain sequence of SEQ ID NO: 108 and variable light chain sequence of SEQ ID NO: 94;

(k) the variable heavy chain sequence of SEQ ID NO: 109 and variable light chain sequence of SEQ ID NO: 95;

(l) the variable heavy chain sequence of SEQ ID NO: 110 and variable light chain sequence of SEQ ID NO: 96;

(m) the variable heavy chain sequence of SEQ ID NO: 111 and variable light chain sequence of SEQ ID NO: 97; or

(n) the variable heavy chain sequence of SEQ ID NO: 112 and variable light chain sequence of SEQ ID NO: 98.

56. A method of treating sporadic inclusion body myositis according to any of aspects 45-55, wherein the antibody comprises:

(a) the heavy chain sequence of SEQ ID NO: 146 and light chain sequence of SEQ ID NO: 141;

(b) the heavy chain sequence of SEQ ID NO: 147 and light chain sequence of SEQ ID NO: 142;

(c) the heavy chain sequence of SEQ ID NO: 148 and light chain sequence of SEQ ID NO: 143;

(d) the heavy chain sequence of SEQ ID NO: 149 and light chain sequence of SEQ ID NO: 144;

(e) the heavy chain sequence of SEQ ID NO: 150 and light chain sequence of SEQ ID NO: 145;

(f) the heavy chain sequence of SEQ ID NO: 156 and light chain sequence of SEQ ID NO: 151;

(g) the heavy chain sequence of SEQ ID NO: 157 and light chain sequence of SEQ ID NO: 152;

(h) the heavy chain sequence of SEQ ID NO: 158 and light chain sequence of SEQ ID NO: 153;

(i) the heavy chain sequence of SEQ ID NO: 159 and light chain sequence of SEQ ID NO: 154; or

(j) the heavy chain sequence of SEQ ID NO: 160 and light chain sequence of SEQ ID NO: 155.

57. A method of treating sporadic inclusion body myositis according to any of aspects 45-56, wherein the antibody comprised in said composition cross-blocks or is cross blocked by at least one antibody of aspect 10 from binding to ActRIIB.

58. A method of treating sporadic inclusion body myositis according to any of aspects 45-57, wherein the antibody comprised in said composition has altered effector function through mutation of the Fc region.

59. A method of treating sporadic inclusion body myositis according to any of aspects 45-58, wherein the antibody comprised in said composition binds to an epitope recognised by an antibody listed in aspects 46-48.

60. A method of treating sporadic inclusion body myositis according to any of aspects 45-59, wherein the antibody is encoded by pBW522 (DSM22873) or pBW524 (DSM22874).

61. A method of treating sporadic inclusion body myositis comprsing administering bimagrumab,

62. A method of treating sporadic inclusion body myositis comprsing administering bimagrumab,wherein bimagrumab is to be administered intraveneously at a dose of about 1-10 mg/kg body weight every four weeks.

63. A method of treating sporadic inclusion body myositis comprsing administering bimagrumab,, wherein bimagrumab is to be administered intraveneously at a dose of about 1 mg/kg body weight every four weeks.

64. A method of treating sporadic inclusion body myositis comprsing administering bimagrumab,, wherein bimagrumab is to be administered intraveneously at a dose of about 3 mg/kg body weight every four weeks.

65. A method of treating sporadic inclusion body myositis comprsing administering bimagrumab,, wherein bimagrumab is to be administered intraveneously at a dose of about 10 mg/kg body weight every four weeks.

66. Bimagrumab for use in treating sporadic inclusion body myositis.

67. A composition comprising 150 mg/ml of bimagrumab for use in a method of treating sporadic inclusion body myositis.

68. A unitary dosage form comprising 150 mg/ml of bimagrumab.

In further aspects the unitary dosage form, i.e., a vial , comprises 100-200 mg/ml of bimagrumab, preferably 100, 105, 110, 115, 120, 125, 130, 135, 140, 145, 150, 155, 160,165, 170, 175, 180, 185, 190, 195, 200 mg/ml of bimagrumab.

69. An infusion bag comprising an appropriate amount of bimagrumab from one or more vials diluted with a solution.

The solution is preferably a dextrose solution.

In some further embodiments, the myostatin antagonist, preferably the AcRII antagonist or anti-ActRII antibody such as bimagrumab is ito be administered at a dose of about 1, 2, 3, 4, 5, 6, 7, 8, 9 or 10 mg/kg body weight.

Various aspects of the disclosure are described in further detail in the following subsections. Standard assays to evaluate the binding ability of the antibodies toward ActRII of various species are known in the art, including for example, ELISAs, western blots and RIAs. Suitable assays are described in detail in the Examples. The binding affinity of the antibodies also can be assessed by standard assays known in the art, such as by Biacore analysis or Solution Equilibrium Titration. Surface plasmon resonance based techniques such as Biacore can determine the binding kinetics which allows the calculation of the binding affinity. Assays to evaluate the effects of the antibodies on functional properties of ActRIIB (*e.g.* receptor binding, preventing or inducing human B cell proliferation or IgG production) are described in further detail in the Examples.

Accordingly, an antibody that "inhibits" one or more of these ActRII functional properties (*e.g.* biochemical, immunochemical, cellular, physiological or other biological activities, or the like) as determined according to methodologies known to the art and described herein, will be understood to relate to a statistically significant decrease in the particular activity relative to that seen in the absence of the antibody (*e.g.* or when a control antibody of irrelevant specificity is present). An antibody that inhibits ActRII activity effects such a statistically significant decrease by at least 10% of the measured parameter, by at least 50%, 80% or 90%, and in certain embodiments an antibody of the disclosure may inhibit greater than 95%, 98% or 99% of ActRIIB functional activity.

The ability or extent to which an antibody or other binding agent is able to interfere with the binding of another antibody or binding molecule to ActRII, and therefore whether it can be said to cross-block according to the disclosure, can be determined using standard competition binding assays. One suitable assay involves the use of the Biacore

technology (*e.g.* by using a BIAcore instrument (Biacore, Uppsala, Sweden)), which can measure the extent of interactions using surface plasmon resonance technology. Another assay for measuring cross-blocking uses an ELISA-based approach. A further assay uses FACS analysis, wherein competition of various antibodies for binding to ActRIIB expressing cells is tested (such as described in the Examples).

According to the disclosure, a cross-blocking antibody or other binding agent according to the disclosure binds to ActRII in the described BIAcore cross-blocking assay such that the recorded binding of the combination (mixture) of the antibodies or binding agents is between 80% and 0.1% (*e.g.* 80% to 4%) of the maximum theoretical binding, specifically between 75% and 0.1% (*e.g.* 75% to 4%) of the maximum theoretical binding, and more specifically between 70% and 0.1% (*e.g.* 70% to 4%), and more specifically between 65% and 0.1% (*e.g.* 65% to 4%) of maximum theoretical binding (as defined above) of the two antibodies or binding agents in combination.

An antibody is defined as cross-blocking an anti-ActRIIB antibody of the disclosure in an ELISA assay, if the test antibody is able to cause a reduction of anti-ActRII antibody binding to ActRIIB of between 60% and 100%, specifically between 70% and 100%, and more specifically between 80% and 100%, when compared to the positive control wells (*i.e.* the same anti-ActRIIB antibody and ActRIIB, but no "test" cross-blocking antibody). Examples of cross blocking antibodies as cited herein are MOR08159 and MOR08213 (disclosed in WO2010/125003). Thus, the disclosure provides compositions comprising antibodies that cross block MOR08159 or MOR08213 for binding to ActRIIB.

Recombinant antibodies

Antibodies, e.g., antagonist antibodies to ActRII, such as bimagrumab, comprised in the compositions used within this disclosure include the human recombinant antibodies, isolated and structurally characterized, as described in the Examples. The V_H amino acid sequences of antibodies comprised in the inventive compositions are shown in SEQ ID NOs: 99-112. The V_L amino acid sequences of antibodies comprised in the inventive compositions are shown in SEQ ID NOs: 85-98 respectively. Examples of preferred full length heavy chain amino acid sequences of antibodies comprised in the inventive compositions are shown in SEQ ID NOs: 146-150 and 156-160. Examples of preferred full length light chain amino acid sequences of antibodies comprised in the inventive compositions are shown in SEQ ID NOs: 141-145 and 151-155 respectively. Other antibodies comprised in the inventive compositions include amino acids that have been

mutated by amino acid deletion, insertion or substitution, yet have at least 60, 70, 80, 90, 95, 97 or 99 percent identity in the CDR regions with the CDR regions depicted in the sequences described above. In some embodiments, it includes mutant amino acid sequences wherein no more than 1, 2, 3, 4 or 5 amino acids have been mutated by amino acid deletion, insertion or substitution in the CDR regions when compared with the CDR regions depicted in the sequence described above.

Further, variable heavy chain parental nucleotide sequences are shown in SEQ ID NOs: 127-140. Variable light chain parental nucleotide sequences are shown in SEQ ID NOs: 113-126. Full length light chain nucleotide sequences optimized for expression in a mammalian cell are shown in SEQ ID NOs: 161-165 and 171-175. Full length heavy chain nucleotide sequences optimized for expression in a mammalian cell are shown in SEQ ID NOs: 166-170 and 176-180. Other antibodies comprised in the inventive compositions include amino acids or are encoded by nucleic acids that have been mutated, yet have at least 60 or more (*i.e.* 80, 90, 95, 97, 99 or more) percent identity to the sequences described above. In some embodiments, it includes mutant amino acid sequences wherein no more than 1, 2, 3, 4 or 5 amino acids have been mutated by amino acid deletion, insertion or substitution in the variable regions when compared with the variable regions depicted in the sequence described above.

Since each of these antibodies binds the same epitope and are progenies from the same parental antibody, the V_H, V_L, full length light chain, and full length heavy chain sequences (nucleotide sequences and amino acid sequences) can be "mixed and matched" to create other anti-ActRIIB binding molecules of the disclosure. ActRIIB binding of such "mixed and matched" antibodies can be tested using the binding assays described above and in the Examples (*e.g.* ELISAs). When these chains are mixed and matched, a V_H sequence from a particular V_H/V_L pairing should be replaced with a structurally similar V_H sequence. Likewise a full length heavy chain sequence from a particular full length heavy chain / full length light chain pairing should be replaced with a structurally similar full length heavy chain sequence. Likewise, a V_L sequence from a particular V_H/V_L pairing should be replaced with a structurally similar V_L sequence. Likewise a full length light chain sequence from a particular full length heavy chain / full length light chain pairing should be replaced with a structurally similar full length light chain sequence. Accordingly, in one aspect, the disclosure provides compositions comprising a recombinant anti-ActRII antibody or antigen binding region thereof having: a heavy chain variable region comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 99-112; and a light chain variable region comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 85-98.

In another aspect, the disclosure provides compositions comprising:

(i) an isolated recombinant anti-ActRII antibody having: a full length heavy chain comprising an amino acid sequence selected from the group consisting of SEQ ID NOs:99-112; and a full length light chain comprising an amino acid sequence selected from the group consisting of SEQ ID NOs:85-98, or

(ii) a functional protein comprising an antigen binding portion thereof.

In another aspect, the disclosure provides compositions comprising:

(i) an isolated recombinant anti-ActRII antibody having a full length heavy chain encoded by a nucleotide sequence that has been optimized for expression in the cell of a mammalian selected from the group consisting of SEQ ID NOs:127-140, and a full length light chain encoded by a nucleotide sequence that has been optimized for expression in the cell of a mammalian selected from the group consisting of SEQ ID NOs:113-126, or

(ii) a functional protein comprising an antigen binding portion thereof.

Examples of amino acid sequences of the V_H CDR1s of the antibodies comprised in the inventive compositions are shown in SEQ ID NOs: 1-14. The amino acid sequences of the V_H CDR2s of the antibodies are shown in SEQ ID NOs: 15-28. The amino acid sequences of the V_H CDR3s of the antibodies are shown in SEQ ID NOs: 29-42. The amino acid sequences of the V_L CDR1s of the antibodies are shown in SEQ ID NOs: 43-56. The amino acid sequences of the V_L CDR2s of the antibodies are shown in SEQ ID NOs: 57-70. The amino acid sequences of the V_L CDR3s of the antibodies are shown in SEQ ID NOs: 71-84. The CDR regions are delineated using the Kabat system (Kabat, E. A., *et al.*, 1991 Sequences of Proteins of Immunological Interest, Fifth Edition, U.S. Department of Health and Human Services, NIH Publication No. 91-3242). An alternative method of determining CDR regions uses the method devised by Chothia (Chothia *et al.* 1989, Nature, 342:877-883). The Chothia definition is based on the location of the structural loop regions. However, due to changes in the numbering system used by Chothia (see e.g. http://www.biochem.ucl.ac.uk/~martin/abs/GeneralInfo.html and http://www.bioinf.org.uk/abs/), this system is now less commonly used. Other systems for defining CDRs exist and are also mentioned in these two websites.

Given that each of these antibodies can bind to ActRII and that antigen-binding specificity is provided primarily by the CDR1, 2 and 3 regions, the V_H CDR1, 2 and 3 sequences and V_L CDR1, 2 and 3 sequences can be "mixed and matched" (*i.e.* CDRs from different antibodies can be mixed and matched, each antibody containing a V_H CDR1, 2 and 3 and a V_L CDR1, 2 and 3 create other anti-ActRII binding molecules of the disclosure. ActRIIB binding of such "mixed and matched" antibodies can be tested using

the binding assays described above and in the Examples (*e.g.* ELISAs). When V_H CDR sequences are mixed and matched, the CDR1, CDR2 and/or CDR3 sequence from a particular V_H sequence should be replaced with a structurally similar CDR sequence(s). Likewise, when V_L CDR sequences are mixed and matched, the CDR1, CDR2 and/or CDR3 sequence from a particular V_L sequence should be replaced with a structurally similar CDR sequence(s). It will be readily apparent to the ordinarily skilled artisan that novel V_H and V_L sequences can be created by substituting one or more V_H and/or V_L CDR region sequences with structurally similar sequences from the CDR sequences shown herein for monoclonal antibodies.

Anti-ActRII antibody comprised in the disclosed compositions, or antigen binding region thereof has: a heavy chain variable region CDR1 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-14; a heavy chain variable region CDR2 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 15-28; a heavy chain variable region CDR3 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 29-42; a light chain variable region CDR1 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 43-56; a light chain variable region CDR2 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 57-70; and a light chain variable region CDR3 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 71-84.

In one embodiment, the antibody comprised in the inventive composition comprises: a heavy chain variable region CDR1 of SEQ ID NO: 1; a heavy chain variable region CDR2 of SEQ ID NO: 15; a heavy chain variable region CDR3 of SEQ ID NO: 29; a light chain variable region CDR1 of SEQ ID NO: 43; a light chain variable region CDR2 of SEQ ID NO: 57; and a light chain variable region CDR3 of SEQ ID NO: 71.

In one embodiment, the antibody comprised in the inventive composition comprises: a heavy chain variable region CDR1 of SEQ ID NO: 2 a heavy chain variable region CDR2 of SEQ ID NO: 16; a heavy chain variable region CDR3 of SEQ ID NO: 30; a light chain variable region CDR1 of SEQ ID NO: 44; a light chain variable region CDR2 of SEQ ID NO: 58; and a light chain variable region CDR3 of SEQ ID NO: 72.

In one embodiment, the antibody comprised in the inventive composition comprises: a heavy chain variable region CDR1 of SEQ ID NO: 3; a heavy chain variable region CDR2 of SEQ ID NO: 17; a heavy chain variable region CDR3 of SEQ ID NO: 31; a light chain

variable region CDR1 of SEQ ID NO: 45; a light chain variable region CDR2 of SEQ ID NO: 59; and a light chain variable region CDR3 of SEQ ID NO: 73.

In one embodiment, the antibody comprised in the inventive composition comprises: a heavy chain variable region CDR1 of SEQ ID NO: 4; a heavy chain variable region CDR2 of SEQ ID NO: 18; a heavy chain variable region CDR3 of SEQ ID NO: 32; a light chain variable region CDR1 of SEQ ID NO: 46; a light chain variable region CDR2 of SEQ ID NO: 60; and a light chain variable region CDR3 of SEQ ID NO: 74.

In one embodiment, the antibody comprised in the inventive composition comprises: a heavy chain variable region CDR1 of SEQ ID NO: 5; a heavy chain variable region CDR2 of SEQ ID NO: 19; a heavy chain variable region CDR3 of SEQ ID NO: 33; a light chain variable region CDR1 of SEQ ID NO: 47; a light chain variable region CDR2 of SEQ ID NO: 61; and a light chain variable region CDR3 of SEQ ID NO: 75.

In one embodiment, the antibody comprised in the inventive composition comprises: a heavy chain variable region CDR1 of SEQ ID NO: 6; a heavy chain variable region CDR2 of SEQ ID NO: 20; a heavy chain variable region CDR3 of SEQ ID NO: 34; a light chain variable region CDR1 of SEQ ID NO: 48; a light chain variable region CDR2 of SEQ ID NO: 62; and a light chain variable region CDR3 of SEQ ID NO: 76.

In one embodiment, the antibody comprised in the inventive composition comprises: a heavy chain variable region CDR1 of SEQ ID NO: 7; a heavy chain variable region CDR2 of SEQ ID NO: 21; a heavy chain variable region CDR3 of SEQ ID NO: 35; a light chain variable region CDR1 of SEQ ID NO: 49; a light chain variable region CDR2 of SEQ ID NO: 63; and a light chain variable region CDR3 of SEQ ID NO: 77.

In one embodiment, the antibody comprised in the inventive composition comprises: a heavy chain variable region CDR1 of SEQ ID NO: 8; a heavy chain variable region CDR2 of SEQ ID NO: 22; a heavy chain variable region CDR3 of SEQ ID NO: 36; a light chain variable region CDR1 of SEQ ID NO: 50 a light chain variable region CDR2 of SEQ ID NO: 64; and a light chain variable region CDR3 of SEQ ID NO: 78.

In one embodiment, the antibody comprised in the inventive composition comprises: a heavy chain variable region CDR1 of SEQ ID NO: 9; a heavy chain variable region CDR2 of SEQ ID NO: 23; a heavy chain variable region CDR3 of SEQ ID NO: 37; a light chain variable region CDR1 of SEQ ID NO: 51; a light chain variable region CDR2 of SEQ ID NO: 65; and a light chain variable region CDR3 of SEQ ID NO: 79.

In one embodiment, the antibody comprised in the inventive composition comprises: a heavy chain variable region CDR1 of SEQ ID NO: 10; a heavy chain variable region CDR2 of SEQ ID NO: 24; a heavy chain variable region CDR3 of SEQ ID NO: 38; a light chain variable region CDR1 of SEQ ID NO: 52; a light chain variable region CDR2 of SEQ ID NO: 66; and a light chain variable region CDR3 of SEQ ID NO: 80.

In one embodiment, the antibody comprised in the inventive composition comprises: a heavy chain variable region CDR1 of SEQ ID NO: 11; a heavy chain variable region CDR2 of SEQ ID NO: 25; a heavy chain variable region CDR3 of SEQ ID NO: 39; a light chain variable region CDR1 of SEQ ID NO: 53; a light chain variable region CDR2 of SEQ ID NO: 67; and a light chain variable region CDR3 of SEQ ID NO: 81.

In one embodiment, the antibody comprised in the inventive composition comprises: a heavy chain variable region CDR1 of SEQ ID NO: 12; a heavy chain variable region CDR2 of SEQ ID NO: 26; a heavy chain variable region CDR3 of SEQ ID NO: 40; a light chain variable region CDR1 of SEQ ID NO: 54; a light chain variable region CDR2 of SEQ ID NO: 68; and a light chain variable region CDR3 of SEQ ID NO: 82.

In one embodiment, the antibody comprised in the inventive composition comprises: a heavy chain variable region CDR1 of SEQ ID NO: 13; a heavy chain variable region CDR2 of SEQ ID NO: 27; a heavy chain variable region CDR3 of SEQ ID NO: 41; a light chain variable region CDR1 of SEQ ID NO: 55; a light chain variable region CDR2 of SEQ ID NO: 69; and a light chain variable region CDR3 of SEQ ID NO: 83.

In one embodiment, the antibody comprised in the inventive composition comprises: a heavy chain variable region CDR1 of SEQ ID NO: 14; a heavy chain variable region CDR2 of SEQ ID NO: 28; a heavy chain variable region CDR3 of SEQ ID NO: 42; a light chain variable region CDR1 of SEQ ID NO: 56; a light chain variable region CDR2 of SEQ ID NO: 70; and a light chain variable region CDR3 of SEQ ID NO: 84.

In one embodiment, the disclosure provides a composition comprising an antibody comprising: (a) the variable heavy chain sequence of SEQ ID NO: 85 and variable light chain sequence of SEQ ID NO: 99; (b) the variable heavy chain sequence of SEQ ID NO: 86 and variable light chain sequence of SEQ ID NO: 100; (c) the variable heavy chain sequence of SEQ ID NO: 87 and variable light chain sequence of SEQ ID NO: 101; (d) the variable heavy chain sequence of SEQ ID NO: 88 and variable light chain sequence of SEQ ID NO: 102; (e) the variable heavy chain sequence of SEQ ID NO: 89 and variable light chain sequence of SEQ ID NO: 103; (f) the variable heavy chain

sequence of SEQ ID NO: 90 and variable light chain sequence of SEQ ID NO: 104; (g) the variable heavy chain sequence of SEQ ID NO: 91 and variable light chain sequence of SEQ ID NO: 105; (h) the variable heavy chain sequence of SEQ ID NO: 92 and variable light chain sequence of SEQ ID NO: 106; (i) the variable heavy chain sequence of SEQ ID NO: 93 and variable light chain sequence of SEQ ID NO: 107; (j) the variable heavy chain sequence of SEQ ID NO: 94 and variable light chain sequence of SEQ ID NO: 108; (k) the variable heavy chain sequence of SEQ ID NO: 95 and variable light chain sequence of SEQ ID NO: 109; (l) the variable heavy chain sequence of SEQ ID NO: 96 and variable light chain sequence of SEQ ID NO: 110; (m) the variable heavy chain sequence of SEQ ID NO: 97 and variable light chain sequence of SEQ ID NO: 111; or (n) the variable heavy chain sequence of SEQ ID NO: 98 and variable light chain sequence of SEQ ID NO: 112.

In one embodiment, the disclosure provides a composition comprising an antibody comprising: (a) the heavy chain sequence of SEQ ID NO: 146 and light chain sequence of SEQ ID NO: 141; (b) the heavy chain sequence of SEQ ID NO: 147 and light chain sequence of SEQ ID NO: 142; (c) the heavy chain sequence of SEQ ID NO: 148 and light chain sequence of SEQ ID NO: 143; (d) the heavy chain sequence of SEQ ID NO: 149 and light chain sequence of SEQ ID NO: 144; (e) the heavy chain sequence of SEQ ID NO: 150 and light chain sequence of SEQ ID NO: 145; (f) the heavy chain sequence of SEQ ID NO: 156 and light chain sequence of SEQ ID NO: 151; (g) the heavy chain sequence of SEQ ID NO: 157 and light chain sequence of SEQ ID NO: 152; (h) the heavy chain sequence of SEQ ID NO: 158 and light chain sequence of SEQ ID NO: 153; (i) the heavy chain sequence of SEQ ID NO: 159 and light chain sequence of SEQ ID NO: 154; or (j) the heavy chain sequence of SEQ ID NO: 160 and light chain sequence of SEQ ID NO: 155.

As used herein, a human antibody comprises heavy or light chain variable regions or full length heavy or light chains that are "the product of" or "derived from" a particular germline sequence if the variable regions or full length chains of the antibody are obtained from a system that uses human germline immunoglobulin genes. Such systems include immunizing a transgenic mouse carrying human immunoglobulin genes with the antigen of interest or screening a human immunoglobulin gene library displayed on phage with the antigen of interest. A human antibody that is "the product of" or "derived from" a human germline immunoglobulin sequence can be identified as such by comparing the amino acid sequence of the human antibody to the amino acid sequences of human germline immunoglobulins and selecting the human germline immunoglobulin sequence that is closest in sequence (*i.e.* greatest % identity) to the sequence of the

human antibody. A human antibody that is "the product of" or "derived from" a particular human germline immunoglobulin sequence may contain amino acid differences as compared to the germline sequence, due to, for example, naturally occurring somatic mutations or intentional introduction of site-directed mutation. However, a selected human antibody typically is at least 90% identical in amino acids sequence to an amino acid sequence encoded by a human germline immunoglobulin gene and contains amino acid residues that identify the human antibody as being human when compared to the germline immunoglobulin amino acid sequences of other species (*e.g.* murine germline sequences). In certain cases, a human antibody may be at least 80%, 90%, or at least 95%, or even at least 96%, 97%, 98%, or 99% identical in amino acid sequence to the amino acid sequence encoded by the germline immunoglobulin gene. Typically, a human antibody derived from a particular human germline sequence will display no more than 10 amino acid differences from the amino acid sequence encoded by the human germline immunoglobulin gene. In certain cases, the human antibody may display no more than 5, or even no more than 4, 3, 2, or 1 amino acid difference from the amino acid sequence encoded by the germline immunoglobulin gene.

In one embodiment the antibody comprised in the compositions of the disclosure is that encoded by pBW522 or pBW524 (deposited at DSMZ, Inhoffenstr. 7B, D-38124 Braunschweig, Germany on 18 August 2009 under deposit numbers DSM22873 and DSM22874, respectively).

Homologous antibodies

In yet another embodiment, an antibody comprised in the inventive composition has full length heavy and light chain amino acid sequences; full length heavy and light chain nucleotide sequences, variable region heavy and light chain nucleotide sequences, or variable region heavy and light chain amino acid sequences that are homologous to the amino acid and nucleotide sequences of the antibodies described herein, and wherein the antibodies retain the desired functional properties of the anti-ActRIIB antibodies of the disclosure.

For example, the disclosure provides a composition comprising an isolated recombinant anti-ActRIIB antibody (or a functional protein comprising an antigen binding portion thereof) comprising a heavy chain variable region and a light chain variable region, wherein: the heavy chain variable region comprises an amino acid sequence that is at least 80%, or at least 90% (preferably at least 95, 97 or 99%) identical to an amino acid sequence selected from the group consisting of SEQ ID NOs: 99-112; the light chain variable region comprises an amino acid sequence that is at least 80%, or at least 90%

(preferably at least 95, 97 or 99%) identical to an amino acid sequence selected from the group consisting of SEQ ID NOs: 85-98; alternatively the compositions comprises a recombinant anti-ActRIIB antibody (or a functional protein comprising an antigen binding portion thereof) comprising a heavy chain variable region and a light chain variable region, wherein: the heavy chain variable region comprises no more than 5 amino acid, or no more than 4 amino acid, or no more than 3 amino acid, or no more than 2 or no more than 1 amino acid change compared to the amino acid sequence selected from the group consisting of SEQ ID NOs: 99-112; the light chain variable region comprises no more than 5 amino acid, or no more than 4 amino acid, or no more than 3 amino acid, or no more than 2 or no more than 1 amino acid change compared to the amino acid sequence selected from the group consisting of SEQ ID NOs: 85-98 and the antibody exhibits at least one of the following functional properties: (i) it inhibits myostatin binding *in vitro* or *in vivo,* (ii) decreases inhibition of muscle differentiation through the Smad-dependent pathway and/or (iii) does not induce hematological changes, in particular no changes in RBC. In this context, the term “change” refers to insertions, deletions and/or substitutions.

In a further example, the disclosure provides a composition comprising an isolated recombinant anti-ActRII antibody, (or a functional protein comprising an antigen binding portion thereof) comprising a full length heavy chain and a full length light chain, wherein: the full length heavy chain comprises an amino acid sequence that is at least 80%, or at least 90% (preferably at least 95, 97 or 99%) identical to an amino acid sequence selected from the group consisting of SEQ ID NOs: 146-150 and 156-160; the full length light chain comprises an amino acid sequence that is at least 80%, or at least 90% (preferably at least 95, 97 or 99%) identical to an amino acid sequence selected from the group consisting of SEQ ID NOs: 141-145 and 151-155; alternatively the compositions comprises a recombinant anti-ActRII antibody (or a functional protein comprising an antigen binding portion thereof) comprising a heavy chain variable region and a light chain variable region, wherein: the heavy chain variable region comprises no more than 5 amino acid, or no more than 4 amino acid, or no more than 3 amino acid, or no more than 2 or no more than 1 amino acid change compared to the amino acid sequence selected from the group consisting of SEQ ID NOs: 146-150 and 156-160; the light chain variable region comprises no more than 5 amino acid, or no more than 4 amino acid, or no more than 3 amino acid, or no more than 2 or no more than 1 amino acid change compared to the amino acid sequence selected from the group consisting of SEQ ID NOs: 141-145 and 151-155 and the antibody exhibits at least one of the following functional properties: (i) it inhibits myostatin binding *in vitro* or *in vivo,* (ii) decreases inhibition of muscle differentiation through the Smad-dependent pathway and/or (iii) does

not induce hematological changes, in particular no changes in RBC. Preferably such an antibody binds to the ligand binding domain of ActRIIB and/or ActRIIA. In this context, the term “change” refers to insertions, deletions and/or substitutions.

In another example, the disclosure provides a composition comprising an isolated recombinant anti-ActRII antibody (or a functional protein comprising an antigen binding portion thereof), comprising a full length heavy chain and a full length light chain, wherein: the full length heavy chain is encoded by a nucleotide sequence that is at least 80%, or at least 90% (preferably at least 95, 97 or 99%) identical to a nucleotide sequence selected from the group consisting of SEQ ID NOs: 166-170 and 176-180; the full length light chain is encoded by a nucleotide sequence that is at least 80%, or at least 90% (preferably at least 95, 97 or 99%) identical to a nucleotide sequence selected from the group consisting of SEQ ID NOs: 161-165 and 171-175; alternatively the compositions comprises a recombinant anti-ActRIIB antibody (or a functional protein comprising an antigen binding portion thereof) comprising a heavy chain variable region and a light chain variable region, wherein: the heavy chain variable region comprises no more than 5 amino acid, or no more than 4 amino acid, or no more than 3 amino acid, or no more than 2 or no more than 1 amino acid change compared to the amino acid sequence selected from the group consisting of SEQ ID NOs: 166-170 and 176-180; the light chain variable region comprises no more than 5 amino acid, or no more than 4 amino acid, or no more than 3 amino acid, or no more than 2 or no more than 1 amino acid change compared to the amino acid sequence selected from the group consisting of SEQ ID NOs: 161-165 and 171-175 and the antibody exhibits at least one of the following functional properties: (i) it inhibits myostatin binding *in vitro* or *in vivo,* (ii) decreases inhibition of muscle differentiation through the Smad-dependent pathway and/or (iii) does not induce hematological changes, in particular no changes in RBC. Preferably such an antibody binds to the ligand binding domain of ActRIIB. In this context, the term “change” refers to insertions, deletions and/or substitutions.

In various embodiments, the antibody comprised in the inventive composition may exhibit one or more, two or more, or three of the functional properties discussed above. The antibody can be, for example, a human antibody, a humanized antibody or a chimeric antibody. Preferably the antibody is a fully human IgG1 antibody.

In other embodiments, the V_H and/or V_L amino acid sequences may be at least 80%, 90%, 95%, 96%, 97%, 98% or 99% identical to the sequences set forth above. In other embodiments, the V_H and/or V_L amino acid sequences may be identical except an amino acid substitution in no more than 1, 2, 3, 4 or 5 amino acid position. An antibody having

V_H and V_L regions having high (*i.e.* 80% or greater) identity to the V_H and V_L regions of SEQ ID NOs 99-112 and SEQ ID NOs: 85-98 respectively, can be obtained by mutagenesis (*e.g.* site-directed or PCR-mediated mutagenesis) of nucleic acid molecules SEQ ID NOs: 127-140 and 113-126 respectively, followed by testing of the encoded altered antibody for retained function (*i.e.* the functions set forth above) using the functional assays described herein.

In other embodiments, the full length heavy chain and/or full length light chain amino acid sequences may be at least 80%, 90%, 95%, 96%, 97%, 98% or 99% identical to the sequences set forth above or may be identical except an amino acid change in no more than 1, 2, 3, 4 or 5 amino acid position. An antibody having a full length heavy chain and full length light chain having high (*i.e.* at least 80% or greater) identity to the full length heavy chains of any of SEQ ID NOs: 146-150 and 156-160 and full length light chains of any of SEQ ID NOs: 141-145 and 151-155 respectively, can be obtained by mutagenesis (*e.g.* site-directed or PCR-mediated mutagenesis) of nucleic acid molecules SEQ ID NOs: 166-170 and 176-180 and SEQ ID NOs: 161-165 and 171-175 respectively, followed by testing of the encoded altered antibody for retained function (*i.e.* the functions set forth above) using the functional assays described herein.

In other embodiments, the full length heavy chain and/or full length light chain nucleotide sequences may be at least 80%, 90%, 95%, 96%, 97%, 98% or 99% identical to the sequences set forth above.

In other embodiments, the variable regions of heavy chain and/or light chain nucleotide sequences may be at least 80%, 90%, 95%, 96%, 97%, 98% or 99% identical to the sequences set forth above or may be identical except an amino acid change in no more than 1, 2, 3, 4 or 5 amino acid position.

As used herein, the percent identity between the two sequences is a function of the number of identical positions shared by the sequences (*i.e.* % identity = # of identical positions/total # of positions x 100), taking into account the number of gaps, and the length of each gap, which need to be introduced for optimal alignment of the two sequences. The comparison of sequences and determination of percent identity between two sequences can be accomplished using a mathematical algorithm, as described below.

The percent identity between two amino acid sequences can be determined using the algorithm of E. Meyers and W. Miller (Comput. Appl. Biosci., 4:11-17, 1988) which has

been incorporated into the ALIGN program (version 2.0), using a PAM120 weight residue table, a gap length penalty of 12 and a gap penalty of 4. In addition, the percent identity between two amino acid sequences can be determined using the Needleman and Wunsch (J. Mol, Biol. 48:444-453, 1970) algorithm which has been incorporated into the GAP program in the GCG software package (available at http://www.gcg.com), using either a Blossom 62 matrix or a PAM250 matrix, and a gap weight of 16, 14, 12, 10, 8, 6, or 4 and a length weight of 1, 2, 3, 4, 5, or 6.

Antibodies with conservative modifications

In certain embodiments, an antibody comprised in the inventive composition has a heavy chain variable region comprising CDR1, CDR2, and CDR3 sequences and a light chain variable region comprising CDR1, CDR2, and CDR3 sequences, wherein one or more of these CDR sequences have specified amino acid sequences based on the antibodies described herein or variant sequences thereof comprising 1, 2, 3, 4 or 5 amino acid changes or conservative modifications thereof, and wherein the antibodies retain the desired functional properties of the anti-ActRIIB antibodies of the disclosure. Accordingly, the disclosure provides compositions comprising an isolated recombinant anti-ActRIIB antibody, or a functional protein comprising an antigen binding portion thereof, consisting of a heavy chain variable region comprising CDR1, CDR2, and CDR3 sequences and a light chain variable region comprising CDR1, CDR2, and CDR3 sequences, wherein: the heavy chain variable region CDR1 amino acid sequences are selected from the group consisting of SEQ ID NOs: 1-14 or variant sequences thereof comprising 1, 2, 3, 4 or 5 amino acid changes, and conservative modifications thereof; the heavy chain variable region CDR2 amino acid sequences are selected from the group consisting of SEQ ID NOs: 15-28 or variant sequences thereof comprising 1, 2, 3, 4 or 5 amino acid changes, and conservative modifications thereof; the heavy chain variable region CDR3 amino acid sequences are selected from the group consisting of SEQ ID NOs: 29-42 or variant sequences thereof comprising 1, 2, 3, 4 or 5 amino acid changes, and conservative modifications thereof; the light chain variable regions CDR1 amino acid sequences are selected from the group consisting of SEQ ID NOs: 43-56 or variant sequences thereof comprising 1, 2, 3, 4 or 5 amino acid changes, and conservative modifications thereof; the light chain variable regions CDR2 amino acid sequences are selected from the group consisting of SEQ ID NOs: 57-70 or variant sequences thereof comprising 1, 2, 3, 4 or 5 amino acid changes, and conservative modifications thereof; the light chain variable regions of CDR3 amino acid sequences are selected from the group consisting of SEQ ID NOs: 71-84 or variant sequences thereof comprising 1, 2, 3, 4 or 5 amino acid changes, and conservative modifications thereof. Preferably the antibody exhibits at least one of the following functional properties: (i) it inhibits myostatin binding *in vitro* or *in vivo,*

(ii) decreases inhibition of muscle differentiation through the Smad-dependent pathway and/or (iii) does not induce hematological changes, in particular no changes in RBC.

In various embodiments, the antibody may exhibit one or both of the functional properties listed above. Such antibodies can be, for example, human antibodies, humanized antibodies or chimeric antibodies.

In other embodiments, an antibody comprised in the inventive composition optimized for expression in a mammalian cell has a full length heavy chain sequence and a full length light chain sequence, wherein one or more of these sequences have specified amino acid sequences based on the antibodies described herein or conservative modifications thereof, and wherein the antibodies retain the desired functional properties of the anti-ActRIIB antibodies of the disclosure. Accordingly, the disclosure provides compositions comprising an isolated monoclonal anti-ActRII antibody optimized for expression in a mammalian cell consisting of a full length heavy chain and a full length light chain wherein: the full length heavy chain has amino acid sequences selected from the group of SEQ ID NOs: 146-150 and 156-160 or variant sequences thereof comprising 1, 2, 3, 4 or 5 amino acid changes, and conservative modifications thereof; and the full length light chain has amino acid sequences selected from the group of SEQ ID NOs: 141-145 and 151-155 or variant sequences thereof comprising 1, 2, 3, 4 or 5 amino acid changes, and conservative modifications thereof; and the antibody exhibits at least one of the following functional properties: (i) it inhibits myostatin binding *in vitro* or *in vivo,* (ii) decreases inhibition of muscle differentiation through the Smad-dependent pathway and/or (iii) does not induce hematological changes, in particular no changes in RBC.

In various embodiments, the antibody may exhibit one or both of the functional properties listed above. Such antibodies can be, for example, human antibodies, humanized antibodies or chimeric antibodies.

As used herein, the term "conservative sequence modifications" is intended to refer to amino acid modifications that do not significantly affect or alter the binding characteristics of the antibody containing the amino acid sequence. Such conservative modifications include amino acid substitutions, additions and deletions. Modifications can be introduced into an antibody of the disclosure by standard techniques known in the art, such as site-directed mutagenesis and PCR-mediated mutagenesis.

Conservative amino acid substitutions are ones in which the amino acid residue is replaced with an amino acid residue having a similar side chain. Families of amino acid

residues having similar side chains have been defined in the art. These families include amino acids with basic side chains (*e.g.* lysine, arginine, histidine), acidic side chains (*e.g.* aspartic acid, glutamic acid), uncharged polar side chains (*e.g.* glycine, asparagine, glutamine, serine, threonine, tyrosine, cysteine, tryptophan), nonpolar side chains (*e.g.* alanine, valine, leucine, isoleucine, proline, phenylalanine, methionine), beta-branched side chains (*e.g.* threonine, valine, isoleucine) and aromatic side chains (*e.g.* tyrosine, phenylalanine, tryptophan, histidine). Thus, one or more amino acid residues within the CDR regions of an antibody of the disclosure can be replaced with other amino acid residues from the same side chain family, and the altered antibody can be tested for retained function using the functional assays described herein.

Antibodies that bind to the same epitope as anti-ActRII antibodies comprised in the disclosed composition

In another embodiment, the disclosure provides compositions comprising antibodies that bind to the same epitope as the various specific anti-ActRII antibodies described herein. All the antibodies described in the examples that are capable of blocking myostatin binding to ActRIIA and ActRIIB bind to one of the epitopes in ActRIIA and ActRIIB with high affinity, said epitope being comprised between amino acids 19-134 of SEQ ID NO:181.

Additional antibodies can therefore be identified based on their ability to cross-compete (*e.g.* to competitively inhibit the binding of, in a statistically significant manner) with other antibodies of the disclosure in standard ActRIIB binding assays. The ability of a test antibody to inhibit the binding of antibodies comprised in the inventive compositions to human ActRIIB demonstrates that the test antibody can compete with said antibody for binding to human ActRIIB; such an antibody may, according to non-limiting theory, bind to the same or a related (*e.g.* a structurally similar or spatially proximal) epitope on human ActRIIB as the antibody with which it competes. In a certain embodiment, the antibody that binds to the same epitope on human ActRIIA and ActRIIA as the antibodies comprised in the inventive compositions is a human recombinant antibody. Such human recombinant antibodies can be prepared and isolated as described in the examples.

Thus, the disclosure provides a composition comprising an antibody that binds to an epitope recognised by and/or that competes for binding with an antibody having the variable heavy chain sequence recited in SEQ ID NO: 85, and the variable light chain sequence recited in SEQ ID NO: 99.

Thus, the disclosure provides a composition comprising an antibody that binds to an epitope recognised by an antibody having the variable heavy chain sequence recited in SEQ ID NO: 86, and the variable light chain sequence recited in SEQ ID NO: 100.

Thus, the disclosure provides a composition comprising an antibody that binds to an epitope recognised by an antibody having the variable heavy chain sequence recited in SEQ ID NO: 87, and the variable light chain sequence recited in SEQ ID NO: 101.

Thus, the disclosure provides a composition comprising an antibody that binds to an epitope recognised by an antibody having the variable heavy chain sequence recited in SEQ ID NO: 88, and the variable light chain sequence recited in SEQ ID NO: 102.

Thus, the disclosure provides a composition comprising an antibody that binds to an epitope recognised by an antibody having the variable heavy chain sequence recited in SEQ ID NO: 89, and the variable light chain sequence recited in SEQ ID NO: 103.

Thus, the disclosure provides a composition comprising an antibody that binds to an epitope recognised by an antibody having the variable heavy chain sequence recited in SEQ ID NO: 90, and the variable light chain sequence recited in SEQ ID NO: 104.

Thus, the disclosure provides a composition comprising an antibody that binds to an epitope recognised by an antibody having the variable heavy chain sequence recited in SEQ ID NO: 91, and the variable light chain sequence recited in SEQ ID NO: 105.

Thus, the disclosure provides a composition comprising an antibody that binds to an epitope recognised by an antibody having the variable heavy chain sequence recited in SEQ ID NO: 92, and the variable light chain sequence recited in SEQ ID NO: 106.

Thus, the disclosure provides a composition comprising an antibody that binds to an epitope recognised by an antibody having the variable heavy chain sequence recited in SEQ ID NO: 93, and the variable light chain sequence recited in SEQ ID NO: 107.

Thus, the disclosure provides a composition comprising an antibody that binds to an epitope recognised by an antibody having the variable heavy chain sequence recited in SEQ ID NO: 94, and the variable light chain sequence recited in SEQ ID NO: 108.

Thus, the disclosure provides a composition comprising an antibody that binds to an epitope recognised by an antibody having the variable heavy chain sequence recited in SEQ ID NO: 95, and the variable light chain sequence recited in SEQ ID NO: 109.

Thus, the disclosure provides a composition comprising an antibody that binds to an epitope recognised by an antibody having the variable heavy chain sequence recited in SEQ ID NO: 96, and the variable light chain sequence recited in SEQ ID NO: 110.

Thus, the disclosure provides a composition comprising an antibody that binds to an epitope recognised by an antibody having the variable heavy chain sequence recited in SEQ ID NO: 97, and the variable light chain sequence recited in SEQ ID NO: 111.

Thus, the disclosure provides a composition comprising an antibody that binds to an epitope recognised by an antibody having the variable heavy chain sequence recited in SEQ ID NO: 98, and the variable light chain sequence recited in SEQ ID NO: 112.

Following more detailed epitope mapping experiments, the binding regions of preferred antibodies of the inventive compositions have been more clearly defined.

Thus, the disclosure provides a composition comprising an antibody that binds to an epitope comprising amino acids 78-83 of SEQ ID NO: 181 (WLDDFN – SEQ ID NO:188). The disclosure also provides a composition comprising an antibody that binds to an epitope comprising amino acids 76-84 of SEQ ID NO: 181 (GCWLDDFNC – SEQ ID NO:186).

The disclosure also provides a composition comprising an antibody that binds to an epitope comprising amino acids 75-85 of SEQ ID NO: 181 (KGCWLDDFNCY – SEQ ID NO:190).

The disclosure also provides a composition comprising an antibody that binds to an epitope comprising amino acids 52-56 of SEQ ID NO: 181 (EQDKR – SEQ ID NO:189). The disclosure also provides a composition comprising an antibody that binds to an epitope comprising amino acids 49-63 of SEQ ID NO: 181 (CEGEQDKRLHCYASW – SEQ ID NO:187).

The disclosure also provides a composition comprising an antibody that binds to an epitope comprising or consisting of amino acids 29-41 of SEQ ID NO: 181 (CIYYNANWELERT– SEQ ID NO:191).

The disclosure also provides a composition comprising an antibody that binds to an epitope comprising or consisting of amino acids 100-110 of SEQ ID NO: 181 (YFCCCEGNFCN – SEQ ID NO:192);

The disclosure also provides a composition comprising an antibody that binds to an epitope comprising or consisting of amino acids 78-83 of SEQ ID NO: 181 (WLDDFN) and amino acids 52-56 of SEQ ID NO: 181 (EQDKR).

The disclosure also provides a composition comprising antibodies that bind to epitopes consisting of these sequences or epitopes comprising combinations of these epitope regions.

Thus, the disclosure also provides a composition comprising an antibody that binds to an epitope comprising or consisting of amino acids 78-83 of SEQ ID NO: 181 (WLDDFN) and amino acids 52-56 of SEQ ID NO: 181 (EQDKR).

Engineered and modified antibodies

An antibody comprised in the inventive compositions further can be prepared using an antibody having one or more of the V_H and/or V_L sequences shown herein as starting material to engineer a modified antibody, which modified antibody may have altered properties from the starting antibody. An antibody can be engineered by modifying one or more residues within one or both variable regions (*i.e.* V_H and/or V_L), for example within one or more CDR regions and/or within one or more framework regions. Additionally or alternatively, an antibody can be engineered by modifying residues within the constant region(s), for example to alter the effector function(s) of the antibody.

One type of variable region engineering that can be performed is CDR grafting. Antibodies interact with target antigens predominantly through amino acid residues that are located in the six heavy and light chain complementarity determining regions (CDRs). For this reason, the amino acid sequences within CDRs are more diverse between individual antibodies than sequences outside of CDRs. Because CDR sequences are responsible for most antibody-antigen interactions, it is possible to express recombinant antibodies that mimic the properties of specific naturally occurring antibodies by constructing expression vectors that include CDR sequences from the specific naturally occurring antibody grafted onto framework sequences from a different antibody with different properties (see, *e.g.* Riechmann, L. *et al.*, 1998 Nature 332:323-327; Jones, P. *et al.*, 1986 Nature 321:522-525; Queen, C. *et al.*, 1989 Proc. Natl. Acad. Sci. U.S.A. 86:10029-10033; U.S. Patent No. 5,225,539 to Winter, and U.S. Patent Nos. 5,530,101; 5,585,089; 5,693,762 and 6,180,370 to Queen *et al.*).

Accordingly, another embodiment of the disclosure pertains to compositions comprising a monoclonal anti- ActRII antibody, or a functional protein comprising an antigen binding portion thereof, comprising a heavy chain variable region comprising CDR1 sequences having an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-14; CDR2 sequences having an amino acid sequence selected from the group consisting of SEQ ID NOs: 15-28; CDR3 sequences having an amino acid sequence selected from the group consisting of SEQ ID NOs: 29-42, respectively; and a light chain variable region having CDR1 sequences having an amino acid sequence selected from the group consisting of SEQ ID NOs: 43-56; CDR2 sequences having an amino acid sequence selected from the group consisting of SEQ ID NOs: 57-70; and CDR3 sequences consisting of an amino acid sequence selected from the group consisting of SEQ ID NOs: 71-84, respectively. Thus, such antibodies contain the V_H and V_L CDR sequences of monoclonal antibodies, yet may contain different framework sequences from these antibodies.

Such framework sequences can be obtained from public DNA databases or published references that include germline antibody gene sequences. For example, germline DNA sequences for human heavy and light chain variable region genes can be found in the "VBase" human germline sequence database (available on the Internet at www.mrc-cpe.cam.ac.uk/vbase), as well as in Kabat, E. A., *et al.*, [*supra*]; Tomlinson, I. M., *et al.*, 1992 J. fol. Biol. 227:776-798; and Cox, J. P. L. *et al.*, 1994 Eur. J Immunol. 24:827-836. An example of framework sequences for use in the antibodies of the disclosure are those that are structurally similar to the framework sequences used by selected antibodies of the disclosure, *e.g.* consensus sequences and/or framework sequences used by monoclonal antibodies of the disclosure. The V_H CDR1, 2 and 3 sequences, and the V_L CDR1, 2 and 3 sequences, can be grafted onto framework regions that have the identical sequence as that found in the germline immunoglobulin gene from which the framework sequence derive, or the CDR sequences can be grafted onto framework regions that contain one or more mutations as compared to the germline sequences. For example, it has been found that in certain instances it is beneficial to mutate residues within the framework regions to maintain or enhance the antigen binding ability of the antibody (see *e.g.* U.S. Patents. 5,530,101; 5,585,089; 5,693,762 and 6,180,370 to Queen *et al*).

Another type of variable region modification is to mutate amino acid residues within the V_H and/or V_L CDR1, CDR2 and/or CDR3 regions to thereby improve one or more binding properties (*e.g.* affinity) of the antibody of interest, known as "affinity maturation." Site-directed mutagenesis or PCR-mediated mutagenesis can be performed to introduce the mutation(s) and the effect on antibody binding, or other functional property of interest,

can be evaluated in *in vitro* or *in vivo* assays as described herein and provided in the Examples. Conservative modifications (as discussed above) can be introduced. The mutations may be amino acid substitutions, additions or deletions. Moreover, typically no more than one, two, three, four or five residues within a CDR region are altered.

Accordingly, in another embodiment, the disclosure provides isolated anti-ActRII monoclonal antibodies, or a functional protein comprising an antigen binding portion thereof, consisting of a heavy chain variable region having: a V_H CDR1 region consisting of an amino acid sequence selected from the group having SEQ ID NOs: 1-14 or an amino acid sequence having one, two, three, four or five amino acid substitutions, deletions or additions as compared to SEQ ID NOs: 1-14; a V_H CDR2 region having an amino acid sequence selected from the group consisting of SEQ ID NOs: 15-28, or an amino acid sequence having one, two, three, four or five amino acid substitutions, deletions or additions as compared to SEQ ID NOs: 15-28; a V_H CDR3 region having an amino acid sequence selected from the group consisting of SEQ ID NOs: 29-42, or an amino acid sequence having one, two, three, four or five amino acid substitutions, deletions or additions as compared to SEQ ID NOs: 29-42; a V_L CDR1 region having an amino acid sequence selected from the group consisting of SEQ ID NOs: 43-56, or an amino acid sequence having one, two, three, four or five amino acid substitutions, deletions or additions as compared to SEQ ID NOs: 43-56; a V_L CDR2 region having an amino acid sequence selected from the group consisting of SEQ ID NOs: 52-70, or an amino acid sequence having one, two, three, four or five amino acid substitutions, deletions or additions as compared to SEQ ID NOs: 52-70; and a V_L CDR3 region having an amino acid sequence selected from the group consisting of SEQ ID NOs: 71-84, or an amino acid sequence having one, two, three, four or five amino acid substitutions, deletions or additions as compared to SEQ ID NOs: 71-84.

Camelid antibodies

Antibody proteins obtained from members of the camel and dromedary family (*Camelus bactrianus* and *Camelus dromaderius*) including new world members such as llama species (*Lama paccos*, *Lama glama and Lama vicugna)* have been characterized with respect to size, structural complexity and antigenicity for human subjects. Certain IgG antibodies from this family of mammals as found in nature lack light chains, and are thus structurally distinct from the typical four chain quaternary structure having two heavy and two light chains, for antibodies from other animals (see WO94/04678).

A region of the camelid antibody which is the small single variable domain identified as V_{HH} can be obtained by genetic engineering to yield a small protein having high affinity

for a target, resulting in a low molecular weight antibody-derived protein known as a "camelid nanobody" (see US5,759,808; Stijlemans, B. *et al.*, 2004 J Biol Chem 279: 1256-1261; Dumoulin, M. *et al.*, 2003 Nature 424: 783-788; Pleschberger, M. *et al.* 2003 Bioconjugate Chem 14: 440-448; Cortez-Retamozo, V. *et al.* 2002 Int J Cancer 89: 456-62; and Lauwereys, M. *et al.* 1998 EMBO J 17: 3512-3520). Engineered libraries of camelid antibodies and antibody fragments are commercially available, for example, from Ablynx, Ghent, Belgium. As with other antibodies of non-human origin, an amino acid sequence of a camelid antibody can be altered recombinantly to obtain a sequence that more closely resembles a human sequence, *i.e.* the nanobody can be "humanized". Thus the natural low antigenicity of camelid antibodies to humans can be further reduced.

The camelid nanobody has a molecular weight approximately one-tenth that of a human IgG molecule, and the protein has a physical diameter of only a few nanometers. One consequence of the small size is the ability of camelid nanobodies to bind to antigenic sites that are functionally invisible to larger antibody proteins, *i.e.* camelid nanobodies are useful as reagents detect antigens that are otherwise cryptic using classical immunological techniques, and as possible therapeutic agents. Thus yet another consequence of small size is that a camelid nanobody can inhibit as a result of binding to a specific site in a groove or narrow cleft of a target protein, and hence can serve in a capacity that more closely resembles the function of a classical low molecular weight drug than that of a classical antibody.

The low molecular weight and compact size further result in camelid nanobodies being extremely thermostable, stable to extreme pH and to proteolytic digestion, and poorly antigenic. Another consequence is that camelid nanobodies readily move from the circulatory system into tissues, and even cross the blood-brain barrier and can treat disorders that affect nervous tissue. Nanobodies can further facilitate drug transport across the blood brain barrier (see US2004/0161738). These features combined with the low antigenicity to humans indicate great therapeutic potential. Further, these molecules can be fully expressed in prokaryotic cells such as *E. coli* and are expressed as fusion proteins with bacteriophage and are functional.

Accordingly, in one embodiment, the present disclosure related to composition comprising a camelid antibody or nanobody having high affinity for ActRIIB. In certain embodiments herein, the camelid antibody or nanobody is naturally produced in the camelid animal, *i.e.* is produced by the camelid following immunization with ActRIIB or a peptide fragment thereof, using techniques described herein for other antibodies.

Alternatively, the anti-ActRIIB camelid nanobody is engineered, *i.e.* produced by selection for example from a library of phage displaying appropriately mutagenized camelid nanobody proteins using panning procedures with ActRIIB as a target as described in the examples herein. Engineered nanobodies can further be customized by genetic engineering to have a half life in a recipient subject of from 45 minutes to two weeks. In a specific embodiment, the camelid antibody or nanobody is obtained by grafting the CDRs sequences of the heavy or light chain of the human antibodies of the disclosure into nanobody or single domain antibody framework sequences, as described for example in WO94/04678.

Non-antibody scaffold

Known non-immunoglobulin frameworks or scaffolds include, but are not limited to, Adnectins (fibronectin) (Compound Therapeutics, Inc., Waltham, MA), ankyrin (Molecular Partners AG, Zurich, Switzerland), domain antibodies (Domantis, Ltd (Cambridge, MA) and Ablynx nv (Zwijnaarde, Belgium)), lipocalin (Anticalin) (Pieris Proteolab AG, Freising, Germany), small modular immuno-pharmaceuticals (Trubion Pharmaceuticals Inc., Seattle, WA), maxybodies (Avidia, Inc. (Mountain View, CA)), Protein A (Affibody AG, Sweden) and affilin (gamma-crystallin or ubiquitin) (Scil Proteins GmbH, Halle, Germany), protein epitope mimetics (Polyphor Ltd, Allschwil, Switzerland).

(i) Fibronectin scaffold

The fibronectin scaffolds are based preferably on fibronectin type III domain (*e.g.* the tenth module of the fibronectin type III (10 Fn3 domain)). The fibronectin type III domain has 7 or 8 beta strands which are distributed between two beta sheets, which themselves pack against each other to form the core of the protein, and further containing loops (analogous to CDRs) which connect the beta strands to each other and are solvent exposed. There are at least three such loops at each edge of the beta sheet sandwich, where the edge is the boundary of the protein perpendicular to the direction of the beta strands (US 6,818,418).

These fibronectin-based scaffolds are not an immunoglobulin, although the overall fold is closely related to that of the smallest functional antibody fragment, the variable region of the heavy chain, which comprises the entire antigen recognition unit in camel and llama IgG. Because of this structure, the non-immunoglobulin antibody mimics antigen binding properties that are similar in nature and affinity to those of antibodies. These scaffolds can be used in a loop randomization and shuffling strategy *in vitro* that is similar to the process of affinity maturation of antibodies *in vivo*. These fibronectin-based molecules

can be used as scaffolds where the loop regions of the molecule can be replaced with CDRs of the disclosure using standard cloning techniques.

(ii) Ankyrin – Molecular Partners

The technology is based on using proteins with ankyrin derived repeat modules as scaffolds for bearing variable regions which can be used for binding to different targets. The ankyrin repeat module is a 33 amino acid polypeptide consisting of two anti-parallel α-helices and a β-turn. Binding of the variable regions is mostly optimized by using ribosome display.

(iii) Maxybodies/Avimers - Avidia

Avimers are derived from natural A-domain containing protein such as LRP-1. These domains are used by nature for protein-protein interactions and in human over 250 proteins are structurally based on A-domains. Avimers consist of a number of different "A-domain" monomers (2-10) linked via amino acid linkers. Avimers can be created that can bind to the target antigen using the methodology described in, for example, US2004/0175756; US2005/0053973; US2005/0048512; and US2006/0008844.

(vi) Protein A – Affibody

Affibody® affinity ligands are small, simple proteins composed of a three-helix bundle based on the scaffold of one of the IgG-binding domains of Protein A. Protein A is a surface protein from the bacterium *Staphylococcus aureus*. This scaffold domain consists of 58 amino acids, 13 of which are randomized to generate Affibody® libraries with a large number of ligand variants (See *e.g.* US 5,831,012). Affibody® molecules mimic antibodies, they have a molecular weight of 6 kDa, compared to the molecular weight of antibodies, which is 150 kDa. In spite of its small size, the binding site of Affibody® molecules is similar to that of an antibody.

(v) Anticalins – Pieris

Anticalins® are products developed by the company Pieris ProteoLab AG. They are derived from lipocalins, a widespread group of small and robust proteins that are usually involved in the physiological transport or storage of chemically sensitive or insoluble compounds. Several natural lipocalins occur in human tissues or body liquids.

The protein architecture is reminiscent of immunoglobulins, with hypervariable loops on top of a rigid framework. However, in contrast with antibodies or their recombinant fragments, lipocalins are composed of a single polypeptide chain with 160 to 180 amino acid residues, being just marginally bigger than a single immunoglobulin domain.

The set of four loops, which makes up the binding pocket, shows pronounced structural plasticity and tolerates a variety of side chains. The binding site can thus be reshaped in a proprietary process in order to recognize prescribed target molecules of different shape with high affinity and specificity.

One protein of lipocalin family, the bilin-binding protein (BBP) of *Pieris brassicae* has been used to develop anticalins by mutagenizing the set of four loops. One example of a patent application describing "anticalins" is WO1999/16873.

(vi) Affilin – Scil Proteins

AFFILIN™ molecules are small non-immunoglobulin proteins which are designed for specific affinities towards proteins and small molecules. New AFFILIN™ molecules can be very quickly selected from two libraries, each of which is based on a different human derived scaffold protein.

AFFILIN™ molecules do not show any structural homology to immunoglobulin proteins. Scil Proteins employs two AFFILIN™ scaffolds, one of which is gamma crystalline, a human structural eye lens protein and the other is "ubiquitin" superfamily proteins. Both human scaffolds are very small, show high temperature stability and are almost resistant to pH changes and denaturing agents. This high stability is mainly due to the expanded beta sheet structure of the proteins. Examples of gamma crystalline derived proteins are described in WO2001/004144 and examples of "ubiquitin-like" proteins are described in WO2004/106368.

(vii) Protein Epitope Mimetics (PEM)

PEM are medium-sized, cyclic, peptide-like molecules (MW 1-2kDa) mimicking beta-hairpin secondary structures of proteins, the major secondary structure involved in protein-protein interactions.

Grafting antigen-binding domains into alternative frameworks or scaffolds

A wide variety of antibody/immunoglobulin frameworks or scaffolds can be employed so long as the resulting polypeptide includes at least one binding region which specifically binds to ActRIIB. Such frameworks or scaffolds include the 5 main idiotypes of human immunoglobulins, or fragments thereof (such as those disclosed elsewhere herein), and include immunoglobulins of other animal species, preferably having humanized aspects. Single heavy-chain antibodies such as those identified in camelids are of particular interest in this regard. Novel frameworks, scaffolds and fragments continue to be discovered and developed by those skilled in the art.

In one aspect, the compositions of the disclosure may comprise non-immunoglobulin based antibodies using non-immunoglobulin scaffolds onto which CDRs of the disclosed antibodies can be grafted. Known or future non-immunoglobulin frameworks and scaffolds may be employed, as long as they comprise a binding region specific for the target protein of SEQ ID NO: 181 (preferably, the ligand binding domain thereof as shown in SEQ ID NO: 182). Such compounds are known herein as “polypeptides comprising a target-specific binding region”. Examples of non-immunoglobulin framework are further described in the sections below (camelid antibodies and non-antibody scaffold).

Framework or Fc engineering

Engineered antibodies comprised in the compositions of the disclosure include those in which modifications have been made to framework residues within V_H and/or V_L, *e.g.* to improve the properties of the antibody. Typically such framework modifications are made to decrease the immunogenicity of the antibody. For example, one approach is to "backmutate" one or more framework residues to the corresponding germline sequence. More specifically, an antibody that has undergone somatic mutation may contain framework residues that differ from the germline sequence from which the antibody is derived. Such residues can be identified by comparing the antibody framework sequences to the germline sequences from which the antibody is derived. To return the framework region sequences to their germline configuration, the somatic mutations can be "backmutated" to the germline sequence by, for example, site-directed mutagenesis or PCR-mediated mutagenesis. Such "backmutated" antibodies can also be comprised in the compositions of the disclosure.

Another type of framework modification involves mutating one or more residues within the framework region, or even within one or more CDR regions, to remove T-cell epitopes to thereby reduce the potential immunogenicity of the antibody. This approach is also referred to as "deimmunization" and is described in further detail in US2003/0153043.

In addition or alternative to modifications made within the framework or CDR regions, antibodies of the disclosure may be engineered to include modifications within the Fc region, typically to alter one or more functional properties of the antibody, such as serum half-life, complement fixation, Fc receptor binding, and/or antigen-dependent cellular cytotoxicity. Furthermore, an antibody comprised in the compositions of the disclosure may be chemically modified (*e.g.* one or more chemical moieties can be attached to the

antibody) or be modified to alter its glycosylation, again to alter one or more functional properties of the antibody. Each of these embodiments is described in further detail below. The numbering of residues in the Fc region is that of the EU index of Kabat.

In one embodiment, the hinge region of CH1 is modified such that the number of cysteine residues in the hinge region is altered, *e.g.* increased or decreased. This approach is described further in US5,677,425. The number of cysteine residues in the hinge region of CH1 is altered to, for example, facilitate assembly of the light and heavy chains or to increase or decrease the stability of the antibody.

In another embodiment, the Fc hinge region of an antibody is mutated to decrease the biological half-life of the antibody. More specifically, one or more amino acid mutations are introduced into the CH2-CH3 domain interface region of the Fc-hinge fragment such that the antibody has impaired Staphylococcyl protein A (SpA) binding relative to native Fc-hinge domain SpA binding. This approach is described in further detail in US 6,165,745.

In another embodiment, the antibody is modified to increase its biological half-life. Various approaches are possible. For example, one or more of the following mutations can be introduced: T252L, T254S, T256F, as described in US6,277,375. Alternatively, to increase the biological half life, the antibody can be altered within the CH1 or CL region to contain a salvage receptor binding epitope taken from two loops of a CH2 domain of an Fc region of an IgG, as described in US5,869,046 and US6,121,022.

In yet other embodiments, the Fc region is altered by replacing at least one amino acid residue with a different amino acid residue to alter the effector functions of the antibody. For example, one or more amino acids can be replaced with a different amino acid residue such that the antibody has an altered affinity for an effector ligand but retains the antigen-binding ability of the parent antibody. The effector ligand to which affinity is altered can be, for example, an Fc receptor or the C1 component of complement. This approach is described in further detail in US5,624,821 and US5,648,260, both by Winter *et al*. In particular, residues 234 and 235 may be mutated. In particular, these mutations may be to alanine. Thus in one embodiment the antibody comprised in the compositions of the disclosure has a mutation in the Fc region at one or both of amino acids 234 and 235. In another embodiment, one or both of amino acids 234 and 235 may be substituted to alanine. Substitution of both amino acids 234 and 235 to alanine results in a reduced ADCC activity.

In another embodiment, one or more amino acids selected from amino acid residues of the described antibodies can be replaced with a different amino acid residue such that the antibody has altered C1q binding and/or reduced or abolished complement dependent cytotoxicity (CDC). This approach is described in further detail in US6,194,551.

In another embodiment, one or more amino acid residues of the described antibodies are altered to thereby alter the ability of the antibody to fix complement. This approach is described further in WO94/29351.

In yet another embodiment, the Fc region of the described antibodies is modified to increase the ability of the antibody to mediate antibody dependent cellular cytotoxicity (ADCC) and/or to increase the affinity of the antibody for an Fcγ receptor by modifying one or more amino acids. This approach is described further in WO00/42072. Moreover, the binding sites on human IgG1 for FcγRI, FcγRII, FcγRIII and FcRn have been mapped and variants with improved binding have been described (see Shields, R.L. *et al.*, 2001 J. Biol. Chen. 276:6591-6604).

In still another embodiment, the glycosylation of an antibody comprised in the compositions of the disclosure is modified. For example, an aglycoslated antibody can be made (*i.e.* the antibody lacks glycosylation). Glycosylation can be altered to, for example, increase the affinity of the antibody for the antigen. Such carbohydrate modifications can be accomplished by; for example, altering one or more sites of glycosylation within the antibody sequence. For example, one or more amino acid substitutions can be made that result in elimination of one or more variable region framework glycosylation sites to thereby eliminate glycosylation at that site. Such aglycosylation may increase the affinity of the antibody for antigen. Such an approach is described in further detail in U.S. Patent Nos. 5,714,350 and 6,350,861 by Co *et al.*

Additionally or alternatively, an antibody can be used that has an altered type of glycosylation, such as a hypofucosylated antibody having reduced amounts of fucosyl residues or an antibody having increased bisecting GlcNac structures. Such altered glycosylation patterns have been demonstrated to increase the ADCC ability of antibodies. Such carbohydrate modifications can be accomplished by, for example, expressing the antibody in a host cell with altered glycosylation machinery. Cells with altered glycosylation machinery have been described in the art and can be used as host cells in which to express the disclosed recombinant antibodies to thereby produce an antibody with altered glycosylation. For example, EP 1,176,195 by Hang *et al.* describes

a cell line with a functionally disrupted FUT8 gene, which encodes a fucosyl transferase, such that antibodies expressed in such a cell line exhibit hypofucosylation. Therefore, in one embodiment, the antibodies comprised in the compositions of the disclosure are produced by recombinant expression in a cell line which exhibit hypofucosylation pattern, for example, a mammalian cell line with deficient expression of the FUT8 gene encoding fucosyltransferase. WO03/035835 describes a variant CHO cell line, Lecl3 cells, with reduced ability to attach fucose to Asn(297)-linked carbohydrates, also resulting in hypofucosylation of antibodies expressed in that host cell (see also Shields, R.L. *et al.*, 2002 J. Biol. Chem. 277:26733-26740). WO99/54342 describes cell lines engineered to express glycoprotein-modifying glycosyl transferases (*e.g.* beta(1,4)-N acetylglucosaminyltransferase III (GnTIII)) such that antibodies expressed in the engineered cell lines exhibit increased bisecting GlcNac structures which results in increased ADCC activity of the antibodies (see also Umana *et al.*, 1999 Nat. Biotech. 17:176-180). Alternatively, the antibodies comprised in the compositions of the disclosure can be produced in a yeast or a filamentous fungi engineered for mammalian-like glycosylation pattern, and capable of producing antibodies lacking fucose as glycosylation pattern (see for example EP1297172B1).

Another modification of the antibodies herein that is contemplated by the disclosure is pegylation. An antibody can be pegylated to, for example, increase the biological (*e.g.* serum) half-life of the antibody. To pegylate an antibody, the antibody, or fragment thereof, typically is reacted with polyethylene glycol (PEG), such as a reactive ester or aldehyde derivative of PEG, under conditions in which one or more PEG groups become attached to the antibody or antibody fragment. The pegylation can be carried out by an acylation reaction or an alkylation reaction with a reactive PEG molecule (or an analogous reactive water-soluble polymer). As used herein, the term "polyethylene glycol" is intended to encompass any of the forms of PEG that have been used to derivatize other proteins, such as mono (C1-C10) alkoxy- or aryloxy-polyethylene glycol or polyethylene glycol-maleimide. In certain embodiments, the used antibody to be pegylated is an aglycosylated antibody. Methods for pegylating proteins are known in the art and can be applied to the disclosed antibodies (see for example, EP0154316 and EP0401384).

Another modification of the antibodies that is contemplated by the disclosure is a conjugate or a protein fusion of at least the antigen-binding region of the antibody comprised in the composition of the disclosure to serum protein, such as human serum albumin or a fragment thereof to increase half-life of the resulting molecule (see, for example, EP0322094).

Another possibility is a fusion of at least the antigen-binding region of the antibody comprised in the composition of the disclosure to proteins capable of binding to serum proteins, such as human serum albumin to increase half life of the resulting molecule (see, for example, EP0486525).

Methods of engineering altered antibodies

As discussed above, the anti-ActRIIB antibodies having CDR sequences, V_H and V_L sequences or full length heavy and light chain sequences shown herein can be used to create new anti-ActRIIB antibodies by modifying the CDR sequences full length heavy chain and/or light chain sequences, V_H and/or V_L sequences, or the constant region(s) attached thereto. Thus, in another aspect of the disclosure, the structural features of an anti-ActRIIB antibody comprised in the compositions of the disclosure are used to create structurally related anti-ActRIIB antibodies that retain at least one functional property of the antibodies comprised in the compositions of the disclosure, such as binding to human ActRIIB but also inhibit one or more functional properties of ActRIIB (for example, the inhibition of Smad activation).

For example, one or more CDR regions of the antibodies comprised in the compositions of the present disclosure, or mutations thereof, can be combined recombinantly with known framework regions and/or other CDRs to create additional, recombinantly-engineered, anti-ActRIIB antibodies comprised in the compositions of the disclosure, as discussed above. Other types of modifications include those described in the previous section. The starting material for the engineering method is one or more of the V_H and/or V_L sequences provided herein, or one or more CDR regions thereof. To create the engineered antibody, it is not necessary to actually prepare (*i.e.* express as a protein) an antibody having one or more of the V_H and/or V_L sequences provided herein, or one or more CDR regions thereof. Rather, the information contained in the sequence(s) is used as the starting material to create a "second generation" sequence(s) derived from the original sequence(s) and then the "second generation" sequence(s) is prepared and expressed as a protein.

The altered antibody sequence can also be prepared by screening antibody libraries having fixed CDR3 sequences selected among the group consisting of SEQ ID NO: 29-42 and SEQ ID NO: 71-84 or minimal essential binding determinants as described in US2005/0255552 and diversity on CDR1 and CDR2 sequences. The screening can be performed according to any screening technology appropriate for screening antibodies from antibody libraries, such as phage display technology.

Standard molecular biology techniques can be used to prepare and express the altered antibody sequence. The antibody encoded by the altered antibody sequence(s) is one that retains one, some or all of the functional properties of the anti-ActRIIB antibodies described herein, which functional properties include, but are not limited to, specifically binding to human ActRIIB and inhibition of Smad activation.

The altered antibody may exhibit one or more, two or more, or three or more of the functional properties discussed above.

The functional properties of the altered antibodies can be assessed using standard assays available in the art and/or described herein, such as those set forth in the Examples (*e.g.* ELISAs).

Mutations can be introduced randomly or selectively along all or part of an anti-ActRIIB antibody coding sequence and the resulting modified anti-ActRIIB antibodies can be screened for binding activity and/or other functional properties as described herein. Mutational methods have been described in the art. For example, WO02/092780 describes methods for creating and screening antibody mutations using saturation mutagenesis, synthetic ligation assembly, or a combination thereof. Alternatively, WO03/074679 describes methods of using computational screening methods to optimize physiochemical properties of antibodies.

Nucleic acid molecules encoding antibodies comprised in the compositions of the disclosure

Examples of full length light chain nucleotide sequences optimized for expression in a mammalian cell are shown in SEQ ID NOs: 161-165 and 171-175. Examples of full length heavy chain nucleotide sequences optimized for expression in a mammalian cell are shown in SEQ ID NOs: 166-170 and 176-180.

The nucleic acids may be present in whole cells, in a cell lysate, or may be nucleic acids in a partially purified or substantially pure form. A nucleic acid is "isolated" or "rendered substantially pure" when purified away from other cellular components or other contaminants, *e.g.* other cellular nucleic acids or proteins, by standard techniques, including alkaline/SDS treatment, CsCl banding, column chromatography, agarose gel electrophoresis and others well known in the art. See, F. Ausubel, *et al.*, ed. 1987 Current Protocols in Molecular Biology, Greene Publishing and Wiley Interscience, New York. Nucleic acids can be obtained using standard molecular biology techniques. For antibodies expressed by hybridomas (*e.g.* hybridomas prepared from transgenic mice

carrying human immunoglobulin genes as described further below), cDNAs encoding the light and heavy chains of the antibody made by the hybridoma can be obtained by standard PCR amplification or cDNA cloning techniques. For antibodies obtained from an immunoglobulin gene library (*e.g.* using phage display techniques), nucleic acid encoding the antibody can be recovered from various phage clones that are members of the library.

Once DNA fragments encoding V_H and V_L segments are obtained, these DNA fragments can be further manipulated by standard recombinant DNA techniques, for example to convert the variable region genes to full-length antibody chain genes, to Fab fragment genes or to an scFv gene. In these manipulations, a V_L- or V_H-encoding DNA fragment is operatively linked to another DNA molecule, or to a fragment encoding another protein, such as an antibody constant region or a flexible linker. The term "operatively linked", as used in this context, is intended to mean that the two DNA fragments are joined in a functional manner, for example, such that the amino acid sequences encoded by the two DNA fragments remain in-frame, or such that the protein is expressed under control of a desired promoter.

The isolated DNA encoding the V_H region can be converted to a full-length heavy chain gene by operatively linking the V_H-encoding DNA to another DNA molecule encoding heavy chain constant regions (CH1, CH2 and CH3). The sequences of human heavy chain constant region genes are known in the art (see *e.g.* Kabat, E. A., *et al.* [*supra*]) and DNA fragments encompassing these regions can be obtained by standard PCR amplification. The heavy chain constant region can be an IgG1, IgG2, IgG3, IgG4, IgA, IgE, IgM or IgD constant region. The heavy chain contstant region can be selected among IgG1 isotypes. For a Fab fragment heavy chain gene, the V_H-encoding DNA can be operatively linked to another DNA molecule encoding only the heavy chain CH1 constant region.

The isolated DNA encoding the V_L region can be converted to a full-length light chain gene (as well as to a Fab light chain gene) by operatively linking the V_L-encoding DNA to another DNA molecule encoding the light chain constant region, CL. The sequences of human light chain constant region genes are known in the art (see *e.g.* Kabat, E. A., *et al.* [*supra*]) and DNA fragments encompassing these regions can be obtained by standard PCR amplification. The light chain constant region can be a kappa or a lambda constant region.

To create an scFv gene, the V_H- and V_L-encoding DNA fragments are operatively linked to another fragment encoding a flexible linker, *e.g.* encoding the amino acid sequence $(Gly4\ -Ser)_3$, such that the V_H and V_L sequences can be expressed as a contiguous single-chain protein, with the V_L and V_H regions joined by the flexible linker (see *e.g.* Bird *et al.*, 1988 Science 242:423-426; Huston *et al.*, 1988 Proc. Natl. Acad. Sci. USA 85:5879-5883; McCafferty *et al.*, 1990 Nature 348:552-554).

Generation of monoclonal antibodies

Monoclonal antibodies (mAbs) can be produced by a variety of techniques, including conventional monoclonal antibody methodology *e.g.* the standard somatic cell hybridization technique of Kohler and Milstein (1975 Nature 256: 495). Many techniques for producing monoclonal antibody can be employed *e.g.* viral or oncogenic transformation of B lymphocytes.

An animal system for preparing hybridomas is the murine system. Hybridoma production in the mouse is a well established procedure. Immunization protocols and techniques for isolation of immunized splenocytes for fusion are known in the art. Fusion partners (*e.g.* murine myeloma cells) and fusion procedures are also known.

Chimeric or humanized antibodies comprised in the compositions of the present disclosure can be prepared based on the sequence of a murine monoclonal antibody prepared as described above. DNA encoding the heavy and light chain immunoglobulins can be obtained from the murine hybridoma of interest and engineered to contain non-murine (*e.g.* human) immunoglobulin sequences using standard molecular biology techniques. For example, to create a chimeric antibody, the murine variable regions can be linked to human constant regions using methods known in the art (see *e.g.* US4,816,567). To create a humanized antibody, the murine CDR regions can be inserted into a human framework using methods known in the art (see *e.g.* U.S. Patent No. 5225539; 5530101; 5585089; 5693762 and 6180370).

In a certain embodiment, the antibodies comprised in the compositions of the disclosure are human monoclonal antibodies. Such human monoclonal antibodies directed against ActRIIB can be generated using transgenic or transchromosomic mice carrying parts of the human immune system rather than the mouse system. These transgenic and transchromosomic mice include mice referred to herein as HuMAb mice and KM mice, respectively, and are collectively referred to herein as "human Ig mice."

The HuMAb mouse® (Medarex, Inc.) contains human immunoglobulin gene miniloci that encode un-rearranged human heavy (μ and γ) and κ light chain immunoglobulin

sequences, together with targeted mutations that inactivate the endogenous μ and κ chain loci (see *e.g.* Lonberg, *et al.*, 1994 Nature 368(6474): 856-859). Accordingly, the mice exhibit reduced expression of mouse IgM or κ, and in response to immunization, the introduced human heavy and light chain transgenes undergo class switching and somatic mutation to generate high affinity human IgGκ monoclonal (Lonberg, N. *et al.*, 1994 [*supra*]; reviewed in Lonberg, N., 1994 Handbook of Experimental Pharmacology 113:49-101; Lonberg, N. and Huszar, D., 1995 Intern. Rev. Immunol.13: 65-93, and Harding, F. and Lonberg, N., 1995 Ann. N. Y. Acad. Sci. 764:536-546). The preparation and use of HuMAb mice, and the genomic modifications carried by such mice, is further described in Taylor, L. *et al.*, 1992 Nucleic Acids Research 20:6287-6295; Chen, J. *et al.*, 1993 International Immunology 5: 647-656; Tuaillon *et al.*, 1993 Proc. Natl. Acad. Sci. USA 94:3720-3724; Choi *et al.*, 1993 Nature Genetics 4:117-123; Chen, J. *et al.*, 1993 EMBO J. 12: 821-830; Tuaillon *et al.*, 1994 J. Immunol. 152:2912-2920; Taylor, L. *et al.*, 1994 International Immunology 579-591; and Fishwild, D. *et al.*, 1996 Nature Biotechnology 14: 845-851, the contents of all of which are hereby specifically incorporated by reference in their entirety. See further, U.S. Patent Nos. 5,545,806; 5,569,825; 5,625,126; 5,633,425; 5,789,650; 5,877,397; 5,661,016; 5,814,318; 5,874,299; 5,770,429; and 5,545,807; as well as WO92/103918, WO93/12227, WO94/25585, WO97/113852, WO98/24884; WO99/45962; and WO01/14424.

In another embodiment, human antibodies comprised in the compositions of the disclosure can be raised using a mouse that carries human immunoglobulin sequences on transgenes and transchomosomes such as a mouse that carries a human heavy chain transgene and a human light chain transchromosome. Such mice, referred to herein as "KM mice", are described in detail in WO02/43478.

Still further, alternative transgenic animal systems expressing human immunoglobulin genes are available in the art and can be used to raise anti-ActRIIB antibodies of the disclosure. For example, an alternative transgenic system referred to as the Xenomouse (Abgenix, Inc.) can be used. Such mice are described in, *e.g.* U.S. Patent Nos. 5,939,598; 6,075,181; 6,114,598; 6, 150,584 and 6,162,963.

Moreover, alternative transchromosomic animal systems expressing human immunoglobulin genes are available in the art and can be used to raise anti-ActRIIB antibodies of the disclosure. For example, mice carrying both a human heavy chain transchromosome and a human light chain tranchromosome, referred to as "TC mice" can be used; such mice are described in Tomizuka *et al.*, 2000 Proc. Natl. Acad. Sci. USA 97:722-727. Furthermore, cows carrying human heavy and light chain

transchromosomes have been described in the art (Kuroiwa *et al.*, 2002 Nature Biotechnology 20:889-894) and can be used to raise anti-ActRIIB antibodies.

Human recombinant antibodies comprised in the compositions of the disclosure can also be prepared using phage display methods for screening libraries of human immunoglobulin genes. Such phage display methods for isolating human antibodies are established in the art or described in the examples below. See for example: U.S. Patent Nos. 5,223,409; 5,403,484; 5,571,698; 5,427,908; 5,580,717; 5,969,108; 6,172,197; 5,885,793; 6,521,404; 6,544,731; 6,555,313; 6,582,915 and 6,593,081.

Human monoclonal antibodies comprised in the compositions of the disclosure can also be prepared using SCID mice into which human immune cells have been reconstituted such that a human antibody response can be generated upon immunization. Such mice are described in, for example, U.S. Patent Nos. 5,476,996 and 5,698,767.

Generation of hybridomas producing human monoclonal antibodies

To generate hybridomas producing human monoclonal antibodies comprised in the compositions of the disclosure, splenocytes and/or lymph node cells from immunized mice can be isolated and fused to an appropriate immortalized cell line, such as a mouse myeloma cell line. The resulting hybridomas can be screened for the production of antigen-specific antibodies. For example, single cell suspensions of splenic lymphocytes from immunized mice can be fused to one-sixth the number of P3X63-Ag8.653 nonsecreting mouse myeloma cells (ATCC, CRL 1580) with 50% PEG. Cells are plated at approximately 2 x 145 in flat bottom microtiter plates, followed by a two week incubation in selective medium containing 20% fetal Clone Serum, 18% "653" conditioned media, 5% origen (IGEN), 4 mM L-glutamine, 1 mM sodium pyruvate, 5mM HEPES, 0:055 mM 2-mercaptoethanol, 50 units/ml penicillin, 50 mg/ml streptomycin, 50 mg/ml gentamycin and 1X HAT (Sigma; the HAT is added 24 hours after the fusion). After approximately two weeks, cells can be cultured in medium in which the HAT is replaced with HT. Individual wells can then be screened by ELISA for human monoclonal IgM and IgG antibodies. Once extensive hybridoma growth occurs, medium can be observed usually after 10-14 days. The antibody secreting hybridomas can be replated, screened again, and if still positive for human IgG, the monoclonal antibodies can be subcloned at least twice by limiting dilution. The stable subclones can then be cultured *in vitro* to generate small amounts of antibody in tissue culture medium for characterization.

To purify human monoclonal antibodies, selected hybridomas can be grown in two-liter spinner-flasks for monoclonal antibody purification. Supernatants can be filtered and concentrated before affinity chromatography with protein A-sepharose (Pharmacia).

Eluted IgG can be checked by gel electrophoresis and high performance liquid chromatography to ensure purity. The buffer solution can be exchanged into PBS, and the concentration can be determined by OD_{280} using 1.43 extinction coefficient. The monoclonal antibodies can be aliquoted and stored at -80°C.

Generation of transfectomas producing monoclonal antibodies

Antibodies comprised in the compositions of the disclosure also can be produced in a host cell transfectoma using, for example, a combination of recombinant DNA techniques and gene transfection methods as is well known in the art (*e.g.* Morrison, S. (1985) Science 229:1202).

For example, to express the antibodies, or antibody fragments thereof, DNAs encoding partial or full-length light and heavy chains, can be obtained by standard molecular biology techniques (*e.g.* PCR amplification or cDNA cloning using a hybridoma that expresses the antibody of interest) and the DNAs can be inserted into expression vectors such that the genes are operatively linked to transcriptional and translational control sequences. In this context, the term "operatively linked" is intended to mean that an antibody gene is ligated into a vector such that transcriptional and translational control sequences within the vector serve their intended function of regulating the transcription and translation of the antibody gene. The expression vector and expression control sequences are chosen to be compatible with the expression host cell used. The antibody light chain gene and the antibody heavy chain gene can be inserted into separate vector or, more typically, both genes are inserted into the same expression vector. The antibody genes are inserted into the expression vector by standard methods (*e.g.* ligation of complementary restriction sites on the antibody gene fragment and vector, or blunt end ligation if no restriction sites are present). The light and heavy chain variable regions of the antibodies described herein can be used to create full-length antibody genes of any antibody isotype by inserting them into expression vectors already encoding heavy chain constant and light chain constant regions of the desired isotype such that the V_H segment is operatively linked to the CH segment(s) within the vector and the V_L segment is operatively linked to the CL segment within the vector. Additionally or alternatively, the recombinant expression vector can encode a signal peptide that facilitates secretion of the antibody chain from a host cell. The antibody chain gene can be cloned into the vector such that the signal peptide is linked in frame to the amino terminus of the antibody chain gene. The signal peptide can be an immunoglobulin signal peptide or a heterologous signal peptide (*i.e.* a signal peptide from a non-immunoglobulin protein).

In addition to the antibody chain genes, the recombinant expression vectors of the disclosure carry regulatory sequences that control the expression of the antibody chain

genes in a host cell. The term "regulatory sequence" is intended to include promoters, enhancers and other expression control elements (*e.g.* polyadenylation signals) that control the transcription or translation of the antibody chain genes. Such regulatory sequences are described, for example, in Goeddel (Gene Expression Technology. Methods in Enzymology 185, Academic Press, San Diego, CA 1990). It will be appreciated by those skilled in the art that the design of the expression vector, including the selection of regulatory sequences, may depend on such factors as the choice of the host cell to be transformed, the level of expression of protein desired, *etc.* Regulatory sequences for mammalian host cell expression include viral elements that direct high levels of protein expression in mammalian cells, such as promoters and/or enhancers derived from cytomegalovirus (CMV), Simian Virus 40 (SV40), adenovirus (*e.g.* the adenovirus major late promoter (AdMLP)), and polyoma. Alternatively, nonviral regulatory sequences may be used, such as the ubiquitin promoter or P-globin promoter. Still further, regulatory elements composed of sequences from different sources, such as the SRa promoter system, which contains sequences from the SV40 early promoter and the long terminal repeat of human T cell leukemia virus type 1 (Takebe, Y. *et al.*, 1988 Mol. Cell. Biol. 8:466-472).

In addition to the antibody chain genes and regulatory sequences, the recombinant expression vectors may carry additional sequences, such as sequences that regulate replication of the vector in host cells (*e.g.* origins of replication) and selectable marker genes. The selectable marker gene facilitates selection of host cells into which the vector has been introduced (see, *e.g.* U.S. Patent Nos. 4,399,216, 4,634,665 and 5,179,017). For example, typically the selectable marker gene confers resistance to drugs, such as G418, hygromycin or methotrexate, on a host cell into which the vector has been introduced. Selectable marker genes include the dihydrofolate reductase (DHFR) gene (for use in dhfr- host cells with methotrexate selection/amplification) and the neo gene (for G418 selection).

For expression of the light and heavy chains, the expression vector(s) encoding the heavy and light chains is transfected into a host cell by standard techniques. The various forms of the term "transfection" are intended to encompass a wide variety of techniques commonly used for the introduction of exogenous DNA into a prokaryotic or eukaryotic host cell, *e.g.* electroporation, calcium-phosphate precipitation, DEAE-dextran transfection and the like. It is theoretically possible to express the antibodies of the disclosure in either prokaryotic or eukaryotic host cells. Expression of antibodies in eukaryotic cells, in particular mammalian host cells, is discussed because such eukaryotic cells, and in particular mammalian cells, are more likely than prokaryotic cells

to assemble and secrete a properly folded and immunologically active antibody. Prokaryotic expression of antibody genes has been reported to be ineffective for production of high yields of active antibody (Boss, M. A. and Wood, C. R., 1985 Immunology Today 6:12-13).

Mammalian host cells for expressing the recombinant antibodies comprised in the compositions of the disclosure include Chinese Hamster Ovary (CHO cells) (including dhfr- CHO cells, described Urlaub and Chasin, 1980 Proc. Natl. Acad. Sci. USA 77:4216-4220 used with a DH FR selectable marker, *e.g.* as described in R.J. Kaufman and P.A. Sharp, 1982 Mol. Biol. 159:601-621), NSO myeloma cells, COS cells and SP2 cells. In one embodiment the host cells are CHO K1PD cells. In particular, for use with NSO myeloma cells, another expression system is the GS gene expression system shown in WO87/04462, WO89/01036 and EP 338,841. Mammalian host cells for expressing the recombinant antibodies comprised in the compositions of the disclosure include mammalian cell lines deficient for FUT8 gene expression, for example as described in US6,946,292B2. When recombinant expression vectors encoding antibody genes are introduced into mammalian host cells, the antibodies are produced by culturing the host cells for a period of time sufficient to allow for expression of the antibody in the host cells or secretion of the antibody into the culture medium in which the host cells are grown. Antibodies can be recovered from the culture medium using standard protein purification methods.

Immunoconjugates

In another aspect, the present disclosure features compositions comprising an anti-ActRIIB antibody, or a fragment thereof, conjugated to a therapeutic moiety, such as a cytotoxin, a drug (*e.g.*, an immunosuppressant) or a radiotoxin. Such conjugates are referred to herein as "immunoconjugates". Immunoconjugates that include one or more cytotoxins are referred to as "immunotoxins." A cytotoxin or cytotoxic agent includes any agent that is detrimental to (*e.g.*, kills) cells.

Cytotoxins can be conjugated to antibodies of the disclosure using linker technology available in the art. Examples of linker types that have been used to conjugate a cytotoxin to an antibody include, but are not limited to, hydrazones, thioethers, esters, disulfides and peptide-containing linkers. A linker can be chosen that is, for example, susceptible to cleavage by low pH within the lysosomal compartment or susceptible to cleavage by proteases, such as proteases preferentially expressed in tumor tissue such as cathepsins (*e.g.* cathepsins B, C, D).

For further discussion of types of cytotoxins, linkers and methods for conjugating therapeutic agents to antibodies, see also Saito, G. *et al.*, 2003 Adv. Drug Deliv. Rev. 55:199-215; Trail, P.A. *et al.*, 2003 Cancer Immunol. Immunother. 52:328-337; Payne, G. 2003 Cancer Cell 3:207-212; Allen, T.M., 2002 Nat. Rev. Cancer 2:750-763; Pastan, I. and Kreitman, R. J., 2002 Curr. Opin. Investig. Drugs 3:1089-1091; Senter, P.D. and Springer, C.J., 2001 Adv. Drug Deliv. Rev. 53:247-264.

Antibodies comprised in the compositions of the present disclosure also can be conjugated to a radioactive isotope to generate cytotoxic radiopharmaceuticals, also referred to as radioimmunoconjugates. Examples of radioactive isotopes that can be conjugated to antibodies for use diagnostically or therapeutically include, but are not limited to, iodine^{131}, indium^{111}, yttrium^{90}, and lutetium^{177}. Methods for preparing radioimmunconjugates are established in the art. Examples of radioimmunoconjugates are commercially available, including Zevalin™ (DEC Pharmaceuticals) and Bexxar™ (Corixa Pharmaceuticals), and similar methods can be used to prepare radioimmunoconjugates using the antibodies of the disclosure.

The antibody conjugates comprised in the compositions of the disclosure can be used to modify a given biological response, and the drug moiety is not to be construed as limited to classical chemical therapeutic agents. For example, the drug moiety may be a protein or polypeptide possessing a desired biological activity. Such proteins may include, for example, an enzymatically active toxin, or active fragment thereof, such as abrin, ricin A, pseudomonas exotoxin, or diphtheria toxin; a protein such as tumor necrosis factor or interferon-γ; or, biological response modifiers such as, for example, lymphokines, interleukin-1 ("IL-1"), interleukin-2 ("IL-2"), interleukin-6 ("IL-6"), granulocyte macrophage colony stimulating factor ("GM-CSF"), granulocyte colony stimulating factor ("G-CSF"), or other growth factors.

Techniques for conjugating such therapeutic moiety to antibodies are well known, see, *e.g.* Amon *et al.*, "Monoclonal Antibodies For Immunotargeting Of Drugs In Cancer Therapy", in Monoclonal Antibodies And Cancer Therapy, Reisfeld *et al.* (eds.), pp. 243-56 (Alan R. Liss, Inc. 1985); Hellstrom *et al.*, "Antibodies For Drug Delivery", in Controlled Drug Delivery (2nd Ed.), Robinson *et al.* (eds.), pp. 623-53 (Marcel Dekker, Inc. 1987); Thorpe, "Antibody Carriers Of Cytotoxic Agents In Cancer Therapy: A Review", in Monoclonal Antibodies '84: Biological And Clinical Applications, Pinchera *et al.* (eds.), pp. 475-506 (1985); "Analysis, Results, And Future Prospective Of The Therapeutic Use Of Radiolabeled Antibody In Cancer Therapy", in Monoclonal Antibodies For Cancer Detection And Therapy, Baldwin *et al.* (eds.), pp. 303-16

(Academic Press 1985), and Thorpe *et al.*, "The Preparation And Cytotoxic Properties Of Antibody-Toxin Conjugates", Inmunol. Rev., 62:119-58 (1982).

Bispecific molecules

In another aspect, the present disclosure features compositions comprising bispecific or multispecific molecules comprising an anti-ActRIIB antibody, or a fragment thereof, of the disclosure. An antibody comprised in the compositions of the disclosure, or antigen-binding regions thereof, can be derivatized or linked to another functional molecule, *e.g.* another peptide or protein (*e.g.* another antibody or ligand for a receptor) to generate a bispecific molecule that binds to at least two different binding sites or target molecules. The antibody of the disclosure may in fact be derivatized or linked to more than one other functional molecule to generate multi-specific molecules that bind to more than two different binding sites and/or target molecules; such multi-specific molecules are also intended to be encompassed by the term "bispecific molecule" as used herein. To create a bispecific molecule of the disclosure, an antibody of the disclosure can be functionally linked (*e.g.* by chemical coupling, genetic fusion, noncovalent association or otherwise) to one or more other binding molecules, such as another antibody, antibody fragment, peptide or binding mimetic, such that a bispecific molecule results.

Accordingly, the present disclosure includes compositions comprising bispecific molecules comprising at least one first binding specificity for ActRIIB and a second binding specificity for a second target epitope. For example, the second target epitope may be another epitope of ActRIIB different from the first target epitope.

Additionally, for the compositions in which the bispecific molecule is multi-specific, the molecule can further include a third binding specificity, in addition to the first and second target epitope.

In one embodiment, the bispecific molecules of the disclosed compositions comprise as a binding specificity at least one antibody, or an antibody fragment thereof, including, *e.g.* an Fab, Fab', $F(ab')_2$, Fv, or a single chain Fv. The antibody may also be a light chain or heavy chain dimer, or any minimal fragment thereof such as a Fv or a single chain construct as described in Ladner *et al.* US4,946,778, the contents of which is expressly incorporated by reference.

Other antibodies which can be employed in the bispecific molecules are murine, chimeric and humanized monoclonal antibodies.

The bispecific molecules comprised in the compositions of the present disclosure can be prepared by conjugating the constituent binding specificities, using methods known in the art. For example, each binding specificity of the bispecific molecule can be generated separately and then conjugated to one another. When the binding specificities are proteins or peptides, a variety of coupling or cross-linking agents can be used for covalent conjugation. Examples of cross-linking agents include protein A, carbodiimide, N-succinimidyl-S-acetyl-thioacetate (SATA), 5,5'-dithiobis(2-nitrobenzoic acid) (DTNB), o-phenylenedimaleimide (oPDM), N-succinimidyl-3-(2-pyridyldithio)propionate (SPDP), and sulfosuccinimidyl 4-(N-maleimidomethyl) cyclohaxane-l-carboxylate (sulfo-SMCC) (see *e.g.* Karpovsky *et al.*, 1984 J. Exp. Med. 160:1686; Liu, MA *et al.*, 1985 Proc. Natl. Acad. Sci. USA 82:8648). Other methods include those described in Paulus, 1985 Behring Ins. Mitt. No. 78,118-132; Brennan *et al.*, 1985 Science 229:81-83), and Glennie *et al.*, 1987 J. Immunol. 139: 2367-2375). Conjugating agents are SATA and sulfo-SMCC, both available from Pierce Chemical Co. (Rockford, IL).

When the binding specificities are antibodies, they can be conjugated by sulfhydryl bonding of the C-terminus hinge regions of the two heavy chains. In a particularly embodiment, the hinge region is modified to contain an odd number of sulfhydryl residues, for example one, prior to conjugation.

Alternatively, both binding specificities can be encoded in the same vector and expressed and assembled in the same host cell. This method is particularly useful where the bispecific molecule is a mAb x mAb, mAb x Fab, Fab x $F(ab')_2$ or ligand x Fab fusion protein. A bispecific molecule comprised in the compositions of the disclosure can be a single chain molecule comprising one single chain antibody and a binding determinant, or a single chain bispecific molecule comprising two binding determinants. Bispecific molecules may comprise at least two single chain molecules. Methods for preparing bispecific molecules are described for example in U.S. Patent Numbers 5,260,203; 5,455,030; 4,881,175; 5,132,405; 5,091,513; 5,476,786; 5,013,653; 5,258,498; and 5,482,858.

Binding of the bispecific molecules to their specific targets can be confirmed by, for example, enzyme-linked immunosorbent assay (ELISA), radioimmunoassay (RIA), FACS analysis, bioassay (*e.g.* growth inhibition), or Western Blot assay. Each of these assays generally detects the presence of protein-antibody complexes of particular interest by employing a labeled reagent (*e.g.* an antibody) specific for the complex of interest.

Multivalent antibodies

In another aspect, the present disclosure relates to compositions comprising multivalent antibodies comprising at least two identical or different antigen-binding portions of the disclosed antibodies binding to ActRIIB. In one embodiment, the multivalent antibodies provide at least two, three or four antigen-binding portions of the antibodies. The antigen-binding portions can be linked together via protein fusion or covalent or non covalent linkage. Alternatively, methods of linkage have been described for the bispecific molecules. In various embodiments, the composition can be mono-, bi- or multi-valent (e.g., capable of binding to one, two or several antigens), and/or mono-, bi- or multi-specific (e.g., having binding region(s) capable of binding to one, two or several different antigens). a composition can be any combination of these, e.g., monovalent and mono-specific (having one binding region that binds to one antigen or epitope); or bi-valent and bi-specific (having two binding regions, each of which bind to a different epitope or antigen); or bi-valent and mono-specific (having two binding regions, each of which bind to the same epitope or antigen); or multi-valent and mono-specific (having several binding regions that all bind to the same antigen or epitope); or multi-valent and multi-specific (having several binding regions that bind to several different antigens or epitopes).

Pharmaceutical compositions

In another aspect, the present disclosure provides a composition, *e.g.* a pharmaceutical composition, containing one or a combination of the above described antibodies/monoclonal antibodies, or antigen-binding portion(s) thereof, formulated together with a pharmaceutically acceptable carrier. Such compositions may include one or a combination of (*e.g.* two or more different) the described antibodies, or immunoconjugates or bispecific molecules. For example, a pharmaceutical composition of the disclosure can comprise a combination of antibodies that bind to different epitopes on the target antigen or that have complementary activities.

Pharmaceutical compositions of the disclosure also can be administered in combination therapy, *i.e.* combined with other agents. For example, the combination therapy can include an anti-ActRII antibody of the present disclosure combined with at least one other muscle mass/strength increasing agent, for example, IGF-1, IGF-2 or variants of IGF-1 or IGF-2, an anti-myostatin antibody, a myostatin propeptide, a myostatin decoy protein that binds ActRIIB but does not activate it, a beta 2 agonist, a Ghrelin agonist, a SARM, GH agonists/mimetics or follistatin. Examples of therapeutic agents that can be used in

combination therapy are described in greater detail below in the section on uses of the antibodies of the disclosure.

As used herein, "pharmaceutically acceptable carrier" includes any and all solvents, dispersion media, coatings, antibacterial and antifungal agents, isotonic and absorption delaying agents, and the like that are physiologically compatible. The carrier should be suitable for intravenous, intramuscular, subcutaneous, parenteral, spinal or epidermal administration (*e.g.* by injection or infusion), preferably for intravenous injection or infusion. Depending on the route of administration, the active compound, *i.e.* antibody, immunoconjuage, or bispecific molecule, may be coated in a material to protect the compound from the action of acids and other natural conditions that may inactivate the compound.

The pharmaceutical compositions of the disclosure may include one or more pharmaceutically acceptable salts. A "pharmaceutically acceptable salt" refers to a salt that retains the desired biological activity of the parent compound and does not impart any undesired toxicological effects (see *e.g.* Berge, S.M., *et al.*, 1977 J. Pharm. Sci. 66:1-19). Examples of such salts include acid addition salts and base addition salts. Acid addition salts include those derived from nontoxic inorganic acids, such as hydrochloric, nitric, phosphoric, sulfuric, hydrobromic, hydroiodic, phosphorous and the like, as well as from nontoxic organic acids such as aliphatic mono- and di-carboxylic acids, phenyl-substituted alkanoic acids, hydroxy alkanoic acids, aromatic acids, aliphatic and aromatic sulfonic acids and the like. Base addition salts include those derived from alkaline earth metals, such as sodium, potassium, magnesium, calcium and the like, as well as from nontoxic organic amines, such as N,N'-dibenzylethylenediamine, N-methylglucamine, chloroprocaine, choline, diethanolamine, ethylenediamine, procaine and the like.

A pharmaceutical composition of the disclosure also may include a pharmaceutically acceptable anti-oxidant. Examples of pharmaceutically acceptable antioxidants include: water soluble antioxidants, such as ascorbic acid, cysteine hydrochloride, sodium bisulfate, sodium metabisulfite, sodium sulfite and the like; oil-soluble antioxidants, such as ascorbyl palmitate, butylated hydroxyanisole (BHA), butylated hydroxytoluene (BHT), lecithin, propyl gallate, alpha-tocopherol, and the like; and metal chelating agents, such as citric acid, ethylenediamine tetraacetic acid (EDTA), sorbitol, tartaric acid, phosphoric acid, and the like.

Examples of suitable aqueous and nonaqueous carriers that may be employed in the pharmaceutical compositions of the disclosure include water, ethanol, polyols (such as

glycerol, propylene glycol, polyethylene glycol, and the like), and suitable mixtures thereof, vegetable oils, such as olive oil, and injectable organic esters, such as ethyl oleate. Proper fluidity can be maintained, for example, by the use of coating materials, such as lecithin, by the maintenance of the required particle size in the case of dispersions, and by the use of surfactants.

These compositions may also contain adjuvants such as preservatives, wetting agents, emulsifying agents and dispersing agents. Prevention of presence of microorganisms may be ensured both by sterilization procedures, *supra*, and by the inclusion of various antibacterial and antifungal agents, for example, paraben, chlorobutanol, phenol sorbic acid, and the like. It may also be desirable to include isotonic agents, such as sugars, sodium chloride, and the like into the compositions. In addition, prolonged absorption of the injectable pharmaceutical form may be brought about by the inclusion of agents which delay absorption such as, aluminum monostearate and gelatin.

Pharmaceutically acceptable carriers include sterile aqueous solutions or dispersions and sterile powders for the extemporaneous preparation of sterile injectable solutions or dispersion. The use of such media and agents for pharmaceutically active substances is known in the art. Except insofar as any conventional media or agent is incompatible with the active compound, use thereof in the pharmaceutical compositions of the disclosure is contemplated. Supplementary active compounds can also be incorporated into the compositions.

Therapeutic compositions typically must be sterile and stable under the conditions of manufacture and storage. The composition can be formulated as a solution, microemulsion, liposome, or other ordered structure suitable to high drug concentration. The carrier can be a solvent or dispersion medium containing, for example, water, ethanol, polyol (for example, glycerol, propylene glycol, and liquid polyethylene glycol, and the like), and suitable mixtures thereof. The proper fluidity can be maintained, for example, by the use of a coating such as lecithin, by the maintenance of the required particle size in the case of dispersion and by the use of surfactants. In many cases, one can include isotonic agents, for example, sugars, polyalcohols such as mannitol, sorbitol, or sodium chloride in the composition. Prolonged absorption of the injectable compositions can be brought about by including in the composition an agent that delays absorption for example, monostearate salts and gelatin.

Sterile injectable solutions can be prepared by incorporating the active compound in the required amount in an appropriate solvent with one or a combination of agents enumerated above, as required, followed by sterilization microfiltration. Generally,

dispersions are prepared by incorporating the active compound into a sterile vehicle that contains a basic dispersion medium and the required other agents from those enumerated above. In the case of sterile powders for the preparation of sterile injectable solutions, the methods of preparation are vacuum drying and freeze-drying (lyophilization) that yield a powder of the active agent plus any additional desired agent from a previously sterile-filtered solution thereof.

The amount of active agent which can be combined with a carrier material to produce a single dosage form will vary depending upon the subject being treated, and the particular mode of administration. The amount of active agent which can be combined with a carrier material to produce a single dosage form will generally be that amount of the composition which produces a therapeutic effect. Generally, out of one hundred percent, this amount will range from about 0.01 per cent to about ninety-nine percent of active agent, from about 0.1 per cent to about 70 per cent, or from about 1 percent to about 30 percent of active agent in combination with a pharmaceutically acceptable carrier.

Dosage regimens are adjusted to provide the optimum desired response (*e.g.* a therapeutic response). For example, a single bolus may be administered, several divided doses may be administered over time or the dose may be proportionally reduced or increased as indicated by the exigencies of the therapeutic situation. It is especially advantageous to formulate parenteral compositions in dosage unit form for ease of administration and uniformity of dosage. Dosage unit form as used herein refers to physically discrete units suited as unitary dosages for the subjects to be treated; each unit contains a predetermined quantity of active compound calculated to produce the desired therapeutic effect in association with the required pharmaceutical carrier. The specification for the dosage unit forms of the disclosure are dictated by and directly dependent on the unique characteristics of the active compound and the particular therapeutic effect to be achieved, and the limitations inherent in the art of compounding such an active compound for the treatment of sensitivity in individuals.

For administration of the antibody comprising composition for use according to the present disclosure, the antibody dosage ranges from about 0.0001 to about 100 mg/kg, and more usually about 0.01 to about 30 mg/kg, of the host body weight. For example dosages are about 1 mg/kg body weight, about 3 mg/kg body weight, about 5 mg/kg body weight or about 10 mg/kg body weight within the ranges of about 1-10 mg/kg e.g., about 1, 2, 3, 4, 5, 6, 7, 8, 9, 10 mg/kg body weight, preferably once every 4 weeks. Such administration is preferably carried out intraveneously. Alternatively, administration is carried out subcutaneously.

Dosage regimens for an anti-ActRII antibody of the disclosure, e.g., bimagrumab, include about 1 mg/kg body weight or about 3 mg/kg body weight or about 10 mg/kg body once every four weeks weight by intravenous administration.

Preferaby the compositions of the disclosure are for use in the treatment of sporadic inclusion body myositis.

In one specific embodiment the physical function and mobility of the patient suffering from sporadic inclusion body myositis are improved.

In one specific embodiment the swallowing difficulties or dysphagia of the patient suffering from sporadic inclusion body myositis are improved.

In one specific embodiment upper extremity strength of the patient suffering from sporadic inclusion body myositis is improved.

In one specific embodiment incidence of falls or risk of fallingof the patient suffering from sporadic inclusion body myositis are reduced.

In some methods, two or more monoclonal antibodies with different binding specificities are comprised in the compositions of the disclosure and, thus, administered simultaneously, in which case the dosage of each antibody administered falls within the ranges indicated. An antibody is usually administered on multiple occasions. Intervals between single dosages can be, for example, weekly, monthly, every three months, every six months or yearly. Intervals can also be irregular as indicated by measuring blood levels of antibody to the target antigen in the patient. In some methods, dosage is adjusted to achieve a plasma antibody concentration of about 1- about 1000 µg/ml and in some methods about 25- about 300 µg/ml. For example, an ActRII antibody of the disclosure could be co-administered with an anti-myostatin antibody.

Dosage and frequency vary depending on the half-life of the antibody in the patient. In general, human antibodies show the longest half-life, followed by humanized antibodies, chimeric antibodies, and nonhuman antibodies. The dosage and frequency of administration can vary depending on whether the treatment is prophylactic or therapeutic. In prophylactic applications, a relatively low dosage is administered at relatively infrequent intervals over a long period of time. Some patients continue to receive treatment for the rest of their lives. In therapeutic applications, a relatively high dosage at relatively short intervals is sometimes required until progression of the disease

is reduced or terminated or until the patient shows partial or complete amelioration of symptoms of disease. Thereafter, the patient can be administered a prophylactic regime.

Administration of a "therapeutically effective dosage" of an anti-ActRII antibody comprised in the compositions of the disclosure can result in a decrease in severity of disease symptoms, an increase in frequency and duration of disease symptom-free periods, or a prevention of impairment or disability due to the disease affliction *i.e.* an increase in muscle mass and/or strength.

The active compounds can be prepared with carriers that will protect the compound against rapid release, such as a controlled release formulation, including implants, transdermal patches, and microencapsulated delivery systems. Biodegradable, biocompatible polymers can be used, such as ethylene vinyl acetate, polyanhydrides, polyglycolic acid, collagen, polyorthoesters, and polylactic acid. Many methods for the preparation of such formulations are patented or generally known to those skilled in the art. See, *e.g.* Sustained and Controlled Release Drug Delivery Systems, J.R. Robinson, ed., Marcel Dekker, Inc., New York, 1978.

Therapeutic compositions can be administered with medical devices known in the art.

Uses and methods of the disclosure

The compositions of the present disclosure and the disclosed antibodies have therapeutic utilities, because they have an impact on the treatment of sporadic inclusion body myositis or on the amelioration of the condition of patients affected by sporadic inclusion body myositis or on the reduction of symptoms associated with sporadic inclusion body myositis.

The term "subject" or “individual” as used herein is intended to include human and non-human animals. Non-human animals include all vertebrates, *e.g.* mammals and non-mammals, such as non-human primates, sheep, dogs, cats, cows, horses, chickens, amphibians, and reptiles.

Hence, the disclosure also relates to methods of treatment in which compositions of the disclosure or the disclosed myostatin antagonists, e.g., myostatin binding molecules or ActRII binding molecules, preferably ActRII binding molecules, more preferably antibodies to ActRII, e.g, bimagrumab or BYM338, inhibit, *i.e.* antagonize, the function of ActRII and thereby resulting in the alleviation of sporadic inclusion body myositis. The disclosure provides a method of treating a patient suffering from sporadic inclusion body

myositis comprising administering a therapeutically effective amount of a myostatin antagonist, e.g., myostatin binding molecule or ActRIIB binding molecule, preferably ActRIIB binding molecule, more preferably an antagonist antibody to ActRIIB, e.g, BYM338 or the disclosed compositions to the patient.

Examples of myostatin antagonists, e.g., myostatin binding molecules or ActRII binding molecules, preferably ActRIIB binding molecules, more preferably antagonist antibodies to ActRIIB, e.g, bimagrumab or BYM338, that can be used in the disclosed methods of treatment are those disclosed or described in detail above. In certain embodiments, the ActRII antibodies (e.g., bimagrumab or BYM338) are comprised in the herein disclosed inventive compositions.

The disclosure also relates to the use of a myostatin antagonist, e.g., myostatin binding molecule or ActRIIB binding molecule, preferably ActRIIB binding molecule, more preferably an antagonist antibody to ActRII, e.g, BYM338, in the manufacture of a medicament for treating sporadic inclusion body myositis.

In a further embodiment, the patient may be one who has not responded to previous treatments. For example, the patient may not have responded to treatment with IGF-1, IGF-2 or variants of IGF-1 or IGF-2, an anti-myostatin antibody, a myostatin propeptide, a myostatin decoy protein that binds ActRIIB but does not activate it, a beta 2 agonist, a Ghrelin agonist, a SARM, GH agonists/mimetics or follistatin. A simple way of measuring a patient's response to treatment may be timing how long it takes for a patient to climb a known height of stairs and comparing the results both before and after treatment.

The myostatin antagonist, e.g., myostatin binding molecule or ActRII binding molecule, preferably ActRII binding molecule, more preferably an antagonist antibody to ActRII, e.g., bimagrumab or BYM338, may be administered as the sole active agent or in conjunction with, *e.g.* as an adjuvant to or in combination to, other drugs *e.g.* IGF-1, IGF-2 or variants of IGF-1 or IGF-2, an anti-myostatin antibody, a myostatin propeptide, a myostatin decoy protein that binds ActRIIB but does not activate it, a beta 2 agonist, a Ghrelin agonist, a SARM, GH agonists/mimetics or follistatin. For example, the antagonists of the disclosure may be used in combination with an IGF-1 mimetic as disclosed in WO2007/146689.

In accordance with the foregoing the present disclosure provides in a yet further aspect:
A method or use as defined above comprising co-administration, *e.g.* concomitantly or in sequence, of a therapeutically effective amount of a myostatin antagonist, e.g., myostatin

binding molecule or ActRII binding molecule, preferably an ActRII or binding molecule, more preferably an antagonist antibody to ActRII, e.g, bimagrumab or BYM338, and at least one second drug substance, said second drug substance being IGF-1, IGF-2 or variants of IGF-1 or IGF-2, an anti-myostatin antibody, a myostatin propeptide, a myostatin decoy protein that binds ActRII but does not activate it, a beta 2 agonist, a Ghrelin agonist, a SARM, GH agonists/mimetics or follistatin.

Kits

The invention also encompasses kits which may comprise a myostatin antagonist, e.g., a myostatin binding molecule (e.g., a myostatin antibody or antigen binding fragment thereof, e.g., bimagrumab or BYM338) or myostatin receptor (i.e., ActRIIB receptor) binding molecule (e.g., anti-ActRIIB antibody or antigen binding fragment thereof) (e.g., in liquid or lyophilized form) or a pharmaceutical composition comprising the myostatin antagonist (described *supra)*. Additionally, such kits may comprise means for administering the myostatin antagonist (e.g., a syringe and vial, a prefilled syringe, a prefilled pen) and instructions for use. These kits may contain additional therapeutic agents (described *supra*), e.g., for delivery in combination with the enclosed myostatin antagonist, e.g., BYM338.

The phrase “means for administering” is used to indicate any available implement for systemically administering a drug top a patient, including, but not limited to, a pre-filled syringe, a vial and syringe, an injection pen, an autoinjector, an i.v. drip and bag, a pump, etc. With such items, a patient may self-administer the drug (i.e., administer the drug on their own behalf) or a physician may administer the drug.

Each component of the kit is usually enclosed within an individual container, and all of the various containers are within a single package along with instructions for use.

SEQUENCES

Table 1: sequence listing

SEQ ID NO	Ab region	Sequence
SEQ ID NO1	HCDR1	GYTFTSSYIN
SEQ ID NO2	HCDR1	GYTFTSSYIN
SEQ ID NO3	HCDR1	GYTFTSSYIN
SEQ ID NO4	HCDR1	GYTFTSSYIN
SEQ ID NO5	HCDR1	GYTFTSSYIN
SEQ ID NO6	HCDR1	GYTFTSSYIN
SEQ ID NO7	HCDR1	GYTFTSSYIN

SEQ ID NO8	HCDR1	GYTFTSSYIN
SEQ ID NO9	HCDR1	GYTFTSSYIN
SEQ ID NO10	HCDR1	GYTFTSSYIN
SEQ ID NO11	HCDR1	GYTFTSSYIN
SEQ ID NO12	HCDR1	GYTFTSSYIN
SEQ ID NO13	HCDR1	GYTFTSSYIN
SEQ ID NO14	HCDR1	GYTFTSSYIN
SEQ ID NO15	HCDR2	TINPVSGNTSYAQKFQG
SEQ ID NO16	HCDR2	TINPVSGNTSYAQKFQG
SEQ ID NO17	HCDR2	TINPVSGNTSYAQKFQG
SEQ ID NO18	HCDR2	TINPVSGNTSYAQKFQG
SEQ ID NO19	HCDR2	MINAPIGTTRYAQKFQG
SEQ ID NO20	HCDR2	QINAASGMTRYAQKFQG
SEQ ID NO21	HCDR2	MINAPIGTTRYAQKFQG
SEQ ID NO22	HCDR2	TINPVSGNTRYAQKFQG
SEQ ID NO23	HCDR2	TINPVSGSTSYAQKFQG
SEQ ID NO24	HCDR2	QINAASGMTRYAQKFQG
SEQ ID NO25	HCDR2	NINAAAGITLYAQKFQG
SEQ ID NO26	HCDR2	TINPPTGGTYYAQKFQG
SEQ ID NO27	HCDR2	GINPPAGTTSYAQKFQG
SEQ ID NO28	HCDR2	NINPATGHADYAQKFQG
SEQ ID NO29	HCDR3	GGWFDY
SEQ ID NO30	HCDR3	GGWFDY
SEQ ID NO31	HCDR3	GGWFDY
SEQ ID NO32	HCDR3	GGWFDY
SEQ ID NO33	HCDR3	GGWFDY
SEQ ID NO34	HCDR3	GGWFDY
SEQ ID NO35	HCDR3	GGWFDY
SEQ ID NO36	HCDR3	GGWFDY
SEQ ID NO37	HCDR3	GGWFDY
SEQ ID NO38	HCDR3	GGWFDY
SEQ ID NO39	HCDR3	GGWFDY
SEQ ID NO40	HCDR3	GGWFDY
SEQ ID NO41	HCDR3	GGWFDY
SEQ ID NO42	HCDR3	GGWFDY
SEQ ID NO43	LCDR1	TGTSSDVGSYNYVN
SEQ ID NO44	LCDR1	TGTSSDVGSYNYVN
SEQ ID NO45	LCDR1	TGTSSDVGSYNYVN
SEQ ID NO46	LCDR1	TGTSSDVGSYNYVN
SEQ ID NO47	LCDR1	TGTSSDVGSYNYVN
SEQ ID NO48	LCDR1	TGTSSDVGSYNYVN
SEQ ID NO49	LCDR1	TGTSSDVGSYNYVN
SEQ ID NO50	LCDR1	TGTSSDVGSYNYVN
SEQ ID NO51	LCDR1	TGTSSDVGSYNYVN
SEQ ID NO52	LCDR1	TGTSSDVGSYNYVN
SEQ ID NO53	LCDR1	TGTSSDVGSYNYVN
SEQ ID NO54	LCDR1	TGTSSDVGSYNYVN
SEQ ID NO55	LCDR1	TGTSSDVGSYNYVN
SEQ ID NO56	LCDR1	TGTSSDVGSYNYVN
SEQ ID NO57	LDCR2	LMIYGVSKRPS

SEQ ID NO58	LDCR2	LMIYGVSKRPS
SEQ ID NO59	LDCR2	LMIYGVSKRPS
SEQ ID NO60	LDCR2	LMIYGVSKRPS
SEQ ID NO61	LDCR2	LMIYGVSKRPS
SEQ ID NO62	LDCR2	LMIYGVSKRPS
SEQ ID NO63	LDCR2	LMIYGVSKRPS
SEQ ID NO64	LDCR2	LMIYGVSKRPS
SEQ ID NO65	LDCR2	LMIYGVSKRPS
SEQ ID NO66	LDCR2	LMIYGVSKRPS
SEQ ID NO67	LDCR2	LMIYGVSKRPS
SEQ ID NO68	LDCR2	LMIYGVSKRPS
SEQ ID NO69	LDCR2	LMIYGVSKRPS
SEQ ID NO70	LDCR2	LMIYGVSKRPS
SEQ ID NO71	LCDR3	QAWTSKMAG
SEQ ID NO72	LCDR3	SSYTRMGHP
SEQ ID NO73	LCDR3	ATYGKGVTPP
SEQ ID NO74	LCDR3	GTFAGGSYYG
SEQ ID NO75	LCDR3	QAWTSKMAG
SEQ ID NO76	LCDR3	QAWTSKMAG
SEQ ID NO77	LCDR3	GTFAGGSYYG
SEQ ID NO78	LCDR3	GTFAGGSYYG
SEQ ID NO79	LCDR3	GTFAGGSYYG
SEQ ID NO80	LCDR3	GTFAGGSYYG
SEQ ID NO81	LCDR3	GTFAGGSYYG
SEQ ID NO82	LCDR3	GTFAGGSYYG
SEQ ID NO83	LCDR3	GTFAGGSYYG
SEQ ID NO84	LCDR3	GTFAGGSYYG
SEQ ID NO85	VL	DIALTQPASVSGSPGQSITISCTGTSSDVGSYNYVNWYQQHPGKAPKLMIYGVSKRPSGV SNRFSGSKSGNTASLTISGLQAEDEADYYCQAWTSKMAGVFGGGTKLTVLGQ
SEQ ID NO86	VL	DIALTQPASVSGSPGQSITISCTGTSSDVGSYNYVNWYQQHPGKAPKLMIYGVSKRPSGV SNRFSGSKSGNTASLTISGLQAEDEADYYCSSYTRMGHPVFGGGTKLTVLGQ
SEQ ID NO87	VL	DIALTQPASVSGSPGQSITISCTGTSSDVGSYNYVNWYQQHPGKAPKLMIYGVSKRPSGV SNRFSGSKSGNTASLTISGLQAEDEADYYCATYGKGVTPPVFGGGTKLTVLGQ
SEQ ID NO88	VL	DIALTQPASVSGSPGQSITISCTGTSSDVGSYNYVNWYQQHPGKAPKLMIYGVSKRPSGV SNRFSGSKSGNTASLTISGLQAEDEADYYCGTFAGGSYYGVFGGGTKLTVLGQ
SEQ ID NO89	VL	DIALTQPASVSGSPGQSITISCTGTSSDVGSYNYVNWYQQHPGKAPKLMIYGVSKRPSGV SNRFSGSKSGNTASLTISGLQAEDEADYYCQAWTSKMAGVFGGGTKLTVLGQ
SEQ ID NO90	VL	DIALTQPASVSGSPGQSITISCTGTSSDVGSYNYVNWYQQHPGKAPKLMIYGVSKRPSGV SNRFSGSKSGNTASLTISGLQAEDEADYYCQAWTSKMAGVFGGGTKLTVLGQ
SEQ ID NO91	VL	DIALTQPASVSGSPGQSITISCTGTSSDVGSYNYVNWYQQHPGKAPKLMIYGVSKRPSGV SNRFSGSKSGNTASLTISGLQAEDEADYYCGTFAGGSYYGVFGGGTKLTVLGQ
SEQ ID NO92	VL	DIALTQPASVSGSPGQSITISCTGTSSDVGSYNYVNWYQQHPGKAPKLMIYGVSKRPSGV SNRFSGSKSGNTASLTISGLQAEDEADYYCGTFAGGSYYGVFGGGTKLTVLGQ
SEQ ID NO93	VL	DIALTQPASVSGSPGQSITISCTGTSSDVGSYNYVNWYQQHPGKAPKLMIYGVSKRPSGV SNRFSGSKSGNTASLTISGLQAEDEADYYCGTFAGGSYYGVFGGGTKLTVLGQ
SEQ ID NO94	VL	DIALTQPASVSGSPGQSITISCTGTSSDVGSYNYVNWYQQHPGKAPKLMIYGVSKRPSGV SNRFSGSKSGNTASLTISGLQAEDEADYYCGTFAGGSYYGVFGGGTKLTVLGQ
SEQ ID NO95	VL	DIALTQPASVSGSPGQSITISCTGTSSDVGSYNYVNWYQQHPGKAPKLMIYGVSKRPSGV SNRFSGSKSGNTASLTISGLQAEDEADYYCGTFAGGSYYGVFGGGTKLTVLGQ
SEQ ID NO96	VL	DIALTQPASVSGSPGQSITISCTGTSSDVGSYNYVNWYQQHPGKAPKLMIYGVSKRPSGV

		SNRFSGSKSGNTASLTISGLQAEDEADYYCGTFAGGSYYGVFGGGTKLTVLGQ
SEQ ID NO97	VL	DIALTQPASVSGSPGQSITISCTGTSSDVGSYNYVNWYQQHPGKAPKLMIYGVSKRPSGV SNRFSGSKSGNTASLTISGLQAEDEADYYCGTFAGGSYYGVFGGGTKLTVLGQ
SEQ ID NO98	VL	DIALTQPASVSGSPGQSITISCTGTSSDVGSYNYVNWYQQHPGKAPKLMIYGVSKRPSGV SNRFSGSKSGNTASLTISGLQAEDEADYYCGTFAGGSYYGVFGGGTKLTVLGQ
SEQ ID NO99	VH	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSSYINWVRQAPGQGLEWMGTINPVSGNT SYAQKFQGRVTMTRDTSISTAYMELSSLRSEDTAVYYCARGGWFDYWGQGTLVTVSS
SEQ ID NO100	VH	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSSYINWVRQAPGQGLEWMGTINPVSGNT SYAQKFQGRVTMTRDTSISTAYMELSSLRSEDTAVYYCARGGWFDYWGQGTLVTVSS
SEQ ID NO101	VH	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSSYINWVRQAPGQGLEWMGTINPVSGNT SYAQKFQGRVTMTRDTSISTAYMELSSLRSEDTAVYYCARGGWFDYWGQGTLVTVSS
SEQ ID NO102	VH	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSSYINWVRQAPGQGLEWMGTINPVSGNT SYAQKFQGRVTMTRDTSISTAYMELSSLRSEDTAVYYCARGGWFDYWGQGTLVTVSS
SEQ ID NO103	VH	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSSYINWVRQAPGQGLEWMGMINAPIGTTR YAQKFQGRVTMTRDTSISTAYMELSSLRSEDTAVYYCARGGWFDYWGQGTLVTVSS
SEQ ID NO104	VH	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSSYINWVRQAPGQGLEWMGQINAASGMT RYAQKFQGRVTMTRDTSISTAYMELSSLRSEDTAVYYCARGGWFDYWGQGTLVTVSS
SEQ ID NO105	VH	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSSYINWVRQAPGQGLEWMGMINAPIGTTR YAQKFQGRVTMTRDTSISTAYMELSSLRSEDTAVYYCARGGWFDYWGQGTLVTVSS
SEQ ID NO106	VH	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSSYINWVRQAPGQGLEWMGTINPVSGNT RYAQKFQGRVTMTRDTSISTAYMELSSLRSEDTAVYYCARGGWFDYWGQGTLVTVSS
SEQ ID NO107	VH	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSSYINWVRQAPGQGLEWMGTINPVSGST SYAQKFQGRVTMTRDTSISTAYMELSSLRSEDTAVYYCARGGWFDYWGQGTLVTVSS
SEQ ID NO108	VH	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSSYINWVRQAPGQGLEWMGQINAASGMT RYAQKFQGRVTMTRDTSISTAYMELSSLRSEDTAVYYCARGGWFDYWGQGTLVTVSS
SEQ ID NO109	VH	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSSYINWVRQAPGQGLEWMGNINAAAGITL YAQKFQGRVTMTRDTSISTAYMELSSLRSEDTAVYYCARGGWFDYWGQGTLVTVSS
SEQ ID NO110	VH	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSSYINWVRQAPGQGLEWMGTINPPTGGT YYAQKFQGRVTMTRDTSISTAYMELSSLRSEDTAVYYCARGGWFDYWGQGTLVTVSS
SEQ ID NO111	VH	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSSYINWVRQAPGQGLEWMGGINPPAGTT SYAQKFQGRVTMTRDTSISTAYMELSSLRSEDTAVYYCARGGWFDYWGQGTLVTVSS
SEQ ID NO112	VH	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSSYINWVRQAPGQGLEWMGNINPATGHA DYAQKFQGRVTMTRDTSISTAYMELSSLRSEDTAVYYCARGGWFDYWGQGTLVTVSS
SEQ ID NO113	DNA VL	GATATCGCACTGACCCAGCCAGCTTCAGTGAGCGGCTCACCAGGTCAGAGCATTACC ATCTCGTGTACGGGTACTAGCAGCGATGTTGGTTCTTATAATTATGTGAATTGGTACC AGCAGCATCCCGGGAAGGCGCCGAAACTTATGATTTATGGTGTTTCTAAGCGTCCCT CAGGCGTGAGCAACCGTTTTAGCGGATCCAAAAGCGGCAACACCGCGAGCCTGACC ATTAGCGGCCTGCAAGCGGAAGACGAAGCGGATTATTATTGCCAGGCTTGGACTTCT AAGATGGCTGGTGTGTTTGGCGGCGGCACGAAGTTAACCGTTCTTGGCCAG
SEQ ID NO114	DNA VL	GATATCGCACTGACCCAGCCAGCTTCAGTGAGCGGCTCACCAGGTCAGAGCATTACC ATCTCGTGTACGGGTACTAGCAGCGATGTTGGTTCTTATAATTATGTGAATTGGTACC AGCAGCATCCCGGGAAGGCGCCGAAACTTATGATTTATGGTGTTTCTAAGCGTCCCT CAGGCGTGAGCAACCGTTTTAGCGGATCCAAAAGCGGCAACACCGCGAGCCTGACC ATTAGCGGCCTGCAAGCGGAAGACGAAGCGGATTATTATTGCTCTTCTTATACTCGTA TGGGTCATCCTGTGTTTGGCGGCGGCACGAAGTTAACCGTTCTTGGCCAG
SEQ ID NO115	DNA VL	GATATCGCACTGACCCAGCCAGCTTCAGTGAGCGGCTCACCAGGTCAGAGCATTACC ATCTCGTGTACGGGTACTAGCAGCGATGTTGGTTCTTATAATTATGTGAATTGGTACC AGCAGCATCCCGGGAAGGCGCCGAAACTTATGATTTATGGTGTTTCTAAGCGTCCCT CAGGCGTGAGCAACCGTTTTAGCGGATCCAAAAGCGGCAACACCGCGAGCCTGACC ATTAGCGGCCTGCAAGCGGAAGACGAAGCGGATTATTATTGCGCTACTTATGGTAAG GGTGTTACTCCTCCTGTGTTTGGCGGCGGCACGAAGTTAACCGTTCTTGGCCAG

SEQ ID NO116	DNA VL	GATATCGCACTGACCCAGCCAGCTTCAGTGAGCGGCTCACCAGGTCAGAGCATTACC ATCTCGTGTACGGGTACTAGCAGCGATGTTGGTTCTTATAATTATGTGAATTGGTACC AGCAGCATCCCGGGAAGGCGCCGAAACTTATGATTTATGGTGTTTCTAAGCGTCCCT CAGGCGTGAGCAACCGTTTTAGCGGATCCAAAAGCGGCAACACCGCGAGCCTGACC ATTAGCGGCCTGCAAGCGGAAGACGAAGCGGATTATTATTGCGGTACTTTTGCTGGT GGTTCTTATTATGGTGTGTTTGGCGGCGGCACGAAGTTAACCGTTCTTGGCCAG
SEQ ID NO117	DNA VL	GATATCGCACTGACCCAGCCAGCTTCAGTGAGCGGCTCACCAGGTCAGAGCATTACC ATCTCGTGTACGGGTACTAGCAGCGATGTTGGTTCTTATAATTATGTGAATTGGTACC AGCAGCATCCCGGGAAGGCGCCGAAACTTATGATTTATGGTGTTTCTAAGCGTCCCT CAGGCGTGAGCAACCGTTTTAGCGGATCCAAAAGCGGCAACACCGCGAGCCTGACC ATTAGCGGCCTGCAAGCGGAAGACGAAGCGGATTATTATTGCCAGGCTTGGACTTCT AAGATGGCTGGTGTGTTTGGCGGCGGCACGAAGTTAACCGTTCTTGGCCAG
SEQ ID NO118	DNA VL	GATATCGCACTGACCCAGCCAGCTTCAGTGAGCGGCTCACCAGGTCAGAGCATTACC ATCTCGTGTACGGGTACTAGCAGCGATGTTGGTTCTTATAATTATGTGAATTGGTACC AGCAGCATCCCGGGAAGGCGCCGAAACTTATGATTTATGGTGTTTCTAAGCGTCCCT CAGGCGTGAGCAACCGTTTTAGCGGATCCAAAAGCGGCAACACCGCGAGCCTGACC ATTAGCGGCCTGCAAGCGGAAGACGAAGCGGATTATTATTGCCAGGCTTGGACTTCT AAGATGGCTGGTGTGTTTGGCGGCGGCACGAAGTTAACCGTTCTTGGCCAG
SEQ ID NO119	DNA VL	GATATCGCACTGACCCAGCCAGCTTCAGTGAGCGGCTCACCAGGTCAGAGCATTACC ATCTCGTGTACGGGTACTAGCAGCGATGTTGGTTCTTATAATTATGTGAATTGGTACC AGCAGCATCCCGGGAAGGCGCCGAAACTTATGATTTATGGTGTTTCTAAGCGTCCCT CAGGCGTGAGCAACCGTTTTAGCGGATCCAAAAGCGGCAACACCGCGAGCCTGACC ATTAGCGGCCTGCAAGCGGAAGACGAAGCGGATTATTATTGCGGTACTTTTGCTGGT GGTTCTTATTATGGTGTGTTTGGCGGCGGCACGAAGTTAACCGTTCTTGGCCAG
SEQ ID NO120	DNA VL	GATATCGCACTGACCCAGCCAGCTTCAGTGAGCGGCTCACCAGGTCAGAGCATTACC ATCTCGTGTACGGGTACTAGCAGCGATGTTGGTTCTTATAATTATGTGAATTGGTACC AGCAGCATCCCGGGAAGGCGCCGAAACTTATGATTTATGGTGTTTCTAAGCGTCCCT CAGGCGTGAGCAACCGTTTTAGCGGATCCAAAAGCGGCAACACCGCGAGCCTGACC ATTAGCGGCCTGCAAGCGGAAGACGAAGCGGATTATTATTGCGGTACTTTTGCTGGT GGTTCTTATTATGGTGTGTTTGGCGGCGGCACGAAGTTAACCGTTCTTGGCCAG
SEQ ID NO121	DNA VL	GATATCGCACTGACCCAGCCAGCTTCAGTGAGCGGCTCACCAGGTCAGAGCATTACC ATCTCGTGTACGGGTACTAGCAGCGATGTTGGTTCTTATAATTATGTGAATTGGTACC AGCAGCATCCCGGGAAGGCGCCGAAACTTATGATTTATGGTGTTTCTAAGCGTCCCT CAGGCGTGAGCAACCGTTTTAGCGGATCCAAAAGCGGCAACACCGCGAGCCTGACC ATTAGCGGCCTGCAAGCGGAAGACGAAGCGGATTATTATTGCGGTACTTTTGCTGGT GGTTCTTATTATGGTGTGTTTGGCGGCGGCACGAAGTTAACCGTTCTTGGCCAG
SEQ ID NO122	DNA VL	GATATCGCACTGACCCAGCCAGCTTCAGTGAGCGGCTCACCAGGTCAGAGCATTACC ATCTCGTGTACGGGTACTAGCAGCGATGTTGGTTCTTATAATTATGTGAATTGGTACC AGCAGCATCCCGGGAAGGCGCCGAAACTTATGATTTATGGTGTTTCTAAGCGTCCCT CAGGCGTGAGCAACCGTTTTAGCGGATCCAAAAGCGGCAACACCGCGAGCCTGACC ATTAGCGGCCTGCAAGCGGAAGACGAAGCGGATTATTATTGCGGTACTTTTGCTGGT GGTTCTTATTATGGTGTGTTTGGCGGCGGCACGAAGTTAACCGTTCTTGGCCAG
SEQ ID NO123	DNA VL	GATATCGCACTGACCCAGCCAGCTTCAGTGAGCGGCTCACCAGGTCAGAGCATTACC ATCTCGTGTACGGGTACTAGCAGCGATGTTGGTTCTTATAATTATGTGAATTGGTACC AGCAGCATCCCGGGAAGGCGCCGAAACTTATGATTTATGGTGTTTCTAAGCGTCCCT CAGGCGTGAGCAACCGTTTTAGCGGATCCAAAAGCGGCAACACCGCGAGCCTGACC ATTAGCGGCCTGCAAGCGGAAGACGAAGCGGATTATTATTGCGGTACTTTTGCTGGT GGTTCTTATTATGGTGTGTTTGGCGGCGGCACGAAGTTAACCGTTCTTGGCCAG
SEQ ID NO124	DNA VL	GATATCGCACTGACCCAGCCAGCTTCAGTGAGCGGCTCACCAGGTCAGAGCATTACC ATCTCGTGTACTGGTACTAGCAGCGATGTTGGTTCTTATAATTATGTGAATTGGTACCA GCAGCATCCCGGGAAGGCGCCGAAACTTATGATTTATGGTGTTTCTAAGCGTCCCTC

		AGGCGTGAGCAACCGTTTTAGCGGATCCAAAAGCGGCAACACCGCGAGCCTGACCAT TAGCGGCCTGCAAGCGGAAGACGAAGCGGATTATTATTGCGGTACTTTTGCTGGTGG TTCTTATTATGGTGTGTTTGGCGGCGGCACGAAGTTAACCGTTCTTGGCCAG
SEQ ID NO125	DNA VL	GATATCGCACTGACCCAGCCAGCTTCAGTGAGCGGCTCACCAGGTCAGAGCATTACC ATCTCGTGTACGGGTACTAGCAGCGATGTTGGTTCTTATAATTATGTGAATTGGTACC AGCAGCATCCCGGGAAGGCGCCGAAACTTATGATTTATGGTGTTTCTAAGCGTCCCT CAGGCGTGAGCAACCGTTTTAGCGGATCCAAAAGCGGCAACACCGCGAGCCTGACC ATTAGCGGCCTGCAAGCGGAAGACGAAGCGGATTATTATTGCGGTACTTTTGCTGGT GGTTCTTATTATGGTGTGTTTGGCGGCGGCACGAAGTTAACCGTTCTTGGCCAG
SEQ ID NO126	DNA VL	GATATCGCACTGACCCAGCCAGCTTCAGTGAGCGGCTCACCAGGTCAGAGCATTACC ATCTCGTGTACGGGTACTAGCAGCGATGTTGGTTCTTATAATTATGTGAATTGGTACC AGCAGCATCCCGGGAAGGCGCCGAAACTTATGATTTATGGTGTTTCTAAGCGTCCCT CAGGCGTGAGCAACCGTTTTAGCGGATCCAAAAGCGGCAACACCGCGAGCCTGACC ATTAGCGGCCTGCAAGCGGAAGACGAAGCGGATTATTATTGCGGTACTTTTGCTGGT GGTTCTTATTATGGTGTGTTTGGCGGCGGCACGAAGTTAACCGTTCTTGGCCAG
SEQ ID NO127	DNA VH	CAGGTGCAATTGGTTCAGAGCGGCGCGGAAGTGAAAAAACCGGGCGCGAGCGTGAA AGTGAGCTGCAAAGCCTCCGGATATACCTTTACTTCTTCTTATATTAATTGGGTCCGCC AAGCCCCTGGGCAGGGTCTCGAGTGGATGGGCACTATCAATCCGGTTTCTGGCAATA CGTCTTACGCGCAGAAGTTTCAGGGCCGGGTGACCATGACCCGTGATACCAGCATTA GCACCGCGTATATGGAACTGAGCAGCCTGCGTAGCGAAGATACGGCCGTGTATTATT GCGCGCGTGGTGGTTGGTTTGATTATTGGGGCCAAGGCACCCTGGTGACGGTTAGCT CA
SEQ ID NO128	DNA VH	CAGGTGCAATTGGTTCAGAGCGGCGCGGAAGTGAAAAAACCGGGCGCGAGCGTGAA AGTGAGCTGCAAAGCCTCCGGATATACCTTTACTTCTTCTTATATTAATTGGGTCCGCC AAGCCCCTGGGCAGGGTCTCGAGTGGATGGGCACTATCAATCCGGTTTCTGGCAATA CGTCTTACGCGCAGAAGTTTCAGGGCCGGGTGACCATGACCCGTGATACCAGCATTA GCACCGCGTATATGGAACTGAGCAGCCTGCGTAGCGAAGATACGGCCGTGTATTATT GCGCGCGTGGTGGTTGGTTTGATTATTGGGGCCAAGGCACCCTGGTGACGGTTAGCT CA
SEQ ID NO129	DNA VH	CAGGTGCAATTGGTTCAGAGCGGCGCGGAAGTGAAAAAACCGGGCGCGAGCGTGAA AGTGAGCTGCAAAGCCTCCGGATATACCTTTACTTCTTCTTATATTAATTGGGTCCGCC AAGCCCCTGGGCAGGGTCTCGAGTGGATGGGCACTATCAATCCGGTTTCTGGCAATA CGTCTTACGCGCAGAAGTTTCAGGGCCGGGTGACCATGACCCGTGATACCAGCATTA GCACCGCGTATATGGAACTGAGCAGCCTGCGTAGCGAAGATACGGCCGTGTATTATT GCGCGCGTGGTGGTTGGTTTGATTATTGGGGCCAAGGCACCCTGGTGACGGTTAGCT CA
SEQ ID NO130	DNA VH	CAGGTGCAATTGGTTCAGAGCGGCGCGGAAGTGAAAAAACCGGGCGCGAGCGTGAA AGTGAGCTGCAAAGCCTCCGGATATACCTTTACTTCTTCTTATATTAATTGGGTCCGCC AAGCCCCTGGGCAGGGTCTCGAGTGGATGGGCACTATCAATCCGGTTTCTGGCAATA CGTCTTACGCGCAGAAGTTTCAGGGCCGGGTGACCATGACCCGTGATACCAGCATTA GCACCGCGTATATGGAACTGAGCAGCCTGCGTAGCGAAGATACGGCCGTGTATTATT GCGCGCGTGGTGGTTGGTTTGATTATTGGGGCCAAGGCACCCTGGTGACGGTTAGCT CA
SEQ ID NO131	DNA VH	CAGGTGCAATTGGTTCAGAGCGGCGCGGAAGTGAAAAAACCGGGCGCGAGCGTGAA AGTGAGCTGCAAAGCCTCCGGATATACCTTTACTTCTTCTTATATTAATTGGGTCCGCC AAGCCCCTGGGCAGGGTCTCGAGTGGATGGGCATGATTAATGCTCCTATTGGTACTA CTCGTTATGCTCAGAAGTTTCAGGGTCGGGTGACCATGACCCGTGATACCAGCATTA GCACCGCGTATATGGAACTGAGCAGCCTGCGTAGCGAAGATACGGCCGTGTATTATT GCGCGCGTGGTGGTTGGTTTGATTATTGGGGCCAAGGCACCCTGGTGACGGTTAGCT CA

SEQ ID NO132	DNA VH	CAGGTGCAATTGGTTCAGAGCGGCGCGGAAGTGAAAAAACCGGGCGCGAGCGTGAA AGTGAGCTGCAAAGCCTCCGGATATACCTTTACTTCTTCTTATATTAATTGGGTCCGCC AAGCCCCTGGGCAGGGTCTCGAGTGGATGGGCCAGATTAATGCTGCTTCTGGTATGA CTCGTTATGCTCAGAAGTTTCAGGGTCGGGTGACCATGACCCGTGATACCAGCATTA GCACCGCGTATATGGAACTGAGCAGCCTGCGTAGCGAAGATACGGCCGTGTATTATT GCGCGCGTGGTGGTTGGTTTGATTATTGGGGCCAAGGCACCCTGGTGACGGTTAGCT CA
SEQ ID NO133	DNA VH	CAGGTGCAATTGGTTCAGAGCGGCGCGGAAGTGAAAAAACCGGGCGCGAGCGTGAA AGTGAGCTGCAAAGCCTCCGGATATACCTTTACTTCTTCTTATATTAATTGGGTCCGCC AAGCCCCTGGGCAGGGTCTCGAGTGGATGGGCATGATTAATGCTCCTATTGGTACTA CTCGTTATGCTCAGAAGTTTCAGGGTCGGGTGACCATGACCCGTGATACCAGCATTA GCACCGCGTATATGGAACTGAGCAGCCTGCGTAGCGAAGATACGGCCGTGTATTATT GCGCGCGTGGTGGTTGGTTTGATTATTGGGGCCAAGGCACCCTGGTGACGGTTAGCT CA
SEQ ID NO134	DNA VH	CAGGTGCAATTGGTTCAGAGCGGCGCGGAAGTGAAAAAACCGGGCGCGAGCGTGAA AGTGAGCTGCAAAGCCTCCGGATATACCTTTACTTCTTCTTATATTAATTGGGTCCGCC AAGCCCCTGGGCAGGGTCTCGAGTGGATGGGCACTATCAATCCGGTTTCTGGCAATA CGCGTTACGCGCAGAAGTTTCAGGGCCGGGTGACCATGACCCGTGATACCAGCATTA GCACCGCGTATATGGAACTGAGCAGCCTGCGTAGCGAAGATACGGCCGTGTATTATT GCGCGCGTGGTGGTTGGTTTGATTATTGGGGCCAAGGCACCCTGGTGACGGTTAGCT CA
SEQ ID NO135	DNA VH	CAGGTGCAATTGGTTCAGAGCGGCGCGGAAGTGAAAAAACCGGGCGCGAGCGTGAA AGTGAGCTGCAAAGCCTCCGGATATACCTTTACTTCTTCTTATATTAATTGGGTCCGCC AAGCCCCTGGGCAGGGTCTCGAGTGGATGGGCACTATCAATCCGGTTTCTGGCTCTA CGTCTTACGCGCAGAAGTTTCAGGGCCGGGTGACCATGACCCGTGATACCAGCATTA GCACCGCGTATATGGAACTGAGCAGCCTGCGTAGCGAAGATACGGCCGTGTATTATT GCGCGCGTGGTGGTTGGTTTGATTATTGGGGCCAAGGCACCCTGGTGACGGTTAGCT CA
SEQ ID NO136	DNA VH	CAGGTGCAATTGGTTCAGAGCGGCGCGGAAGTGAAAAAACCGGGCGCGAGCGTGAA AGTGAGCTGCAAAGCCTCCGGATATACCTTTACTTCTTCTTATATTAATTGGGTCCGCC AAGCCCCTGGGCAGGGTCTCGAGTGGATGGGCCAGATTAATGCTGCTTCTGGTATGA CTCGTTATGCTCAGAAGTTTCAGGGTCGGGTCACCATGACCCGTGATACCAGCATTA GCACCGCGTATATGGAACTGAGCAGCCTGCGTAGCGAAGATACGGCCGTGTATTATT GCGCGCGTGGTGGTTGGTTTGATTATTGGGGCCAAGGCACCCTGGTGACGGTTAGCT CA
SEQ ID NO137	DNA VH	CAGGTGCAATTGGTTCAGAGCGGCGCGGAAGTGAAAAAACCGGGCGCGAGCGTGAA AGTGAGCTGCAAAGCCTCCGGATATACCTTTACTTCTTCTTATATTAATTGGGTCCGCC AAGCCCCTGGGCAGGGTCTCGAGTGGATGGGCAATATTAATGCTGCTGCTGGTATTA CTCTTTATGCTCAGAAGTTTCAGGGTCGGGTCACCATGACCCGTGATACCAGCATTAG CACCGCGTATATGGAACTGAGCAGCCTGCGTAGCGAAGATACGGCCGTGTATTATTG CGCGCGTGGTGGTTGGTTTGATTATTGGGGCCAAGGCACCCTGGTGACGGTTAGCTC A
SEQ ID NO138	DNA VH	CAGGTGCAATTGGTTCAGAGCGGCGCGGAAGTGAAAAAACCGGGCGCGAGCGTGAA AGTGAGCTGCAAAGCCTCCGGATATACCTTTACTTCTTCTTATATTAATTGGGTCCGCC AAGCCCCTGGGCAGGGTCTCGAGTGGATGGGCACTATTAATCCTCCTACTGGAGGTA CTTATTATGCTCAGAAGTTTCAGGGTCGGGTGACCATGACCCGTGATACCAGCATTAG CACCGCGTATATGGAACTGAGCAGCCTGCGTAGCGAAGATACGGCCGTGTATTATTG CGCGCGTGGTGGTTGGTTTGATTATTGGGGCCAAGGCACCCTGGTGACGGTTAGCTC A

SEQ ID NO139	DNA VH	CAGGTGCAATTGGTTCAGAGCGGCGCGGAAGTGAAAAAACCGGGCGCGAGCGTGAA AGTGAGCTGCAAAGCCTCCGGATATACCTTTACTTCTTCTTATATTAATTGGGTCCGCC AAGCCCCTGGGCAGGGTCTCGAGTGGATGGGCGGTATTAATCCTCCTGCTGGTACTA CTTCTTATGCTCAGAAGTTTCAGGGTCGGGTCACCATGACCCGTGATACCAGCATTAG CACCGCGTATATGGAACTGAGCAGCCTGCGTAGCGAAGATACGGCCGTGTATTATTG CGCGCGTGGTGGTTGGTTTGATTATTGGGGCCAAGGCACCCTGGTGACGGTTAGCTC A
SEQ ID NO140	DNA VH	CAGGTGCAATTGGTTCAGAGCGGCGCGGAAGTGAAAAAACCGGGCGCGAGCGTGAA AGTGAGCTGCAAAGCCTCCGGATATACCTTTACTTCTTCTTATATTAATTGGGTCCGCC AAGCCCCTGGGCAGGGTCTCGAGTGGATGGGCAATATTAATCCTGCTACTGGTCATG CTGATTATGCTCAGAAGTTTCAGGGTCGGGTGACCATGACCCGTGATACCAGCATTA GCACCGCGTATATGGAACTGAGCAGCCTGCGTAGCGAAGATACGGCCGTGTATTATT GCGCGCGTGGTGGTTGGTTTGATTATTGGGGCCAAGGCACCCTGGTGACGGTTAGCT CA
SEQ ID NO141	Light Chain	QSALTQPASVSGSPGQSITISCTGTSSDVGSYNYVNWYQQHPGKAPKLMIYGVSKRPSG VSNRFSGSKSGNTASLTISGLQAEDEADYYCGTFAGGSYYGVFGGGTKLTVLGQPKAAP SVTLFPPSSEELQANKATLVCLISDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYA ASSYLSLTPEQWKSHRSYSCQVTHEGSTVEKTVAPTECS
SEQ ID NO142	Light Chain	QSALTQPASVSGSPGQSITISCTGTSSDVGSYNYVNWYQQHPGKAPKLMIYGVSKRPSG VSNRFSGSKSGNTASLTISGLQAEDEADYYCGTFAGGSYYGVFGGGTKLTVLGQPKAAP SVTLFPPSSEELQANKATLVCLISDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYA ASSYLSLTPEQWKSHRSYSCQVTHEGSTVEKTVAPTECS
SEQ ID NO143	Light Chain	QSALTQPASVSGSPGQSITISCTGTSSDVGSYNYVNWYQQHPGKAPKLMIYGVSKRPSG VSNRFSGSKSGNTASLTISGLQAEDEADYYCGTFAGGSYYGVFGGGTKLTVLGQPKAAP SVTLFPPSSEELQANKATLVCLISDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYA ASSYLSLTPEQWKSHRSYSCQVTHEGSTVEKTVAPTECS
SEQ ID NO144	Light Chain	QSALTQPASVSGSPGQSITISCTGTSSDVGSYNYVNWYQQHPGKAPKLMIYGVSKRPSG VSNRFSGSKSGNTASLTISGLQAEDEADYYCGTFAGGSYYGVFGGGTKLTVLGQPKAAP SVTLFPPSSEELQANKATLVCLISDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYA ASSYLSLTPEQWKSHRSYSCQVTHEGSTVEKTVAPTECS
SEQ ID NO145	Light Chain	QSALTQPASVSGSPGQSITISCTGTSSDVGSYNYVNWYQQHPGKAPKLMIYGVSKRPSG VSNRFSGSKSGNTASLTISGLQAEDEADYYCGTFAGGSYYGVFGGGTKLTVLGQPKAAP SVTLFPPSSEELQANKATLVCLISDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYA ASSYLSLTPEQWKSHRSYSCQVTHEGSTVEKTVAPTECS
SEQ ID NO146	Heavy Chain	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSSYINWVRQAPGQGLEWMGTINPVSGST SYAQKFQGRVTMTRDTSISTAYMELSRLRSDDTAVYYCARGGWFDYWGQGTLVTVSSA STKGPSVFPLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSS GLYSLSSVVTVPSSSLGTQTYICNVNHKPSNTKVDKRVEPKSCDKTHTCPPCPAPEAAGG PSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPREEQY NSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREPQVYTLPPSRE EMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKSR WQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
SEQ ID NO147	Heavy Chain	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSSYINWVRQAPGQGLEWMGQINAASGMT RYAQKFQGRVTMTRDTSISTAYMELSRLRSDDTAVYYCARGGWFDYWGQGTLVTVSSA STKGPSVFPLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSS GLYSLSSVVTVPSSSLGTQTYICNVNHKPSNTKVDKRVEPKSCDKTHTCPPCPAPEAAGG PSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPREEQY NSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREPQVYTLPPSRE EMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKSR WQQGNVFSCSVMHEALHNHYTQKSLSLSPGK

SEQ ID NO148	Heavy Chain	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSSYINWVRQAPGQGLEWMGNINAAAGITL YAQKFQGRVTMTRDTSISTAYMELSRLRSDDTAVYYCARGGWFDYWGQGTLVTVSSAS TKGPSVFPLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGL YSLSSVVTVPSSSLGTQTYICNVNHKPSNTKVDKRVEPKSCDKTHTCPPCPAPEAAGGPS VFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPREEQYNST YRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREPQVYTLPPSREEMT KNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKSRWQ QGNVFSCSVMHEALHNHYTQKSLSLSPGK
SEQ ID NO149	Heavy Chain	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSSYINWVRQAPGQGLEWMGGINPPAGTT SYAQKFQGRVTMTRDTSISTAYMELSRLRSDDTAVYYCARGGWFDYWGQGTLVTVSSA STKGPSVFPLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSS GLYSLSSVVTVPSSSLGTQTYICNVNHKPSNTKVDKRVEPKSCDKTHTCPPCPAPEAAGG PSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPREEQY NSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREPQVYTLPPSRE EMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKSR WQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
SEQ ID NO150	Heavy Chain	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSSYINWVRQAPGQGLEWMGNINPATGHA DYAQKFQGRVTMTRDTSISTAYMELSRLRSDDTAVYYCARGGWFDYWGQGTLVTVSSA STKGPSVFPLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSS GLYSLSSVVTVPSSSLGTQTYICNVNHKPSNTKVDKRVEPKSCDKTHTCPPCPAPEAAGG PSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPREEQY NSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREPQVYTLPPSRE EMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKSR WQQGNVFSCSVMHEALHNHYTQKSLSLSPGK
SEQ ID NO151	Light Chain	QSALTQPASVSGSPGQSITISCTGTSSDVGSYNYVNWYQQHPGKAPKLMIYGVSKRPSG VSNRFSGSKSGNTASLTISGLQAEDEADYYCGTFAGGSYYGVFGGGTKLTVLGQPKAAP SVTLFPPSSEELQANKATLVCLISDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYA ASSYLSLTPEQWKSHRSYSCQVTHEGSTVEKTVAPTECS
SEQ ID NO152	Light Chain	QSALTQPASVSGSPGQSITISCTGTSSDVGSYNYVNWYQQHPGKAPKLMIYGVSKRPSG VSNRFSGSKSGNTASLTISGLQAEDEADYYCGTFAGGSYYGVFGGGTKLTVLGQPKAAP SVTLFPPSSEELQANKATLVCLISDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYA ASSYLSLTPEQWKSHRSYSCQVTHEGSTVEKTVAPTECS
SEQ ID NO153	Light Chain	QSALTQPASVSGSPGQSITISCTGTSSDVGSYNYVNWYQQHPGKAPKLMIYGVSKRPSG VSNRFSGSKSGNTASLTISGLQAEDEADYYCGTFAGGSYYGVFGGGTKLTVLGQPKAAP SVTLFPPSSEELQANKATLVCLISDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYA ASSYLSLTPEQWKSHRSYSCQVTHEGSTVEKTVAPTECS
SEQ ID NO154	Light Chain	QSALTQPASVSGSPGQSITISCTGTSSDVGSYNYVNWYQQHPGKAPKLMIYGVSKRPSG VSNRFSGSKSGNTASLTISGLQAEDEADYYCGTFAGGSYYGVFGGGTKLTVLGQPKAAP SVTLFPPSSEELQANKATLVCLISDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYA ASSYLSLTPEQWKSHRSYSCQVTHEGSTVEKTVAPTECS
SEQ ID NO155	Light Chain	QSALTQPASVSGSPGQSITISCTGTSSDVGSYNYVNWYQQHPGKAPKLMIYGVSKRPSG VSNRFSGSKSGNTASLTISGLQAEDEADYYCGTFAGGSYYGVFGGGTKLTVLGQPKAAP SVTLFPPSSEELQANKATLVCLISDFYPGAVTVAWKADSSPVKAGVETTTPSKQSNNKYA ASSYLSLTPEQWKSHRSYSCQVTHEGSTVEKTVAPTECS
SEQ ID NO156	Heavy Chain	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSSYINWVRQAPGQGLEWMGTINPVSGST SYAQKFQGRVTMTRDTSISTAYMELSRLRSDDTAVYYCARGGWFDYWGQGTLVTVSSA STKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSG LYSLSSVVTVPSSNFGTQTYTCNVDHKPSNTKVDKTVERKCCVECPPCPAPPVAGPSVFL FPPKPKDTLMISRTPEVTCVVVDVSHEDPEVQFNWYVDGVEVHNAKTKPREEQFNSTFR VVSVLTVVHQDWLNGKEYKCKVSNKGLPAPIEKTISKTKGQPREPQVYTLPPSREEMTKN QVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPMLDSDGSFFLYSKLTVDKSRWQQG

		NVFSCSVMHEALHNHYTQKSLSLSPGK
SEQ ID NO157	Heavy Chain	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSSYINWVRQAPGQGLEWMGQINAASGMT RYAQKFQGRVTMTRDTSISTAYMELSRLRSDDTAVYYCARGGWFDYWGQGTLVTVSSA STKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSG LYSLSSVVTVPSSNFGTQTYTCNVDHKPSNTKVDKTVERKCCVECPPCPAPPVAGPSVFL FPPKPKDTLMISRTPEVTCVVVDVSHEDPEVQFNWYVDGVEVHNAKTKPREEQFNSTFR VVSVLTVVHQDWLNGKEYKCKVSNKGLPAPIEKTISKTKGQPREPQVYTLPPSREEMTKN QVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPMLDSDGSFFLYSKLTVDKSRWQQG NVFSCSVMHEALHNHYTQKSLSLSPGK
SEQ ID NO158	Heavy Chain	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSSYINWVRQAPGQGLEWMGNINAAAGITL YAQKFQGRVTMTRDTSISTAYMELSRLRSDDTAVYYCARGGWFDYWGQGTLVTVSSAS TKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGL YSLSSVVTVPSSNFGTQTYTCNVDHKPSNTKVDKTVERKCCVECPPCPAPPVAGPSVFLF PPKPKDTLMISRTPEVTCVVVDVSHEDPEVQFNWYVDGVEVHNAKTKPREEQFNSTFRV VSVLTVVHQDWLNGKEYKCKVSNKGLPAPIEKTISKTKGQPREPQVYTLPPSREEMTKNQ VSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPMLDSDGSFFLYSKLTVDKSRWQQGN VFSCSVMHEALHNHYTQKSLSLSPGK
SEQ ID NO159	Heavy Chain	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSSYINWVRQAPGQGLEWMGGINPPAGTT SYAQKFQGRVTMTRDTSISTAYMELSRLRSDDTAVYYCARGGWFDYWGQGTLVTVSSA STKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSG LYSLSSVVTVPSSNFGTQTYTCNVDHKPSNTKVDKTVERKCCVECPPCPAPPVAGPSVFL FPPKPKDTLMISRTPEVTCVVVDVSHEDPEVQFNWYVDGVEVHNAKTKPREEQFNSTFR VVSVLTVVHQDWLNGKEYKCKVSNKGLPAPIEKTISKTKGQPREPQVYTLPPSREEMTKN QVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPMLDSDGSFFLYSKLTVDKSRWQQG NVFSCSVMHEALHNHYTQKSLSLSPGK
SEQ ID NO160	Heavy Chain	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSSYINWVRQAPGQGLEWMGNINPATGHA DYAQKFQGRVTMTRDTSISTAYMELSRLRSDDTAVYYCARGGWFDYWGQGTLVTVSSA STKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSG LYSLSSVVTVPSSNFGTQTYTCNVDHKPSNTKVDKTVERKCCVECPPCPAPPVAGPSVFL FPPKPKDTLMISRTPEVTCVVVDVSHEDPEVQFNWYVDGVEVHNAKTKPREEQFNSTFR VVSVLTVVHQDWLNGKEYKCKVSNKGLPAPIEKTISKTKGQPREPQVYTLPPSREEMTKN QVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPMLDSDGSFFLYSKLTVDKSRWQQG NVFSCSVMHEALHNHYTQKSLSLSPGK
SEQ ID NO161	DNA Light Chain	CAGAGCGCCCTGACCCAGCCCGCCAGCGTGTCCGGCAGCCCAGGCCAGTCTATCAC AATCAGCTGCACCGGCACCTCCAGCGACGTGGGCAGCTACAACTACGTGAACTGGTA TCAGCAGCACCCCGGCAAGGCCCCCAAGCTGATGATCTACGGCGTGAGCAAGAGGC CCAGCGGCGTGTCCAACAGGTTCAGCGGCAGCAAGAGCGGCAACACCGCCAGCCTG ACAATCAGTGGGCTGCAGGCTGAGGACGAGGCCGACTACTACTGCGGCACCTTTGC CGGCGGATCATACTACGGCGTGTTCGGCGGAGGGACCAAGCTGACCGTGCTGGGCC AGCCTAAGGCTGCCCCCAGCGTGACCCTGTTCCCCCCCAGCAGCGAGGAGCTGCAG GCCAACAAGGCCACCCTGGTGTGCCTGATCAGCGACTTCTACCCAGGCGCCGTGAC CGTGGCCTGGAAGGCCGACAGCAGCCCCGTGAAGGCCGGCGTGGAGACCACCACC CCCAGCAAGCAGAGCAACAACAAGTACGCCGCCAGCAGCTACCTGAGCCTGACCCC CGAGCAGTGGAAGAGCCACAGGTCCTACAGCTGCCAGGTGACCCACGAGGGCAGCA CCGTGGAAAAGACCGTGGCCCCAACCGAGTGCAGC
SEQ ID NO162	DNA Light Chain	CAGAGCGCCCTGACCCAGCCCGCCAGCGTGTCCGGCAGCCCAGGCCAGTCTATCAC AATCAGCTGCACCGGCACCTCCAGCGACGTGGGCAGCTACAACTACGTGAACTGGTA TCAGCAGCACCCCGGCAAGGCCCCCAAGCTGATGATCTACGGCGTGAGCAAGAGGC CCAGCGGCGTGTCCAACAGGTTCAGCGGCAGCAAGAGCGGCAACACCGCCAGCCTG

		ACAATCAGTGGGCTGCAGGCTGAGGACGAGGCCGACTACTACTGCGGCACCTTTGC CGGCGGATCATACTACGGCGTGTTCGGCGGAGGGACCAAGCTGACCGTGCTGGGCC AGCCTAAGGCTGCCCCCAGCGTGACCCTGTTCCCCCCCAGCAGCGAGGAGCTGCAG GCCAACAAGGCCACCCTGGTGTGCCTGATCAGCGACTTCTACCCAGGCGCCGTGAC CGTGGCCTGGAAGGCCGACAGCAGCCCCGTGAAGGCCGGCGTGGAGACCACCACC CCCAGCAAGCAGAGCAACAACAAGTACGCCGCCAGCAGCTACCTGAGCCTGACCCC CGAGCAGTGGAAGAGCCACAGGTCCTACAGCTGCCAGGTGACCCACGAGGGCAGCA CCGTGGAAAAGACCGTGGCCCCAACCGAGTGCAGC
SEQ ID NO163	DNA Light Chain	CAGAGCGCACTGACCCAGCCAGCTTCAGTGAGCGGCTCACCAGGTCAGAGCATTAC CATCTCGTGTACGGGTACTAGCAGCGATGTTGGTTCTTATAATTATGTGAATTGGTAC CAGCAGCATCCCGGGAAGGCGCCGAAACTTATGATTTATGGTGTTTCTAAGCGTCCC TCAGGCGTGAGCAACCGTTTTAGCGGATCCAAAAGCGGCAACACCGCGAGCCTGAC CATTAGCGGCCTGCAAGCGGAAGACGAAGCGGATTATTATTGCGGTACTTTTGCTGG TGGTTCTTATTATGGTGTGTTTGGCGGCGGCACGAAGTTAACCGTCCTAGGTCAGCC CAAGGCTGCCCCCTCGGTCACTCTGTTCCCGCCCTCCTCTGAGGAGCTTCAAGCCAA CAAGGCCACACTGGTGTGTCTCATAAGTGACTTCTACCCGGGAGCCGTGACAGTGGC CTGGAAGGCAGATAGCAGCCCCGTCAAGGCGGGAGTGGAGACCACCACACCCTCCA AACAAAGCAACAACAAGTACGCGGCCAGCAGCTATCTGAGCCTGACGCCTGAGCAGT GGAAGTCCCACAGAAGCTACAGCTGCCAGGTCACGCATGAAGGGAGCACCGTGGAG AAGACAGTGGCCCCTACAGAATGTTCA
SEQ ID NO164	DNA Light Chain	CAGAGCGCACTGACCCAGCCAGCTTCAGTGAGCGGCTCACCAGGTCAGAGCATTAC CATCTCGTGTACGGGTACTAGCAGCGATGTTGGTTCTTATAATTATGTGAATTGGTAC CAGCAGCATCCCGGGAAGGCGCCGAAACTTATGATTTATGGTGTTTCTAAGCGTCCC TCAGGCGTGAGCAACCGTTTTAGCGGATCCAAAAGCGGCAACACCGCGAGCCTGAC CATTAGCGGCCTGCAAGCGGAAGACGAAGCGGATTATTATTGCGGTACTTTTGCTGG TGGTTCTTATTATGGTGTGTTTGGCGGCGGCACGAAGTTAACCGTCCTAGGTCAGCC CAAGGCTGCCCCCTCGGTCACTCTGTTCCCGCCCTCCTCTGAGGAGCTTCAAGCCAA CAAGGCCACACTGGTGTGTCTCATAAGTGACTTCTACCCGGGAGCCGTGACAGTGGC CTGGAAGGCAGATAGCAGCCCCGTCAAGGCGGGAGTGGAGACCACCACACCCTCCA AACAAAGCAACAACAAGTACGCGGCCAGCAGCTATCTGAGCCTGACGCCTGAGCAGT GGAAGTCCCACAGAAGCTACAGCTGCCAGGTCACGCATGAAGGGAGCACCGTGGAG AAGACAGTGGCCCCTACAGAATGTTCA
SEQ ID NO165	DNA Light Chain	CAGAGCGCACTGACCCAGCCAGCTTCAGTGAGCGGCTCACCAGGTCAGAGCATTAC CATCTCGTGTACGGGTACTAGCAGCGATGTTGGTTCTTATAATTATGTGAATTGGTAC CAGCAGCATCCCGGGAAGGCGCCGAAACTTATGATTTATGGTGTTTCTAAGCGTCCC TCAGGCGTGAGCAACCGTTTTAGCGGATCCAAAAGCGGCAACACCGCGAGCCTGAC CATTAGCGGCCTGCAAGCGGAAGACGAAGCGGATTATTATTGCGGTACTTTTGCTGG TGGTTCTTATTATGGTGTGTTTGGCGGCGGCACGAAGTTAACCGTCCTAGGTCAGCC CAAGGCTGCCCCCTCGGTCACTCTGTTCCCGCCCTCCTCTGAGGAGCTTCAAGCCAA CAAGGCCACACTGGTGTGTCTCATAAGTGACTTCTACCCGGGAGCCGTGACAGTGGC CTGGAAGGCAGATAGCAGCCCCGTCAAGGCGGGAGTGGAGACCACCACACCCTCCA AACAAAGCAACAACAAGTACGCGGCCAGCAGCTATCTGAGCCTGACGCCTGAGCAGT GGAAGTCCCACAGAAGCTACAGCTGCCAGGTCACGCATGAAGGGAGCACCGTGGAG AAGACAGTGGCCCCTACAGAATGTTCA
SEQ ID NO166	DNA Heavy Chain	CAGGTGCAGCTGGTGCAGAGCGGAGCTGAGGTGAAGAAGCCAGGCGCCAGCGTCAA GGTGTCCTGCAAGGCCAGCGGCTACACCTTCACCAGCAGCTACATCAACTGGGTCCG CCAGGCTCCTGGGCAGGGACTGGAGTGGATGGGCACCATCAACCCCGTGTCCGGCA GCACCAGCTACGCCCAGAAGTTCCAGGGCAGAGTCACCATGACCAGGGACACCAGC ATCAGCACCGCCTACATGGAGCTGTCCAGGCTGAGAAGCGACGACACCGCCGTGTA CTACTGCGCCAGGGGCGGCTGGTTCGACTACTGGGGCCAGGGCACCCTGGTGACCG TGTCCTCAGCTAGCACCAAGGGCCCCAGCGTGTTCCCCCTGGCCCCCAGCAGCAAG

		AGCACCTCCGGCGGCACAGCCGCCCTGGGCTGCCTGGTGAAGGACTACTTCCCCGA GCCCGTGACCGTGTCCTGGAACAGCGGAGCCCTGACCAGCGGCGTGCACACCTTCC CCGCCGTGCTGCAGAGCAGCGGCCTGTACAGCCTGTCCAGCGTGGTGACAGTGCCC AGCAGCAGCCTGGGCACCCAGACCTACATCTGCAACGTGAACCACAAGCCCAGCAAC ACCAAGGTGGACAAGAGAGTGGAGCCCAAGAGCTGCGACAAGACCCACACCTGCCC CCCCTGCCCAGCCCCCGAAGCTGCAGGCGGCCCTTCCGTGTTCCTGTTCCCCCCCA AGCCCAAGGACACCCTGATGATCAGCAGGACCCCCGAGGTGACCTGCGTGGTGGTG GACGTGAGCCACGAGGACCCAGAGGTGAAGTTCAACTGGTACGTGGACGGCGTGGA GGTGCACAACGCCAAGACCAAGCCCAGAGAGGAGCAGTACAACAGCACCTACAGGG TGGTGTCCGTGCTGACCGTGCTGCACCAGGACTGGCTGAACGGCAAAGAATACAAGT GCAAGGTCTCCAACAAGGCCCTGCCTGCCCCCATCGAAAAGACCATCAGCAAGGCCA AGGGCCAGCCACGGGAGCCCCAGGTGTACACCCTGCCCCCTTCTCGGGAGGAGATG ACCAAGAACCAGGTGTCCCTGACCTGTCTGGTGAAGGGCTTCTACCCCAGCGACATC GCCGTGGAGTGGGAGAGCAACGGCCAGCCCGAGAACAACTACAAGACCACCCCCCC AGTGCTGGACAGCGACGGCAGCTTCTTCCTGTACAGCAAGCTGACCGTGGACAAGAG CAGGTGGCAGCAGGGCAACGTGTTCAGCTGCAGCGTGATGCACGAGGCCCTGCACA ACCACTACACCCAGAAGAGCCTGAGCCTGTCACCCGGCAAG
SEQ ID NO167	DNA Heavy Chain	CAGGTGCAGCTGGTGCAGAGCGGAGCTGAGGTGAAGAAGCCAGGCGCCAGCGTCAA GGTGTCCTGCAAGGCCAGCGGCTACACCTTCACCAGCAGCTACATCAACTGGGTGCG CCAGGCTCCAGGGCAGGGACTGGAGTGGATGGGCCAGATCAACGCCGCCAGCGGC ATGACCAGATACGCCCAGAAGTTCCAGGGCAGAGTCACAATGACCAGGGACACCTCT ATCAGCACCGCCTACATGGAGCTGTCCAGGCTGAGAAGCGACGACACCGCCGTGTA CTACTGCGCCAGGGGCGGCTGGTTCGACTACTGGGGCCAGGGCACCCTGGTGACCG TGTCCTCAGCTAGCACCAAGGGCCCCAGCGTGTTCCCCCTGGCCCCCAGCAGCAAG AGCACCTCCGGCGGCACAGCCGCCCTGGGCTGCCTGGTGAAGGACTACTTCCCCGA GCCCGTGACCGTGTCCTGGAACAGCGGAGCCCTGACCAGCGGCGTGCACACCTTCC CCGCCGTGCTGCAGAGCAGCGGCCTGTACAGCCTGTCCAGCGTGGTGACAGTGCCC AGCAGCAGCCTGGGCACCCAGACCTACATCTGCAACGTGAACCACAAGCCCAGCAAC ACCAAGGTGGACAAGAGAGTGGAGCCCAAGAGCTGCGACAAGACCCACACCTGCCC CCCCTGCCCAGCCCCCGAAGCTGCAGGCGGCCCTTCCGTGTTCCTGTTCCCCCCCA AGCCCAAGGACACCCTGATGATCAGCAGGACCCCCGAGGTGACCTGCGTGGTGGTG GACGTGAGCCACGAGGACCCAGAGGTGAAGTTCAACTGGTACGTGGACGGCGTGGA GGTGCACAACGCCAAGACCAAGCCCAGAGAGGAGCAGTACAACAGCACCTACAGGG TGGTGTCCGTGCTGACCGTGCTGCACCAGGACTGGCTGAACGGCAAAGAATACAAGT GCAAGGTCTCCAACAAGGCCCTGCCTGCCCCCATCGAAAAGACCATCAGCAAGGCCA AGGGCCAGCCACGGGAGCCCCAGGTGTACACCCTGCCCCCTTCTCGGGAGGAGATG ACCAAGAACCAGGTGTCCCTGACCTGTCTGGTGAAGGGCTTCTACCCCAGCGACATC GCCGTGGAGTGGGAGAGCAACGGCCAGCCCGAGAACAACTACAAGACCACCCCCCC AGTGCTGGACAGCGACGGCAGCTTCTTCCTGTACAGCAAGCTGACCGTGGACAAGAG CAGGTGGCAGCAGGGCAACGTGTTCAGCTGCAGCGTGATGCACGAGGCCCTGCACA ACCACTACACCCAGAAGAGCCTGAGCCTGTCACCCGGCAAG
SEQ ID NO168	DNA Heavy Chain	CAGGTGCAATTGGTTCAGAGCGGCGCGGAAGTGAAAAAACCGGGGCGCGAGCGTGAA AGTGAGCTGCAAAGCCTCCGGATATACCTTTACTTCTTCTTATATTAATTGGGTCCGCC AAGCCCCTGGGCAGGGTCTCGAGTGGATGGGCAATATTAATGCTGCTGCTGGTATTA CTCTTTATGCTCAGAAGTTTCAGGGTCGGGTCACCATGACCCGTGATACCAGCATTAG CACCGCGTATATGGAACTGAGCCGCCTGCGTAGCGATGATACGGCCGTGTATTATTG CGCGCGTGGTGGTTGGTTTGATTATTGGGGCCAAGGCACCCTGGTGACGGTTAGCTC AGCCTCCACCAAGGGTCCATCGGTCTTCCCCCTGGCACCCTCCTCCAAGAGCACCTC TGGGGGCACAGCGGCCCTGGGCTGCCTGGTCAAGGACTACTTCCCCGAACCGGTGA CGGTGTCGTGGAACTCAGGCGCCCTGACCAGCGGCGTGCACACCTTCCCGGCTGTC CTACAGTCCTCAGGACTCTACTCCCTCAGCAGCGTGGTGACCGTGCCCTCCAGCAGC

		TTGGGCACCCAGACCTACATCTGCAACGTGAATCACAAGCCCAGCAACACCAAGGTG GACAAGAGAGTTGAGCCCAAATCTTGTGACAAAACTCACACATGCCCACCGTGCCCA GCACCTGAAGCAGCGGGGGGACCGTCAGTCTTCCTCTTCCCCCCAAAACCCAAGGA CACCCTCATGATCTCCCGGACCCCTGAGGTCACATGCGTGGTGGTGGACGTGAGCC ACGAAGACCCTGAGGTCAAGTTCAACTGGTACGTGGACGGCGTGGAGGTGCATAATG CCAAGACAAAGCCGCGGGAGGAGCAGTACAACAGCACGTACCGGGTGGTCAGCGTC CTCACCGTCCTGCACCAGGACTGGCTGAATGGCAAGGAGTACAAGTGCAAGGTCTCC AACAAAGCCCTCCCAGCCCCCATCGAGAAAACCATCTCCAAAGCCAAAGGGCAGCCC CGAGAACCACAGGTGTACACCCTGCCCCCATCCCGGGAGGAGATGACCAAGAACCA GGTCAGCCTGACCTGCCTGGTCAAAGGCTTCTATCCCAGCGACATCGCCGTGGAGTG GGAGAGCAATGGGCAGCCGGAGAACAACTACAAGACCACGCCTCCCGTGCTGGACT CCGACGGCTCCTTCTTCCTCTACAGCAAGCTCACCGTGGACAAGAGCAGGTGGCAGC AGGGGAACGTCTTCTCATGCTCCGTGATGCATGAGGCTCTGCACAACCACTACACGC AGAAGAGCCTCTCCCTGTCTCCGGGTAAA
SEQ ID NO169	DNA Heavy Chain	CAGGTGCAATTGGTTCAGAGCGGCGCGGAAGTGAAAAAACCGGGCGCGAGCGTGAA AGTGAGCTGCAAAGCCTCCGGATATACCTTTACTTCTTCTTATATTAATTGGGTCCGCC AAGCCCCTGGGCAGGGTCTCGAGTGGATGGGCGGTATTAATCCTCCTGCTGGTACTA CTTCTTATGCTCAGAAGTTTCAGGGTCGGGTCACCATGACCCGTGATACCAGCATTAG CACCGCGTATATGGAACTGAGCCGCCTGCGTAGCGATGATACGGCCGTGTATTATTG CGCGCGTGGTGGTTGGTTTGATTATTGGGGCCAAGGCACCCTGGTGACGGTTAGCTC AGCCTCCACCAAGGGTCCATCGGTCTTCCCCCTGGCACCCTCCTCCAAGAGCACCTC TGGGGGCACAGCGGCCCTGGGCTGCCTGGTCAAGGACTACTTCCCCGAACCGGTGA CGGTGTCGTGGAACTCAGGCGCCCTGACCAGCGGCGTGCACACCTTCCCGGCTGTC CTACAGTCCTCAGGACTCTACTCCCTCAGCAGCGTGGTGACCGTGCCCTCCAGCAGC TTGGGCACCCAGACCTACATCTGCAACGTGAATCACAAGCCCAGCAACACCAAGGTG GACAAGAGAGTTGAGCCCAAATCTTGTGACAAAACTCACACATGCCCACCGTGCCCA GCACCTGAAGCAGCGGGGGGACCGTCAGTCTTCCTCTTCCCCCCAAAACCCAAGGA CACCCTCATGATCTCCCGGACCCCTGAGGTCACATGCGTGGTGGTGGACGTGAGCC ACGAAGACCCTGAGGTCAAGTTCAACTGGTACGTGGACGGCGTGGAGGTGCATAATG CCAAGACAAAGCCGCGGGAGGAGCAGTACAACAGCACGTACCGGGTGGTCAGCGTC CTCACCGTCCTGCACCAGGACTGGCTGAATGGCAAGGAGTACAAGTGCAAGGTCTCC AACAAAGCCCTCCCAGCCCCCATCGAGAAAACCATCTCCAAAGCCAAAGGGCAGCCC CGAGAACCACAGGTGTACACCCTGCCCCCATCCCGGGAGGAGATGACCAAGAACCA GGTCAGCCTGACCTGCCTGGTCAAAGGCTTCTATCCCAGCGACATCGCCGTGGAGTG GGAGAGCAATGGGCAGCCGGAGAACAACTACAAGACCACGCCTCCCGTGCTGGACT CCGACGGCTCCTTCTTCCTCTACAGCAAGCTCACCGTGGACAAGAGCAGGTGGCAGC AGGGGAACGTCTTCTCATGCTCCGTGATGCATGAGGCTCTGCACAACCACTACACGC AGAAGAGCCTCTCCCTGTCTCCGGGTAAA
SEQ ID NO170	DNA Heavy Chain	CAGGTGCAATTGGTTCAGAGCGGCGCGGAAGTGAAAAAACCGGGCGCGAGCGTGAA AGTGAGCTGCAAAGCCTCCGGATATACCTTTACTTCTTCTTATATTAATTGGGTCCGCC AAGCCCCTGGGCAGGGTCTCGAGTGGATGGGCAATATTAATCCTGCTACTGGTCATG CTGATTATGCTCAGAAGTTTCAGGGTCGGGTGACCATGACCCGTGATACCAGCATTA GCACCGCGTATATGGAACTGAGCCGCCTGCGTAGCGATGATACGGCCGTGTATTATT GCGCGCGTGGTGGTTGGTTTGATTATTGGGGCCAAGGCACCCTGGTGACGGTTAGCT CAGCCTCCACCAAGGGTCCATCGGTCTTCCCCCTGGCACCCTCCTCCAAGAGCACCT CTGGGGGCACAGCGGCCCTGGGCTGCCTGGTCAAGGACTACTTCCCCGAACCGGTG ACGGTGTCGTGGAACTCAGGCGCCCTGACCAGCGGCGTGCACACCTTCCCGGCTGT CCTACAGTCCTCAGGACTCTACTCCCTCAGCAGCGTGGTGACCGTGCCCTCCAGCAG CTTGGGCACCCAGACCTACATCTGCAACGTGAATCACAAGCCCAGCAACACCAAGGT GGACAAGAGAGTTGAGCCCAAATCTTGTGACAAAACTCACACATGCCCACCGTGCCC AGCACCTGAAGCAGCGGGGGGACCGTCAGTCTTCCTCTTCCCCCCAAAACCCAAGGA

		CACCCTCATGATCTCCCGGACCCCTGAGGTCACATGCGTGGTGGTGGACGTGAGCC ACGAAGACCCTGAGGTCAAGTTCAACTGGTACGTGGACGGCGTGGAGGTGCATAATG CCAAGACAAAGCCGCGGGAGGAGCAGTACAACAGCACGTACCGGGTGGTCAGCGTC CTCACCGTCCTGCACCAGGACTGGCTGAATGGCAAGGAGTACAAGTGCAAGGTCTCC AACAAAGCCCTCCCAGCCCCCATCGAGAAAACCATCTCCAAAGCCAAAGGGCAGCCC CGAGAACCACAGGTGTACACCCTGCCCCCATCCCGGGAGGAGATGACCAAGAACCA GGTCAGCCTGACCTGCCTGGTCAAAGGCTTCTATCCCAGCGACATCGCCGTGGAGTG GGAGAGCAATGGGCAGCCGGAGAACAACTACAAGACCACGCCTCCCGTGCTGGACT CCGACGGCTCCTTCTTCCTCTACAGCAAGCTCACCGTGGACAAGAGCAGGTGGCAGC AGGGGAACGTCTTCTCATGCTCCGTGATGCATGAGGCTCTGCACAACCACTACACGC AGAAGAGCCTCTCCCTGTCTCCGGGTAAA
SEQ ID NO171	DNA Light Chain	CAGAGCGCCCTGACCCAGCCCGCCAGCGTGTCCGGCAGCCCAGGCCAGTCTATCAC AATCAGCTGCACCGGCACCTCCAGCGACGTGGGCAGCTACAACTACGTGAACTGGTA TCAGCAGCACCCCGGCAAGGCCCCCAAGCTGATGATCTACGGCGTGAGCAAGAGGC CCAGCGGCGTGTCCAACAGGTTCAGCGGCAGCAAGAGCGGCAACACCGCCAGCCTG ACAATCAGTGGGCTGCAGGCTGAGGACGAGGCCGACTACTACTGCGGCACCTTTGC CGGCGGATCATACTACGGCGTGTTCGGCGGAGGGACCAAGCTGACCGTGCTGGGCC AGCCTAAGGCTGCCCCCAGCGTGACCCTGTTCCCCCCCAGCAGCGAGGAGCTGCAG GCCAACAAGGCCACCCTGGTGTGCCTGATCAGCGACTTCTACCCAGGCGCCGTGAC CGTGGCCTGGAAGGCCGACAGCAGCCCCGTGAAGGCCGGCGTGGAGACCACCACC CCCAGCAAGCAGAGCAACAACAAGTACGCCGCCAGCAGCTACCTGAGCCTGACCCC CGAGCAGTGGAAGAGCCACAGGTCCTACAGCTGCCAGGTGACCCACGAGGGCAGCA CCGTGGAAAAGACCGTGGCCCCAACCGAGTGCAGC
SEQ ID NO172	DNA Light Chain	CAGAGCGCCCTGACCCAGCCCGCCAGCGTGTCCGGCAGCCCAGGCCAGTCTATCAC AATCAGCTGCACCGGCACCTCCAGCGACGTGGGCAGCTACAACTACGTGAACTGGTA TCAGCAGCACCCCGGCAAGGCCCCCAAGCTGATGATCTACGGCGTGAGCAAGAGGC CCAGCGGCGTGTCCAACAGGTTCAGCGGCAGCAAGAGCGGCAACACCGCCAGCCTG ACAATCAGTGGGCTGCAGGCTGAGGACGAGGCCGACTACTACTGCGGCACCTTTGC CGGCGGATCATACTACGGCGTGTTCGGCGGAGGGACCAAGCTGACCGTGCTGGGCC AGCCTAAGGCTGCCCCCAGCGTGACCCTGTTCCCCCCCAGCAGCGAGGAGCTGCAG GCCAACAAGGCCACCCTGGTGTGCCTGATCAGCGACTTCTACCCAGGCGCCGTGAC CGTGGCCTGGAAGGCCGACAGCAGCCCCGTGAAGGCCGGCGTGGAGACCACCACC CCCAGCAAGCAGAGCAACAACAAGTACGCCGCCAGCAGCTACCTGAGCCTGACCCC CGAGCAGTGGAAGAGCCACAGGTCCTACAGCTGCCAGGTGACCCACGAGGGCAGCA CCGTGGAAAAGACCGTGGCCCCAACCGAGTGCAGC
SEQ ID NO173	DNA Light Chain	CAGAGCGCACTGACCCAGCCAGCTTCAGTGAGCGGCTCACCAGGTCAGAGCATTAC CATCTCGTGTACGGGTACTAGCAGCGATGTTGGTTCTTATAATTATGTGAATTGGTAC CAGCAGCATCCCGGGAAGGCGCCGAAACTTATGATTTATGGTGTTTCTAAGCGTCCC TCAGGCGTGAGCAACCGTTTTAGCGGATCCAAAAGCGGCAACACCGCGAGCCTGAC CATTAGCGGCCTGCAAGCGGAAGACGAAGCGGATTATTATTGCGGTACTTTTGCTGG TGGTTCTTATTATGGTGTGTTTGGCGGCGGCACGAAGTTAACCGTCCTAGGTCAGCC CAAGGCTGCCCCCTCGGTCACTCTGTTCCCGCCCTCCTCTGAGGAGCTTCAAGCCAA CAAGGCCACACTGGTGTGTCTCATAAGTGACTTCTACCCGGGAGCCGTGACAGTGGC CTGGAAGGCAGATAGCAGCCCCGTCAAGGCGGGAGTGGAGACCACCACACCCTCCA AACAAAGCAACAACAAGTACGCGGCCAGCAGCTATCTGAGCCTGACGCCTGAGCAGT GGAAGTCCCACAGAAGCTACAGCTGCCAGGTCACGCATGAAGGGAGCACCGTGGAG AAGACAGTGGCCCCTACAGAATGTTCA
SEQ ID NO174	DNA Light Chain	CAGAGCGCACTGACCCAGCCAGCTTCAGTGAGCGGCTCACCAGGTCAGAGCATTAC CATCTCGTGTACGGGTACTAGCAGCGATGTTGGTTCTTATAATTATGTGAATTGGTAC CAGCAGCATCCCGGGAAGGCGCCGAAACTTATGATTTATGGTGTTTCTAAGCGTCCC TCAGGCGTGAGCAACCGTTTTAGCGGATCCAAAAGCGGCAACACCGCGAGCCTGAC

		CATTAGCGGCCTGCAAGCGGAAGACGAAGCGGATTATTATTGCGGTACTTTTGCTGG TGGTTCTTATTATGGTGTGTTTGGCGGCGGCACGAAGTTAACCGTCCTAGGTCAGCC CAAGGCTGCCCCCTCGGTCACTCTGTTCCCGCCCTCCTCTGAGGAGCTTCAAGCCAA CAAGGCCACACTGGTGTGTCTCATAAGTGACTTCTACCCGGGAGCCGTGACAGTGGC CTGGAAGGCAGATAGCAGCCCCGTCAAGGCGGGAGTGGAGACCACCACACCCTCCA AACAAAGCAACAACAAGTACGCGGCCAGCAGCTATCTGAGCCTGACGCCTGAGCAGT GGAAGTCCCACAGAAGCTACAGCTGCCAGGTCACGCATGAAGGGAGCACCGTGGAG AAGACAGTGGCCCCTACAGAATGTTCA
SEQ ID NO175	DNA Light Chain	CAGAGCGCACTGACCCAGCCAGCTTCAGTGAGCGGCTCACCAGGTCAGAGCATTAC CATCTCGTGTACGGGTACTAGCAGCGATGTTGGTTCTTATAATTATGTGAATTGGTAC CAGCAGCATCCCGGGAAGGCGCCGAAACTTATGATTTATGGTGTTTCTAAGCGTCCC TCAGGCGTGAGCAACCGTTTTAGCGGATCCAAAAGCGGCAACACCGCGAGCCTGAC CATTAGCGGCCTGCAAGCGGAAGACGAAGCGGATTATTATTGCGGTACTTTTGCTGG TGGTTCTTATTATGGTGTGTTTGGCGGCGGCACGAAGTTAACCGTCCTAGGTCAGCC CAAGGCTGCCCCCTCGGTCACTCTGTTCCCGCCCTCCTCTGAGGAGCTTCAAGCCAA CAAGGCCACACTGGTGTGTCTCATAAGTGACTTCTACCCGGGAGCCGTGACAGTGGC CTGGAAGGCAGATAGCAGCCCCGTCAAGGCGGGAGTGGAGACCACCACACCCTCCA AACAAAGCAACAACAAGTACGCGGCCAGCAGCTATCTGAGCCTGACGCCTGAGCAGT GGAAGTCCCACAGAAGCTACAGCTGCCAGGTCACGCATGAAGGGAGCACCGTGGAG AAGACAGTGGCCCCTACAGAATGTTCA
SEQ ID NO176	DNA Heavy Chain	CAGGTGCAGCTGGTGCAGAGCGGAGCTGAGGTGAAGAAGCCAGGCGCCAGCGTCAA GGTGTCCTGCAAGGCCAGCGGCTACACCTTCACCAGCAGCTACATCAACTGGGTCCG CCAGGCTCCTGGGCAGGGACTGGAGTGGATGGGCACCATCAACCCCGTGTCCGGCA GCACCAGCTACGCCCAGAAGTTCCAGGGCAGAGTCACCATGACCAGGGACACCAGC ATCAGCACCGCCTACATGGAGCTGTCCAGGCTGAGAAGCGACGACACCGCCGTGTA CTACTGCGCCAGGGGCGGCTGGTTCGACTACTGGGGCCAGGGCACCCTGGTGACCG TGTCCTCAGCTAGCACCAAGGGCCCCAGCGTGTTCCCCCTGGCCCCCTGCAGCAGA AGCACCAGCGAGAGCACAGCCGCCCTGGGCTGCCTGGTGAAGGACTACTTCCCCGA GCCAGTGACCGTGTCCTGGAACAGCGGAGCCCTGACCAGCGGCGTGCACACCTTCC CCGCCGTGCTGCAGAGCAGCGGCCTGTACAGCCTGTCCAGCGTGGTGACCGTGCCC AGCAGCAACTTCGGCACCCAGACCTACACCTGCAACGTGGACCACAAGCCCAGCAAC ACCAAGGTGGACAAGACCGTGGAGAGGAAGTGCTGCGTGGAGTGCCCCCCCTGCCC AGCCCCCCCAGTGGCCGGACCCTCCGTGTTCCTGTTCCCCCCCAAGCCCAAGGACA CCCTGATGATCAGCAGGACCCCCGAGGTGACCTGCGTGGTGGTGGACGTGAGCCAC GAGGACCCAGAGGTGCAGTTCAACTGGTACGTGGACGGCGTGGAGGTGCACAACGC CAAGACCAAGCCCAGAGAGGAACAGTTTAACAGCACCTTCAGGGTGGTGTCCGTGCT GACCGTGGTGCACCAGGACTGGCTGAACGGCAAAGAGTACAAGTGCAAGGTCTCCA ACAAGGGCCTGCCAGCCCCCATCGAGAAAACCATCAGCAAGACCAAGGGCCAGCCA CGGGAGCCCCAGGTGTACACCCTGCCCCCCAGCCGGGAGGAAATGACCAAGAACCA GGTGTCCCTGACCTGTCTGGTGAAGGGCTTCTACCCCAGCGACATCGCCGTGGAGT GGGAGAGCAACGGCCAGCCCGAGAACAACTACAAGACCACCCCCCCCCATGCTGGAC AGCGACGGCAGCTTCTTCCTGTACAGCAAGCTGACAGTGGACAAGAGCAGGTGGCA GCAGGGCAACGTGTTCAGCTGCAGCGTGATGCACGAGGCCCTGCACAACCACTACA CCCAGAAGAGCCTGAGCCTGTCCCCCGGCAAG
SEQ ID NO177	DNA Heavy Chain	CAGGTGCAGCTGGTGCAGAGCGGAGCTGAGGTGAAGAAGCCAGGCGCCAGCGTCAA GGTGTCCTGCAAGGCCAGCGGCTACACCTTCACCAGCAGCTACATCAACTGGGTGCG CCAGGCTCCAGGGCAGGGACTGGAGTGGATGGGCCAGATCAACGCCGCCAGCGGC ATGACCAGATACGCCCAGAAGTTCCAGGGCAGAGTCACAATGACCAGGGACACCTCT ATCAGCACCGCCTACATGGAGCTGTCCAGGCTGAGAAGCGACGACACCGCCGTGTA CTACTGCGCCAGGGGCGGCTGGTTCGACTACTGGGGCCAGGGCACCCTGGTGACCG TGTCCTCAGCTAGCACCAAGGGCCCCAGCGTGTTCCCCCTGGCCCCCTGCAGCAGA

		AGCACCAGCGAGAGCACAGCCGCCCTGGGCTGCCTGGTGAAGGACTACTTCCCCGA GCCAGTGACCGTGTCCTGGAACAGCGGAGCCCTGACCAGCGGCGTGCACACCTTCC CCGCCGTGCTGCAGAGCAGCGGCCTGTACAGCCTGTCCAGCGTGGTGACCGTGCCC AGCAGCAACTTCGGCACCCAGACCTACACCTGCAACGTGGACCACAAGCCCAGCAAC ACCAAGGTGGACAAGACCGTGGAGAGGAAGTGCTGCGTGGAGTGCCCCCCCTGCCC AGCCCCCCCAGTGGCCGGACCCTCCGTGTTCCTGTTCCCCCCCAAGCCCAAGGACA CCCTGATGATCAGCAGGACCCCCGAGGTGACCTGCGTGGTGGTGGACGTGAGCCAC GAGGACCCAGAGGTGCAGTTCAACTGGTACGTGGACGGCGTGGAGGTGCACAACGC CAAGACCAAGCCCAGAGAGGAACAGTTTAACAGCACCTTCAGGGTGGTGTCCGTGCT GACCGTGGTGCACCAGGACTGGCTGAACGGCAAAGAGTACAAGTGCAAGGTCTCCA ACAAGGGCCTGCCAGCCCCCATCGAGAAAACCATCAGCAAGACCAAGGGCCAGCCA CGGGAGCCCCAGGTGTACACCCTGCCCCCCAGCCGGGAGGAAATGACCAAGAACCA GGTGTCCCTGACCTGTCTGGTGAAGGGCTTCTACCCCAGCGACATCGCCGTGGAGT GGGAGAGCAACGGCCAGCCCGAGAACAACTACAAGACCACCCCCCCCCATGCTGGAC AGCGACGGCAGCTTCTTCCTGTACAGCAAGCTGACAGTGGACAAGAGCAGGTGGCA GCAGGGCAACGTGTTCAGCTGCAGCGTGATGCACGAGGCCCTGCACAACCACTACA CCCAGAAGAGCCTGAGCCTGTCCCCCGGCAAG
SEQ ID NO178	DNA Heavy Chain	CAGGTGCAATTGGTTCAGAGCGGCGCGGAAGTGAAAAAACCGGGGCGCGAGCGTGAA AGTGAGCTGCAAAGCCTCCGGATATACCTTTACTTCTTCTTATATTAATTGGGTCCGCC AAGCCCCTGGGCAGGGTCTCGAGTGGATGGGCAATATTAATGCTGCTGCTGGTATTA CTCTTTATGCTCAGAAGTTTCAGGGTCGGGTCACCATGACCCGTGATACCAGCATTAG CACCGCGTATATGGAACTGAGCCGCCTGCGTAGCGATGATACGGCCGTGTATTATTG CGCGCGTGGTGGTTGGTTTGATTATTGGGGCCAAGGCACCCTGGTGACGGTTAGCTC AGCTTCCACCAAGGGCCCCAGCGTGTTCCCCCTGGCCCCCTGCAGCAGAAGCACCA GCGAGAGCACAGCCGCCCTGGGCTGCCTGGTGAAGGACTACTTCCCCGAGCCCGTG ACCGTGAGCTGGAACAGCGGAGCCCTGACCAGCGGCGTGCACACCTTCCCCGCCGT GCTGCAGAGCAGCGGCCTGTACAGCCTGAGCAGCGTGGTGACCGTGCCCAGCAGCA ACTTCGGCACCCAGACCTACACCTGCAACGTGGACCACAAGCCCAGCAACACCAAGG TGGACAAGACCGTGGAGCGGAAGTGCTGCGTGGAGTGCCCCCCCCTGCCCTGCCCCT CCTGTGGCCGGACCCTCCGTGTTCCTGTTCCCCCCCAAGCCCAAGGACACCCTGATG ATCAGCCGGACCCCCGAGGTGACCTGCGTGGTGGTGGACGTGAGCCACGAGGACCC CGAGGTGCAGTTCAACTGGTACGTGGACGGCGTGGAGGTGCACAACGCCAAGACCA AGCCCCGGGAGGAACAGTTCAACAGCACCTTCCGGGTGGTGTCCGTGCTGACCGTG GTGCACCAGGACTGGCTGAACGGCAAAGAATACAAGTGCAAGGTGTCCAACAAGGG CCTGCCTGCCCCCATCGAGAAAACCATCAGCAAGACAAAGGGCCAGCCCAGGGAAC CCCAGGTGTACACCCTGCCCCCCAGCCGGGAGGAAATGACCAAGAACCAGGTGTCC CTGACCTGTCTGGTGAAGGGCTTCTACCCCAGCGACATCGCCGTGGAGTGGGAGAG CAACGGCCAGCCCGAGAACAACTACAAGACCACCCCCCCCCATGCTGGACAGCGACG GCAGCTTCTTCCTGTACAGCAAGCTGACAGTGGACAAGAGCCGGTGGCAGCAGGGC AACGTGTTCAGCTGCAGCGTGATGCACGAGGCCCTGCACAACCACTACACCCAGAAG AGCCTGAGCCTGTCCCCCGGCAAA
SEQ ID NO179	DNA Heavy Chain	CAGGTGCAATTGGTTCAGAGCGGCGCGGAAGTGAAAAAACCGGGGCGCGAGCGTGAA AGTGAGCTGCAAAGCCTCCGGATATACCTTTACTTCTTCTTATATTAATTGGGTCCGCC AAGCCCCTGGGCAGGGTCTCGAGTGGATGGGCGGTATTAATCCTCCTGCTGGTACTA CTTCTTATGCTCAGAAGTTTCAGGGTCGGGTCACCATGACCCGTGATACCAGCATTAG CACCGCGTATATGGAACTGAGCCGCCTGCGTAGCGATGATACGGCCGTGTATTATTG CGCGCGTGGTGGTTGGTTTGATTATTGGGGCCAAGGCACCCTGGTGACGGTTAGCTC AGCTTCCACCAAGGGCCCCAGCGTGTTCCCCCTGGCCCCCTGCAGCAGAAGCACCA GCGAGAGCACAGCCGCCCTGGGCTGCCTGGTGAAGGACTACTTCCCCGAGCCCGTG ACCGTGAGCTGGAACAGCGGAGCCCTGACCAGCGGCGTGCACACCTTCCCCGCCGT GCTGCAGAGCAGCGGCCTGTACAGCCTGAGCAGCGTGGTGACCGTGCCCAGCAGCA

		ACTTCGGCACCCAGACCTACACCTGCAACGTGGACCACAAGCCCAGCAACACCAAGG TGGACAAGACCGTGGAGCGGAAGTGCTGCGTGGAGTGCCCCCCCTGCCCTGCCCCT CCTGTGGCCGGACCCTCCGTGTTCCTGTTCCCCCCCAAGCCCAAGGACACCCTGATG ATCAGCCGGACCCCCGAGGTGACCTGCGTGGTGGTGGACGTGAGCCACGAGGACCC CGAGGTGCAGTTCAACTGGTACGTGGACGGCGTGGAGGTGCACAACGCCAAGACCA AGCCCCGGGAGGAACAGTTCAACAGCACCTTCCGGGTGGTGTCCGTGCTGACCGTG GTGCACCAGGACTGGCTGAACGGCAAAGAATACAAGTGCAAGGTGTCCAACAAGGG CCTGCCTGCCCCCATCGAGAAAACCATCAGCAAGACAAAGGGCCAGCCCAGGGAAC CCCAGGTGTACACCCTGCCCCCCAGCCGGGAGGAAATGACCAAGAACCAGGTGTCC CTGACCTGTCTGGTGAAGGGCTTCTACCCCAGCGACATCGCCGTGGAGTGGGAGAG CAACGGCCAGCCCGAGAACAACTACAAGACCACCCCCCCCCATGCTGGACAGCGACG GCAGCTTCTTCCTGTACAGCAAGCTGACAGTGGACAAGAGCCGGTGGCAGCAGGGC AACGTGTTCAGCTGCAGCGTGATGCACGAGGCCCTGCACAACCACTACACCCAGAAG AGCCTGAGCCTGTCCCCCGGCAAA
SEQ ID NO180	DNA Heavy Chain	CAGGTGCAATTGGTTCAGAGCGGCGCGGAAGTGAAAAAACCGGGCGCGAGCGTGAA AGTGAGCTGCAAAGCCTCCGGATATACCTTTACTTCTTCTTATATTAATTGGGTCCGCC AAGCCCCTGGGCAGGGTCTCGAGTGGATGGGCAATATTAATCCTGCTACTGGTCATG CTGATTATGCTCAGAAGTTTCAGGGTCGGGTGACCATGACCCGTGATACCAGCATTA GCACCGCGTATATGGAACTGAGCCGCCTGCGTAGCGATGATACGGCCGTGTATTATT GCGCGCGTGGTGGTTGGTTTGATTATTGGGGCCAAGGCACCCTGGTGACGGTTAGCT CAGCTTCCACCAAGGGCCCCAGCGTGTTCCCCCTGGCCCCCTGCAGCAGAAGCACC AGCGAGAGCACAGCCGCCCTGGGCTGCCTGGTGAAGGACTACTTCCCCGAGCCCGT GACCGTGAGCTGGAACAGCGGAGCCCTGACCAGCGGCGTGCACACCTTCCCCGCCG TGCTGCAGAGCAGCGGCCTGTACAGCCTGAGCAGCGTGGTGACCGTGCCCAGCAGC AACTTCGGCACCCAGACCTACACCTGCAACGTGGACCACAAGCCCAGCAACACCAAG GTGGACAAGACCGTGGAGCGGAAGTGCTGCGTGGAGTGCCCCCCCTGCCCTGCCCC TCCTGTGGCCGGACCCTCCGTGTTCCTGTTCCCCCCCAAGCCCAAGGACACCCTGAT GATCAGCCGGACCCCCGAGGTGACCTGCGTGGTGGTGGACGTGAGCCACGAGGAC CCCGAGGTGCAGTTCAACTGGTACGTGGACGGCGTGGAGGTGCACAACGCCAAGAC CAAGCCCCGGGAGGAACAGTTCAACAGCACCTTCCGGGTGGTGTCCGTGCTGACCG TGGTGCACCAGGACTGGCTGAACGGCAAAGAATACAAGTGCAAGGTGTCCAACAAGG GCCTGCCTGCCCCCATCGAGAAAACCATCAGCAAGACAAAGGGCCAGCCCAGGGAA CCCCAGGTGTACACCCTGCCCCCCAGCCGGGAGGAAATGACCAAGAACCAGGTGTC CCTGACCTGTCTGGTGAAGGGCTTCTACCCCAGCGACATCGCCGTGGAGTGGGAGA GCAACGGCCAGCCCGAGAACAACTACAAGACCACCCCCCCCCATGCTGGACAGCGAC GGCAGCTTCTTCCTGTACAGCAAGCTGACAGTGGACAAGAGCCGGTGGCAGCAGGG CAACGTGTTCAGCTGCAGCGTGATGCACGAGGCCCTGCACAACCACTACACCCAGAA GAGCCTGAGCCTGTCCCCCGGCAAA
SEQ ID NO181	ActRIIB	MTAPWVALALLWGSLCAGSGRGEAETRECIYYNANWELERTNQSGLERCEGEQDKRLH CYASWRNSSGTIELVKKGCWLDDFNCYDRQECVATEENPQVYFCCCEGNFCNERFTHL PEAGGPEVTYEPPPTAPTLLTVLAYSLLPIGGLSLIVLLAFWMYRHRKPPYGHVDIHEDPG PPPPSPLVGLKPLQLLEIKARGRFGCVWKAQLMNDFVAVKIFPLQDKQSWQSEREIFSTP GMKHENLLQFIAAEKRGSNLEVELWLITAFHDKGSLTDYLKGNIITWNELCHVAETMSRGL SYLHEDVPWCRGEGHKPSIAHRDFKSKNVLLKSDLTAVLADFGLAVRFEPGKPPGDTHG QVGTRRYMAPEVLEGAINFQRDAFLRIDMYAMGLVLWELVSRCKAADGPVDEYMLPFEE EIGQHPSLEELQEVVVHKKMRPTIKDHWLKHPGLAQLCVTIEACWDHDAEARLSAGCVEE RVSLIRRSVNGTTSDCLVSLVTSVTNVDLPPKESSI
SEQ ID NO182	ActRIIB ligand binding domain	SGRGEAETRECIYYNANWELERTNQSGLERCEGEQDKRLHCYASWRNSSGTIELVKKGC WLDDFNCYDRQECVATEENPQVYFCCCEGNFCNERFTHLPEAGGPEVTYEPPPTAPT

	(aa19-134)	
SEQ ID NO183	Antibody binding region	IELVKKGSWLDDFNS
SEQ ID NO184	Antibody binding region	VKKGSWLDDFNSYDR
SEQ ID NO185	Antibody binding region	GSWLDDFNSYDRQES
SEQ ID NO186	Antibody binding region	GCWLDDFNC
SEQ ID NO187	Antibody binding region	CEGEQDKRLHCYASW
SEQ ID NO188	Antibody binding region	WLDDFN
SEQ ID NO189	Antibody binding region	EQDKR
SEQ ID NO190	Antibody binding region	KGCWLDDFNCY
SEQ ID NO191	Antibody binding region	CIYYNANWELERT
SEQ ID NO192	Antibody binding region	YFCCCEGNFCN
SEQ ID NO193	Light – h/mIgG2aLALA Chain	DIALTQPASVSGSPGQSITISCTGTSSDVGSYNYVNWYQQHPGKAPKLMIYGVSKRPSGVSNRFSGSKSGNTASLTISGLQAEDEADYYCGTFAGGSYYGVFGGGTKLTVLGQPKSTPTLTVFPPSSEELKENKATLVCLISNFSPSGVTVAWKANGTPITQGVDTSNPTKEGNKFMASSFLHLTSDQWRSHNSFTCQVTHEGDTVEKSLSPAECL
SEQ ID NO194	Heavy-h/mIgG2aLALA chain	QVQLVQSGAEVKKPGASVKVSCKASGYTFT SSYINWVRQAPGQGLEWMGTINPVSGSTSYAQKFQGRVTMTRDTSISTAYMELSSLRSEDTAVYYCARGGWFDYWGQGTLVTVSSAKTTAPSVYPLAPVCGDTTGSSVTLGCLVKGYFPEPVTLTWNSGSLSSGVHTFPAVLQSDLYTLSSSVTVTSSTWPSQSITCNVAHPASSTKVDKKIEPRGPTIKPCPPCKCPAPNAAGGPSVFIFPPKIKDVLMISLSPIVTCVVVDVSEDDPDVQISWFVNNVEVHTAQTQTHREDYNSTLRVVSALPIQHQDWMSGKEFKCKVNNKDLPAPIERTISKPKGSVRAPQVYVLPPPEEEMTKKQVTLTCMVTDFMPEDIYVEWTNNGKTELNYKNTEPVLDSDGSYFMYSKLRVEKKNWVERNSYSCSVVHEGLHNHHTTKSFSRTPGK

Some embodiments of the disclosed methods, treatments, regimens, uses and kits employ a myostatin antagonist, e.g., a myostatin binding molecule or an ActRIIB

binding molecule. In further embodiments, the ActRIIB binding molecule is an antagonist antibody to ActRIIB.

In some embodiments of the disclosed methods, treatments, regimens, uses and kits, the anti-ActRIIB antibody is selected from the group consisting of: a) an anti-ActRIIB antibody that binds to an epitope of ActRIIB comprising SEQ ID NO: amino acids 78-83 of SEQ ID NO: 181 (WLDDFN – SEQ ID NO:188);

(b) amino acids 76-84 of SEQ ID NO: 181 (GCWLDDFNC – SEQ ID NO:186);

(c) amino acids 75-85 of SEQ ID NO: 181 (KGCWLDDFNCY – SEQ ID NO:190);

(d) amino acids 52-56 of SEQ ID NO: 181 (EQDKR – SEQ ID NO:189);

(e) amino acids 49-63 of SEQ ID NO: 181 (CEGEQDKRLHCYASW – SEQ ID NO:187);

(f) amino acids 29-41 of SEQ ID NO: 181 (CIYYNANWELERT– SEQ ID NO:191);

(g) amino acids 100-110 of SEQ ID NO: 181 (YFCCCEGNFCN – SEQ ID NO:192); or

(h) amino acids 78-83 of SEQ ID NO: 181 (WLDDFN) and amino acids 52-56 of SEQ ID NO: 181 (EQDKR).

; and b) an antagonist antibody to ActRIIB that binds to an epitope of ActRIIB comprising amino acids 78-83 of SEQ ID NO: 181 (WLDDFN – SEQ ID NO:188);

(b) amino acids 76-84 of SEQ ID NO: 181 (GCWLDDFNC – SEQ ID NO:186);

(c) amino acids 75-85 of SEQ ID NO: 181 (KGCWLDDFNCY – SEQ ID NO:190);

(d) amino acids 52-56 of SEQ ID NO: 181 (EQDKR – SEQ ID NO:189);

(e) amino acids 49-63 of SEQ ID NO: 181 (CEGEQDKRLHCYASW – SEQ ID NO:187);

(f) amino acids 29-41 of SEQ ID NO: 181 (CIYYNANWELERT– SEQ ID NO:191);

(g) amino acids 100-110 of SEQ ID NO: 181 (YFCCCEGNFCN – SEQ ID NO:192); or

(h) amino acids 78-83 of SEQ ID NO: 181 (WLDDFN) and amino acids 52-56 of SEQ ID NO: 181 (EQDKR), wherein the antibody has a K_D of about 2 pM.

In some embodiments of the disclosed methods, treatments, regimens, uses and kits, the antagonist antibody to ActRIIB is a human antibody.

In some embodiments of the disclosed methods, treatments, regimens, uses and kits, the antibody is bimagrumab or BYM338.

The details of one or more embodiments of the disclosure are set forth in the accompanying description above. Although any methods and materials similar or

equivalent to those described herein can be used in the practice or testing of the present disclosure, the preferred methods and materials are now described. Other features, objects, and advantages of the disclosure will be apparent from the description and from the claims. In the specification and the appended claims, the singular forms include plural referents unless the context clearly dictates otherwise. Unless defined otherwise, all technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which this disclosure belongs. All patents and publications cited in this specification are incorporated by reference. The following Examples are presented in order to more fully illustrate the preferred embodiments of the disclosure. These examples should in no way be construed as limiting the scope of the disclosed patient matter, as defined by the appended claims.

EXAMPLES

General Methodology

ActRIIB antibodies, their characterisation and methods related thereto like (i) Functional Assays, (ii) REPORTER GENE ASSAYs (RGA), (iii) Cultivation of HEK293T/17 Cell Lines, (iv) Myostatin-Induced Luciferase Reporter Gene Assays,(v) SPECIFICITY ELISAs, (vi) ActRIIB/Fc-Myostatin Binding Interaction ELISA, (vii) FACS titration on hActRIIB- and hActRIIA-Expressing Cells, (viii) Binding to primary human skeletal muscle cells, (ix) affinity Determination of Selected Anti-Human ActRIIB Fabs Using Surface Plasmon Resonance (Biacore), (x) CK ASSAY, (xi) Animal Models, (xii) TREATMENT PROTOCOLs, (xiii) Statistical Analysis, (xiiii) Pannings, (xv)antibody identification and characterization, (xvi) Optimization of antibodies derived from first affinity maturation, (xvii) IgG2 Conversion of Affinity Matured Fabs (1st Maturation), (xviiii) Second Affinity Maturation, (xx) IgG2 Conversion and Characterization of IgG2 (2nd Maturation), (xxi) Characterization of anti-ActRIIB antibodies in in vivo murine studies, (xxii) Confirmation of affinity by SET, (xxiii) Cross Blocking Studies and (xxiv) Epiotpe mapping details and technologies have been disclosed in the WO 2010/125003.

Study design

This is a multi-center, randomized, double-blind, placebo-controlled, pivotal, 4 arm dose-finding phase IIb/III trial. A total of approximately 240 subjects with sporadic Inclusion Body Myositis (sIBM) are randomized into one of the 4 trial arms (A, B, C, or D) in a 1:1:1:1 ratio (**Figure 2**).

A maximum of 20% of patients with baseline 6 minute walk distance ≥400 meters will be randomized.

For each subject, the study consists of a maximum 28 day screening period which includes a baseline visit up to 5 days, a 52 week treatment period, a variable 0-52 week maintenance treatment period, and a 28 day treatment-free follow up period.

The study includes 4 study epochs:

1. Screening epoch: With duration up to 28 days (Day -28 to Day -1), during which study eligibility will be confirmed and if applicable, to taper subjects off disallowed medications. The screening epoch consists of a screening visit and a baseline visit. Subjects do not receive any study medication during this epoch.
   Screening visit: Visit may occur 28 to 6 days prior to randomization (Day -28 to Day -6). Informed consent must be obtained prior to implementing any study specific procedure. Subjects that meet the inclusion/exclusion criteria will undergo safety assessments that will include physical examinations, ECG, vital signs, standard clinical laboratory evaluations (hematology, blood chemistry, urinalysis).
   Baseline visit: All required assessments prior to beginning study drug administration will be performed during this 5 day timeframe (Day -5 to Day -1). Subjects that meet the safety requirements and inclusion/ exclusion criteria at the screening visit will be asked to return to the site for baseline assessments. Once baseline evaluations (including the 6MWD) have been checked and the patient again confirmed as eligible for the study, the patient will be asked to return to the site at Day 1 to be randomized.
2. Treatment epoch: The treatment period is defined as Day 1 to Week 52 (pre-dose administration). At the Day 1 visit, subjects will be randomized into one of four parallel treatment arms:
   - Arm A: BYM338 10 mg/kg i.v. infusion every 4 weeks
   - Arm B: BYM338 3 mg/kg i.v. infusion every 4 weeks
   - Arm C: BYM338 1 mg/kg i.v. infusion every 4 weeks
   - Arm D: Placebo, administered as i.v. infusion every 4 weeks

   Depending on the study arm the subject will receive either 13 doses of BYM338 or matching placebo administered as intravenous infusion every 4 weeks. The first study drug administration will occur at Day 1 and all baseline safety evaluation results must be available prior to dosing. For all subsequent visits including Week 52, all evaluations must be performed prior to drug administration. Final dose administration for this treatment epoch will occur at the Week 48 visit. Subjects will return to the site every four weeks for safety, pharmacodynamics, pharmacokinetic, biomarker and efficacy evaluations.

All subjects will perform their End of Treatment assessments at Week 52 and then move into the Maintenance Treatment epoch until the last subject continuing in this Treatment epoch has completed their Week 48 dose.

Once the last subject continuing in this Treatment epoch has completed their Week 48 dose, no other subjects will receive study drug, and:

- Subjects that have not yet completed this Treatment epoch should complete an End of Treatment (EOT) visit approximately 28 days following their last study drug administration.
- After the EOT visit is completed, these subjects will enter directly into the treatment-free follow up period (they will not be eligible to enter into the Maintenance Treatment epoch).
- For subjects that have already entered the Maintenance epoch, see below.

3. Maintenance Treatment epoch: This epoch will be defined by the duration of patient enrolment, but will be limited to a maximum duration of 52 weeks (in case the enrolment period exceeds 52 weeks). Thus, with variable duration of 0 up to 52 weeks, the maintenance treatment period is defined as Week 52 post-dose administration up to a maximum of Week 104.

   During this Maintenance Treatment epoch, subjects will continue in the treatment arm assigned at randomization. The dose given at Week 52 is the first dose of the maintenance treatment period and is followed by doses of BYM338 or placebo every four weeks up to Week 100. Subjects will return to the site every four weeks for safety and efficacy evaluations.

   The Week 104 visit is the end of maintenance treatment (EOMT) visit. The EOMT visit is considered complete when all visit assessments are completed and the visit has been registered into Interactive Response Technology (IRT).

   Treatment duration for all subjects beyond Week 52 will be determined by the date the last subject continuing in the Treatment epoch receives their Week 48 dose, but will be limited to 52 weeks (i.e. a maximum total treatment duration in the study of 2 years).

   Once the last subject has received their Week 48 dose, no other subjects will receive study drug, and:

   - All subjects in the Maintenance Treatment epoch should complete an End of Maintenance Treatment (EOMT) visit approximately 28 days following their last study drug administration.
   - After EOMT visit is completed, all subjects will enter the treatment-free follow up period.

4. Post-Treatment Follow Up epoch: With duration of 28 days, subjects do not receive any study medication during this epoch. The End of Follow Up (EOF) visit will occur 4 weeks after completion of the respective subject's treatment epoch (Treatment epoch or Maintenance epoch) and 8 weeks after the last study drug administration. The EOF visit should be completed for all subjects, regardless of whether they complete the entire study as planned or discontinue prematurely). Subjects that complete this study may be eligible for entry into an open-label study to collect additional long-term efficacy and safety data of BYM338 in sIBM patients.

An outline of the study design is presented in Figure 3-1, while the Maintenance Treatment epoch variable treatment duration is represented in Figure 3-2. A detailed visit and assessment schedule can be found in Table 6-1 and Table 6-2.

Rationale of study design

Preliminary clinical data are available for a single dose administration of BYM338 30 mg/kg in patients with sIBM. No multiple dose data and no data on other doses of BYM338 are available yet in patients with sIBM. A dose-ranging study is proposed to study different BYM338 doses on the basis of efficacy, pharmacodynamics, pharmacokinetics, and key safety data in the targeted patient population. The design of this study addresses these needs, focusing in particular on subject safety and the effect of BYM338 on lean body mass, muscle strength, muscle function, and mobility. The primary endpoint of this trial measures a change in physical function and mobility via the 6-minute walk distance test at Week 52.

This double-blind, placebo-controlled study is designed to allow dose ranging assessment between doses of 1 mg/kg, 3 mg/kg, and 10 mg/kg BYM338 administered every 4 weeks. Currently, there is no approved treatment for sIBM. In addition, there is no pharmacological therapy available that would help sIBM patients to reverse the steady course of the disease progression (Greenberg 2009, Aggarwal and Oddis 2012). The finding of elevated expression of TGFβ signaling activation through pSMAD suggests that myostatin over-expression could be an important part of the patho-physiology of sIBM, and therefore that BYM338 could be beneficial in this disease.
Given that there is no drug approved for the treatment of sIBM and no current standard of care, a placebo-control is proposed. Placebo-controlled trials provide the maximum ability to distinguish adverse effects caused by a drug from those resulting from underlying disease or “background noise.” Placebo-controlled trials also provide an optimal setting to demonstrate the efficacy of an investigational therapy, and to demonstrate a dose-response relationship, and may increase the ability of the study to detect true drug effects by decreasing the amount of improvement resulting from subject or investigator expectations. It is particularly important to be able to distinguish the adverse effect profile of BYM338 in sIBM patients, as sIBM is the first indication for BYM338 and only limited human data with BYM338 are available to date.
Subjects randomized to any of the three non-placebo arms will receive at least 13 doses of BYM338 every 4 weeks. Study drug treatment duration beyond 52 weeks will vary per subject with a minimum total treatment duration in this study of 52 weeks and a maximum of 104 weeks, followed in all cases by a 4 week treatment-free follow up period, to allow for evaluation of long-term efficacy and safety in this population. This study is designed to gather data regarding onset of therapeutic effect as measured by

the 6 minute walk test. The trial also includes measurements of the effects of sIBM on quality of life, including swallowing difficulties that may result from this disease.

The blinding is maintained until database lock to ensure reliable efficacy and safety measures. Ideally, a standard exercise program would be performed by all subjects, to avoid variability in assessing the efficacy of various BYM338 doses. Due to the wide variation of physical function within the sIBM population (i.e. asymmetrical neuromuscular pathology), subjects will be provided with a customized, subject-specific exercise recommendation by a qualified physical therapist that will assess the subject's abilities and severity of weakness at each visit. Subjects will be encouraged to engage in the physical activity that is determined to be safe for them by the site physical therapist. A video describing a suggested home-based exercise program will be provided to the physical therapist to assist in the development of an exercise recommendation to the subject. The physical therapist can choose to provide a copy of the video to the subject, if he/she judges it to be appropriate to be used by the subject. The effect of an exercise program alone remains to be defined in a placebo group and may synergize with a group receiving active drug.

The study population will consist of ambulatory sIBM patients fulfilling the sIBM diagnostic criteria (adapted from the proposed Medical Research Council criteria – refer to Appendix 1) at the time of screening. Eligible subjects are required to have pathologically defined or clinically defined sIBM diagnosis as per the sIBM diagnostic criteria (Hohlfeld 2011 adapted from Hilton-Jones et al 2010).

Patients with 6MWD ≥ 400 m at baseline will represent no more than 20% of the study sample size as many publications argue for a threshold of 400 m as a marker of subnormal physical function and mobility in healthy volunteers and patients under various disease conditions (Chetta et al 2006, Troosters et al 1999, Simonsick et al 2008, Chang et al 2004, Enright et al 2003, Enright et al 1998, Kruis et al 2010, Morley et al 2011, Kommuri et al 2010, Villalba et al 2007).

3.3 Rationale of dose/regimen, route of administration and duration of treatment

The proposed doses have been chosen based on inferences made from available clinical data, including consideration of the recently available on-treatment results from the ongoing study CBYM338X2205.

Data from study CBYM338X2205 on 14 patients (11 active, 3 placebo) demonstrated that administration of a single dose of BYM338 30 mg/kg i.v. showed a significant increase from baseline in thigh muscle volume (TMV) via MRI as well as total and appendicular LBM via DXA as compared to placebo. In addition, this dose also induced sustained strength and functional performance and mobility improvements as per the Quantitative Muscle Testing and the 6-minute walking distance tests 16 weeks after drug

administration. As the single 30 mg/kg dose showed an effect of at least 8 weeks duration, repeated treatment with 10 mg/kg may achieve similar efficacy. Therefore, 10 mg/kg has been selected as the highest dose in this study. However, the minimum efficacious dose remains to be identified.

In this study, the efficacy of long-term repeated dosing will be investigated with the objective to understand the dose-response relationship of BYM338. All doses of BYM338 chosen for this study (10, 3, and 1 mg/kg i.v.) have been tested and found to be well-tolerated in normal healthy volunteers.

The objective for the doses is to evaluate whether repeated doses of 10, 3, or 1 mg/kg i.v. can offer additional muscle structure and functional benefit to patients, and whether lower doses can do so with fewer side-effects such as acne or involuntary muscle contractions.

In healthy volunteers (CBYM338X2101 and CBYM338X2102), thigh muscle MRI volume increased comparably for single doses of 10 mg/kg and 30 mg/kg, but the effect of 30 mg/kg lasted longer. With 3 repeat monthly doses of BYM338, there was a comparable increase in TMV in healthy adults at 3 mg/kg and 10 mg/kg, even though it is thought that 3 mg/kg causes complete receptor occupancy for approximately half the duration of 10 mg/kg. It is not yet clear if 10 mg/kg would have fewer adverse effects than 30 mg/kg, presumably due to a lower Cmax, even if it were given more frequently than the 30 mg/kg dose.

In summary 3 mg/kg and 10 mg/kg q4 were effective in increasing TMV in healthy volunteers, thus are expected to be effective as well in sIBM patients.

The lower dose of 1 mg/kg did not demonstrate a persistent increase in TMV beyond 2 weeks after a single dose was administered to healthy volunteers. However, it is being tested on the premise that patients with sIBM could have an increased sensitivity to BYM338 and that this dose could prove effective when administered multiple times.

A consensus paper arising from a conference of international experts in neuromuscular diseases in 2010 recommends that treatment trials in sIBM must exceed 6 months in length in order to adequately assess the effect of a treatment on the disease (Hilton-Jones et al 2010). Both EMA and FDA agreed that a 12-month duration for sIBM trials was adequate, thus all subjects will receive at least 52 weeks of treatment. Additional data up to a maximum of 3 years from randomization is being generated to evaluate the long-term safety and efficacy in this population.

Although it is preferable to maintain subjects on placebo for the shortest possible duration, there is currently no approved treatment for sIBM, and no standard of care.

Furthermore, given that sIBM is the first indication planned for registration with BYM338, and because sIBM is expected to require chronic treatment, it is important to develop a robust database with sufficient long-term safety data. Therefore, it is reasonable to conduct a 1 year placebo-controlled study with an additional maintenance period (for a total treatment duration of up to 2 years). Regular assessments of disease activity ensure that subjects who are experiencing worsening of disease in any of the treatment groups can exit the study upon their own wish or based on the advice of the investigator at any time. It should be noted that as soon as efficacy and safety are determined based on the results of the study, all subjects will be invited to participate in a separate open label extension study with active drug. Thus, no subject will be on placebo after the 52 week safety and efficacy has been determined. Though currently available data are insufficient to demonstrate convincing efficacy of BYM338 in the sIBM population, this study is designed to collect this data and inform on both efficacy and safety of BYM338 in the sIBM population at the study conclusion.

Rationale for choice of comparator

Given that there is no drug approved for the treatment of sIBM, a placebo-control is proposed. Placebo-controlled trials also provide the maximum ability to distinguish adverse effects caused by a drug from those resulting from underlying disease or "background noise." Placebo-controlled trials also provide an optimal setting to demonstrate the efficacy of an investigational therapy, and to demonstrate a dose-response relationship, and may increase the ability of the study to detect true drug effects by decreasing the amount of improvement resulting from subject or investigator expectations.

Risks and benefits

The safety data from the completed Phase I studies do not highlight any specific safety risk or concern or any particular pattern of major event clustering. From the standpoint of the overall risk benefit assessment the current study design and its trial-related risks (infusion, blood draws, exposure to radiation during imaging) with BYM338 is justified.

To date 213 patients and healthy volunteers have been enrolled in studies with bimagrumab with 150 having received active drug at doses ranging from single and multiple doses of 0.1 to 30 mg/kg i.v. Early phase clinical data support a favorable benefit-risk profile for BYM338 in healthy volunteers and patients.

Preliminary data from the first studies in healthy volunteers and older patients indicate observations of diffuse acne on the face and less frequently on the back and chest may be present in some adults receiving a single dose of 30 mg/kg. A possible mechanistic link between BYM338 and skin reactivity remains unclear.

In approximately 20% - 48% of normal healthy volunteers and patients with sIBM treated with a single 30 mg/kg i.v. dose of BYM338, involuntary muscle contractions, referred to as 'muscle twitching' were reported. The reported muscle twitches were mild in intensity, transient, of short duration, self-limiting and did not require medical treatment. While twitching could theoretically be more pronounced in aging and atrophying muscle, this has not been observed in sIBM patients with age up to 78 years. The biological explanation of twitching is being pursued by the BYM338 Project Team using both internal and external resources.

Preclinical studies in rats demonstrated a marked cardiac hypertrophy deemed compensatory to the substantial increase in skeletal muscle mass. Although inconsistently seen in monkey studies, the observed increase in ventricular muscle mass without any effect on left ventricular chamber size and, importantly, on global left ventricular systolic function in female monkeys is also supportive of a compensatory effect rather than a direct effect of BYM338. It is not clear if cardiac hypertrophy may occur with BYM338 treatment in patients with small increases in skeletal muscle mass. Echocardiography and EKG Holter monitor data showed no evidence of significant change in posterior wall thickness, interventricular septum thickness, left ventricular diastolic diameter, left ventricular mass or cardiac conduction when compared with placebo after single and multiple doses up to and including 30 mg/kg. ECGs and electrocardiography will be conducted in study CBYM338B2203 to monitor this potential risk.

Weekly doses of 1, 10 and 100 mg/kg of BYM338 in the 4- and 13-week toxicity studies resulted in reversible changes in the uterus, vagina and ovaries of the rat and to a much lesser extent to the ovaries of the cynomolgus monkey. At 100 mg/kg given weekly, BYM338 was shown to be developmentally toxic in the rat, with evidence of teratogenicity. These findings are not unexpected based on the pharmacology of BYM338 and the role of activins in organogenesis (Xia and Schneyer 2009). Suppression of FSH levels in postmenopausal and premenopausal women of non-childbearing potential, increasing with the dose of BYM338, has been identified and plans are in place to monitor data closely in ongoing and future studies. The clinical meaning of reduced FSH in postmenopausal women is unknown; in premenopausal women it is expected that such a reduction could interfere with menstrual cycling, fecundity and potentially cause symptoms of low estrogen in the body (e.g. hot flushes, vaginal dryness, accelerated bone loss), which could require additional treatment. Only, women of non-child bearing potential have been recruited in all healthy volunteer studies. For subjects who are premenopausal, this study plans to enroll women of childbearing potential (WOCBP) who consent to use two forms of highly effective contraception from

the day written consent is given through 6 months after the last study drug administration. Pregnancy testing will be administered at screening, throughout the study period and 30 days after the last visit.

Body weight may increase with BYM338 treatment, but it is expected that the increase will be primarily in LBM, with stable or reduced body fat mass.

It is possible that anti-drug antibodies could develop against BYM338, which could neutralize the drug and attenuate its efficacy. However, as of the publication date of the IB version 4.0, no neutralizing antibodies have been observed in the samples evaluated from single dose studies. Analysis is planned for the multiple dose study and other single dose studies.

Infusion related reactions can occur with monoclonal antibodies. Hypersensitivity reactions can manifest as fever, chills, urticaria, dyspnea, headaches, myalgia and/or hypotension. A serious infusion reaction that results in anaphylaxis is a rare event in monoclonal antibody therapy. If a severe hypersensitivity reaction occurs, administration of BYM338 should be discontinued and appropriate therapy initiated.

As with any other investigational drug, there are unknown risks to BYM338 which may be serious and unforeseen.

Population

The study population will consist of ambulatory (i.e. not wheel-chair bound or bed-ridden at screening and baseline) male and female patients with a pathologically defined or clinically defined diagnosis of sIBM as per the 2010 definition of the European Neuromuscular Center. It is aimed to randomize a total of 240 patients in approximately 30 centers worldwide.

Inclusion criteria

Patients eligible for inclusion in this study have to fulfill all of the following criteria at screening and baseline visits:

Male and female patients age 35 to 85 years of age (inclusive)

Diagnosed with pathologically-defined or clinically-defined sporadic inclusion body myositis as per the sIBM diagnostic criteria (adapted from the proposed 2010 MRC criteria), including evaluation of a pathology report.

Must not be wheel-chair bound (intermittent use of wheelchair is allowed) at both screening and baseline visits, as defined by a 6MWD score >1 meter.

Able to communicate well with the investigator.

Willing to participate for the entire duration of the study with commitment to follow study requirements and procedures.

Exclusion criteria

Patients fulfilling any of the following criteria at screening and baseline visits are not eligible for inclusion in this study.

1. Use of other investigational drugs at the time of enrollment or within 30 days or 5 half-lives of enrollment, whichever is longer, or longer if required by any other limitation of participation in an investigational trial based on local regulations. Current or planned participation to another clinical trial during this study. Participation in observational or registry studies not involving drug therapy is allowed;
2. History of an allergic reaction or anaphylactic reaction to the study medication or history of hypersensitivity to any component of the study medication. History of hypersensitivity to biologics;
3. Reasons that preclude adequate intake of protein, defined as at least 0.8 g protein/kg/day and/or diseases known to cause malabsorption of protein or energy from the gastrointestinal tract, such as inflammatory bowel disease, celiac disease, short bowel syndrome, pancreatic insufficiency;
4. Any non-sIBM conditions or other neurologic (congenital or acquired) or neuromuscular diseases that cause significant lower leg muscular or joint pain or muscle weakness that significantly limit mobility, including polymyositis or dermatomyositis, polymyalgia rheumatica, fibromyalgia, significant dementia/Alzheimer's disease, Parkinson's disease, amyotrophic lateral sclerosis, stroke, cerebral palsy, epilepsy, multiple sclerosis, spinal cord injury, muscular dystrophy, myasthenia gravis, and weakness related to degenerative joint disease of the spine. Note: Previous misdiagnosis of these is not exclusionary. Patients with conditions such as osteoarthritis can participate unless their pain limits them from performing study procedures;
5. Any active chronic non-sIBM condition associated with cachexia or muscle atrophy or that limits mobility as a result of respiratory function, including: severe chronic obstructive pulmonary disease (FEV1 < 50% predicted or functional dyspnea ≥ Grade 3 on the Medical Research Council Dyspnea Scale), advanced organ failure, chronic kidney disease (estimated GFR < 30 mL/min using the MDRD equation), hyperthyroidism, rheumatoid arthritis;
6. Uncontrolled hypothyroidism. Hypothyroid patients who have changed their dose of hormone replacement therapy in the past 6 weeks prior to screening are not eligible for the study;

7. Uncontrolled diabetes mellitus (i.e. HbA1C ≥9.0 mmol/l) and/or any other uncontrolled endocrine disease;
8. Potential subjects who have recently donated blood in the past 60 days, or plasma donation in the past 2 weeks;
9. History of a hip fracture in the last 6 months or has undergone surgery for a hip or knee prosthesis in the last 6 months;
10. Experienced an acute illness within the 30 days prior to screening (Screening Visit) that, in the opinion of the investigator, affects lower extremity function or the patient's ability to participate in the study;
11. Score "yes" on item 4 or item 5 of the Suicidal Ideation section of the Columbia Suicide Severity Rating Scale (CSSRS), if this ideation occurred in the past 6 months, or "yes" on any item of the Suicidal Behavior section, except for the "Non-Suicidal Self-Injurious Behavior" (item also included in the Suicidal Behavior section), if this behavior occurred in the past 2 years;
12. Pregnant or nursing (lactating) women, where pregnancy is defined as the state of a female after conception and until the termination of gestation, confirmed by a positive hCG laboratory test;
13. Women of child-bearing potential, defined as all women physiologically capable of becoming pregnant, UNLESS they are using highly effective methods of contraception during dosing and for 6 months after the last BYM338 dose.

Highly effective contraception methods include:

1. Total abstinence (when this is in line with the preferred and usual lifestyle of the subject). Periodic abstinence (e.g., calendar, ovulation, symptothermal, post-ovulation methods) and withdrawal are not acceptable methods of contraception.
2. Female Sterilization (have had surgical bilateral oophorectomy with or without hysterectomy) or tubal ligation at least six weeks before taking study treatment. In case of oophorectomy alone, only when the reproductive status of the woman has been confirmed by follow up hormone level assessment.

Male sterilization (at least 6 months prior to screening). For female subjects in the study, the vasectomised male partner should be the sole partner for that subject.

4. Combination of any two of the following methods (a+b or a+c or b+c):

a. Use of oral, injected or implanted hormonal methods of contraception or other forms of hormonal contraception that have comparable efficacy (failure rate <1%), for example hormone vaginal ring or transdermal hormone contraception.

b. Placement of an intrauterine device (IUD) or intrauterine system (IUS).

c. Barrier methods of contraception: Condom or Occlusive cap (diaphragm or cervical/vault caps) with spermicidal foam/gel/film/cream/vaginal suppository.

In case of use of oral contraception women should have been stable on the same pill for a minimum of 3 months before taking study treatment.

Women are considered post-menopausal and not of child bearing potential if they have had 12 months of natural (spontaneous) amenorrhea with an appropriate clinical profile (e.g. age appropriate, history of vasomotor symptoms) or have had surgical bilateral oophorectomy (with or without hysterectomy) or tubal ligation at least six weeks ago. In

the case of oophorectomy alone, only when the reproductive status of the woman has been confirmed by follow up hormone level assessment is she considered not of child bearing potential;

14. Any anticipated scheduled in-patient surgery within 12 months following randomization;

15. Severe Vitamin D deficiency defined as 25-OH-vitamin D levels <9.2 ng/mL or <23 nmol/mL at screening;

16. Significant psychiatric disorder, clinically manifest peripheral vascular disease or disorder, or systemic disorder which could affect any of the efficacy assessments (e.g. diabetic neuropathy, chronic fatigue syndrome, schizophrenia, bipolar disorder, severe depression, intermittent claudication);

17. Lack of peripheral venous access;

18. Known heart failure classified as New York Heart Association Class III and IV or a history of chronic hypotension (systolic blood pressure <100 mmHg);

19. Systolic blood pressure >180 or <90 mm Hg or diastolic blood pressure >100 or <50 mmHg at screening, or malignant hypertension;

20. History of unstable angina, myocardial infarction, coronary artery bypass graft surgery, or percutaneous coronary intervention (such as angioplasty or stent placement) within 180 days of screening;

21. Prolonged QT syndrome or QTcF > 450 msec (Fridericia Correction) for males and > 470 msec for females at screening or baseline at repeated assessment;

22. ECG showing clinically significant abnormalities including any current supra-ventricular arrhythmia with an uncontrolled ventricular response (mean heart rate >100 beats per minute [bpm]) at rest despite medical or device therapy, or any history of spontaneous or induced sustained ventricular tachycardia (heart rate >100 bpm for 30 sec) despite medical or device therapy, or any history of resuscitated cardiac arrest or presence of an automated internal cardioverter-defibrillator;

23. Significant coagulopathy, platelet count less than 75,000/mm3, hemoglobin less than 11.0 g/dL;

24. Liver disease or liver injury as indicated by abnormal liver function tests such as SGOT (AST), SGPT (ALT), alkaline phosphatase, or serum bilirubin (except Gilbert's Disease).

The Investigator should be guided by the following criteria:

- Any single transaminase may not exceed 3x times upper limit of normal (ULN). A single parameter elevated up to and including 3 x ULN should be re-checked as soon as possible, and in all cases, at least prior to randomization, to rule out any lab error.
- If the total bilirubin concentration is increased above 1.5x ULN, total bilirubin should be differentiated into the direct and indirect reacting bilirubin. In any case, serum bilirubin should not exceed the value of 1.6 mg/dL (27 µmol/L);

14. Known history or presence of severe acute or chronic liver disease (compensated or decompensated), known gallbladder or bile duct disease, acute or chronic pancreatitis, renal failure or chronic treatment with medication which has a hepatotoxic potential;
15. History of malignancy of any organ system (other than localized basal cell carcinoma or squamous cell carcinoma of the skin that has been definitively treated), treated or untreated, within the past 5 years, regardless of whether there is evidence of local recurrence or metastases. Participants with carcinoma in situ of the uterine cervix treated definitively more than 1 year prior to screening may enter the study;
16. Use of prohibited systemic treatments, including VEGF inhibitors (bevacizumab), within past 6 months prior to randomization or any therapies known to affect muscle mass, including androgen supplements (including OTC DHEA), GnRH analogues, anti-androgens, anti-estrogens (tamoxifen), progestins with known androgenic component (e.g. NETA), recombinant human growth hormone, oral beta agonists, insulin, megestrol acetate, or dronabinol within the past 3 months prior to randomization;
17. Ongoing chronic corticosteroid use or history of systemic corticosteroid use for at least 90 days prior to randomization at a daily dose greater than or equal to 10 mg prednisone equivalent;
18. Ongoing immunosuppressive therapy OR antibody immunosuppressive therapy (rituximab or i.v. immunoglobulin, TNF-alpha inhibitors) within the past 6 months prior to randomization OR non-antibody therapy for autoimmune diseases (e.g. cyclosporine, methotrexate, tacrolimus, cyclophosphamide, azathioprine) within the past 3 months (90 days) prior to randomization);
19. Currently active alcohol or drug abuse or history of alcohol or drug abuse within the last 24 weeks prior to randomization;
20. Known active infection of any kind (excluding fungal infection of nail beds) or any major episode of infection requiring hospitalization or treatment with i.v. anti-infectives within 8 weeks prior to randomization;
21. Any chronic active infection (e.g., HIV, hepatitis B or C, tuberculosis, etc). Patients receiving chemoprophylaxis for latent tuberculosis infection are eligible for the study;
22. Patient has any medical condition or laboratory finding during screening, which, in the investigator's opinion may interfere with participation, confound the results, or pose any additional risk to the patient when administering BYM338;
23. A maximum of 20% of patients with baseline 6 minute walk distance $\geq$400 meters will be randomized.

No additional exclusions may be applied by the investigator, in order to ensure that the study population will be representative of all eligible patients.

Treatment

Protocol requested treatment

1. Investigational treatment

Novartis will supply the following investigational drug:

- BYM338: 150 mg liquid in vial, provided in colorless glass vials with rubber stopper and aluminum flip-off caps.
- Placebo: Provided in colorless glass vials with rubber stopper and aluminum flip-off caps.

Note: To maintain a blind, open-label investigational treatment will be prepared by an unblinded pharmacist/designee and administered only by blinded study personnel.

2. Additional study treatment

No additional treatment beyond investigational treatment is required for this trial.

However, subjects may receive sufficient elemental calcium and vitamin D treatment as per local guidelines and under the guidance of the treating physician.

Treatment arms

This is a four arm study. 240 subjects with sIBM will be randomized at Day 1 to one of the following 4 arms in a ratio of 1:1:1:1:

- Arm A: BYM338 at 10 mg/kg i.v. infusion every 4 weeks (n=60)
- Arm B: BYM338 at 3 mg/kg i.v. infusion every 4 weeks (n=60)
- Arm C: BYM338 at 1 mg/kg i.v. infusion every 4 weeks (n=60)
- Arm D: Placebo, administered as i.v. infusion every 4 weeks for (n=60)

Depending on the study arm the subject receives at minimum 13 doses of BYM338 or matching placebo administered as intravenous infusion every 4 weeks. The first administration will occur at Day 1 and the final dose administration during the Treatment Epoch will occur at the Week 48 visit, defining the minimum treatment duration of 52 weeks. Subjects that enter the Maintenance Treatment epoch will continue to receive BYM338 or matching placebo i.v. infusions every 4 weeks according to their Day 1 treatment arm randomization assignment. Since all subjects will continue to receive study medication until the last subject reaches the Week 48 treatment dose, the final study drug administration is variable. However, the maximum total treatment duration for an individual subject in this study will be limited to 2 years (104 weeks),; in this maximal treatment duration case, the final study drug administration would occur at the Week 100 visit.

Claims

1. A myostatin antagonist for use in treating sporadic inclusion body myositis.

2. A myostatin antagonist for use according to claim 1, wherein said myostatin antagonist is to be administered to a patient in need thereof at a dose of about 1-10 mg/kg.

3. A myostatin antagonist for use according to claim 1 or 2, wherein said myostatin antagonist is to be administered at a dose of about 1, about 3 or about 10 mg/kg body weight.

4. A myostatin antagonist for use according to claim 1-3, wherein said myostatin antagonist is to be administered intraveneously.

5. A myostatin antagonist for use according to anyone of claims 1-4, wherein said myostatin anatagonis is to be administred every four weeks.

6. A myostatin antagonist for use according to anyone of claims 1-5, wherein said patient is ambulatory.

7. A myostatin antagonist for use according to anyone of claims 1-6, wherein treating sporadic inclusion body myositis comprises slowing down the progression of the disease or improving physical function and mobility.

8. A myostatin antagonist for use according to anyone of claims 1-6, wherein treating sporadic inclusion body myositis comprises improving dysphagia or swallowing difficulties.

9. A myostatin antagonist for use according to anyone of claims 1-6, wherein treating sporadic inclusion body myositis comprises improving upper extremity strength.

10. A myostatin antagonist for use according to anyone of claims 1-6, wherein treating sporadic inclusion body myositis comprises reducing incidence of falls or preventing falls.

11. A myostatin antagonist for use according to anyone of claims 1-10, wherein the myostatin antagonist is a myostatin receptor binding molecule.

12. A myostatin antagonist for use according to anyone of claims 1-11, wherein the myostatin antagonist is an ActRII receptor antagonist.

13. A myostatin antagonist for use according to anyone of claims 1-12, wherein the myostatin antagonist is an anti-ActRII receptor antibody.

14. A myostatin antagonist for use according to anyone of claims 1-13, wherein the anti-ActRII receptor antibody is bimagrumab.

15. A myostatin antagonist for use according to any one of claims 13-14, wherein the myostatin antagonist is an anti-ActRII antibody that binds to an epitope of ActRIIB consisting of amino acids 19-134 of SEQ ID NO: 181 (SEQ ID NO: 182).

16. A myostatin antagonist for use according to anyone of claims 13-15, wherein the anti-ActRII antibody binds to an epitope of ActRIIB comprising or consisting of:

(a) amino acids 78-83 of SEQ ID NO: 181 (WLDDFN – SEQ ID NO:188);

(b) amino acids 76-84 of SEQ ID NO: 181 (GCWLDDFNC – SEQ ID NO:186);

(c) amino acids 75-85 of SEQ ID NO: 181 (KGCWLDDFNCY – SEQ ID NO:190);

(d) amino acids 52-56 of SEQ ID NO: 181 (EQDKR – SEQ ID NO:189);

(e) amino acids 49-63 of SEQ ID NO: 181 (CEGEQDKRLHCYASW – SEQ ID NO:187);

(f) amino acids 29-41 of SEQ ID NO: 181 (CIYYNANWELERT– SEQ ID NO:191);

(g) amino acids 100-110 of SEQ ID NO: 181 (YFCCCEGNFCN – SEQ ID NO:192); or

(h) amino acids 78-83 of SEQ ID NO: 181 (WLDDFN) and amino acids 52-56 of SEQ ID NO: 181 (EQDKR).

17. A myostatin antagonist for use according to any of claims 13-16, wherein the anti-ActRIIB antibody is selected from the group consisting of:

a) an anti-ActRIIB antibody that binds to an epitope of ActRIIB comprising :

(a) amino acids 78-83 of SEQ ID NO: 181 (WLDDFN – SEQ ID NO:188);

(b) amino acids 76-84 of SEQ ID NO: 181 (GCWLDDFNC – SEQ ID NO:186);

(c) amino acids 75-85 of SEQ ID NO: 181 (KGCWLDDFNCY – SEQ ID NO:190);

(d) amino acids 52-56 of SEQ ID NO: 181 (EQDKR – SEQ ID NO:189);

(e) amino acids 49-63 of SEQ ID NO: 181 (CEGEQDKRLHCYASW – SEQ ID NO:187);
(f) amino acids 29-41 of SEQ ID NO: 181 (CIYYNANWELERT– SEQ ID NO:191);
(g) amino acids 100-110 of SEQ ID NO: 181 (YFCCCEGNFCN – SEQ ID NO:192); or
(h) amino acids 78-83 of SEQ ID NO: 181 (WLDDFN) and amino acids 52-56 of SEQ ID NO: 181 (EQDKR).
; and b) an antagonist antibody to ActRIIB that binds to an epitope of ActRIIB comprising amino acids 78-83 of SEQ ID NO: 181 (WLDDFN – SEQ ID NO:188);
(b) amino acids 76-84 of SEQ ID NO: 181 (GCWLDDFNC – SEQ ID NO:186);
(c) amino acids 75-85 of SEQ ID NO: 181 (KGCWLDDFNCY – SEQ ID NO:190);
(d) amino acids 52-56 of SEQ ID NO: 181 (EQDKR – SEQ ID NO:189);
(e) amino acids 49-63 of SEQ ID NO: 181 (CEGEQDKRLHCYASW – SEQ ID NO:187);
(f) amino acids 29-41 of SEQ ID NO: 181 (CIYYNANWELERT– SEQ ID NO:191);
(g) amino acids 100-110 of SEQ ID NO: 181 (YFCCCEGNFCN – SEQ ID NO:192); or
(h) amino acids 78-83 of SEQ ID NO: 181 (WLDDFN) and amino acids 52-56 of SEQ ID NO: 181 (EQDKR), wherein the antibody has a K_D of about 2 pM.

18. A myostatin antagonist for use according to any of claims 13-17, wherein the antibody binds to ActRIIB with a 10-fold or greater affinity than it binds to ActRIIA.

19. A myostatin antagonist for use according to anyone of claims 13-18, wherein the antibody comprises a heavy chain variable region CDR1 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-14; a heavy chain variable region CDR2 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 15-28; a heavy chain variable region CDR3 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 29-42; a light chain variable region CDR1 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 43-56; a light chain variable region CDR2 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 57-70; and a light chain variable region CDR3 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 71-84.

20. A myostatin antagonist for use according to any of claims 13-19 wherein the antibody comprises:

(a) a heavy chain variable region CDR1 of SEQ ID NO: 1; a heavy chain variable region CDR2 of SEQ ID NO: 15; a heavy chain variable region CDR3 of SEQ ID NO: 29; a light chain variable region CDR1 of SEQ ID NO: 43; a light chain variable region CDR2 of SEQ ID NO: 57; and a light chain variable region CDR3 of SEQ ID NO: 71,

(b) a heavy chain variable region CDR1 of SEQ ID NO: 2; a heavy chain variable region CDR2 of SEQ ID NO: 16; a heavy chain variable region CDR3 of SEQ ID NO: 30; a light chain variable region CDR1 of SEQ ID NO: 44; a light chain variable region CDR2 of SEQ ID NO: 58; and a light chain variable region CDR3 of SEQ ID NO: 72,

(c) a heavy chain variable region CDR1 of SEQ ID NO: 3; a heavy chain variable region CDR2 of SEQ ID NO: 17; a heavy chain variable region CDR3 of SEQ ID NO: 31; a light chain variable region CDR1 of SEQ ID NO: 45; a light chain variable region CDR2 of SEQ ID NO: 59; and a light chain variable region CDR3 of SEQ ID NO: 73,

(d) a heavy chain variable region CDR1 of SEQ ID NO: 4; a heavy chain variable region CDR2 of SEQ ID NO: 18; a heavy chain variable region CDR3 of SEQ ID NO: 32; a light chain variable region CDR1 of SEQ ID NO: 46; a light chain variable region CDR2 of SEQ ID NO: 60; and a light chain variable region CDR3 of SEQ ID NO: 74,

(e) a heavy chain variable region CDR1 of SEQ ID NO: 5; a heavy chain variable region CDR2 of SEQ ID NO: 19; a heavy chain variable region CDR3 of SEQ ID NO: 33; a light chain variable region CDR1 of SEQ ID NO: 47; a light chain variable region CDR2 of SEQ ID NO: 61; and a light chain variable region CDR3 of SEQ ID NO: 75,

(f) a heavy chain variable region CDR1 of SEQ ID NO: 6; a heavy chain variable region CDR2 of SEQ ID NO: 20; a heavy chain variable region CDR3 of SEQ ID NO: 34; a light chain variable region CDR1 of SEQ ID NO: 48; a light chain variable region CDR2 of SEQ ID NO: 62; and a light chain variable region CDR3 of SEQ ID NO: 76,

(g) a heavy chain variable region CDR1 of SEQ ID NO: 7; a heavy chain variable region CDR2 of SEQ ID NO: 21; a heavy chain variable region CDR3 of SEQ ID NO: 35; a light chain variable region CDR1 of SEQ ID NO: 49; a light chain

variable region CDR2 of SEQ ID NO: 63; and a light chain variable region CDR3 of SEQ ID NO: 77,

(h) a heavy chain variable region CDR1 of SEQ ID NO: 8; a heavy chain variable region CDR2 of SEQ ID NO: 22; a heavy chain variable region CDR3 of SEQ ID NO: 36; a light chain variable region CDR1 of SEQ ID NO: 50 a light chain variable region CDR2 of SEQ ID NO: 64; and a light chain variable region CDR3 of SEQ ID NO: 78,

(i) a heavy chain variable region CDR1 of SEQ ID NO: 9; a heavy chain variable region CDR2 of SEQ ID NO: 23; a heavy chain variable region CDR3 of SEQ ID NO: 37; a light chain variable region CDR1 of SEQ ID NO: 51; a light chain variable region CDR2 of SEQ ID NO: 65; and a light chain variable region CDR3 of SEQ ID NO: 79,

(j) a heavy chain variable region CDR1 of SEQ ID NO: 10; a heavy chain variable region CDR2 of SEQ ID NO: 24; a heavy chain variable region CDR3 of SEQ ID NO: 38; a light chain variable region CDR1 of SEQ ID NO: 52; a light chain variable region CDR2 of SEQ ID NO: 66; and a light chain variable region CDR3 of SEQ ID NO: 80,

(k) a heavy chain variable region CDR1 of SEQ ID NO: 11; a heavy chain variable region CDR2 of SEQ ID NO: 25; a heavy chain variable region CDR3 of SEQ ID NO: 39; a light chain variable region CDR1 of SEQ ID NO: 53; a light chain variable region CDR2 of SEQ ID NO: 67; and a light chain variable region CDR3 of SEQ ID NO: 81,

(l) a heavy chain variable region CDR1 of SEQ ID NO: 12; a heavy chain variable region CDR2 of SEQ ID NO: 26; a heavy chain variable region CDR3 of SEQ ID NO: 40; a light chain variable region CDR1 of SEQ ID NO: 54; a light chain variable region CDR2 of SEQ ID NO: 68; and a light chain variable region CDR3 of SEQ ID NO: 82,

(m) a heavy chain variable region CDR1 of SEQ ID NO: 13; a heavy chain variable region CDR2 of SEQ ID NO: 27; a heavy chain variable region CDR3 of SEQ ID NO: 41; a light chain variable region CDR1 of SEQ ID NO: 55; a light chain variable region CDR2 of SEQ ID NO: 69; and a light chain variable region CDR3 of SEQ ID NO: 83, or

(n) a heavy chain variable region CDR1 of SEQ ID NO: 14; a heavy chain variable region CDR2 of SEQ ID NO: 28; a heavy chain variable region CDR3 of SEQ ID NO: 42; a light chain variable region CDR1 of SEQ ID NO: 56; a light

chain variable region CDR2 of SEQ ID NO: 70; and a light chain variable region CDR3 of SEQ ID NO: 84.

21. A myostatin antagonist for use according to according to any of claims 13-20, wherein the antibody comprises a full length heavy chain amino acid sequence having at least 95% sequence identity to at least one sequence selected from the group consisting of SEQ ID NOs:146-150 and 156-160.

22. A myostatin antagonist for use according to according to any of claims 13-21, wherein the antibody comprises a full length light chain amino acid sequence having at least 95% sequence identity to at least one sequence selected from the group consisting of SEQ ID NOs:141-145 and 151-155.

23. A myostatin antagonist for use according to according to any of claims 13-22, wherein the antibody comprises:

(a) the variable heavy chain sequence of SEQ ID NO: 99 and variable light chain sequence of SEQ ID NO: 85;
(b) the variable heavy chain sequence of SEQ ID NO: 100 and variable light chain sequence of SEQ ID NO: 86;
(c) the variable heavy chain sequence of SEQ ID NO: 101 and variable light chain sequence of SEQ ID NO: 87;
(d) the variable heavy chain sequence of SEQ ID NO: 102 and variable light chain sequence of SEQ ID NO: 88;
(e) the variable heavy chain sequence of SEQ ID NO: 103 and variable light chain sequence of SEQ ID NO: 89;
(f) the variable heavy chain sequence of SEQ ID NO: 104 and variable light chain sequence of SEQ ID NO: 90;
(g) the variable heavy chain sequence of SEQ ID NO: 105 and variable light chain sequence of SEQ ID NO: 91;
(h) the variable heavy chain sequence of SEQ ID NO: 106 and variable light chain sequence of SEQ ID NO: 92;
(i) the variable heavy chain sequence of SEQ ID NO: 107 and variable light chain sequence of SEQ ID NO: 93;
(j) the variable heavy chain sequence of SEQ ID NO: 108 and variable light chain sequence of SEQ ID NO: 94;
(k) the variable heavy chain sequence of SEQ ID NO: 109 and variable light chain sequence of SEQ ID NO: 95;
(l) the variable heavy chain sequence of SEQ ID NO: 110 and variable light chain sequence of SEQ ID NO: 96;

(m) the variable heavy chain sequence of SEQ ID NO: 111 and variable light chain sequence of SEQ ID NO: 97; or
(n) the variable heavy chain sequence of SEQ ID NO: 112 and variable light chain sequence of SEQ ID NO: 98.

24. A myostatin antagonist for use according to according to any of claims 13-23, wherein the antibody comprises:

(a) the heavy chain sequence of SEQ ID NO: 146 and light chain sequence of SEQ ID NO: 141;
(b) the heavy chain sequence of SEQ ID NO: 147 and light chain sequence of SEQ ID NO: 142;
(c) the heavy chain sequence of SEQ ID NO: 148 and light chain sequence of SEQ ID NO: 143;
(d) the heavy chain sequence of SEQ ID NO: 149 and light chain sequence of SEQ ID NO: 144;
(e) the heavy chain sequence of SEQ ID NO: 150 and light chain sequence of SEQ ID NO: 145;
(f) the heavy chain sequence of SEQ ID NO: 156 and light chain sequence of SEQ ID NO: 151;
(g) the heavy chain sequence of SEQ ID NO: 157 and light chain sequence of SEQ ID NO: 152;
(h) the heavy chain sequence of SEQ ID NO: 158 and light chain sequence of SEQ ID NO: 153;
(i) the heavy chain sequence of SEQ ID NO: 159 and light chain sequence of SEQ ID NO: 154; or
(j) the heavy chain sequence of SEQ ID NO: 160 and light chain sequence of SEQ ID NO: 155.

25. A myostatin antagonist for use according to according to any of claims 13-24, wherein the antibody comprised in said composition cross-blocks or is cross blocked by at least one antibody of claim 10 from binding to ActRIIB.

26. A myostatin antagonist for use according to according to any of claims 13-25, wherein the antibody comprised in said composition has altered effector function through mutation of the Fc region.

27. A myostatin antagonist for use according to according to any of claims 13-26, wherein the antibody comprised in said composition binds to an epitope recognised by an antibody listed in claims 15-17.

28. A myostatin antagonist for use according to any of claims 13-27, wherein the antibody is encoded by pBW522 (DSM22873) or pBW524 (DSM22874).

29. Bimaghumab for use in treating sporadic inclusion body myositis, wherein bimagrumab is to be administered intraveneously at a dose of about 1-10 mg/kg body weight every four weeks.

30. Bimagrumab for use in treating sporadic inclusion body myositis, wherein bimagrumab is to be administered intraveneously at a dose of about 1 mg/kg body weight every four weeks.

31. Bimagrumab for use in treating sporadic inclusion body myositis, wherein bimagrumab is to be administered intraveneously at a dose of about 3 mg/kg body weight every four weeks.

32. Bimagrumab for use in treating sporadic inclusion body myositis, wherein bimagrumab is to be administered intraveneously at a dose of about 10 mg/kg body weight every four weeks.

33. A method of treating sporadic inclusion body myositis comprising administering a therapeutically effective amount of myostatin antagonist to a patient in need thereof.

34. A method of treating sporadic inclusion body myositis according to claim 33, comprising adminstering said myostatin antagonist at a dose of about 1-10 mg/kg.

35. A method of treating sporadic inclusion body myositis according to any of claims 33-34, comprising administering at a dose of about 1, about 3 or about 10 mg/kg body weight.

36. A method of treating sporadic inclusion body myositis according to any of claims 33-35, comprising administering said myostatin antagonist intraveneously.

37. A method of treating sporadic inclusion body myositis according to any of claims 33-36, comprising administering said myostatin anatagonist every four weeks.

38. A method of treating sporadic inclusion body myositis according to any of claims 33-37, wherein said patient is ambulatory.

39. A method of treating sporadic inclusion body myositis according to any of claims 33-38, wherein treating sporadic inclusion body myositis comprises slowing down the progression of the disease or improving physical function and mobility.

40. A method of treating sporadic inclusion body myositis according to any of claims 33-38, wherein treating sporadic inclusion body myositis comprises improving dysphagia/swallowing difficulties.

41. A method of treating sporadic inclusion body myositis according to any of claims 33-38, wherein treating sporadic inclusion body myositis comprises improving upper extremity strength.

42. A method of treating sporadic inclusion body myositis according to any of claims 33-38, wherein treating sporadic inclusion body myositis comprises reducing incidence of falls or preventing falls.

43. A method of treating sporadic inclusion body myositis according to any of claims 33-42, wherein the myostatin antagonist is a myostatin receptor binding molecule.

44. A method of treating sporadic inclusion body myositis according to any of claims 33-43, wherein the myostatin antagonist is an ActRII receptor antagonist.

45. A method of treating sporadic inclusion body myositis according to any of claims 33-44, wherein the myostatin antagonist is an anti-ActRII receptor antibody.

46. A method of treating sporadic inclusion body myositis according to any of claims 33-45, wherein the anti-ActRII receptor antibody is bimagrumab.

47. A method of treating sporadic inclusion body myositis according to any of claims 45 or 46, wherein the myostatin antagonist is an anti-ActRII antibody that binds to an epitope of ActRIIB consisting of amino acids 19-134 of SEQ ID NO: 181 (SEQ ID NO: 182).

48. A method of treating sporadic inclusion body myositis according to any of claims 45-47, wherein the anti-ActRII antibody binds to an epitope of ActRIIB comprising or consisting of:

(a) amino acids 78-83 of SEQ ID NO: 181 (WLDDFN – SEQ ID NO:188);

(b) amino acids 76-84 of SEQ ID NO: 181 (GCWLDDFNC – SEQ ID NO:186);

(c) amino acids 75-85 of SEQ ID NO: 181 (KGCWLDDFNCY – SEQ ID NO:190);

(d) amino acids 52-56 of SEQ ID NO: 181 (EQDKR – SEQ ID NO:189);

(e) amino acids 49-63 of SEQ ID NO: 181 (CEGEQDKRLHCYASW – SEQ ID NO:187);

(f) amino acids 29-41 of SEQ ID NO: 181 (CIYYNANWELERT– SEQ ID NO:191);

(g) amino acids 100-110 of SEQ ID NO: 181 (YFCCCEGNFCN – SEQ ID NO:192); or

(h) amino acids 78-83 of SEQ ID NO: 181 (WLDDFN) and amino acids 52-56 of SEQ ID NO: 181 (EQDKR).

49. A method of treating sporadic inclusion body myositis according to any of claims 45-48, wherein the anti-ActRIIB antibody is selected from the group consisting of:

a) an anti-ActRIIB antibody that binds to an epitope of ActRIIB comprising :

(a) amino acids 78-83 of SEQ ID NO: 181 (WLDDFN – SEQ ID NO:188);

(b) amino acids 76-84 of SEQ ID NO: 181 (GCWLDDFNC – SEQ ID NO:186);

(c) amino acids 75-85 of SEQ ID NO: 181 (KGCWLDDFNCY – SEQ ID NO:190);

(d) amino acids 52-56 of SEQ ID NO: 181 (EQDKR – SEQ ID NO:189);

(e) amino acids 49-63 of SEQ ID NO: 181 (CEGEQDKRLHCYASW – SEQ ID NO:187);

(f) amino acids 29-41 of SEQ ID NO: 181 (CIYYNANWELERT– SEQ ID NO:191);

(g) amino acids 100-110 of SEQ ID NO: 181 (YFCCCEGNFCN – SEQ ID NO:192); or

(h) amino acids 78-83 of SEQ ID NO: 181 (WLDDFN) and amino acids 52-56 of SEQ ID NO: 181 (EQDKR).

; and b) an antagonist antibody to ActRIIB that binds to an epitope of ActRIIB comprising amino acids 78-83 of SEQ ID NO: 181 (WLDDFN – SEQ ID NO:188);

(b) amino acids 76-84 of SEQ ID NO: 181 (GCWLDDFNC – SEQ ID NO:186);

(c) amino acids 75-85 of SEQ ID NO: 181 (KGCWLDDFNCY – SEQ ID NO:190);

(d) amino acids 52-56 of SEQ ID NO: 181 (EQDKR – SEQ ID NO:189);

(e) amino acids 49-63 of SEQ ID NO: 181 (CEGEQDKRLHCYASW – SEQ ID NO:187);

(f) amino acids 29-41 of SEQ ID NO: 181 (CIYYNANWELERT– SEQ ID NO:191);

(g) amino acids 100-110 of SEQ ID NO: 181 (YFCCCEGNFCN – SEQ ID NO:192); or

(h) amino acids 78-83 of SEQ ID NO: 181 (WLDDFN) and amino acids 52-56 of SEQ ID NO: 181 (EQDKR), wherein the antibody has a K_D of about 2 pM.

50. A method of treating sporadic inclusion body myositis according to any of claims 45-49, wherein the antibody binds to ActRIIB with a 10-fold or greater affinity than it binds to ActRIIA.

51. A method of treating sporadic inclusion body myositis to any of claims 45-50, wherein the antibody comprises a heavy chain variable region CDR1 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 1-14; a heavy chain variable region CDR2 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 15-28; a heavy chain variable region CDR3 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 29-42; a light chain variable region CDR1 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 43-56; a light chain variable region CDR2 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 57-70; and a light chain variable region CDR3 comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 71-84.

52. A method of treating sporadic inclusion body myositis according to any of claims 45-51, wherein the antibody comprises:

(a) a heavy chain variable region CDR1 of SEQ ID NO: 1; a heavy chain variable region CDR2 of SEQ ID NO: 15; a heavy chain variable region CDR3 of SEQ ID NO: 29; a light chain variable region CDR1 of SEQ ID NO: 43; a light chain variable region CDR2 of SEQ ID NO: 57; and a light chain variable region CDR3 of SEQ ID NO: 71,

(b) a heavy chain variable region CDR1 of SEQ ID NO: 2; a heavy chain variable region CDR2 of SEQ ID NO: 16; a heavy chain variable region CDR3 of SEQ ID NO: 30; a light chain variable region CDR1 of SEQ ID NO: 44; a light chain variable region CDR2 of SEQ ID NO: 58; and a light chain variable region CDR3 of SEQ ID NO: 72,

(c) a heavy chain variable region CDR1 of SEQ ID NO: 3; a heavy chain variable region CDR2 of SEQ ID NO: 17; a heavy chain variable region CDR3 of SEQ ID

NO: 31; a light chain variable region CDR1 of SEQ ID NO: 45; a light chain variable region CDR2 of SEQ ID NO: 59; and a light chain variable region CDR3 of SEQ ID NO: 73,

(d) a heavy chain variable region CDR1 of SEQ ID NO: 4; a heavy chain variable region CDR2 of SEQ ID NO: 18; a heavy chain variable region CDR3 of SEQ ID NO: 32; a light chain variable region CDR1 of SEQ ID NO: 46; a light chain variable region CDR2 of SEQ ID NO: 60; and a light chain variable region CDR3 of SEQ ID NO: 74,

(e) a heavy chain variable region CDR1 of SEQ ID NO: 5; a heavy chain variable region CDR2 of SEQ ID NO: 19; a heavy chain variable region CDR3 of SEQ ID NO: 33; a light chain variable region CDR1 of SEQ ID NO: 47; a light chain variable region CDR2 of SEQ ID NO: 61; and a light chain variable region CDR3 of SEQ ID NO: 75,

(f) a heavy chain variable region CDR1 of SEQ ID NO: 6; a heavy chain variable region CDR2 of SEQ ID NO: 20; a heavy chain variable region CDR3 of SEQ ID NO: 34; a light chain variable region CDR1 of SEQ ID NO: 48; a light chain variable region CDR2 of SEQ ID NO: 62; and a light chain variable region CDR3 of SEQ ID NO: 76,

(g) a heavy chain variable region CDR1 of SEQ ID NO: 7; a heavy chain variable region CDR2 of SEQ ID NO: 21; a heavy chain variable region CDR3 of SEQ ID NO: 35; a light chain variable region CDR1 of SEQ ID NO: 49; a light chain variable region CDR2 of SEQ ID NO: 63; and a light chain variable region CDR3 of SEQ ID NO: 77,

(h) a heavy chain variable region CDR1 of SEQ ID NO: 8; a heavy chain variable region CDR2 of SEQ ID NO: 22; a heavy chain variable region CDR3 of SEQ ID NO: 36; a light chain variable region CDR1 of SEQ ID NO: 50 a light chain variable region CDR2 of SEQ ID NO: 64; and a light chain variable region CDR3 of SEQ ID NO: 78,

(i) a heavy chain variable region CDR1 of SEQ ID NO: 9; a heavy chain variable region CDR2 of SEQ ID NO: 23; a heavy chain variable region CDR3 of SEQ ID NO: 37; a light chain variable region CDR1 of SEQ ID NO: 51; a light chain variable region CDR2 of SEQ ID NO: 65; and a light chain variable region CDR3 of SEQ ID NO: 79,

(j) a heavy chain variable region CDR1 of SEQ ID NO: 10; a heavy chain variable region CDR2 of SEQ ID NO: 24; a heavy chain variable region CDR3 of SEQ ID

NO: 38; a light chain variable region CDR1 of SEQ ID NO: 52; a light chain variable region CDR2 of SEQ ID NO: 66; and a light chain variable region CDR3 of SEQ ID NO: 80,

(k) a heavy chain variable region CDR1 of SEQ ID NO: 11; a heavy chain variable region CDR2 of SEQ ID NO: 25; a heavy chain variable region CDR3 of SEQ ID NO: 39; a light chain variable region CDR1 of SEQ ID NO: 53; a light chain variable region CDR2 of SEQ ID NO: 67; and a light chain variable region CDR3 of SEQ ID NO: 81,

(l) a heavy chain variable region CDR1 of SEQ ID NO: 12; a heavy chain variable region CDR2 of SEQ ID NO: 26; a heavy chain variable region CDR3 of SEQ ID NO: 40; a light chain variable region CDR1 of SEQ ID NO: 54; a light chain variable region CDR2 of SEQ ID NO: 68; and a light chain variable region CDR3 of SEQ ID NO: 82,

(m) a heavy chain variable region CDR1 of SEQ ID NO: 13; a heavy chain variable region CDR2 of SEQ ID NO: 27; a heavy chain variable region CDR3 of SEQ ID NO: 41; a light chain variable region CDR1 of SEQ ID NO: 55; a light chain variable region CDR2 of SEQ ID NO: 69; and a light chain variable region CDR3 of SEQ ID NO: 83, or

(n) a heavy chain variable region CDR1 of SEQ ID NO: 14; a heavy chain variable region CDR2 of SEQ ID NO: 28; a heavy chain variable region CDR3 of SEQ ID NO: 42; a light chain variable region CDR1 of SEQ ID NO: 56; a light chain variable region CDR2 of SEQ ID NO: 70; and a light chain variable region CDR3 of SEQ ID NO: 84.

53. A method of treating sporadic inclusion body myositis according to any of claims 45-52, wherein the antibody comprises a full length heavy chain amino acid sequence having at least 95% sequence identity to at least one sequence selected from the group consisting of SEQ ID NOs:146-150 and 156-160.

54. A method of treating sporadic inclusion body myositis to according to any of claims 45-53, wherein the antibody comprises a full length light chain amino acid sequence having at least 95% sequence identity to at least one sequence selected from the group consisting of SEQ ID NOs:141-145 and 151-155.

55. A method of treating sporadic inclusion body myositis according to any of claims 45-54, wherein the antibody comprises:

(a) the variable heavy chain sequence of SEQ ID NO: 99 and variable light chain sequence of SEQ ID NO: 85;

(b) the variable heavy chain sequence of SEQ ID NO: 100 and variable light chain sequence of SEQ ID NO: 86;

(c) the variable heavy chain sequence of SEQ ID NO: 101 and variable light chain sequence of SEQ ID NO: 87;

(d) the variable heavy chain sequence of SEQ ID NO: 102 and variable light chain sequence of SEQ ID NO: 88;

(e) the variable heavy chain sequence of SEQ ID NO: 103 and variable light chain sequence of SEQ ID NO: 89;

(f) the variable heavy chain sequence of SEQ ID NO: 104 and variable light chain sequence of SEQ ID NO: 90;

(g) the variable heavy chain sequence of SEQ ID NO: 105 and variable light chain sequence of SEQ ID NO: 91;

(h) the variable heavy chain sequence of SEQ ID NO: 106 and variable light chain sequence of SEQ ID NO: 92;

(i) the variable heavy chain sequence of SEQ ID NO: 107 and variable light chain sequence of SEQ ID NO: 93;

(j) the variable heavy chain sequence of SEQ ID NO: 108 and variable light chain sequence of SEQ ID NO: 94;

(k) the variable heavy chain sequence of SEQ ID NO: 109 and variable light chain sequence of SEQ ID NO: 95;

(l) the variable heavy chain sequence of SEQ ID NO: 110 and variable light chain sequence of SEQ ID NO: 96;

(m) the variable heavy chain sequence of SEQ ID NO: 111 and variable light chain sequence of SEQ ID NO: 97; or

(n) the variable heavy chain sequence of SEQ ID NO: 112 and variable light chain sequence of SEQ ID NO: 98.

56. A method of treating sporadic inclusion body myositis according to any of claims 45-55, wherein the antibody comprises:

(a) the heavy chain sequence of SEQ ID NO: 146 and light chain sequence of SEQ ID NO: 141;

(b) the heavy chain sequence of SEQ ID NO: 147 and light chain sequence of SEQ ID NO: 142;

(c) the heavy chain sequence of SEQ ID NO: 148 and light chain sequence of SEQ ID NO: 143;
(d) the heavy chain sequence of SEQ ID NO: 149 and light chain sequence of SEQ ID NO: 144;
(e) the heavy chain sequence of SEQ ID NO: 150 and light chain sequence of SEQ ID NO: 145;
(f) the heavy chain sequence of SEQ ID NO: 156 and light chain sequence of SEQ ID NO: 151;
(g) the heavy chain sequence of SEQ ID NO: 157 and light chain sequence of SEQ ID NO: 152;
(h) the heavy chain sequence of SEQ ID NO: 158 and light chain sequence of SEQ ID NO: 153;
(i) the heavy chain sequence of SEQ ID NO: 159 and light chain sequence of SEQ ID NO: 154; or
(j) the heavy chain sequence of SEQ ID NO: 160 and light chain sequence of SEQ ID NO: 155.

57. A method of treating sporadic inclusion body myositis according to any of claims 45-56, wherein the antibody comprised in said composition cross-blocks or is cross blocked by at least one antibody of aspect 10 from binding to ActRIIB.

58. A method of treating sporadic inclusion body myositis according to any of claims 45-57, wherein the antibody comprised in said composition has altered effector function through mutation of the Fc region.

59. A method of treating sporadic inclusion body myositis according to any of claims 45-58, wherein the antibody comprised in said composition binds to an epitope recognised by an antibody listed in claims 46-48.

60. A method of treating sporadic inclusion body myositis according to any of claims 45-59, wherein the antibody is encoded by pBW522 (DSM22873) or pBW524 (DSM22874).

61. A method of treating sporadic inclusion body myositis comprsing administering bimagrumab,

62. A method of treating sporadic inclusion body myositis comprsing administering bimagrumab,wherein bimagrumab is to be administered intraveneously at a dose of about 1-10 mg/kg body weight every four weeks.

63. A method of treating sporadic inclusion body myositis comprsing administering bimagrumab,, wherein bimagrumab is to be administered intraveneously at a dose of about 1 mg/kg body weight every four weeks.

64. A method of treating sporadic inclusion body myositis comprsing administering bimagrumab,, wherein bimagrumab is to be administered intraveneously at a dose of about 3 mg/kg body weight every four weeks.

65. A method of treating sporadic inclusion body myositis comprsing administering bimagrumab,, wherein bimagrumab is to be administered intraveneously at a dose of about 10 mg/kg body weight every four weeks.

66. Bimagrumab for use in treating sporadic inclusion body myositis.

67. A composition comprising 150 mg/ml of bimagrumab for use in a method of treating sporadic inclusion body myositis.

68. A unitary dosage form comprising 150 mg/ml of bimagrumab.

69. An infusion bag comprising an appropriate amount of bimagrumab from one or more vials diluted with a solution.