ALPHA-4-BETA-7 HETERODIMER SPECIFIC ANTAGONIST ANTIBODY

CROSS-REFERENCE TO RELATED APPLICATIONS

This application claims the benefit under 35 U.S.C. 119(e) of United States patent application number 61/162,154, filed March 20, 2009 and United States patent application number 61/306,829, filed February 22, 2010, which are incorporated herein by reference.

FIELD OF THE INVENTION

This application provides compositions and methods relating to alpha4beta7 heterodimer-specific antigen binding proteins.

BACKGROUND

Integrins are heterodimeric Type I transmembrane proteins formed of two subunits (one alpha subunit and one beta subunit), and mediate many different cell-cell and cell-extracellular matrix interactions. Functionally, integrins have been shown to be involved in diverse biological processes, including leukocyte migration and recirculation and the immune response. In mammals, there are 18 known alpha subunits and eight known beta subunits, which combine to form 24 distinct integrins. Ligand specificity is determined in large part by the particular combinations of alpha and beta subunits expressed, while affinity for ligand is modulated by integrin conformational changes and is divalent-cation dependent.

The ligands for integrins form a structurally diverse group that includes extracellular matrix proteins such as collagens, fibronection, vitronectin and laminins; counter-receptors such as the cellular adhesion molecules (for example, vascular cellular adhesion molecule or VCAM), and plasma proteins. Numerous pathogenic microorganisms also utilize integrins to initiate infection or as sites for toxin binding. The structurally diverse ligands share an exposed glutamic or aspartic acid residue, usually present in an extended, flexible loop, which is important for recognition by integrins.

The alpha4 integrins (alpha 4 partnered with either the beta1 or beta7 subunit) play an important role in the immune system. Alpha4beta1 is expressed on lymphocytes and myeloid cells; it appears to be the major binding partner for vascular cell adhesion molecule (VCAM). VCAM is ubiquitously expressed on vascular endothelium, is up regulated during inflammation, and binds alpha4beta7 as well as alpha4beta1 (albeit weakly to alpha4beta7). Though also detected on d peripheral T cells, B cells, NK cells and eosinphils, alpha4beta7 is most highly expressed on a subpopulation of CD4+CD45RA- memory T cells which has been shown to preferentially home to the gut. The primary ligand for the alpha4beta7 heterodimer is mucosal addressin cell adhesion molecule 1 (MAdCAM-1 or MAdCAM), which is expressed in gut endothelium.

In addition to pairing with the alpha4 chain, the beta7 subunit also partners with alphaE to form alphaEbeta7, which is primarily expressed on intraepithelial lymphocytes (IEL) in intestine, lung and genitourinary tract. AlphaEbeta7 is also expressed on dendritic cells in the gut. The alphaEbeta7 heterodimer binds to E-cadherin, which is expressed on epithelial cells. The IEL cells are thought to provide a mechanism for immunosurveillance within the epithelial compartment.

Antibodies that bind alpha4 and inhibit binding of alpha4beta1 to VCAM-1 and fibronection mapped to a 52-amino acid region of alpha4, between residues 152 and 203 (Schiffer et al., J. Biol. Chem. 270:14270; 1995). Tidswell et al. (J. Immuno 159:1497; 1997) identified domains of beta7 that are important in binding to MAdCAM-1, utilizing a panel of antibodies that bind beta7 in a mouse/human chimeric beta7 subunit approach. They found that six of seven antibodies that inhibited binding to MAdCAM-1 and E-cadherin mapped to a region comprising amino acids 176 through 250, which appears to have homology to the metal-ion dependent adhesion site (MIDAS) of other integrin subunits. One of the antibodies used by Tidswell et al. was an alpha4beta7 heterodimer specific antibody referred to as ACT-1.

The ACT-1 antibody was originally described by Lazarovitz et al. (J. Immunol. 133:1857; 1984) as an antibody developed by immunizing mice with human tetanus toxoid-specific T lymphocyte line from PBMC. Later it was shown that ACT-1 binds to the alpha4beta7 heterodimer specifically (Schweighoffer et al., J. Immunol. 151:717, 1993). While ACT-1 does not bind murine alpha4beta7, it does bind alpha4beta7 from least some non-human primate species, and has been shown to attenuate spontaneous colitis in captive cotton-top tamarins (Hesterberg et al., Gastroenterology 111:1373; 1996)

ACT-1 has been humanized and evaluated as a human therapeutic in ulcerative colitis (Feagan et al., N Engl J Med. 352:2499; 2005), and recently in Crohn's disease (Feagan et al, Clinical Gastroenterology and Hepatology, 6:1370, 2008). Humanized ACT-1, also known as vedolizumab, is described in WO 98/06248 and US patent 7,147,85, as well as WO 07/061679 and US 2007-0122404. Another humanized antibody, natalizumab (Tysabri®), has been used to treat Crohn's disease. Natalizumab is a humanized version of an alpha4-specific murine antibody. Vedolizumab has been shown to lead to a neutralizing anti-humanized antibody response in a portion of patients, and natalizumab has been associated with progressive multifocal leukoencephalopathy (PML), a neurological disorder that is associated with reactivation of prior infection with JC virus in immunocompromised individuals. Accordingly, there is a need for a therapeutic agent that ameliorates these disadvantages while disrupting the alpha4beta7/MAdCAM-1 pathway.

SUMMARY OF THE INVENTION

In a first aspect, the present invention refers to an isolated, alpha4beta7 heterodimer specific antigen binding protein

having a heavy chain variable region comprising CDR1, CDR2 and CDR3 from SEQ ID NO:37, and a light chain variable region comprising CDR1, CDR2 and CDR3 from SEQ ID NO:9.

In a second aspect, the present invention refers to an isolated, alpha4beta7 heterodimer specific antigen binding

protein having a heavy chain variable region and a light chain variable region, wherein the heavy chain variable region is encoded by a nucleic acid that encodes SEQ ID NO:37, and the light chain variable region is encoded by a nucleic acid that encodes SEQ ID NO:9.

In a third aspect, the present invention refers to a composition for treating inflammatory bowel disease, pancreatitis,

insulin-dependent diabetes mellitus, mastitis, cholecystitis, cholangitis, pericholangitis, chronic bronchitis, chronic sinusitis, asthma, or graft versus host disease, comprising the protein as defined in anyone of claims 1 to 18 and a physiologically acceptable diluent, excipient or carrier.

In a fourth aspect, the present invention refers to an isolated nucleic acid that encodes the antigen binding protein of the present invention.

In a fifth aspect, the present invention refers to a vector comprising the nucleic acid of the present invention.

In a sixth aspect, the present invention refers to an isolated host cell transfected or transformed with the vector of the present invention.

In a seventh aspect, the present invention refers to a method for the production of an antigen binding protein comprising culturing the host cell of the present invention to express the protein and recovering the protein from the culture medium.

In one example, it is described an isolated antigen binding protein that specifically binds to human alpha4beta7 (i.e., an alpha4beta7 heterodimer specific antigen binding protein). In another example, the antigen binding protein specifically binds to the alpha4beta7 of a non-human primate, a cynomologous monkey, a chimpanzee, a non-primate mammal, a rodent, a mouse, a rat, a hamster, a guinea pig, a cat, or a dog. In another example, the isolated antigen binding protein comprises a human antibody; a chimeric antibody; a monoclonal antibody; a recombinant antibody; an antigen-binding antibody fragment; a single chain antibody; a diabody; a triabody; a tetrabody; a Fab fragment; a $F(ab')_2$ fragment; a domain antibody; an IgD antibody; an IgE antibody; an IgM antibody; an IgG1 antibody; an IgG2 antibody; an IgG3 antibody; an IgG4 antibody; or an IgG4 antibody having at least one mutation in a hinge region that alleviates a tendency to form intra-H chain disulfide bond. In another example, the isolated antigen binding protein comprises a heavy chain constant region from one of the aforementioned antibodies; in another example, the constant region is a polypeptide comprising SEQ ID NO:72; a polypeptide at least 90% identical to SEQ ID NO:72; a

polypeptide having an amino acid sequence as set forth in SEQ ID NO:72 from which one, two, three, four or five N-terminal and/or C-terminal amino acids have been removed; or one of the aforementioned polypeptides which incorporates one or more post-translational modifications. In one example, the isolated antigen binding protein comprises a kappa light chain constant region, in another it comprises a lambda light chain region. In one example, the light chain constant region is a polypeptide comprising SEQ ID NO:70; a polypeptide at least 90% identical to SEQ ID NO:70; a polypeptide having an amino acid sequence as set forth in SEQ ID NO:70 from which one, two, three, four or five N-terminal and/or C-terminal amino acids have been removed; or one of the aforementioned polypeptides which incorporates one or more post-translational modifications

One example, it is described an alpha4beta7 heterodimer specific antigen binding protein having a heavy chain and a light chain, each of which comprise one or more complementarity determining regions, or CDRs. In another example, the heavy chain variable region comprises CDR1, CDR2 and CDR3 and a light chain variable region comprises CDR1, CDR2 and CDR3, wherein each respective CDR is selected from the group consisting of the light chain CDR1, CDR2 and CDR3 from SEQ ID NO:55, and the heavy chain CDR1, CDR2 and CDR3 from SEQ ID NO:58; the light chain CDR1, CDR2 and CDR3 from SEQ ID NO:56, and the heavy chain CDR1, CDR2 and CDR3 from SEQ ID NO:59; and the light chain CDR1, CDR2 and CDR3 from SEQ ID NO:57, and the heavy chain CDR1, CDR2 and CDR3 from SEQ ID NO:60.

In another example, the heavy chain variable region further comprises four framework regions (FRs) designated FR1, FR2, FR3 andFR4, and the light chain variable region further comprises four framework regions (FRs) designated FR1, FR2, FR3 andFR4. In one aspect, the FRs are selected from the same SEQ ID NO as the CDRs; in another, the FRs are selected from a different SEQ ID NO. In a further example, it is described an alpha4beta7 heterodimer specific antigen binding protein wherein the light chain variable region comprises SEQ ID NO:55, and the heavy chain variable region comprises SEQ ID NO:58; the light chain variable region comprises SEQ ID NO:56, and the heavy chain variable region comprises SEQ ID NO:59; or the light chain variable region comprises SEQ ID NO:57, and the heavy chain variable region comprises SEQ ID NO:60.

In another example, it is described an isolated alpha4beta7 heterodimer specific antigen binding protein, having a heavy chain and a light chain, each of which comprise one or more complementarity determining regions, or CDRs. In another example, the heavy chain variable region comprises CDR1, CDR2 and CDR3 and the light chain variable region comprises CDR1, CDR2 and CDR3. In one example, the light chain CDRs are selected from the group consisting of a CDR1, CDR2 and CDR3 at least 90% identical to a CDR1, CDR2 and CDR3, respectively, of SEQ ID NO: 3; a CDR1, CDR2 and CDR3 at least 90% identical to a CDR1, CDR2 and CDR3, respectively, of SEQ ID NO: 5; a CDR1, CDR2 and CDR3 at least 90% identical to a CDR1, CDR2 and CDR3, respectively, of SEQ ID NO: 7; a CDR1, CDR2 and CDR3 at least 90% identical to a CDR1, CDR2 and CDR3, respectively, of SEQ ID NO: 22; and a CDR1, CDR2 and CDR3 at least 90% identical to a CDR1, CDR2 and CDR3, respectively, of SEQ ID NO: 24; and the heavy chain variable CDR1, CDR2 and CDR3 are from SEQ ID NO:58.

In another example, the heavy chain variable region further comprises four framework regions (FRs) designated FR1, FR2, FR3 andFR4, and the light chain variable region further comprises four framework regions (FRs) designated FR1, FR2, FR3 andFR4. In one example, the FRs are selected from the same SEQ ID NO as the CDRs; in another, the FRs are selected from a different SEQ ID NO. In a further example, it is described an alpha4beta7 heterodimer specific antigen binding protein wherein the light chain variable region is selected from the group consisting of a light chain variable region at least 90% identical to SEQ ID NO:3; a light chain variable region at least 90% identical to SEQ ID NO:5; a light chain variable region at least 90% identical to SEQ ID NO:7; a light chain variable region at least 90% identical to SEQ ID NO:22; and a light chain variable region at least 90% identical to SEQ ID NO:24; and the heavy chain variable region comprises SEQ ID NO:58.

Another example describes an isolated, alpha4beta7 heterodimer specific antigen binding protein having a heavy chain variable region comprising CDR1, CDR2 and CDR3 and a light chain variable region comprising CDR1, CDR2 and CDR3, wherein the light chain CDR1, CDR2 and CDR3 are selected from the group consisting of a CDR1, CDR2 and CDR3 at least 90% identical to a CDR1, CDR2 and CDR3, respectively, of SEQ ID NO:12; a CDR1, CDR2 and CDR3 at least 90% identical to a CDR1, CDR2 and CDR3, respectively, of SEQ ID NO: 25; and a CDR1, CDR2 and CDR3 at least 90% identical to a CDR1, CDR2 and CDR3, respectively, of SEQ ID NO: 26; and the heavy chain CDR1, CDR2 and CDR3 are selected from the group consisting of a CDR1, CDR2 and CDR3 at least 90% identical to a CDR1, CDR2 and CDR3, respectively, of SEQ ID NO:41; and a CDR1, CDR2 and CDR3 at least 90% identical to a CDR1, CDR2 and CDR3, respectively, of SEQ ID NO:54. In one example, the light chain variable region is selected from the group consisting of variable regions that are at least 90% identical to any one of SEQ ID NOs: 12, 25 and 26, and the heavy variable region is selected from the group consisting of variable regions that are at least 90% identical to any one of SEQ ID NOs:41 and 54. In another example, the heavy chain variable region further comprises four framework regions (FRs) designated FR1, FR2, FR3 andFR4, and the light chain variable region further comprises four framework regions (FRs) designated FR1, FR2, FR3 andFR4. In one example, the FRs are selected from the same SEQ ID NO as the CDRs; in another, the FRs are selected from a different SEQ ID NO.

In one example, it is described an isolated, alpha4beta7 heterodimer specific antigen binding protein having a heavy chain variable region comprising CDR1, CDR2 and CDR3 and a light chain variable region comprising CDR1, CDR2 and CDR3, wherein each respective CDR is at least 90% identical to a CDR selected from the group consisting of a light chain CDR1, CDR2 and CDR3 from SEQ ID NO:10, and a heavy chain CDR1, CDR2 and CDR3 from SEQ ID NO:38; a light chain CDR1, CDR2 and CDR3 from SEQ ID NO:2, and a heavy chain CDR1, CDR2 and CDR3 from SEQ ID NO:30; a light chain CDR1, CDR2 and CDR3 from SEQ ID NO:20, and a heavy chain CDR1, CDR2 and CDR3 from SEQ ID NO:51; a light chain CDR1, CDR2 and CDR3 from SEQ ID NO:11, and a heavy chain CDR1, CDR2 and CDR3 from SEQ ID NO:39; a light chain CDR1, CDR2 and CDR3 from SEQ ID NO:13, and a heavy chain CDR1, CDR2 and CDR3 from SEQ ID NO:42; a light chain CDR1, CDR2 and CDR3 from SEQ ID NO:17, and a heavy chain CDR1, CDR2 and CDR3 from

SEQ ID NO:46; a light chain CDR1, CDR2 and CDR3 from SEQ ID NO:8, and a heavy chain CDR1, CDR2 and CDR3 from SEQ ID NO:36; a light chain CDR1, CDR2 and CDR3 from SEQ ID NO:19, and a heavy chain CDR1, CDR2 and CDR3 from SEQ ID NO:49; a light chain CDR1, CDR2 and CDR3 from SEQ ID NO:18, and a heavy chain CDR1, CDR2 and CDR3 from SEQ ID NO:47; a light chain CDR1, CDR2 and CDR3 from SEQ ID NO:21, and a heavy chain CDR1, CDR2 and CDR3 from SEQ ID NO:52; a light chain CDR1, CDR2 and CDR3 from SEQ ID NO:3, and a heavy chain CDR1, CDR2 and CDR3 from SEQ ID NO:31; a light chain CDR1, CDR2 and CDR3 from SEQ ID NO:7, and a heavy chain CDR1, CDR2 and CDR3 from SEQ ID NO:35; a light chain CDR1, CDR2 and CDR3 from SEQ ID NO:6, and a heavy chain CDR1, CDR2 and CDR3 from SEQ ID NO:34; a light chain CDR1, CDR2 and CDR3 from SEQ ID NO:1, and a heavy chain CDR1, CDR2 and CDR3 from SEQ ID NO:29; a light chain CDR1, CDR2 and CDR3 from SEQ ID NO:22, and a heavy chain CDR1, CDR2 and CDR3 from SEQ ID NO:50; a light chain CDR1, CDR2 and CDR3 from SEQ ID NO:24, and a heavy chain CDR1, CDR2 and CDR3 from SEQ ID NO:40; a light chain CDR1, CDR2 and CDR3 from SEQ ID NO:9, and a heavy chain CDR1, CDR2 and CDR3 from SEQ ID NO:37; a light chain CDR1, CDR2 and CDR3 from SEQ ID NO:4, and a heavy chain CDR1, CDR2 and CDR3 from SEQ ID NO:32; a light chain CDR1, CDR2 and CDR3 from SEQ ID NO:28, and a heavy chain CDR1, CDR2 and CDR3 from SEQ ID NO:53; a light chain CDR1, CDR2 and CDR3 from SEQ ID NO:16, and a heavy chain CDR1, CDR2 and CDR3 from SEQ ID NO:45; a light chain CDR1, CDR2 and CDR3 from SEQ ID NO:15, and a heavy chain CDR1, CDR2 and CDR3 from SEQ ID NO:44; a light chain CDR1, CDR2 and CDR3 from SEQ ID NO:14, and a heavy chain CDR1, CDR2 and CDR3 from SEQ ID NO:43; a light chain CDR1, CDR2 and CDR3 from SEQ ID NO:27, and a heavy chain CDR1, CDR2 and CDR3 from SEQ ID NO:43; a light chain CDR1, CDR2 and CDR3 from SEQ ID NO:5, and a heavy chain CDR1, CDR2 and CDR3 from SEQ ID NO:33; a light chain CDR1, CDR2 and CDR3 from SEQ ID NO:12, and a heavy chain CDR1, CDR2 and CDR3 from SEQ ID NO:41; a light chain CDR1, CDR2 and CDR3 from SEQ ID NO:23, and a heavy chain CDR1, CDR2 and CDR3 from SEQ ID NO:48; a light chain CDR1, CDR2 and CDR3 from SEQ ID NO:25, and a heavy chain CDR1, CDR2 and CDR3 from SEQ ID NO:54; and a light chain CDR1, CDR2 and CDR3 from SEQ ID NO:26, and a heavy chain CDR1, CDR2 and CDR3 from SEQ ID NO:54. In another example, the heavy chain and light chain CDRs are identical to the respective CDRs of the recited SEQ ID NOs. In one example, the heavy chain variable region further comprises four framework regions (FRs) designated FR1, FR2, FR3 andFR4, and the light chain variable region further comprises four framework regions (FRs) designated FR1, FR2, FR3 andFR4. In one example, the FRs are selected from the same SEQ ID NO as the CDRs; in another, the FRs are selected from a different SEQ ID NO.

In another example, an alpha4beta7 heterodimer specific antigen binding protein comprises a light chain variable region and a heavy chain variable region, wherein the light chain variable region is at least 90% identical to SEQ ID NO:10, and the heavy chain variable region is at least 90% identical to SEQ ID NO:38; the light chain variable region is at least 90% identical to SEQ ID NO:2, and the heavy chain variable region is at least 90% identical to SEQ ID NO:30; the light chain variable region is at least 90% identical to SEQ ID NO:20, and the heavy chain variable region is at least 90% identical to SEQ ID NO:51; the light chain variable region is at least 90% identical to SEQ

ID NO:11, and the heavy chain variable region is at least 90% identical to SEQ ID NO:39; the light chain variable region is at least 90% identical to SEQ ID NO:13, and the heavy chain variable region is at least 90% identical to SEQ ID NO:42; the light chain variable region is at least 90% identical to SEQ ID NO:17, and the heavy chain variable region is at least 90% identical to SEQ ID NO:46; the light chain variable region is at least 90% identical to SEQ ID NO:8, and the heavy chain variable region is at least 90% identical to SEQ ID NO:36; the light chain variable region is at least 90% identical to SEQ ID NO:19, and the heavy chain variable region is at least 90% identical to SEQ ID NO:49; the light chain variable region is at least 90% identical to SEQ ID NO:18, and the heavy chain variable region is at least 90% identical to SEQ ID NO:47; the light chain variable region is at least 90% identical to SEQ ID NO:21, and the heavy chain variable region is at least 90% identical to SEQ ID NO:52; the light chain variable region is at least 90% identical to SEQ ID NO:3, and the heavy chain variable region is at least 90% identical to SEQ ID NO:31; the light chain variable region is at least 90% identical to SEQ ID NO:7, and the heavy chain variable region is at least 90% identical to SEQ ID NO:35; the light chain variable region is at least 90% identical to SEQ ID NO:6, and the heavy chain variable region is at least 90% identical to SEQ ID NO:34; the light chain variable region is at least 90% identical to SEQ ID NO:1, and the heavy chain variable region is at least 90% identical to SEQ ID NO:29; the light chain variable region is at least 90% identical to SEQ ID NO:22, and the heavy chain variable region is at least 90% identical to SEQ ID NO:50; the light chain variable region is at least 90% identical to SEQ ID NO:24, and the heavy chain variable region is at least 90% identical to SEQ ID NO:40; the light chain variable region is at least 90% identical to SEQ ID NO:9, and the heavy chain variable region is at least 90% identical to SEQ ID NO:37; the light chain variable region is at least 90% identical to SEQ ID NO:4, and the heavy chain variable region is at least 90% identical to SEQ ID NO:32; the light chain variable region is at least 90% identical to SEQ ID NO:28, and the heavy chain variable region is at least 90% identical to SEQ ID NO:53; the light chain variable region is at least 90% identical to SEQ ID NO:16, and the heavy chain variable region is at least 90% identical to SEQ ID NO:45; the light chain variable region is at least 90% identical to SEQ ID NO:15, and the heavy chain variable region is at least 90% identical to SEQ ID NO:44; the light chain variable region is at least 90% identical to SEQ ID NO:14, and the heavy chain variable region is at least 90% identical to SEQ ID NO:43; the light chain variable region is at least 90% identical to SEQ ID NO:27, and the heavy chain variable region is at least 90% identical to SEQ ID NO:43; the light chain variable region is at least 90% identical to SEQ ID NO:5, and the heavy chain variable region is at least 90% identical to SEQ ID NO:33; the light chain variable region is at least 90% identical to SEQ ID NO:12, and the heavy chain variable region is at least 90% identical to SEQ ID NO:41; the light chain variable region is at least 90% identical to SEQ ID NO:23, and the heavy chain variable region is at least 90% identical to SEQ ID NO:48; the light chain variable region is at least 90% identical to SEQ ID NO:25, and the heavy chain variable region is at least 90% identical to SEQ ID NO:54; or the light chain variable region is at least 90% identical to SEQ ID NO:26, and the heavy chain variable region is at least 90% identical to SEQ ID NO:54. In another example, the heavy chain and light chain variable regions are identical to the respective variable regions of the recited SEQ ID NOs.

One example describes an isolated, alpha4beta7 heterodimer specific antigen binding protein having an EC50 of less than 35 ng/ml in a CD4+ memory T cell binding assay; another describes an isolated, alpha4beta7 heterodimer specific antigen binding which has an EC50 of less than 10 ng/ml in a CD4+ memory T cell binding assay. In another example, it is described an isolated, alpha4beta7 heterodimer specific antigen binding protein having an IC50 in a MAdCAM competition assay of less than 30 ng/m; in another is provided an isolated, alpha4beta7 heterodimer specific antigen binding which has an IC50 of less than 10 ng/ml in a MAdCAM competition assay. One example describes an isolated, alpha4beta7 heterodimer specific antigen binding protein that binds an S250N mutant of alha4beta7.

In one example, it is described nucleic acids encoding the aforementioned polypeptides. In another example the nucleic acid is a vector. In another example, host cells transformed or transfected with the inventive nucleic acids are described. In another example, there is described a method of preparing a polypeptide comprising incubating the host cells under conditions promoting expression of the polypeptides and harvesting the polypeptides.

In another example, it is described an isolated cell that secretes an antigen binding protein that binds alpha4beta7. In another example, the cell is a hybridoma. In another example, it is described a method of making an antigen binding protein that specifically binds alpha4beta7 (i.e., human alpha4beta7), comprising incubating said isolated cell under conditions that allow it to express said antigen binding protein.

In one example, it is described an isolated antigen binding protein that specifically binds to an alpha4beta7 heterodimer. In another example, the isolated antigen binding protein, when bound to a human alpha4beta7, inhibits binding of alpha4beta7 to MAdCAM-1. Accordingly, one example describes a method of inhibiting at least one activity of alpha4beta7, comprising contacting a cell expressing alpha4beta7 with an alpha4beta7 heterodimer-specific antigen binding protein such that the activity is partially or fully inhibited. In one example, such method is carried out in vivo. In one example, the isolated antigen binding protein inhibits adhesion of cells expressing alpha4beta7 to cells expressing MAdCAM-1. In yet another example, the isolated antigen binding protein inhibits trafficking of cells expressing alpha4beta7 to areas or tissues populated by cells expressing MAdCAM-1; in one example, the isolated antigen binding proteins inhibit trafficking of lymphocytes to the gut.

In another example, it is described a pharmaceutical composition comprising the antigen binding protein. In one example, it is described a method of treating a condition in a subject comprising administering the pharmaceutical composition to the subject, wherein the condition is treatable by reducing the activity (partially or fully) of alpha4beta7 in the subject. In another example, the subject is a human being. In another example, the condition is an inflammatory condition of the gastrointestinal system. Thus, there is described a method of treating an individual afflicted with a condition characterized by inappropriate trafficking of cells expressing alpha4beta7 to tissues comprising cells expressing MAdCAM, comprising administering to the individual an alpha4beta7 heterodimer specific antigen binding protein in am amount sufficient to

inhibit (partially or fully) the trafficking of cells expressing alpha4beta7 to tissues comprising cells expressing MAdCAM. In one example, the condition is inflammatory bowel disease, for example, ulcerative colitis, Crohn's disease, Celiac disease (nontropical Sprue), enteropathy associated with seronegative arthropathies, microscopic or collagenous colitis, eosinophilic gastroenteritis, or pouchitis resulting after proctocolectomy and ileoanal anastomosis. In another example, the condition is s pancreatitis, insulin-dependent diabetes mellitus, mastitis, cholecystitis, cholangitis, pericholangitis, chronic bronchitis, chronic sinusitis, asthma or graft versus host disease.

In another example, the method further comprises administering to the subject a second treatment. In another example, the second treatment is administered to the subject before and/or simultaneously with and/or after the pharmaceutical composition is administered to the subject. In another example, the second treatment comprises an anti-inflammatory agent. In another example, the second pharmaceutical composition comprises an agent selected from the group consisting of non-steroidal anti-inflammatory drugs, steroids, and immunomodulating agents. In another example, the method comprises administering to the subject a third treatment.

In another example, it is described a method of increasing the longevity of a subject comprising administering to the subject the pharmaceutical composition. In another example, it is described a method of decreasing alpha4beta7 activity in a subject in need thereof comprising administering to the subject the pharmaceutical composition. In another example, it is described a method of decreasing alpha4beta7-mediated trafficking (for example, alpha4beta7mediated gut homing) in a subject in need thereof comprising administering to the subject the pharmaceutical composition.

DETAILED DESCRIPTION OF THE INVENTION

The present invention provides compositions, kits, and methods relating to molecules that bind to the integrin alpha4beta7 ("alpha4beta7"), including molecules that agonize or antagonize alpha4beta7, such as anti-alpha4beta7 antibodies, antibody fragments, and antibody derivatives, *e.g.*, antagonistic anti-alpha4beta7 antibodies, antibody fragments, or antibody derivatives. Also provided are nucleic acids, and derivatives and fragments thereof, comprising a sequence of nucleotides that encodes all or a portion of a polypeptide that binds to alpha4beta7, *e.g.*, a nucleic acid encoding all or part of an anti-alpha4beta7 antibody, antibody fragment, or antibody derivative, plasmids and vectors comprising such nucleic acids, and cells or cell lines comprising such nucleic acids and/or vectors and plasmids. The provided methods include, for example, methods of making, identifying, or isolating molecules that bind to alpha4beta7, such as anti-alpha4beta7 antibodies, methods of determining whether a molecule binds to alpha4beta7, methods of determining whether a molecule agonizes or antagonizes alpha4beta7, methods of making compositions, such as pharmaceutical compositions, comprising a molecule that binds to alpha4beta7, and methods for administering a molecule that binds alpha4beta7 to a subject, for example, methods for treating a condition mediated by alpha4beta7, and for agonizing or antagonizing a biological activity of alpha4beta7, *in vivo* or *in vitro*.

Polynucleotide and polypeptide sequences are indicated using standard one- or three-letter abbreviations. Unless otherwise indicated, each polypeptide sequence has amino terminus at the left

and a carboxy terminus at the right; each single-stranded nucleic acid sequence, and the top strand of each double-stranded nucleic acid sequence has a 5' terminus at the left and a 3' terminus at the right. A particular polypeptide or polynucleotide sequence also can be described by explaining how it differs from a reference sequence.

Unless otherwise defined herein, scientific and technical terms used in connection with the present invention shall have the meanings that are commonly understood by those of ordinary skill in the art. Further, unless otherwise required by context, singular terms shall include pluralities and plural terms shall include the singular. Generally, nomenclatures used in connection with, and techniques of, cell and tissue culture, molecular biology, immunology, microbiology, genetics and protein and nucleic acid chemistry and hybridization described herein are those well known and commonly used in the art. The methods and techniques of the present invention are generally performed according to conventional methods well known in the art and as described in various general and more specific references that are cited and discussed throughout the present specification unless otherwise indicated. See, *e.g.*, Sambrook *et al.* Molecular Cloning: A Laboratory Manual, 2d ed., Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N.Y. (1989) and Ausubel *et al.*, Current Protocols in Molecular Biology, Greene Publishing Associates (1992), and Harlow and Lane Antibodies: A Laboratory Manual Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N.Y. (1990), which are incorporated herein by reference. Enzymatic reactions and purification techniques are performed according to manufacturer's specifications, as commonly accomplished in the art or as described herein. The terminology used in connection with, and the laboratory procedures and techniques of, analytical chemistry, synthetic organic chemistry, and medicinal and pharmaceutical chemistry described herein are those well known and commonly used in the art. Standard techniques can be used for chemical syntheses, chemical analyses, pharmaceutical preparation, formulation, and delivery, and treatment of patients.

The following terms, unless otherwise indicated, shall be understood to have the following meanings:

The term "isolated molecule" (where the molecule is, for example, a polypeptide, a polynucleotide, or an antibody) is a molecule that by virtue of its origin or source of derivation (1) is not associated with naturally associated components that accompany it in its native state, (2) is substantially free of other molecules from the same species (3) is expressed by a cell from a different species, or (4) does not occur in nature without human intervention. Thus, a molecule that is chemically synthesized, or synthesized in a cellular system different from the cell from which it naturally originates, will be "isolated" from its naturally associated components. A molecule also may be rendered substantially free of naturally associated components by isolation, using purification techniques well known in the art. Molecule purity or homogeneity may be assayed by a number of means well known in the art. For example, the purity of a polypeptide sample may be assayed using polyacrylamide gel electrophoresis and staining of the gel to visualize the polypeptide using techniques well known in the art. For certain purposes, higher resolution may be provided by using HPLC or other means well known in the art for purification.

The terms "alpha4beta7 inhibitor" and "alpha4beta7 antagonist" are used interchangeably. Each is a molecule that detectably inhibits at least one function of alpha4beta7. Conversely, an "alpha4beta7 agonist" is a molecule that detectably increases at least one function of alpha4beta7. The inhibition caused by an alpha4beta7 inhibitor need not be complete so long as it is detectable, for example by using an assay. Any assay of a function of alpha4beta7 can be used, examples of which are provided herein. Examples of functions of alpha4beta7 that can be inhibited by an alpha4beta7 inhibitor (or increased by an alpha4beta7 agonist) include ligand binding (i.e., binding to MAdCAM-1), adhesion to ligand-expressing cells, trafficking to a particular compartment such as the gut, release of cytokines, chemokines and other mediators, enhancing or exacerbating inflammatory response and tissue damage, and so on. Examples of types of alpha4beta7 inhibitors and alpha4beta7 agonists include, but are not limited to, alpha4beta7 binding polypeptides such as antigen binding proteins (e.g., alpha4beta7 antigen binding proteins), antibodies, antibody fragments, and antibody derivatives.

The terms "peptide," "polypeptide" and "protein" each refers to a molecule comprising two or more amino acid residues joined to each other by peptide bonds. These terms encompass, *e.g.*, native and artificial proteins, protein fragments and polypeptide analogs (such as muteins, variants, and fusion proteins) of a protein sequence as well as post-translationally, or otherwise covalently or non-covalently, modified proteins. A peptide, polypeptide, or protein may be monomeric or polymeric.

The term "polypeptide fragment" as used herein refers to a polypeptide that has an amino-terminal and/or carboxy-terminal deletion as compared to a corresponding full-length protein. Fragments can be, for example, at least 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 20, 50, 70, 80, 90, 100, 150 or 200 amino acids in length. Fragments can also be, for example, at most 1,000, 750, 500, 250, 200, 175, 150, 125, 100, 90, 80, 70, 60, 50, 40, 30, 20, 15, 14, 13, 12, 11, or 10 amino acids in length. Fragments can also result from proteolytic (or other) processing, which, for example, results in variation in the amino and/or carboxy terminus of from one to five amino acids from that predicted. A fragment can further comprise, at either or both of its ends, one or more additional amino acids, for example, a sequence of amino acids from a different naturally-occurring protein (*e.g.*, an Fc or leucine zipper domain) or an artificial amino acid sequence (*e.g.*, an artificial linker sequence or a tag protein).

Polypeptides of the invention include polypeptides that have been modified in any way and for any reason, for example, to: (1) reduce susceptibility to proteolysis, (2) reduce susceptibility to oxidation, (3) alter binding affinity for forming protein complexes, (4) alter binding affinities, and (4) confer or modify other physicochemical or functional properties. Analogs include muteins of a polypeptide. For example, single or multiple amino acid substitutions (*e.g.*, conservative amino acid substitutions) may be made in the naturally occurring sequence (*e.g.*, in the portion of the polypeptide outside the domain(s) forming intermolecular contacts). Consensus sequences can be used to select amino acid residues for substitution; those of skill in the art recognize that additional amino acid residues may also be substituted.

A "conservative amino acid substitution" is one that does not substantially change the structural characteristics of the parent sequence (*e.g.*, a replacement amino acid should not tend to break a helix that occurs in the parent sequence, or disrupt other types of secondary structure that characterize the parent sequence or are necessary for its functionality). Examples of art-recognized

polypeptide secondary and tertiary structures are described in Proteins, Structures and Molecular Principles (Creighton, Ed., W. H. Freeman and Company, New York (1984)); Introduction to Protein Structure (C. Branden and J. Tooze, eds., Garland Publishing, New York, N.Y. (1991)); and Thornton *et at.* Nature 354:105 (1991), which are each incorporated herein by reference.

The present invention also provides non-peptide analogs of alpha4beta7 binding polypeptides. Non-peptide analogs are commonly used in the pharmaceutical industry as drugs with properties analogous to those of the template peptide. These types of non-peptide compound are termed "peptide mimetics" or "peptidomimetics," see, for example, Fauchere, J. Adv. Drug Res. 15:29 (1986); Veber and Freidinger TINS p.392 (1985); and Evans *et al.* J. Med. Chem. 30:1229 (1987), which are incorporated herein by reference. Peptide mimetics that are structurally similar to therapeutically useful peptides may be used to produce an equivalent therapeutic or prophylactic effect. Generally, peptidomimetics are structurally similar to a paradigm polypeptide (*i.e.*, a polypeptide that has a desired biochemical property or pharmacological activity), such as a human antibody, but have one or more peptide linkages optionally replaced by a linkage selected from the group consisting of: --CH_2NH--, --CH_2S--, --CH_2--CH_2--, --CH=CH-(*cis* and *trans*), --$COCH_2$--, --$CH(OH)CH_2$--, and --CH_2SO--, by methods well known in the art. Systematic substitution of one or more amino acids of a consensus sequence with a D-amino acid of the same type (*e.g.*, D-lysine in place of L-lysine) may also be used to generate more stable peptides. In addition, constrained peptides comprising a consensus sequence or a substantially identical consensus sequence variation may be generated by methods known in the art (Rizo and Gierasch Ann. Rev. Biochem. 61:387 (1992), incorporated herein by reference), for example, by adding internal cysteine residues capable of forming intramolecular disulfide bridges which cyclize the peptide.

A "variant" of a polypeptide (*e.g.*, an antibody) comprises an amino acid sequence wherein one or more amino acid residues are inserted into, deleted from and/or substituted into the amino acid sequence relative to another polypeptide sequence. Variants of the invention include fusion proteins.

A "derivative" of a polypeptide is a polypeptide (*e.g.*, an antibody) that has been chemically modified, *e.g.*, via conjugation to another chemical moiety (such as, for example, polyethylene glycol or albumin, *e.g.*, human serum albumin), phosphorylation, and/or glycosylation. Unless otherwise indicated, the term "antibody" includes, in addition to antibodies comprising two full-length heavy chains and two full-length light chains, derivatives, variants, fragments, and muteins thereof, examples of which are described below.

An "antigen binding protein" is a protein comprising a portion that binds to an antigen and, optionally, a scaffold or framework portion that allows the antigen binding portion to adopt a conformation that promotes binding of the antigen binding protein to the antigen. Examples of antigen binding proteins include antibodies, antibody fragments (e.g., an antigen binding portion of an antibody), antibody derivatives, and antibody analogs. The antigen binding protein can comprise, for example, an alternative protein scaffold or artificial scaffold with grafted CDRs or CDR derivatives. Such scaffolds include, but are not limited to, antibody-derived scaffolds comprising mutations introduced to, for example, stabilize the three-dimensional structure of the antigen binding protein as well as wholly synthetic scaffolds comprising, for example, a biocompatible polymer. See, for

example, Korndorfer et al., 2003, Proteins: Structure, Function, and Bioinformatics, Volume 53, Issue 1:121-129; Roque et al., 2004, Biotechnol. Prog. 20:639-654. In addition, peptide antibody mimetics ("PAMs") can be used, as well as scaffolds based on antibody mimetics utilizing fibronection components as a scaffold.

An antigen binding protein can have, for example, the structure of a naturally occurring immunoglobulin. An "immunoglobulin" is a tetrameric molecule. In a naturally occurring immunoglobulin, each tetramer is composed of two identical pairs of polypeptide chains, each pair having one "light" (about 25 kDa) and one "heavy" chain (about 50-70 kDa). The amino-terminal portion of each chain includes a variable region of about 100 to 110 or more amino acids primarily responsible for antigen recognition. The carboxy-terminal portion of each chain defines a constant region primarily responsible for effector function. Human light chains are classified as kappa or lambda light chains. Heavy chains are classified as mu, delta, gamma, alpha, or epsilon, and define the antibody's isotype as IgM, IgD, IgG, IgA, and IgE, respectively. Within light and heavy chains, the variable and constant regions are joined by a "J" region of about 12 or more amino acids, with the heavy chain also including a "D" region of about 10 more amino acids. See generally, Fundamental Immunology Ch. 7 (Paul, W., ed., 2nd ed. Raven Press, N.Y. (1989)) (incorporated by reference in its entirety for all purposes). The variable regions of each light/heavy chain pair form the antibody binding site such that an intact immunoglobulin has two binding sites.

The variable regions of naturally occurring immunoglobulin chains exhibit the same general structure of relatively conserved framework regions (FR) joined by three hypervariable regions, also called complementarity determining regions or CDRs. From N-terminus to C-terminus, both light and heavy chains comprise the domains FR1, CDR1, FR2, CDR2, FR3, CDR3 and FR4. The assignment of amino acids to each domain is in accordance with the definitions of Kabat *et al.* in *Sequences of Proteins of Immunological Interest*, 5th Ed., US Dept. of Health and Human Services, PHS, NIH, NIH Publication no. 91-3242, 1991. Other numbering systems for the amino acids in immunoglobulin chains include IMGT® (the international ImMunoGeneTics information system; Lefranc et al, *Dev. Comp. Immunol.* 29:185-203; 2005) and AHo (Honegger and Pluckthun, *J. Mol. Biol.* 309(3):657-670; 2001).

Antibodies can be obtained from sources such as serum or plasma that contain immunoglobulins having varied antigenic specificity. If such antibodies are subjected to affinity purification, they can be enriched for a particular antigenic specificity. Such enriched preparations of antibodies usually are made of less than about 10% antibody having specific binding activity for the particular antigen. Subjecting these preparations to several rounds of affinity purification can increase the proportion of antibody having specific binding activity for the antigen. Antibodies prepared in this manner are often referred to as "monospecific." Monospecfic antibody preparations can be made up of about 10 %, 20%, 30%, 40%, 50%, 60%, 70%, 75%, 80%, 85%, 90%, 95%, 97%, 99%, or 99.9% antibody having specific binding activity for the particular antigen.

An "antibody" refers to an intact immunoglobulin or to an antigen binding portion thereof that competes with the intact antibody for specific binding, unless otherwise specified. Antigen binding portions may be produced by recombinant DNA techniques or by enzymatic or chemical cleavage of

intact antibodies. Antigen binding portions include, *inter alia*, Fab, Fab', $F(ab')_2$, Fv, domain antibodies (dAbs), and complementarity determining region (CDR) fragments, variable region fragments, single-chain antibodies (scFv), chimeric antibodies, diabodies, triabodies, tetrabodies, and polypeptides that contain at least a portion of an immunoglobulin that is sufficient to confer specific antigen binding to the polypeptide.

A Fab fragment is a monovalent fragment having the V_L, V_H, C_L and C_H1 domains; a $F(ab')_2$ fragment is a bivalent fragment having two Fab fragments linked by a disulfide bridge at the hinge region; a Fd fragment has the V_H and C_H1 domains; an Fv fragment has the V_L and V_H domains of a single arm of an antibody; and a dAb fragment has a V_H domain, a V_L domain, or an antigen-binding fragment of a V_H or V_L domain (US Pat. No. 6,846,634, 6,696,245, US App. Pub. No. 05/0202512, 04/0202995, 04/0038291, 04/0009507, 03/0039958, Ward *et al.*, Nature 341:544-546, 1989).

A single-chain antibody (scFv) is an antibody in which a V_L and a V_H region are joined via a linker (*e.g.*, a synthetic sequence of amino acid residues) to form a continuous protein chain wherein the linker is long enough to allow the protein chain to fold back on itself and form a monovalent antigen binding site (see, *e.g.*, Bird *et al.*, 1988, Science 242:423-26 and Huston *et al.*, 1988, Proc. Natl. Acad. Sci. USA 85:5879-83). Diabodies are bivalent antibodies comprising two polypeptide chains, wherein each polypeptide chain comprises V_H and V_L domains joined by a linker that is too short to allow for pairing between two domains on the same chain, thus allowing each domain to pair with a complementary domain on another polypeptide chain (see, *e.g.*, Holliger *et al.*, 1993, Proc. Natl. Acad. Sci. USA 90:6444-48, and Poljak *et al.*, 1994, Structure 2:1121-23). If the two polypeptide chains of a diabody are identical, then a diabody resulting from their pairing will have two identical antigen binding sites. Polypeptide chains having different sequences can be used to make a diabody with two different antigen binding sites. Similarly, triabodies and tetrabodies are antibodies comprising three and four polypeptide chains, respectively, and forming three and four antigen binding sites, respectively, which can be the same or different.

Complementarity determining regions (CDRs) and framework regions (FR) of a given antibody may be identified using the system described by Kabat *et al. supra*; Lefranc et al., *supra* and/or Honegger and Pluckthun, *supra*. One or more CDRs may be incorporated into a molecule either covalently or noncovalently to make it an antigen binding protein. An antigen binding protein may incorporate the CDR(s) as part of a larger polypeptide chain, may covalently link the CDR(s) to another polypeptide chain, or may incorporate the CDR(s) noncovalently. The CDRs permit the antigen binding protein to specifically bind to a particular antigen of interest.

An antigen binding protein may have one or more binding sites. If there is more than one binding site, the binding sites may be identical to one another or may be different. For example, a naturally occurring human immunoglobulin typically has two identical binding sites, while a "bispecific" or "bifunctional" antibody has two different binding sites.

The term "human antibody" includes all antibodies that have one or more variable and constant regions derived from human immunoglobulin sequences. In one embodiment, all of the variable and constant domains are derived from human immunoglobulin sequences (a fully human antibody). These antibodies may be prepared in a variety of ways, examples of which are described

below, including through the immunization with an antigen of interest of a mouse that is genetically modified to express antibodies derived from human heavy and/or light chain-encoding genes.

A humanized antibody has a sequence that differs from the sequence of an antibody derived from a non-human species by one or more amino acid substitutions, deletions, and/or additions, such that the humanized antibody is less likely to induce an immune response, and/or induces a less severe immune response, as compared to the non-human species antibody, when it is administered to a human subject. In one embodiment, certain amino acids in the framework and constant domains of the heavy and/or light chains of the non-human species antibody are mutated to produce the humanized antibody. In another embodiment, the constant domain(s) from a human antibody are fused to the variable domain(s) of a non-human species. In another embodiment, one or more amino acid residues in one or more CDR sequences of a non-human antibody are changed to reduce the likely immunogenicity of the non-human antibody when it is administered to a human subject, wherein the changed amino acid residues either are not critical for immunospecific binding of the antibody to its antigen, or the changes to the amino acid sequence that are made are conservative changes, such that the binding of the humanized antibody to the antigen is not significantly worse than the binding of the non-human antibody to the antigen. Examples of how to make humanized antibodies may be found in U.S. Pat. Nos. 6,054,297, 5,886,152 and 5,877,293.

The term "chimeric antibody" refers to an antibody that contains one or more regions from one antibody and one or more regions from one or more other antibodies. In one embodiment, one or more of the CDRs are derived from a human anti-alpha4beta7 antibody. In another embodiment, all of the CDRs are derived from a human anti-alpha4beta7 antibody. In another embodiment, the CDRs from more than one human anti-alpha4beta7 antibodies are mixed and matched in a chimeric antibody. For instance, a chimeric antibody may comprise a CDR1 from the light chain of a first human anti-alpha4beta7 antibody, a CDR2 and a CDR3 from the light chain of a second human anti-alpha4beta7 antibody, and the CDRs from the heavy chain from a third anti-alpha4beta7 antibody. Other combinations are possible and are included within the embodiments of the invention.

Further, the framework regions may be derived from one of the same anti-alpha4beta7 antibodies, from one or more different antibodies, such as a human antibody, or from a humanized antibody. In one example of a chimeric antibody, a portion of the heavy and/or light chain is identical with, homologous to, or derived from an antibody from a particular species or belonging to a particular antibody class or subclass, while the remainder of the chain(s) is/are identical with, homologous to, or derived from an antibody (-ies) from another species or belonging to another antibody class or subclass. Also included are fragments of such antibodies that exhibit the desired biological activity (*i.e.*, the ability to specifically bind alpha4beta7). See, *e.g.*, U.S. Patent No. 4,816,567 and Morrison, 1985, Science 229:1202-07.

A "neutralizing antibody" or an "inhibitory antibody" is an antibody that inhibits the interaction of alpha4beta7 with MAdCAM-1 when an excess of the anti-alpha4beta7 antibody reduces the amount of interaction by at least about 20% using an assay such as those described herein in the Examples. In various embodiments, the antigen binding protein reduces the interaction of alpha4beta7

with MAdCAM-1 alpha4beta7 by at least 30%, 40%, 50%, 60%, 70%, 75%, 80%, 85%, 90%, 95%, 97%, 99%, and 99.9%.

Fragments or analogs of antibodies can be readily prepared by those of ordinary skill in the art following the teachings of this specification and using techniques well-known in the art. Amino- and carboxy-termini of fragments or analogs occur near boundaries of functional domains. Structural and functional domains can be identified by comparison of the nucleotide and/or amino acid sequence data to public or proprietary sequence databases. Computerized comparison methods can be used to identify sequence motifs or predicted protein conformation domains that occur in other proteins of known structure and/or function. Methods to identify protein sequences that fold into a known three-dimensional structure are known. See, *e.g.*, Bowie *et al.*, 1991, Science 253:164.

A "CDR grafted antibody" is an antibody comprising one or more CDRs derived from an antibody of a particular species or isotype and the framework of another antibody of the same or different species or isotype.

A "multi-specific antibody" is an antibody that recognizes more than one epitope on one or more antigens. A subclass of this type of antibody is a "bi-specific antibody" which recognizes two distinct epitopes on the same or different antigens.

An antigen binding protein "specifically binds" to an antigen (*e.g.*, human alpha4beta7) if it binds to the antigen with a dissociation constant of 1 nanomolar or less. As used herein, an antigen binding protein is "heterodimer specific" if it binds to a first heterodimeric integrin but not to other integrins that share one chain with the first integrin. For example, an antibody that is alpha4beta7 heterodimer specific will bind to alpha4beta7 but not to alpha4beta1 or alphaEbeta7.

Integrins are known to adapt different conformations, depending on the activation state of the cell(s) expressing them and on the presence or absence of certain metal ions. An integrin in "active" conformation binds to its cognate ligand with higher affinity than the same integrin in "inactive" conformation. An antigen binding protein may bind to an integrin in only its active conformation, in only its inactive conformation, or in both or either conformations. For example, an alpha4beta7 heterodimer specific antigen binding protein may bind alpha4beta7 in the presence or absence of the divalent cation manganese^{2+} (Mn^{2+}), indicating that the antigen binding protein binds both active and inactive alpah4beta7.

An "antigen binding domain," "antigen binding region," or "antigen binding site" is a portion of an antigen binding protein that contains amino acid residues (or other moieties) that interact with an antigen and contribute to the antigen binding protein's specificity and affinity for the antigen. For an antibody that specifically binds to its antigen, this will include at least part of at least one of its CDR domains.

An "epitope" is the portion of a molecule that is bound by an antigen binding protein (e.g., by an antibody). An epitope can comprise non-contiguous portions of the molecule (*e.g.*, in a polypeptide, amino acid residues that are not contiguous in the polypeptide's primary sequence but that, in the context of the polypeptide's tertiary and quaternary structure, are near enough to each other to be bound by an antigen binding protein).

The "percent identity" of two polynucleotide or two polypeptide sequences is determined by comparing the sequences using the GAP computer program (a part of the GCG Wisconsin Package, version 10.3 (Accelrys, San Diego, CA)) using its default parameters.

The terms "polynucleotide," "oligonucleotide" and "nucleic acid" are used interchangeably throughout and include DNA molecules (*e.g.*, cDNA or genomic DNA), RNA molecules (*e.g.*, mRNA), analogs of the DNA or RNA generated using nucleotide analogs (*e.g.*, peptide nucleic acids and non-naturally occurring nucleotide analogs), and hybrids thereof. The nucleic acid molecule can be single-stranded or double-stranded. In one embodiment, the nucleic acid molecules of the invention comprise a contiguous open reading frame encoding an antibody, or a fragment, derivative, mutein, or variant thereof, of the invention.

Two single-stranded polynucleotides are "the complement" of each other if their sequences can be aligned in an anti-parallel orientation such that every nucleotide in one polynucleotide is opposite its complementary nucleotide in the other polynucleotide, without the introduction of gaps, and without unpaired nucleotides at the 5' or the 3' end of either sequence. A polynucleotide is "complementary" to another polynucleotide if the two polynucleotides can hybridize to one another under moderately stringent conditions. Thus, a polynucleotide can be complementary to another polynucleotide without being its complement.

A "vector" is a nucleic acid that can be used to introduce another nucleic acid linked to it into a cell. One type of vector is a "plasmid," which refers to a linear or circular double stranded DNA molecule into which additional nucleic acid segments can be ligated. Another type of vector is a viral vector (*e.g.*, replication defective retroviruses, adenoviruses and adeno-associated viruses), wherein additional DNA segments can be introduced into the viral genome. Certain vectors are capable of autonomous replication in a host cell into which they are introduced (*e.g.*, bacterial vectors comprising a bacterial origin of replication and episomal mammalian vectors). Other vectors (*e.g.*, non-episomal mammalian vectors) are integrated into the genome of a host cell upon introduction into the host cell, and thereby are replicated along with the host genome. An "expression vector" is a type of vector that can direct the expression of a chosen polynucleotide.

A nucleotide sequence is "operably linked" to a regulatory sequence if the regulatory sequence affects the expression (*e.g.*, the level, timing, or location of expression) of the nucleotide sequence. A "regulatory sequence" is a nucleic acid that affects the expression (*e.g.*, the level, timing, or location of expression) of a nucleic acid to which it is operably linked. The regulatory sequence can, for example, exert its effects directly on the regulated nucleic acid, or through the action of one or more other molecules (*e.g.*, polypeptides that bind to the regulatory sequence and/or the nucleic acid). Examples of regulatory sequences include promoters, enhancers and other expression control elements (*e.g.*, polyadenylation signals). Further examples of regulatory sequences are described in, for example, Goeddel, 1990, Gene Expression Technology: Methods in Enzymology 185, Academic Press, San Diego, CA and Baron *et al.*, 1995, Nucleic Acids Res. 23:3605 06.

A "host cell" is a cell that can be used to express a nucleic acid, *e.g.*, a nucleic acid of the invention. A host cell can be a prokaryote, for example, *E. coli*, or it can be a eukaryote, for example, a single-celled eukaryote (*e.g.*, a yeast or other fungus), a plant cell (*e.g.*, a tobacco or tomato plant cell),

an animal cell (*e.g.*, a human cell, a monkey cell, a hamster cell, a rat cell, a mouse cell, or an insect cell) or a hybridoma. Examples of host cells include the COS-7 line of monkey kidney cells (ATCC CRL 1651) (see Gluzman *et al.*, 1981, Cell 23:175), L cells, C127 cells, 3T3 cells (ATCC CCL 163), Chinese hamster ovary (CHO) cells or their derivatives such as Veggie CHO and related cell lines which grow in serum-free media (see Rasmussen *et al.*, 1998, Cytotechnology 28:31) or CHO strain DX-B11, which is deficient in DHFR (see Urlaub *et al.*, 1980, Proc. Natl. Acad. Sci. USA 77:4216-20), HeLa cells, BHK (ATCC CRL 10) cell lines, the CV1/EBNA cell line derived from the African green monkey kidney cell line CV1 (ATCC CCL 70) (see McMahan *et al.*, 1991, EMBO J. 10:2821), human embryonic kidney cells such as 293, 293 EBNA or MSR 293, human epidermal A431 cells, human Colo205 cells, other transformed primate cell lines, normal diploid cells, cell strains derived from in vitro culture of primary tissue, primary explants, HL-60, U937, HaK or Jurkat cells. Typically, a host cell is a cultured cell that can be transformed or transfected with a polypeptide-encoding nucleic acid, which can then be expressed in the host cell. The phrase "recombinant host cell" can be used to denote a host cell that has been transformed or transfected with a nucleic acid to be expressed. A host cell also can be a cell that comprises the nucleic acid but does not express it at a desired level unless a regulatory sequence is introduced into the host cell such that it becomes operably linked with the nucleic acid. It is understood that the term host cell refers not only to the particular subject cell but also to the progeny or potential progeny of such a cell. Because certain modifications may occur in succeeding generations due to, *e.g.*, mutation or environmental influence, such progeny may not, in fact, be identical to the parent cell, but are still included within the scope of the term as used herein.

Antigen binding proteins

In one aspect, the present invention provides antigen binding proteins (*e.g.*, antibodies, antibody fragments, antibody derivatives, antibody muteins, and antibody variants) that bind to alpha4beta7, *e.g.*, human alpha4beta7.

Antigen binding proteins in accordance with the present invention include antigen binding proteins that inhibit a biological activity of alpha4beta7. Examples of such biological activities include binding of alpha4beta7 to MAdCAM-1, and adhesion between cells expressing alpha4beta7 and those expressing MAdCAM-1. Other biological activities include those mediated by alpha4beta7 in vivo, such as trafficking or homing; in particular, alpha4beta7 is involved in the trafficking of lymphocytes to the gut, Increased MAdCAM-1 expression in the inflamed gut enhances recruitment of alpha4beta7 expressing lymphocytes to the gut, where aberrant lymphocyte activation augments inflammatory response and tissue damage.

Different antigen binding proteins may bind to different domains or epitopes of alpha4beta7 or act by different mechanisms of action. Examples include but are not limited to antigen binding proteins that interfere with the ability of alpha4beta7 to bind MAdCAM-1 or that inhibit cellular interactions such as adhesion between cells expressing alpha4beta7 and cells expressing MAdCAM-1. The site of action may be, for example, intracellular (*e.g.*, by interfering with an intracellular signaling cascade) or extracellular. An antigen binding protein need not completely inhibit alpha4beta7 induced activity to find use in the present invention; rather, antigen binding proteins that reduce a particular

activity of alpha4beta7 are contemplated for use as well. (Discussions herein of particular mechanisms of action for alpha4beta7-binding antigen binding proteins in treating particular diseases are illustrative only, and the methods presented herein are not bound thereby.)

Other derivatives of anti- alpha4beta7 antibodies within the scope of this invention include covalent or aggregative conjugates of anti-alpha4beta7 antibodies, or fragments thereof, with other proteins or polypeptides, such as by expression of recombinant fusion proteins comprising heterologous polypeptides fused to the N-terminus or C-terminus of an anti- alpha4beta7 antibody polypeptide. For example, the conjugated peptide may be a heterologous signal (or leader) polypeptide, *e.g.*, the yeast alpha-factor leader, or a peptide such as an epitope tag. Antigen binding protein-containing fusion proteins can comprise peptides added to facilitate purification or identification of antigen binding protein (*e.g.*, poly-His). An antigen binding protein also can be linked to the FLAG® peptide Asp-Tyr-Lys-Asp-Asp-Asp-Asp-Lys (DYKDDDDK) (SEQ ID NO:62) as described in Hopp *et al.*, *Bio/Technology* 6:1204, 1988, and U.S. Patent 5,011,912. The FLAG® peptide is highly antigenic and provides an epitope reversibly bound by a specific monoclonal antibody (mAb), enabling rapid assay and facile purification of expressed recombinant protein. Reagents useful for preparing fusion proteins in which the FLAG® peptide is fused to a given polypeptide are commercially available (Sigma-Aldrich, St. Louis MO).

Oligomers that contain one or more antigen binding proteins may be employed as alpha4beta7 antagonists. Oligomers may be in the form of covalently-linked or non-covalently-linked dimers, trimers, or higher oligomers. Oligomers comprising two or more antigen binding protein are contemplated for use, with one example being a homodimer. Other oligomers include heterodimers, homotrimers, heterotrimers, homotetramers, heterotetramers, *etc.*

One embodiment is directed to oligomers comprising multiple antigen binding proteins joined *via* covalent or non-covalent interactions between peptide moieties fused to the antigen binding proteins. Such peptides may be peptide linkers (spacers), or peptides that have the property of promoting oligomerization. Leucine zippers and certain polypeptides derived from antibodies are among the peptides that can promote oligomerization of antigen binding proteins attached thereto, as described in more detail below.

In particular embodiments, the oligomers comprise from two to four antigen binding proteins. The antigen binding proteins of the oligomer may be in any form, such as any of the forms described above, *e.g.*, variants or fragments. Preferably, the oligomers comprise antigen binding proteins that have alpha4beta7 binding activity.

In one embodiment, an oligomer is prepared using polypeptides derived from immunoglobulins. Preparation of fusion proteins comprising certain heterologous polypeptides fused to various portions of antibody-derived polypeptides (including the Fc domain) has been described, *e.g.*, by Ashkenazi *et al.*, 1991, PNAS USA 88:10535; Byrn *et al.*, 1990, Nature 344:677; and Hollenbaugh *et al.*, 1992 "Construction of Immunoglobulin Fusion Proteins", in *Current Protocols in Immunology*, Suppl. 4, pages 10.19.1 - 10.19.11.

One embodiment of the present invention is directed to a dimer comprising two fusion proteins created by fusing an alpha4beta7 binding fragment of an anti- alpha4beta7 antibody to the Fc

region of an antibody. The dimer can be made by, for example, inserting a gene fusion encoding the fusion protein into an appropriate expression vector, expressing the gene fusion in host cells transformed with the recombinant expression vector, and allowing the expressed fusion protein to assemble much like antibody molecules, whereupon interchain disulfide bonds form between the Fc moieties to yield the dimer.

The term "Fc polypeptide" as used herein includes native and mutein forms of polypeptides derived from the Fc region of an antibody. Truncated forms of such polypeptides containing the hinge region that promotes dimerization also are included. Fusion proteins comprising Fc moieties (and oligomers formed therefrom) offer the advantage of facile purification by affinity chromatography over Protein A or Protein G columns.

One suitable Fc polypeptide, described in PCT application WO 93/10151 (hereby incorporated by reference), is a single chain polypeptide extending from the N-terminal hinge region to the native C-terminus of the Fc region of a human IgG1 antibody. Another useful Fc polypeptide is the Fc mutein described in U.S. Patent 5,457,035 and in Baum *et al.*, 1994, EMBO J. 13:3992-4001. The amino acid sequence of this mutein is identical to that of the native Fc sequence presented in WO 93/10151, except that amino acid 19 has been changed from Leu to Ala, amino acid 20 has been changed from Leu to Glu, and amino acid 22 has been changed from Gly to Ala. The mutein exhibits reduced affinity for Fc receptors.

In other embodiments, the variable portion of the heavy and/or light chains of an anti-alpha4beta7 antibody may be substituted for the variable portion of an antibody heavy and/or light chain.

Alternatively, the oligomer is a fusion protein comprising multiple antigen binding proteins, with or without peptide linkers (spacer peptides). Among the suitable peptide linkers are those described in U.S. Patents 4,751,180 and 4,935,233.

Another method for preparing oligomeric antigen binding proteins involves use of a leucine zipper. Leucine zipper domains are peptides that promote oligomerization of the proteins in which they are found. Leucine zippers were originally identified in several DNA-binding proteins (Landschulz *et al.*, 1988, Science 240:1759), and have since been found in a variety of different proteins. Among the known leucine zippers are naturally occurring peptides and derivatives thereof that dimerize or trimerize. Examples of leucine zipper domains suitable for producing soluble oligomeric proteins are described in PCT application WO 94/10308, and the leucine zipper derived from lung surfactant protein D (SPD) described in Hoppe *et al.*, 1994, FEBS Letters 344:191, hereby incorporated by reference. The use of a modified leucine zipper that allows for stable trimerization of a heterologous protein fused thereto is described in Fanslow *et al.*, 1994, Semin. Immunol. 6:267-78. In one approach, recombinant fusion proteins comprising an anti- alpha4beta7 antibody fragment or derivative fused to a leucine zipper peptide are expressed in suitable host cells, and the soluble oligomeric anti- alpha4beta7 antibody fragments or derivatives that form are recovered from the culture supernatant.

In one aspect, the present invention provides antigen binding proteins that interfere with the binding of alpha4beta7 to MAdCAM-1. Such antigen binding proteins can be made against alpha4beta7, or a fragment, variant or derivative thereof, and screened in conventional assays for the

ability to interfere with the binding of alpha4beta7 to MAdCAM-1. Examples of suitable assays are assays that test the antigen binding proteins for the ability to inhibit binding of MAdCAM-1 (i.e., soluble MAdCAM-1) to cells expressing alpha4beta7, or that test antigen binding proteins for the ability to reduce a biological or cellular response that results from the interaction of MAdCAM-1 and alpha4beta7 (i.e., adhesion of cells expressing alpha4beta7 to MAdCAM-1, or MAdCAM-1-expressing cells). Additional assays that test the antigen binding proteins include those that qualitatively or quantitatively compare the binding of an antigen binding protein to a alpha4beta7 polypeptide to the binding of a known antigen binding protein to a alpha4beta7 polypeptide, several examples of which are disclosed herein.

In another aspect, the present invention provides an antigen binding protein that demonstrates species selectivity. In one embodiment, the antigen binding protein binds to one or more mammalian alpha4beta7, for example, to human alpha4beta7 and one or more of mouse, rat, guinea pig, hamster, gerbil, cat, rabbit, dog, goat, sheep, cow, horse, camel, and non-human primate alpha4beta7. In another embodiment, the antigen binding protein binds to one or more primate alpha4beta7, for example, to human alpha4beta7 and one or more of cynomologous, marmoset, rhesus, tamarin and chimpanzee alpha4beta7. In another embodiment, the antigen binding protein binds specifically to human, cynomologous, marmoset, rhesus, tamarin or chimpanzee alpha4beta7. In another embodiment, the antigen binding protein does not bind to one or more of mouse, rat, guinea pig, hamster, gerbil, cat, rabbit, dog, goat, sheep, cow, horse, camel, and non-human primate alpha4beta7. In another embodiment, the antigen binding protein does not bind to a New World monkey species such as a marmoset.

In another embodiment, the antigen binding protein does not exhibit specific binding to any naturally occurring protein other than alpha4beta7. In another embodiment, the antigen binding protein does not exhibit specific binding to any naturally occurring protein other than mammalian alpha4beta7. In another embodiment, the antigen binding protein does not exhibit specific binding to any naturally occurring protein other than primate alpha4beta7. In another embodiment, the antigen binding protein does not exhibit specific binding to any naturally occurring protein other than human alpha4beta7. In another embodiment, the antigen binding protein specifically binds to alpha4beta7 from at least one non-human primate, for example, cynomologous monkey, and human alpha4beta7. In another embodiment, the antigen binding protein specifically binds to non-human primate, cynomologous monkey, and human alpha4beta7 with a similar binding affinity. In another embodiment, the antigen binding protein blocks an activity of non-human primate, cynomologous monkey, and human alpha4beta7. In another embodiment, the antigen binding protein has a similar IC_{50} or EC_{50} against non-human primate, cynomologous monkey, and human alpha4beta7 in an assay as described herein.

One may determine the selectivity of an antigen binding protein for an alpha4beta7 using methods well known in the art and following the teachings of the specification. For example, one may determine the selectivity using Western blot, FACS, ELISA or RIA.

In another aspect, the present invention provides an alpha4beta7 binding antigen binding protein (for example, an anti-alpha4beta7 antibody), that has one or more of the following characteristics: binds to both human and non-human primate alpha4beta7, inhibits binding of

MAdCAM-1 to alpha4beta7, inhibits the adhesion of cells expressing alpha4beta7 to MAdCAM-1, inhibits the adhesion of cells expressing alpha4beta7 to cells expressing MAdCAM-1, inhibits trafficking of cells expressing alpha4beta7 to tissues comprising cells expressing MAdCAM-1, binds both active and inactive forms of alpha4beta7, causes relatively little down-regulation of cell-surface expressed alpha4beta7.

Antigen-binding fragments of antigen binding proteins of the invention may be produced by conventional techniques. Examples of such fragments include, but are not limited to, Fab and $F(ab')_2$ fragments. Antibody fragments and derivatives produced by genetic engineering techniques also are contemplated.

Additional embodiments include chimeric antibodies, *e.g.*, humanized versions of non-human (*e.g.*, murine) monoclonal antibodies. Such humanized antibodies may be prepared by known techniques, and offer the advantage of reduced immunogenicity when the antibodies are administered to humans. In one embodiment, a humanized monoclonal antibody comprises the variable domain of a murine antibody (or all or part of the antigen binding site thereof) and a constant domain derived from a human antibody. Alternatively, a humanized antibody fragment may comprise the antigen binding site of a murine monoclonal antibody and a variable domain fragment (lacking the antigen-binding site) derived from a human antibody. Procedures for the production of chimeric and further engineered monoclonal antibodies include those described in Riechmann *et al.*, 1988, Nature 332:323, Liu *et al.*, 1987, Proc. Nat. Acad. Sci. USA 84:3439, Larrick *et al.*, 1989, Bio/Technology 7:934, and Winter *et al.*, 1993, TIPS 14:139. In one embodiment, the chimeric antibody is a CDR grafted antibody. Techniques for humanizing antibodies are discussed in, *e.g.*, U.S. Pat. App. No. 10/194,975 (published February 27, 2003), U.S. Pat. No.s 5,869,619, 5,225,539, 5,821,337, 5,859,205, Padlan *et al.*, 1995, FASEB J. 9:133-39, and Tamura *et al.*, 2000, J. Immunol. 164:1432-41.

Procedures have been developed for generating human or partially human antibodies in non-human animals. For example, mice in which one or more endogenous immunoglobulin genes have been inactivated by various means have been prepared. Human immunoglobulin genes have been introduced into the mice to replace the inactivated mouse genes. Antibodies produced in the animal incorporate human immunoglobulin polypeptide chains encoded by the human genetic material introduced into the animal. In one embodiment, a non-human animal, such as a transgenic mouse, is immunized with an alpha4beta7 polypeptide, such that antibodies directed against the alpha4beta7 polypeptide are generated in the animal. One example of a suitable immunogen is a soluble human alpha4beta7, such as a polypeptide comprising a portion of alpha4beta7, or other immunogenic fragment alpha4beta7. Another example of a suitable immunogen is cells expressing high levels of alpha4beta7, or cell membrane preparations therefrom.

Examples of techniques for production and use of transgenic animals for the production of human or partially human antibodies are described in U.S. Patents 5,814,318, 5,569,825, and 5,545,806, Davis *et al.*, 2003, *Production of human antibodies from transgenic mice* in Lo, ed. Antibody Engineering: Methods and Protocols, Humana Press, NJ:191-200, Kellermann *et al.*, 2002, Curr Opin Biotechnol. 13:593-97, Russel *et al.*, 2000, Infect Immun. 68:1820-26, Gallo *et al.*, 2000, Eur J Immun. 30:534-40, Davis *et al.*, 1999, Cancer Metastasis Rev. 18:421-25, Green, 1999, J

Immunol Methods. 231:11-23, Jakobovits, 1998, Adv Drug Deliv Rev 31:33-42, Green *et al.*, 1998, J Exp Med. 188:483-95, Jakobovits A, 1998, Exp. Opin. Invest. Drugs. 7:607-14, Tsuda *et al.*, 1997, Genomics 42:413-21, Mendez *et al.*, 1997, Nat Genet. 15:146-56, Jakobovits, 1994, Curr Biol. 4:761-63, Arbones *et al.*, 1994, Immunity. 1:247-60, Green *et al.*, 1994, Nat Genet. 7:13-21, Jakobovits *et al.*, 1993, Nature 362:255-58, Jakobovits *et al.*, 1993, Proc Natl Acad Sci U S A. 90:2551-55. Chen, J. et al., 1993, Int Immunol 5: 647-656, Choi *et al.*, 1993, Nature Genetics 4: 117-23, Fishwild *et al.*, 1996, Nat Biotechnol 14: 845-51, Harding *et al.*, 1995, Ann NY Acad Sci, Lonberg *et al.*, 1994, Nature 368: 856-59, Lonberg, 1994, *Transgenic Approaches to Human Monoclonal Antibodies* in Handbook of Experimental Pharmacology 113: 49-101, Lonberg *et al.*, 1995, Int Rev Immunol 13: 65-93, Neuberger, 1996, Nat Biotechnol 14: 826, Taylor *et al.*, 1992, Nucleic Acids Research 20: 6287-95, Taylor *et al.*, 1994, Int Immunol 6: 579-91, Tomizuka *et al.*, 1997, Nat Gen 16: 133-43, Tomizuka *et al.*, 2000, Proc Natl Acad Sci U S A. 97: 722-27, Tuaillon *et al.*, 1993, Proc Natl Acad Sci U S A. 90: 3720-24, and Tuaillon *et al.*, 1994, J Immunol 152: 2912-20. These and other examples are also discussed in U.S. Patent application publication 2007-0098715, published May 3, 2007.

In another aspect, the present invention provides monoclonal antibodies that bind to alpha4beta7. Monoclonal antibodies may be produced using any technique known in the art, *e.g.*, by immortalizing spleen cells harvested from the transgenic animal after completion of the immunization schedule. The spleen cells can be immortalized using any technique known in the art, *e.g.*, by fusing them with myeloma cells to produce hybridomas. Myeloma cells for use in hybridoma-producing fusion procedures preferably are non-antibody-producing, have high fusion efficiency, and enzyme deficiencies that render them incapable of growing in certain selective media which support the growth of only the desired fused cells (hybridomas). Examples of suitable cell lines for use in mouse fusions include Sp-20, P3-X63/Ag8, P3-X63-Ag8.653, NS1/1.Ag 4 1, Sp210-Ag14, FO, NSO/U, MPC-11, MPC11-X45-GTG 1.7 and S194/5XX0 Bul; examples of cell lines used in rat fusions include R210.RCY3, Y3-Ag 1.2.3, IR983F and 4B210. Other cell lines useful for cell fusions are U-266, GM1500-GRG2, LICR-LON-HMy2 and UC729-6.

In one embodiment, a hybridoma cell line is produced by immunizing an animal (*e.g.*, a transgenic animal having human immunoglobulin sequences) with an alpha4beta7 immunogen; harvesting spleen cells from the immunized animal; fusing the harvested spleen cells to a myeloma cell line, thereby generating hybridoma cells; establishing hybridoma cell lines from the hybridoma cells, and identifying a hybridoma cell line that produces an antibody that binds an alpha4beta7 polypeptide. Such hybridoma cell lines, and anti-alpha4beta7 monoclonal antibodies produced by them, are encompassed by the present invention.

Monoclonal antibodies secreted by a hybridoma cell line can be purified using any technique known in the art. Hybridomas or mAbs may be further screened to identify mAbs with particular properties, such as the ability to block an alpha4beta7 induced activity. Examples of such screens are provided in the examples below.

Monoclonal antibodies can also be produced using a process referred to as genetic immunization. For example, a nucleic acid encoding the antigen of interest can be incorporated into a viral vector (such as an adenoviral vector). The resulting vector is then used to develop an immune

response against the antigen of interest in a suitable host animal (for example, a non-obese diabetic, or NOD, mouse). This techniques is substantially described by Ritter et al., Biodrugs16(1): 3 – 10 (2002), the disclosure of which is incorporated by reference herein.

Molecular evolution of the complementarity determining regions (CDRs) in the center of the antibody binding site also has been used to isolate antibodies with increased affinity, for example, antibodies having increased affinity for c-erbB-2, as described by Schier *et al.*, 1996, J. Mol. Biol. 263:551. Accordingly, such techniques are useful in preparing antibodies to alpha4beta7.

Antigen binding proteins directed against an alpha4beta7 can be used, for example, in assays to detect the presence of alpha4beta7 polypeptides or cells expressing alpha4beta7, either *in vitro* or *in vivo*. The antigen binding proteins also may be employed in purifying alpha4beta7 proteins by immunoaffinity chromatography. Those antigen binding proteins that additionally can block the interaction of MAdCAM-1 and alpha4beta7 may be used to inhibit a biological activity that results from such interaction. Blocking antigen binding proteins can be used in the methods of the present invention. Such antigen binding proteins that function as alpha4beta7 antagonists may be employed in treating any alpha4beta7-induced condition, including but not limited to inflammatory conditions. In one embodiment, a human anti- alpha4beta7 monoclonal antibody generated by procedures involving immunization of transgenic mice is employed in treating such conditions.

Antigen binding proteins may be employed in an *in vitro* procedure, or administered *in vivo* to inhibit an alpha4beta7-induced biological activity. Disorders caused or exacerbated (directly or indirectly) by alpha4beta7 and its interaction with MAdCAM-1, examples of which are provided herein, thus may be treated. In one embodiment, the present invention provides a therapeutic method comprising *in vivo* administration of an alpha4beta7 blocking antigen binding protein to a mammal in need thereof in an amount effective for reducing an alpha4beta7-induced biological activity.

Antigen binding proteins of the invention include partially human and fully human monoclonal antibodies that inhibit a biological activity of alpha4beta7. One embodiment is directed to a human monoclonal antibody that at least partially blocks the interaction of human alpha4beta7 with MAdCAM-1. In one embodiment, the antibodies are generated by immunizing a transgenic mouse with an alpha4beta7 immunogen. In another embodiment, the immunogen is a human alpha4beta7 polypeptide (e.g., a cell transformed or transfected to express alpha4beta7, or a cell that naturally expresses alpha4beta7). Hybridoma cell lines derived from such immunized mice, wherein the hybridoma secretes a monoclonal antibody that binds alpha4beta7, also are provided herein.

Although human, partially human, or humanized antibodies will be suitable for many applications, particularly those involving administration of the antibody to a human subject, other types of antigen binding proteins will be suitable for certain applications. The non-human antibodies of the invention can be, for example, derived from any antibody-producing animal, such as mouse, rat, rabbit, goat, donkey, or non-human primate (such as monkey (*e.g.*, cynomologous or rhesus monkey) or ape (*e.g.*, chimpanzee)). Non-human antibodies of the invention can be used, for example, in *in vitro* and cell-culture based applications, or any other application where an immune response to the antibody of the invention does not occur, is insignificant, can be prevented, is not a concern, or is desired. In one embodiment, a non-human antibody of the invention is administered to a non-human subject. In

another embodiment, the non-human antibody does not elicit an immune response in the non-human subject. In another embodiment, the non-human antibody is from the same species as the non-human subject, *e.g.*, a mouse antibody of the invention is administered to a mouse. An antibody from a particular species can be made by, for example, immunizing an animal of that species with the desired immunogen (*e.g.*, cells expressing alph4beta7, or a soluble alpha4beta7 polypeptide) or using an artificial system for generating antibodies of that species (*e.g.*, a bacterial or phage display-based system for generating antibodies of a particular species), or by converting an antibody from one species into an antibody from another species by replacing, *e.g.*, the constant region of the antibody with a constant region from the other species, or by replacing one or more amino acid residues of the antibody so that it more closely resembles the sequence of an antibody from the other species. In one embodiment, the antibody is a chimeric antibody comprising amino acid sequences derived from antibodies from two or more different species.

Antigen binding proteins may be prepared by any of a number of conventional techniques. For example, they may be purified from cells that naturally express them (e.g., an antibody can be purified from a hybridoma that produces it), or produced in recombinant expression systems, using any technique known in the art. See, for example, *Monoclonal Antibodies, Hybridomas: A New Dimension in Biological Analyses*, Kennet *et al.* (eds.), Plenum Press, New York (1980); and *Antibodies: A Laboratory Manual*, Harlow and Land (eds.), Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY, (1988).

Any expression system known in the art can be used to make the recombinant polypeptides of the invention. In general, host cells are transformed with a recombinant expression vector that comprises DNA encoding a desired polypeptide. Among the host cells that may be employed are prokaryotes, yeast or higher eukaryotic cells. Prokaryotes include gram negative or gram positive organisms, for example *E. coli* or bacilli. Higher eukaryotic cells include insect cells and established cell lines of mammalian origin. Examples of suitable mammalian host cell lines include the COS-7 line of monkey kidney cells (ATCC CRL 1651) (Gluzman *et al.*, 1981, Cell 23:175), L cells, 293 cells, C127 cells, 3T3 cells (ATCC CCL 163), Chinese hamster ovary (CHO) cells, HeLa cells, BHK (ATCC CRL 10) cell lines, and the CVI/EBNA cell line derived from the African green monkey kidney cell line CVI (ATCC CCL 70) as described by McMahan *et al.*, 1991, EMBO J. 10: 2821. Appropriate cloning and expression vectors for use with bacterial, fungal, yeast, and mammalian cellular hosts are described by Pouwels *et al.* (*Cloning Vectors: A Laboratory Manual*, Elsevier, New York, 1985).

The transformed cells can be cultured under conditions that promote expression of the polypeptide, and the polypeptide recovered by conventional protein purification procedures. One such purification procedure includes the use of affinity chromatography, *e.g.*, over a matrix having all or a portion (*e.g.*, the extracellular domain) of alpha4beta7 bound thereto. Polypeptides contemplated for use herein include substantially homogeneous recombinant mammalian anti- alpha4beta7 antibody polypeptides substantially free of contaminating endogenous materials.

The amino acid sequence of the polypeptides may be verified by any means known in the art, and may be identical to the sequences disclosed herein in the Sequence Listing, or may differ from those sequences at one or more amino acid residues as result of processing. For example, on all or a

portion of the substantially homogenous polypeptides, a C-terminal amino acid from either the light chain or the heavy chain (or relevant single-chain molecule) may be removed, by proteolytic processing or other processing that occurs during culture, for example, processing of C-terminal Lys residues. Alternatively, more than one C-terminal amino acid residue is removed, for example two C-terminal amino acids, or three, four or five C-terminal amino acids. For example, a C-terminal might be truncated to amidated proline of the heavy chain of an antibody as disclosed. Similarly, N-terminal amino acids may be absent, for example, one, two, three, four or five N-terminal amino acids may be absent.

Alternatively, or additionally, amino acid residues may undergo post-translational modifications, for example but not limited to, glutamine (in particular, glutamine at the N-terminus) may be cyclized or converted to pyroglutamic acid; additionally or alternatively, amino acids may undergo deamidation, isomerization, glycation and/or oxidation. The polypeptides of the invention may undergo additional post-translational modification, including glycosylation, for example N-linked or O-linked glycosylation, at sites that are well-known in the art. As described previously, changes may be made in the amino acid sequence of a polypeptide to preclude or minimize such alterations, or to facilitate them in circumstances where such processing is beneficial.

Preparations of substantially homogenous polypeptides may comprise about 1%, 5%, 10 %, 20%, 30%, 40%, 50%, 60%, 70%, 75%, 80%, 85%, 90%, 95%, 97%, 98%, or 99% polypeptide having undergone a particular form (or forms) of processing. Preparations of substantially homogenous polypeptides may comprise some (less than or equal to 50%), most (more than 50% but less than 90 %) or substantially all (more than 90%) of a particular form(s) of processed polypeptide. Moreover, such preparations may comprise polypeptides that have varying levels of more than one type of processing-related modification, for example, a polypeptide may have some, most or substantially all of a C-terminal lysine removed (for example, the C-terminal lysine in SEQ ID NO: 72) and some, most or substantially all of an N-terminal amino acid converted to pyroglutamic acid (for example, any polypeptide shown in Table 1 and/or 2 or in the consensus sequences).

Antigen binding proteins may be prepared, and screened for desired properties, by any of a number of known techniques. Certain of the techniques involve isolating a nucleic acid encoding a polypeptide chain (or portion thereof) of an antigen binding protein of interest (e.g., an anti-alpha4beta7 antibody), and manipulating the nucleic acid through recombinant DNA technology. The nucleic acid may be fused to another nucleic acid of interest, or altered (*e.g.*, by mutagenesis or other conventional techniques) to add, delete, or substitute one or more amino acid residues, for example.

In one aspect, the present invention provides antigen-binding fragments of an anti-alpha4beta7 antibody of the invention. Such fragments can consist entirely of antibody-derived sequences or can comprise additional sequences. Examples of antigen-binding fragments include Fab, F(ab')2, single chain antibodies, diabodies, triabodies, tetrabodies, and domain antibodies. Other examples are provided in Lunde *et al.*, 2002, Biochem. Soc. Trans. 30:500-06.

Single chain antibodies may be formed by linking heavy and light chain variable domain (Fv region) fragments via an amino acid bridge (short peptide linker), resulting in a single polypeptide chain. Such single-chain Fvs (scFvs) have been prepared by fusing DNA encoding a peptide linker

between DNAs encoding the two variable domain polypeptides (V_L and V_H). The resulting polypeptides can fold back on themselves to form antigen-binding monomers, or they can form multimers (*e.g.*, dimers, trimers, or tetramers), depending on the length of a flexible linker between the two variable domains (Kortt *et al.*, 1997, Prot. Eng. 10:423; Kortt *et al.*, 2001, Biomol. Eng. 18:95-108). By combining different V_L and V_H-comprising polypeptides, one can form multimeric scFvs that bind to different epitopes (Kriangkum *et al.*, 2001, Biomol. Eng. 18:31-40). Techniques developed for the production of single chain antibodies include those described in U.S. Patent No. 4,946,778; Bird, 1988, Science 242:423; Huston *et al.*, 1988, Proc. Natl. Acad. Sci. USA 85:5879; Ward *et al.*, 1989, Nature 334:544, de Graaf *et al.*, 2002, Methods Mol Biol. 178:379-87.

Antigen binding proteins (e.g., antibodies, antibody fragments, and antibody derivatives) of the invention can comprise any constant region known in the art. The light chain constant region can be, for example, a kappa- or lambda-type light chain constant region, *e.g.*, a human kappa- or lambda-type light chain constant region. The heavy chain constant region can be, for example, an alpha-, delta-, epsilon-, gamma-, or mu-type heavy chain constant regions, *e.g.*, a human alpha-, delta-, epsilon-, gamma-, or mu-type heavy chain constant region. In one embodiment, the light or heavy chain constant region is a fragment, derivative, variant, or mutein of a naturally occurring constant region.

Techniques are known for deriving an antibody of a different subclass or isotype from an antibody of interest, *i.e.*, subclass switching. Thus, IgG antibodies may be derived from an IgM antibody, for example, and *vice versa*. Such techniques allow the preparation of new antibodies that possess the antigen-binding properties of a given antibody (the parent antibody), but also exhibit biological properties associated with an antibody isotype or subclass different from that of the parent antibody. Recombinant DNA techniques may be employed. Cloned DNA encoding particular antibody polypeptides may be employed in such procedures, *e.g.*, DNA encoding the constant domain of an antibody of the desired isotype. See also Lantto *et al.*, 2002, Methods Mol. Biol.178:303-16. Moreover, if an IgG4 is desired, it may also be desired to introduce a point mutation (CPSCP -> CPPCP) in the hinge region as described in Bloom *et al.*, 1997, Protein Science 6:407, incorporated by reference herein) to alleviate a tendency to form intra-H chain disulfide bonds that can lead to heterogeneity in the IgG4 antibodies.

Moreover, techniques for deriving antigen binding proteins having different properties (*i.e.*, varying affinities for the antigen to which they bind) are also known. One such technique, referred to as chain shuffling, involves displaying immunoglobulin variable domain gene repertoires on the surface of filamentous bacteriophage, often referred to as phage display. Chain shuffling has been used to prepare high affinity antibodies to the hapten 2-phenyloxazol-5-one, as described by Marks *et al.*, 1992, BioTechnology, 10:779.

In another embodiment, the present invention provides an antigen binding protein that has a low dissociation constant from alpha4beta7. In one embodiment, the antigen binding protein has a K_d of 100 pM or lower. In another embodiment, the K_d is 10 pM or lower; in another embodiment, it is 5 pM or lower, or it is 1 pM or lower. In another embodiment, the K_d is substantially the same as an antibody described herein in the Examples. In another embodiment, the antigen binding protein binds to alpha4beta7 with substantially the same K_d as an antibody described herein in the Examples.

In another aspect, the present invention provides an antigen binding protein that inhibits an activity of alpha4beta7, for example binding (or adhesion) to MAdCAM-1, binding to cells expressing MAdCAM-1, or adhesion between cells expressing alpha4beta 7 and cells expressing MAdCAM-1. In one embodiment, the antigen binding protein has an IC_{50} of 1000pM or lower. In another embodiment, the IC_{50} is 500pM or lower; in another embodiment, the IC_{50} is 100pM or lower. In another embodiment, the IC_{50} is substantially the same as that of an antibody described herein in the Examples. In another embodiment, the antigen binding protein inhibits an activity of alpha4beta7 with substantially the same IC_{50} as an antibody described herein in the Examples.

In one embodiment, antigen binding proteins of the present invention have an apparent affinity for alpha4beta7 (or cells expressing alpha4beta7) of 1000 pM or lower. In other embodiments, the antigen binding proteins exhibit an apparent affinity of 500 pM or lower, 200 pM or lower, 100 pM or lower, 80 pM or lower, 40 pM or lower, or 15 pM or lower. In another embodiment, the antigen binding protein exhibits an apparent affinity substantially the same as that of an antibody described herein in the Examples. In another embodiment, the antigen binding protein has an apparent affinity substantially the same that of an antibody described herein in the Examples.

In another aspect, the present invention provides an antigen binding protein that binds both active and inactive forms of alpha4beta7. In another embodiment, an antigen binding protein binds only one form, or preferentially binds one form, of alpha4beta7. For example, an antigen binding protein may bind alpha4beta7 in the presence or absence of Mn^{2+} (i.e., it binds both active and inactive forms). Alternatively, an antigen binding protein may bind alpha4beta7 only in the presence of Mn^{2+} or only in the absence of Mn^{2+}, or it may bind with higher affinity under one such condition than another, indicating preferential binding to a particular form of alpha4beta7.

In another embodiment, the present invention provides an antigen binding protein that competes for binding to alpha4beta7 with an antibody disclosed herein. Such competitive ability can be determined by methods that are well-known in the art, for example by competition in binding to alpha4beta7-expressing cells as observed using fluorescence activate cells sorting (FACS) techniques or other, similar assays, by competition in an assay such as an adhesion assay (i.e., between cells expressing alpha4beta7 and cells expressing MAdCAM-1), or by competition in another assay described herein. In one aspect, an antigen binding protein that competes for binding to alpha4beta7 with an antibody disclosed herein binds the same epitope or an overlapping (or adjacent) epitope as the antibody. In another aspect, the antigen binding protein that competes for binding to alpha4beta7 with an antibody disclosed herein inhibits an activity of alpha4beta7.

In another aspect, the present invention provides an antigen binding protein that binds to human alpha4beta7 expressed on the surface of a cell and, when so bound, inhibits alpha4beta7 interaction with MAdCAM-1 without causing a significant reduction in the amount of alpha4beta7 on the surface of the cell. Any method for determining or estimating the amount of alpha4beta7 on the surface and/or in the interior of the cell can be used. In one embodiment, the present invention provides an antigen binding protein that binds to alpha4beta7 expressed on the surface of a cell and, when so bound, inhibits alpha4beta7 interaction with MAdCAM-1 without significantly increasing the rate of internalization of the alpha4beta7 from the surface of the cell. In other embodiments, binding of

the antigen binding protein to the alpha4beta7-expressing cell causes less than about 75%, 50%, 40%, 30%, 20%, 15%, 10%, 5%, 1%, or 0.1% of the cell-surface alpha4beta7 to be internalized.

In another aspect, the present invention provides an antigen binding protein having a half-life of at least one day *in vitro* or *in vivo* (*e.g.*, when administered to a human subject). In one embodiment, the antigen binding protein has a half-life of at least three days. In another embodiment, the antigen binding protein has a half-life of four days or longer. In another embodiment, the antigen binding protein has a half-life of eight days or longer. In another embodiment, the antigen binding protein is derivatized or modified such that it has a longer half-life as compared to the underivatized or unmodified antigen binding protein. In another embodiment, the antigen binding protein contains one or more point mutations to increase serum half life, such as described in WO 00/09560, published Feb.24, 2000, incorporated by reference.

The present invention further provides multi-specific antigen binding proteins, for example, bispecific antigen binding protein, *e.g.*, antigen binding protein that bind to two different epitopes of alpha4beta7, or to an epitope of alpha4beta7 and an epitope of another molecule, via two different antigen binding sites or regions. Moreover, bispecific antigen binding protein as disclosed herein can comprise an alpha4beta7 binding site from one of the herein-described antibodies and a second alpha4beta7 binding region from another of the herein-described antibodies, including those described herein by reference to other publications. Alternatively, a bispecific antigen binding protein may comprise an antigen binding site from one of the herein described antibodies and a second antigen binding site from another alpha4beta7 antibody that is known in the art, or from an antibody that is prepared by known methods or the methods described herein.

Numerous methods of preparing bispecific antibodies are known in the art, and discussed in US Patent Application 09/839,632, filed April 20, 2001 (incorporated by reference herein). Such methods include the use of hybrid-hybridomas as described by Milstein *et al.*, 1983, Nature 305:537, and others (U.S. Patent 4,474,893, U.S. Patent 6,106,833), and chemical coupling of antibody fragments (Brennan *et al.*,1985, Science 229:81; Glennie *et al.*,1987, J. Immunol. 139:2367; U.S. Patent 6,010,902). Moreover, bispecific antibodies can be produced via recombinant means, for example by using leucine zipper moieties *(i.e.*, from the Fos and Jun proteins, which preferentially form heterodimers; Kostelny *et al.*, 1992, J. Immnol. 148:1547) or other lock and key interactive domain structures as described in U.S. Patent 5,582,996. Additional useful techniques include those described in Kortt *et al.*, 1997, *supra*; U.S. Patent 5,959,083; and U.S. Patent 5,807,706.

In another aspect, the antigen binding protein of the present invention comprises a derivative of an antibody. The derivatized antibody can comprise any molecule or substance that imparts a desired property to the antibody, such as increased half-life in a particular use. The derivatized antibody can comprise, for example, a detectable (or labeling) moiety (*e.g.*, a radioactive, colorimetric, antigenic or enzymatic molecule, a detectable bead (such as a magnetic or electrodense (*e.g.*, gold) bead), or a molecule that binds to another molecule (*e.g.*, biotin or streptavidin)), a therapeutic or diagnostic moiety (*e.g.*, a radioactive, cytotoxic, or pharmaceutically active moiety), or a molecule that increases the suitability of the antibody for a particular use (*e.g.*, administration to a subject, such as a human subject, or other *in vivo* or *in vitro* uses). Examples of molecules that can be used to derivatize

an antibody include albumin (*e.g.*, human serum albumin) and polyethylene glycol (PEG). Albumin-linked and PEGylated derivatives of antibodies can be prepared using techniques well known in the art. In one embodiment, the antibody is conjugated or otherwise linked to transthyretin (TTR) or a TTR variant. The TTR or TTR variant can be chemically modified with, for example, a chemical selected from the group consisting of dextran, poly(n-vinyl pyurrolidone), polyethylene glycols, propropylene glycol homopolymers, polypropylene oxide/ethylene oxide co-polymers, polyoxyethylated polyols and polyvinyl alcohols. US Pat. App. No. 20030195154.

In another aspect, the present invention provides methods of screening for a molecule that binds to alpha4beta7 using the antigen binding proteins of the present invention. Any suitable screening technique can be used. In one embodiment, an alpha4beta7 molecule, or a fragment thereof to which an antigen binding protein of the present invention binds, is contacted with the antigen binding protein of the invention and with another molecule, wherein the other molecule binds to alpha4beta7 if it reduces the binding of the antigen binding protein to alpha4beta7. Binding of the antigen binding protein can be detected using any suitable method, *e.g.*, an ELISA. Detection of binding of the antigen binding protein to alpha4beta7 can be simplified by detectably labeling the antigen binding protein, as discussed above. In another embodiment, the alpha4beta7-binding molecule is further analyzed to determine whether it inhibits alpha4beta7 activation and/or signaling.

Nucleic acids

In one aspect, the present invention provides isolated nucleic acid molecules. The nucleic acids comprise, for example, polynucleotides that encode all or part of an antigen binding protein , for example, one or both chains of an antibody of the invention, or a fragment, derivative, mutein, or variant thereof, polynucleotides sufficient for use as hybridization probes, PCR primers or sequencing primers for identifying, analyzing, mutating or amplifying a polynucleotide encoding a polypeptide, anti-sense nucleic acids for inhibiting expression of a polynucleotide, and complementary sequences of the foregoing. The nucleic acids can be any length. They can be, for example, 5, 10, 15, 20, 25, 30, 35, 40, 45, 50, 75, 100, 125, 150, 175, 200, 250, 300, 350, 400, 450, 500, 750, 1,000, 1,500, 3,000, 5,000 or more nucleotides in length, and/or can comprise one or more additional sequences, for example, regulatory sequences, and/or be part of a larger nucleic acid, for example, a vector. The nucleic acids can be single-stranded or double-stranded and can comprise RNA and/or DNA nucleotides, and artificial variants thereof (*e.g.*, peptide nucleic acids).

Nucleic acids encoding antibody polypeptides (*e.g.*, heavy or light chain, variable domain only, or full length) may be isolated from B-cells of mice that have been immunized with alpha4beta7. The nucleic acid may be isolated by conventional procedures such as polymerase chain reaction (PCR).

The invention further provides nucleic acids that hybridize to other nucleic acids under particular hybridization conditions. Methods for hybridizing nucleic acids are well-known in the art. See, *e.g.*, Current Protocols in Molecular Biology, John Wiley & Sons, N.Y. (1989), 6.3.1-6.3.6. As defined herein, a moderately stringent hybridization condition uses a prewashing solution containing 5X sodium chloride/sodium citrate (SSC), 0.5% SDS, 1.0 mM EDTA (pH 8.0), hybridization buffer of about 50% formamide, 6X SSC, and a hybridization temperature of 55° C (or other similar

hybridization solutions, such as one containing about 50% formamide, with a hybridization temperature of 42° C), and washing conditions of 60° C, in 0.5X SSC, 0.1% SDS. A stringent hybridization condition hybridizes in 6X SSC at 45° C, followed by one or more washes in 0.1X SSC, 0.2% SDS at 68° C. Furthermore, one of skill in the art can manipulate the hybridization and/or washing conditions to increase or decrease the stringency of hybridization such that nucleic acids comprising nucleotide sequences that are at least 65, 70, 75, 80, 85, 90, 95, 98 or 99% identical to each other typically remain hybridized to each other. The basic parameters affecting the choice of hybridization conditions and guidance for devising suitable conditions are set forth by, for example, Sambrook, Fritsch, and Maniatis (1989, Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N.Y., chapters 9 and 11; and Current Protocols in Molecular Biology, 1995, Ausubel *et al.*, eds., John Wiley & Sons, Inc., sections 2.10 and 6.3-6.4), and can be readily determined by those having ordinary skill in the art based on, for example, the length and/or base composition of the DNA.

Changes can be introduced by mutation into a nucleic acid, thereby leading to changes in the amino acid sequence of a polypeptide (*e.g.*, an antigen binding protein) that it encodes. Mutations can be introduced using any technique known in the art. In one embodiment, one or more particular amino acid residues are changed using, for example, a site-directed mutagenesis protocol. In another embodiment, one or more randomly selected residues is changed using, for example, a random mutagenesis protocol. However it is made, a mutant polypeptide can be expressed and screened for a desired property (*e.g.*, binding to alpha4beta7 or blocking the binding of alpha4beta7 to an addressin such as MAdCAM).

Mutations can be introduced into a nucleic acid without significantly altering the biological activity of a polypeptide that it encodes. For example, one can make nucleotide substitutions leading to amino acid substitutions at non-essential amino acid residues. In one embodiment, a nucleotide sequence, or a desired fragment, variant, or derivative thereof, is mutated such that it encodes an amino acid sequence comprising one or more deletions or substitutions of amino acid residues. In another embodiment, the mutagenesis inserts an amino acid adjacent to one or more amino acid residues. Alternatively, one or more mutations can be introduced into a nucleic acid that selectively change the biological activity (*e.g.*, binding of alpha4beta7, inhibiting binding of alpha4beta7 to an addressin such as MAdCAM, *etc.*) of a polypeptide that it encodes. For example, the mutation can quantitatively or qualitatively change the biological activity. Examples of quantitative changes include increasing, reducing or eliminating the activity. Examples of qualitative changes include changing the antigen specificity of an antigen binding protein.

In another aspect, the present invention provides nucleic acid molecules that are suitable for use as primers or hybridization probes for the detection of nucleic acid sequences of the invention. A nucleic acid molecule of the invention can comprise only a portion of a nucleic acid sequence encoding a full-length polypeptide of the invention, for example, a fragment that can be used as a probe or primer or a fragment encoding an active portion (*e.g.*, an alpha4beta7 binding portion) of a polypeptide of the invention.

Probes based on the sequence of a nucleic acid of the invention can be used to detect the nucleic acid or similar nucleic acids, for example, transcripts encoding a polypeptide of the invention. The probe can comprise a label group, *e.g.*, a radioisotope, a fluorescent compound, an enzyme, or an enzyme co-factor. Such probes can be used to identify a cell that expresses the polypeptide.

In another aspect, the present invention provides vectors comprising a nucleic acid encoding a polypeptide of the invention or a portion thereof. Examples of vectors include, but are not limited to, plasmids, viral vectors, non-episomal mammalian vectors and expression vectors, for example, recombinant expression vectors.

The recombinant expression vectors of the invention can comprise a nucleic acid of the invention in a form suitable for expression of the nucleic acid in a host cell. The recombinant expression vectors include one or more regulatory sequences, selected on the basis of the host cells to be used for expression, which is operably linked to the nucleic acid sequence to be expressed. Regulatory sequences include those that direct constitutive expression of a nucleotide sequence in many types of host cells (*e.g.*, SV40 early gene enhancer, Rous sarcoma virus promoter and cytomegalovirus promoter), those that direct expression of the nucleotide sequence only in certain host cells (*e.g.*, tissue-specific regulatory sequences, see Voss *et al.*, 1986, Trends Biochem. Sci. 11:287, Maniatis *et al.*, 1987, Science 236:1237, incorporated by reference herein in their entireties), and those that direct inducible expression of a nucleotide sequence in response to particular treatment or condition (*e.g.*, the metallothionin promoter in mammalian cells and the tet-responsive and/or streptomycin responsive promoter in both prokaryotic and eukaryotic systems (see *id.*). It will be appreciated by those skilled in the art that the design of the expression vector can depend on such factors as the choice of the host cell to be transformed, the level of expression of protein desired, *etc.* The expression vectors of the invention can be introduced into host cells to thereby produce proteins or peptides, including fusion proteins or peptides, encoded by nucleic acids as described herein.

In another aspect, the present invention provides host cells into which a recombinant expression vector of the invention has been introduced. A host cell can be any prokaryotic cell (for example, *E. coli*) or eukaryotic cell (for example, yeast, insect, or mammalian cells (*e.g.*, CHO cells)). Vector DNA can be introduced into prokaryotic or eukaryotic cells via conventional transformation or transfection techniques. For stable transfection of mammalian cells, it is known that, depending upon the expression vector and transfection technique used, only a small fraction of cells may integrate the foreign DNA into their genome. In order to identify and select these integrants, a gene that encodes a selectable marker (*e.g.*, for resistance to antibiotics) is generally introduced into the host cells along with the gene of interest. Preferred selectable markers include those that confer resistance to drugs, such as G418, hygromycin and methotrexate. Cells stably transfected with the introduced nucleic acid can be identified by drug selection (*e.g.*, cells that have incorporated the selectable marker gene will survive, while the other cells die), among other methods.

Indications

In one aspect, the present invention provides methods of treating a subject. The method can, for example, have a generally salubrious effect on the subject, *e.g.*, it can increase the subject's

expected longevity. Alternatively, the method can, for example, treat, prevent, cure, relieve, or ameliorate ("treat") a disease, disorder, condition, or illness ("a condition"). Among the conditions to be treated in accordance with the present invention are conditions characterized by inappropriate expression or activity of alpha4beta7. Such conditions include those that are associated with inappropriate trafficking of cells, for example, the trafficking of leukocytes (such as lymphocytes or monocytes) to the gastrointestinal tract or other tissues comprising cells that express MAdCAM-1 (as a result of binding of the leukocytes to the cells that express MAdCAM-1). Diseases which can be treated accordingly include inflammatory bowel disease, such as ulcerative colitis, Crohn's disease, Celiac disease (nontropical Sprue), enteropathy associated with seronegative arthropathies, microscopic or collagenous colitis, eosinophilic gastroenteritis, or pouchitis resulting after proctocolectomy and ileoanal anastomosis. Additional conditions that may be treated in accordance with the present invention include pancreatitis, insulin-dependent diabetes mellitus, mastitis, cholecystitis, cholangitis, pericholangitis, chronic bronchitis, chronic sinusitis, asthma and graft versus host disease.

Therapeutic methods and administration of antigen binding proteins

Certain methods provided herein comprise administering an alpha4beta7 heterodimer specific antigen binding protein to a subject, thereby reducing an alpha4beta7-induced biological response that plays a role in a particular condition. In particular embodiments, methods of the invention involve contacting endogenous alpha4beta7 with an alpha4beta7 antigen binding protein, *e.g.*, via administration to a subject or in an *ex vivo* procedure.

The term "treatment" encompasses alleviation or prevention of at least one symptom or other aspect of a disorder, or reduction of disease severity, and the like. An antigen binding protein need not effect a complete cure, or eradicate every symptom or manifestation of a disease, to constitute a viable therapeutic agent. As is recognized in the pertinent field, drugs employed as therapeutic agents may reduce the severity of a given disease state, but need not abolish every manifestation of the disease to be regarded as useful therapeutic agents. Similarly, a prophylactically administered treatment need not be completely effective in preventing the onset of a condition in order to constitute a viable prophylactic agent. Simply reducing the impact of a disease (for example, by reducing the number or severity of its symptoms, or by increasing the effectiveness of another treatment, or by producing another beneficial effect), or reducing the likelihood that the disease will occur or worsen in a subject, is sufficient. One embodiment of the invention is directed to a method comprising administering to a patient an alpha4beta7 antagonist in an amount and for a time sufficient to induce a sustained improvement over baseline of an indicator that reflects the severity of the particular disorder.

As is understood in the pertinent field, pharmaceutical compositions comprising the molecules of the invention are administered to a subject in a manner appropriate to the indication. Pharmaceutical compositions may be administered by any suitable technique, including but not limited to parenterally, topically, or by inhalation. If injected, the pharmaceutical composition can be administered, for example, via intra-articular, intravenous, intramuscular, intralesional, intraperitoneal or subcutaneous routes, by bolus injection, or continuous infusion. Localized administration, *e.g.* at a site of disease or injury is contemplated, as are transdermal delivery and sustained release from implants. Delivery by

inhalation includes, for example, nasal or oral inhalation, use of a nebulizer, inhalation of the antagonist in aerosol form, and the like. Other alternatives include eyedrops; oral preparations including pills, syrups, lozenges or chewing gum; and topical preparations such as lotions, gels, sprays, and ointments.

Use of antigen binding proteins in *ex vivo* procedures also is contemplated. For example, a patient's blood or other bodily fluid may be contacted with an antigen binding protein that binds alpha4beta7 *ex vivo*. The antigen binding protein may be bound to a suitable insoluble matrix or solid support material.

Advantageously, antigen binding proteins are administered in the form of a composition comprising one or more additional components such as a physiologically acceptable carrier, excipient or diluent. Optionally, the composition additionally comprises one or more physiologically active agents, for example, a second inflammation- or immune-inhibiting substance, an anti-angiogenic substance, an analgesic substance, *etc.*, non-exclusive examples of which are provided herein. In various particular embodiments, the composition comprises one, two, three, four, five, or six physiologically active agents in addition to an alpha4beta7 binding antigen binding protein

In one embodiment, the pharmaceutical composition comprise an antigen binding protein of the invention together with one or more substances selected from the group consisting of a buffer, an antioxidant such as ascorbic acid, a low molecular weight polypeptide (such as those having fewer than 10 amino acids), a protein, an amino acid, a carbohydrate such as glucose, sucrose or dextrins, a chelating agent such as EDTA, glutathione, a stabilizer, and an excipient. Neutral buffered saline or saline mixed with conspecific serum albumin are examples of appropriate diluents. In accordance with appropriate industry standards, preservatives such as benzyl alcohol may also be added. The composition may be formulated as a lyophilizate using appropriate excipient solutions (*e.g.*, sucrose) as diluents. Suitable components are nontoxic to recipients at the dosages and concentrations employed. Further examples of components that may be employed in pharmaceutical formulations are presented in Remington's Pharmaceutical Sciences, 16th Ed. (1980) and 20th Ed. (2000), Mack Publishing Company, Easton, PA.

Kits for use by medical practitioners include an alpha4beta7-inhibiting substance of the invention and a label or other instructions for use in treating any of the conditions discussed herein. In one embodiment, the kit includes a sterile preparation of one or more alpha4beta7 binding antigen binding proteins, which may be in the form of a composition as disclosed above, and may be in one or more vials.

Dosages and the frequency of administration may vary according to such factors as the route of administration, the particular antigen binding proteins employed, the nature and severity of the disease to be treated, whether the condition is acute or chronic, and the size and general condition of the subject. Appropriate dosages can be determined by procedures known in the pertinent art, *e.g.* in clinical trials that may involve dose escalation studies.

An alpha4beta7 inhibiting substance of the invention may be administered, for example, once or more than once, *e.g.*, at regular intervals over a period of time. In particular embodiments, an antigen binding protein is administered over a period of at least a month or more, *e.g.*, for one, two, or

three months or even indefinitely. For treating chronic conditions, long-term treatment is generally most effective. However, for treating acute conditions, administration for shorter periods, *e.g.* from one to six weeks, may be sufficient. In general, the antigen binding protein is administered until the patient manifests a medically relevant degree of improvement over baseline for the chosen indicator or indicators.

Particular embodiments of the present invention involve administering an antigen binding protein at a dosage of from about 1 ng of antigen binding protein per kg of subject's weight per day ("1ng/kg/day") to about 10 mg/kg/day, more preferably from about 500 ng/kg/day to about 5 mg/kg/day, and most preferably from about 5 μg/kg/day to about 2 mg/kg/day, to a subject. In additional embodiments, an antigen binding protein is administered to adults one time per week, two times per week, or three or more times per week, to treat an alpha4beta7 mediated disease, condition or disorder, *e.g.*, a medical disorder disclosed herein. If injected, the effective amount of antigen binding protein per adult dose may range from 1-20 mg/m^2, and preferably is about 5-12 mg/m^2. Alternatively, a flat dose may be administered; the amount may range from 5-100 mg/dose. One range for a flat dose is about 20-30 mg per dose. In one embodiment of the invention, a flat dose of 25 mg/dose is repeatedly administered by injection. If a route of administration other than injection is used, the dose is appropriately adjusted in accordance with standard medical practices. One example of a therapeutic regimen involves injecting a dose of about 20-30 mg of antigen binding protein to one to three times per week over a period of at least three weeks, though treatment for longer periods may be necessary to induce the desired degree of improvement. For pediatric subjects (age 4-17), one exemplary suitable regimen involves the subcutaneous injection of 0.4 mg/kg, up to a maximum dose of 25 mg of antigen binding protein administered two or three times per week.

Particular embodiments of the methods provided herein involve subcutaneous injection of from 0.5 mg to 10 mg, preferably from 3 to 5 mg, of an antigen binding protein, once or twice per week. Another embodiment is directed to pulmonary administration (*e.g.*, by nebulizer) of 3 or more mg of antigen binding protein once a week.

Examples of therapeutic regimens provided herein comprise subcutaneous injection of an antigen binding protein once a week, at a dose of 1.5 to 3 mg, to treat a condition in which alpha4beta7 plays a role. Examples of such conditions are provided herein and include, for example, rheumatic conditions as previously described, and other conditions in which excessive or inappropriate trafficking of alpha4beta7-expressing cells plays a role (described herein; for example, inflammatory bowel disease, pancreatitis, etc). Weekly administration of antigen binding protein is continued until a desired result is achieved, *e.g.*, the subject's symptoms subside. Treatment may resume as needed, or, alternatively, maintenance doses may be administered.

Other examples of therapeutic regimens provided herein comprise subcutaneous or intravenous administration of a dose of 1, 3, 5, 6, 7, 8, 9, 10, 11, 12, 15, or 20 milligrams of an alpha4beta7 inhibitor of the present invention per kilogram body mass of the subject (mg/kg). The dose can be administered once to the subject, or more than once at a certain interval, for example, once a day, three times a week, twice a week, once a week, three times a month, twice a month, once a month, once every two months, once every three months, once every six months, or once a year. The

duration of the treatment, and any changes to the dose and/or frequency of treatment, can be altered or varied during the course of treatment in order to meet the particular needs of the subject.

In another embodiment, an antigen binding protein is administered to the subject in an amount and for a time sufficient to induce an improvement, preferably a sustained improvement, in at least one indicator that reflects the severity of the disorder that is being treated. Various indicators that reflect the extent of the subject's illness, disease or condition may be assessed for determining whether the amount and time of the treatment is sufficient. Such indicators include, for example, clinically recognized indicators of disease severity, symptoms, or manifestations of the disorder in question. In one embodiment, an improvement is considered to be sustained if the subject exhibits the improvement on at least two occasions separated by two to four weeks. The degree of improvement generally is determined by a physician, who may make this determination based on signs, symptoms, biopsies, or other test results, and who may also employ questionnaires that are administered to the subject, such as quality-of-life questionnaires developed for a given disease.

Alteration of alpha4beta7 expression and/or activation of alpha4beta7, and or its binding partner MAdCAM-1, are associated with a number of disorders, including, for example, inflammatory conditions of the gastrointestinal system. Subjects with a given disorder may be screened, to identify those individuals who have altered alpha4beta7 or MAdCAM-1 expression and/or activation, thereby identifying the subjects who may benefit most from treatment with an alpha4beta7 binding antigen binding protein. Thus, treatment methods provided herein optionally comprise a first step of measuring a subject's alpha4beta7 or MAdCAM-1activation or expression levels. An antigen binding protein may be administered to a subject in whom alpha4beta7 and/or MAdCAM-1 expression and/or activation is elevated above normal.

A subject's levels of alpha4beta7 or MAdCAM-1 activity may be monitored before, during and/or after treatment with an antigen binding protein, to detect changes, if any, in alpha4beta7 or MAdCAM-1 activity. For some disorders, the incidence of elevated alpha4beta7 and/or MAdCAM-1 activity may vary according to such factors as the stage of the disease or the particular form of the disease. Known techniques may be employed for measuring such activity, *e.g.*, in a subject's blood or tissue samples. Alpha4beta7or MAdCAM1 activity may be measured using any suitable technique.

Particular embodiments of methods and compositions of the invention involve the use of an antigen binding protein and one or more additional alpha4beta7 antagonists, for example, two or more antigen binding proteins of the invention, or an antigen binding protein of the invention and one or more other alpha4beta7 antagonists. In further embodiments, antigen binding protein are administered alone or in combination with other agents useful for treating the condition with which the patient is afflicted. Examples of such agents include both proteinaceous and non-proteinaceous drugs. When multiple therapeutics are co-administered, dosages may be adjusted accordingly, as is recognized in the pertinent art. "Co-administration" and combination therapy are not limited to simultaneous administration, but also include treatment regimens in which an antigen binding protein is administered at least once during a course of treatment that involves administering at least one other therapeutic agent to the patient.

Examples of other agents that may be co-administered with an antigen binding protein are other antigen binding proteins or therapeutic polypeptides that are chosen according to the particular condition to be treated. Alternatively, non-proteinaceous drugs that are useful in treating one of the particular conditions discussed above may be co-administered with an alpha4beta7 antagonist.

Combination therapy

In another aspect, the present invention provides a method of treating a subject with an alpha4beta7 inhibiting antigen binding protein and one or more other treatments. In one embodiment, such a combination therapy achieves synergy or an additive effect by, for example, attacking multiple sites or molecular targets in a tumor. Types of combination therapies that can be used in connection with the present invention include inhibiting or activating (as appropriate) multiple nodes in a single disease-related pathway, multiple pathways in a target cell, and multiple cell types within a target tissue.

In another embodiment, a combination therapy method comprises administering to the subject two, three, four, five, six, or more of the alpha4beta7 agonists or antagonists described herein. In another embodiment, the method comprises administering to the subject two or more treatments that together inhibit or activate (directly or indirectly) alpha4beta7-mediated signal transduction. Examples of such methods include using combinations of two or more alpha4beta7 inhibiting antigen binding proteins, of an alpha4beta7 inhibiting antigen binding protein and one or more other therapeutic moiety having anti-inflammatory properties (for example, non-steroidal anti-inflammatory agents, steroids, and/or immunomodulators), or of an alpha4beta7 inhibiting antigen binding protein and one or more other treatments (*e.g.*, surgery, ultrasound, or treatment effective to reduce inflammation). Useful agents that may be combined with alpha4beta7 inhibitors include those used to treat, for example, Crohn's disease or ulcerative colitis, such as aminosalicylate (for example, mesalamine), corticosteroids (including predisone), antibiotics such as metronidazole or ciprofloxacin (or other antibiotics useful for treating, for example, patients afflicted with fistulas), and immunosupporessives such as azathioprine, 6-mercaptopurine, methotrexate, tacrolimus and cyclosporine. Combinations of such agents are also contemplated for use with the inventive alpha4beta7 inhibitors. Such agent(s) may be administered orally or by another route, for example via suppository or enema.

Furthermore, one or more anti-alpha4beta7 antibodies or antibody derivatives can be used in combination with one or more molecules or other treatments, wherein the other molecule(s) and/or treatment(s) do not directly bind to or affect alpha4beta7, but which combination is effective for treating or preventing the condition being treated. For example, an alpha4eta7 inhibitor can be used in combination with probiotic therapy, or other therapy used to restore or maintain normal gut flora. In one embodiment, one or more of the molecule(s) and/or treatment(s) treats or prevents a condition that is caused by one or more of the other molecule(s) or treatment(s) in the course of therapy, *e.g.*, nausea, fatigue, alopecia, cachexia, insomnia, *etc.* In every case where a combination of molecules and/or other treatments is used, the individual molecule(s) and/or treatment(s) can be administered in any order, over any length of time, which is effective, *e.g.*, simultaneously, consecutively, or alternately. In one embodiment, the method of treatment comprises completing a first course of treatment with one

molecule or other treatment before beginning a second course of treatment. The length of time between the end of the first course of treatment and beginning of the second course of treatment can be any length of time that allows the total course of therapy to be effective, *e.g.*, seconds, minutes, hours, days, weeks, months, or even years.

In another embodiment, the method comprises administering one or more of the alpha4beta7 antagonists described herein and one or more other treatments (*e.g.*, a therapeutic or palliative treatment). Where a method comprises administering more than one treatment to a subject, it is to be understood that the order, timing, number, concentration, and volume of the administrations is limited only by the medical requirements and limitations of the treatment, *i.e.*, two treatments can be administered to the subject, *e.g.*, simultaneously, consecutively, alternately, or according to any other regimen.

The following examples, both actual and prophetic, are provided for the purpose of illustrating specific embodiments or features of the instant invention and do not limit its scope.

EXAMPLE 1: PREPARATION OF ANTIBODIES

Monoclonal antibodies against human alpha4beta7 were developed by immunizing XenoMouse™ XG2kappalambda (kl) and XG4kl mice (transgenic mice that express human IgG2 or IgG4, and human kappa and lambda light chains, respectively; Abgenix Inc., Fremont CA) with cells expressing human alpha4beta7, either transiently transfected human embryonic kidney (HEK) 293 cells (293-a4b7) or stably transfected Chinese hamster ovary (CHO) cells (CHO-a4b7). Serum titer was monitored by fluorescence activated cell sorter (FACS) analysis comparing alpha4beta7 transfected cells to the respective parental control cells. Hyperimmune animals from either immunization campaign were sacrificed and spleen and lymph node tissues were subjected to hybridoma fusion.

Alpha4beta7 heterodimer specific antibodies were identified using a series of assays. Hybridoma supernatants were first screened by Fluorometric Microvolume Assay Technology (FMAT™ Applera Corporation, Foster City CA; a high-throughput screening cellular detection system) for binding to alpha4beta7 transfected cells as compared to mock-transfected cells. Supernatants identified as positive for binding to alpha4beta7 (1001 positive binding supernatants from CHO-a4b7 cell immunization campaign and 1143 positive binding supernatants from 293-a4b7 cell immunization campaign) were evaluated for the ability to inhibit HUT78 cell adhesion to MAdCAM-1-Fc in a similar fashion as described (Erle, J. Immunol, (1994) 153:517). In this assay, 60 supernatants from CHO-a4b7 campaign and 174 supernatants from 293-a4b7 campaign showed greater than 90% inhibition (n=2) and were subject to further specificity and potency analysis.

Alpha4beta7-transfected, alpha4beta1-transfected, and alphaEbeta7-transfected 293 cells were prepared and used in FACS analysis with the hybridoma supernatants that were identified in the inhibition assay. Supernatants that demonstrated binding to only the alpha4beta7 transfected cells were classified as heterodimers-specific, since antibodies to the alpha4 subunit of this integrin would also bind the alpha4beta1-transfected cells, and antibodies that bound the beta7 chain would bind alphaEbeta7-transfected cells. The hybridoma supernatants were also analyzed for binding activity to cynomologous monkey alpha4beta7-transfected 293 cells by FACS analysis. Seven lines from the

CHO-a4b7 campaign and 25 lines from the 293-a4b7 campaign were selected for sub-cloning and further analysis.

EXAMPLE 2: ANALYSIS OF ANTIBODIES

The antibody-secreting cells obtained were cloned, and the antibody-encoding nucleic acids were isolated and sequenced. Site directed mutagenesis was used to prepared variants that differed from the isolated sequences at one or more amino acid residues. The amino acid sequence of the light and heavy chains of the antibodies and variants are shown in Tables 1 and 2 below. It is recognized that the boundaries of the CDR and FR regions can vary from that shown below, as discussed previously herein.

Table 1: Sequence analysis of light chains

Light chain	FR1	CDR1	FR2
1A10K	DIQMTQSPSSVSASVGDRVTITC	RASQGVSSWLA	WYQQKPGMAPKLLIY
11E7K1	EIVMTQSPATLSVSPGETATLSC	RASQTVSSNLA	WYQQKPGQAPRLLIY
11E7K2	DIQMTQSPSSLSASIGDRVTITC	RASQGIRNYLA	WYQRKPGKVPKLLIY
2F12K	DIQMTQSPSSVFASVGDRVTITC	RASQGISSWLA	WYQQKPGKAPNLLIY
14E4L	QSVLTQPPSVSAAPGQKVTISC	SGSSSNIGNNYVS	WYQQLPGTAPKLLIY
3A5K	DIQMTQSPSSVSASVGDRVTITC	RASQGVISWLA	WYQQKPGMAPKLLIY
10D7K	DIQMTQSPSSVSASVGDRVTITC	RASQGVNNWLA	WYQQKPGKAPKLLIF
27D8K	EIVMMQSPATLSVSPGERATLSC	RASQSVSTNLA	WYQQKPGQAPRLLIY
18A11K	DIQMTQSPSSVSASVGDRVTITC	RASQGISSWLA	WYQQKPGKAPKLLIY
20D7K	EIVLTQSPGTLSLSPGERATLSC	RASQSVSSSYLA	WYQQKPGQAPRLLIY
23H6K	EIVMTQSPATLSVSPGERATLSC	RASQSVNSNLA	WYQQKPGQAPRLLIY
27G8L	QSVLTQPPSVSEAPRQRVTISC	SGSNSNIGNNPVN	WYQLFPGRAPKLLIY
26C7K	EIVMTQSPATLSVSPGERATLSC	RASQSVSDNLA	WYQQKPGQPPRLLIY
26H3K	DIQMTQSPSSLSASVGDRVTITC	QASQDISNYLN	WYQQKPGKAPKLLIY
19G6K	DIQMTQSPSSLSASVGDRVTISC	QASQDINTYLN	WYQQKPGKVPKLLIY
22B2K	DVQMTQSPSSLSASVGDRVTITC	QASQDITDYLN	WYQQKPGKAPKLLIY
24A2K	EVMMTQSPATLSVSPGERATLSC	RASQSVSSNLA	WYQQKPGQAPRLLIF
26E9K	ELVMTQSPATLSVSPGERATVSC	RASQSVSSDLA	WYQQKPGQAPRLLIY
22F5K	EIVMTQSPATLSVFPGEGATLSC	RASQSVSSDLA	WYQQKPGQAPRLLIY
26C10K	EIVLTQSPGTLSLSPGEGATLSC	RASQTVTSSYLA	WYQQSPSQSPRLLIY
17C8K	EIVMTQSPATLSVSPGERATLSC	RASQSVSSNLV	WYQQKPGQAPRLLIY
25C9k	DIQMTQSPSSVSASVGDRVTITC	RASQDISSWLA	WYQRKPGKAPKVLIY
19E6L	SYELTQPPSVSVSPGQTASITC	SGDKLGDKYAC	WYQQKPGQSPVLVIY
26G2k	DIQMTQSPSSVSASVGDRVTITC	RASQDISSWLA	WYQQKPGTAPKVLIY
27G8L (a)	QSVLTQPPSVSGAPRQRVTISC	SGSNSNIGNNPVN	WYQLFPGRAPKLLIY
27G8L (b)	QSVLTQPRSVSGAPRQRVTISC	SGSNSNIGNNPVN	WYQLFPGRAPKLLIY
26H3K (c)	DIQMTQSPSSLSASVGDRVTITC	QASQDISNYLN	WYQQKPGKAPKLLIY
1A10K (d)	DIQMTQSPSSVSASVGDRVTITC	RASQGVSSWLA	WYQQKPGKAPKLLIY

Table 1 (cont'd)

Light chain	CDR2	FR3
1A10K	AASILQS	GVPSRFSGSGSGTDFTLTISSLQPEDFATYYC
11E7K1	GASTRAT	GIPARFSGSGSGTEFTLTISSLQSEDFAVYYC
11E7K2	AASTLQS	GVPSRFSGSGSGTDFTLTISSLQPEDVATYCC
2F12K	GASSLQN	GVPLRFSGSGSGTDFTLTISSLQPEDFATYYC
14E4L	DNNKRPS	GIPDRFSGSKSGTSAILDITGLQTGDEADYYC
3A5K	AASILQS	GVPSRFSGSGSGTDFTLTISSLQPEDFATYYC

10D7K	ATSSLQS	GVPSRFSGSGSGTDFTLTINSLQPEDFATYYC
27D8K	GASTRAT	GIPARFSGSGSGTEFTLTISSLQSEDFAVYFC
18A11K	GASNLES	GVPSRFSGSGSGTDFTLTISSLQPEDFANYYC
20D7K	GASSRAT	GIPDRFSGSGSGTDFTLTISRLEPEDFAVYYC
23H6K	GASTRAT	GIPARFSGSGSGTEFTLTISSLQSEDFAVYYC
27G8L	HDDLLPS	GVSDRFSGSRSGTSASLAISGLQSEDETDYYC
26C7K	GASTRAT	GIPARFSGSGSGTEFTLTISSLQSEDFAVYYC
26H3K	DASNLET	GVPSRFSGSGSGTDFTFTINSLQPEDIATYFC
19G6K	DASNLET	GVPSRFSGSGSGTDFTFTISGLQPEDIATYYC
22B2K	DTSNLEA	GVPSRFSGSGSGTDFTFTISSLQPEDIATYYC
24A2K	GASTRAT	GIPARFSGSGSGTEFTLTISSLQSEDFAVYCC
26E9K	GASSRAT	GIPARFSGSGSGTEFTLTISSLQSEDFAVYYC
22F5K	GASARAT	GIPARFSGSGSGTEFTLTISSLQSEDFAVYYC
26C10K	GASTRAT	GIPDRFSGSGSGTDFTLTISRLEPEDFAVYYC
17C8K	GASTRAT	GIPARFSGSGSGTDFTLTISSLQSEDFAVYYC
25C9k	SASSLQS	GVPSRFSGSGSGTDFTLTISSLQPEDFATYYC
19E6L	QDSKRPS	GIPERFSGSNSGNTATLTISGTQAMDEADYYC
26G2k	SASSLQN	GVPSRFSGRGSGTDFALTISSLQPEDFATYYC
27G8L (a)	HDDLLPS	GVSDRFSGSRSGTSASLAISGLQSADETDYYC
27G8L (b)	HDDLLPS	GVSDRFSGSRSGTSASLAISGLRSADETDYYC
26H3K (c)	DASNLET	GVPSRFSGSGSGTDFTFTINSLQPEDIATYFC
1A10K (d)	AASILQS	GVPSRFSGSGSGTDFTLTISSLQPEDFATYYC

Table 1 (cont'd)

Light chain	CDR3	FR4
1A10K	QQANSFPWT	FGQGTKVEIK
11E7K1	QQYDYWPPLT	FGGGTRVEIK
11E7K2	QKYDSAPFT	FGPGTKVDIK
2F12K	QQANSFPWT	FGQGTKVEIK
14E4L	GTWDSSLSAGRV	FGGGTKLTVL
3A5K	QQANSFPWT	FGQGTNVEIK
10D7K	QQVNSFPGT	FGQGTKVEIK
27D8K	QQYNDWPT	FGGGTKVEIK
18A11K	QQANSFPWT	FGQGTKVEIK
20D7K	QQYDSSPPT	FGGGTKVAIK
23H6K	QQYDDWPPVT	FGQGTRLEIK
27G8L	TAWDDSLNGWV	FGGGTKLTVL
26C7K	QQYDDWPT	FGGGTRVEIK
26H3K	QQYDNLPCS	FGQGTKLEIK
19G6K	QQFDNLPIT	FGQGTRLEIK
22B2K	QQYDILPYS	FGQGTDLEIK
24A2K	QQYDDWPT	FGGGTKVEIK
26E9K	QQYNNWPPLT	FGGGTKVEIK
22F5K	QQYHDWPPLS	FGGGTKVEIK
26C10K	QQYDSSPPT	FGGGTKVEIK
17C8K	QQYDDWPPLT	FGGGTTVEIK
25C9k	QQADSFPWT	FGQGTKVEIK
19E6L	QAWDSSTVV	FGGGTKLTVL
26G2k	QQADSFPWT	FGRGTKVEIK
27G8L (a)	TAWDDSLNGWV	FGGGTKLTVL
27G8L (b)	TAWDDSLNGWV	FGGGTKLTVL
26H3K (c)	QQYDNLPSS	FGQGTKLEIK
1A10K (d)	QQANSFPWT	FGQGTKVEIK

Table 2: Sequence analysis of heavy chains

Heavy	FR1	CDR1	FR2

chain			
1A10H	QVQLVQSGAEVKKPGASVKVSCKVSGYTLN	DLSMH	WVRQAPGKGLEWMG
11E7H1	QVQLVESGGGLVKPGGSLRLSCVASGFTFS	DYYMS	WIRQAPGKGLEWVS
11E7H2	QVQLVESGGGVVQPGRSLRLSCAASGFTFS	SYGMH	WVRQAPGKGLEWVA
2F12H	QVQLVQSGAEVKKPGASVKVSCKVSGYTVT	DLSMH	WVRQAPGKGLEWMG
14E4H	QVQLVESGGGLVKPGGSLRLSCAASGFTFS	DYYMS	WIRQAPGKGLEWVS
3A5H	QVQLVQSGAEVKKPGASVKVSCKVSGYTLN	DLSMH	WVRQAPGKGLEWMG
10D7H	QVQLVESGGGLVKPGGSLRLSCAASGFTFS	DYYMS	WIRQAPGKGLEWVS
27D8H	QVQLVESGGGLVKPGGSLRLSCAASGFTFS	DNYMS	WIRQAPGKGLEWVS
18A11H	QVQLVQSGAEVKKPGASVKVSCKVSGYTLS	DLSIH	WVRQAPGKGLEWMG
20D7H	QVQLVESGGGLVKPGGSLRLSCTASGFTFS	DYYMS	WIRQAPGKGLEWVS
23H6H	QVQLVESGGGLVKPGGSLRLSCAASGFTFS	DYYMS	WIRQAPGKGLEWVS
26G2H	QVQLVESGGGLVKPGGSLRLSCAASGFTFS	DYYMS	WIRQAPGKGLEWVS
27G8H	EVQLVESGGGLVQPGGSLRLSCAASGFTFS	SYWMS	WVRQASGKGLEWVA
26C7H	QVQLVESGGGLVKPGGSLRLSCAASGFTFS	DYYMS	WIRQAPGKGLEWVS
26H3H	EVQLVQSGAEVKKPGESLKISCKGSGYSFT	GYWIG	WVRQMPGKGLEWMG
19G6H	QVQLVESGGDLVKPGGSLRLSCAASGFTFS	DYYMS	WIRQAPGKGLEWIS
22B2H	EVQLVQSGAEVKEPGESLKISCKGSGYIFT	SYWIA	WVRQLPGKGLEWMG
24A2H	QVQLVESGGDLVEPGGSLRLSCAASGFTFR	DYYMS	WIRQAPGKGLEWVS
26E9H	QVQLVESGGGLVKPGGSLRLSCAASGFTFR	DYYMS	WIRQAPGKGLEWVS
19E6H	EVQLLESGGGLVQPGGSLRLSCAASGFTFS	SYAMS	WVRQAPGKGLEWVS
22F5H	QVQLVESGGGLVKPGGSLRLSCAASGFTFS	DYYMS	WIRQAPGKGLEWVS
25C9H	QVQLVESGGGLVKPGGSLRLSCAASGFTFN	DYYMS	WIRQAPGKGLEWVS
26C10H	QVQLVESGGGLVKPGGSLRLSCVASGFTFS	DYYMS	WIRQTPGKGLEWVS
17C8H	QVQLVESGGGLVKPGGSLRLSCAASGFTFS	DYYMS	WIRQAPGKGLEWLS
1A10H(a)	QVQLVQSGAEVKKPGASVKVSCKVSGYTLN	DLSMH	WVRQAPGKGLEWMG
27G8H(b)	EVQLVESGGGLVKPGRSLRLSCAASGFTFS	SYWMS	WVRQASGKGLEWVA

Table 2 (cont'd)

Heavy chain	CDR2	FR3
1A10H	GFDPAEGKIISAQKFQD	RVTMTDDTSTDTAYMELSSLRSEDSAVYYCAT
11E7H1	YISSSGSAIYYADSVKG	RFTISRDNAKNSLYLQLNSLRAEDTAVYYCAR
11E7H2	VIWYDGSNKYYADSVKG	RFTISRDNSKNTLHLQMNSLRAEDTAVYYCAR
2F12H	GFDPQDGETIYAQKFQG	RVTMTEDTSTDTAYMELRSLRSEDTAVYYCTT
14E4H	YISNSGSVVYYADSVKG	RFTISRHNAKNSLYLQMNSLRADDTAVYYCAR
3A5H	GFDPAEGKIISAQKFQD	RVTMTDDTSTDTAYMELSSLRSEDSAVYYCAT
10D7H	YISSTGSAMYDADSVKG	RFTISRDNAKNSLYLQMNSLRAEDTAVYYCAR
27D8H	YISSSGSATYYADSVKG	RFTISRDNAKNSLYLQMSSLRAEDTAVYYCAR
18A11H	GFDPQDGETIYAQKFQG	RVTMTEDTSTDTAYMELSSLKSEDTAVYYCAT
20D7H	YISSSGSAIYYADSVKG	RFTISRDNAKNSLYLQMDSLRAEDTAVFYCAR
23H6H	YISSSGSAMYSADSVKG	RFTISRDNAKNSLYLQMNSLRAEDTAVYYCAR
26G2H	YISSIGSAIHYADSVKG	RFTISRDNAKNSLYLQMNSLRAEDTAVYYCAR
27G8H	NIKQDGSEKYYVDSVKG	RFTISRDNAKNSLYLQMNSLRAEDTAVYYCAR
26C7H	YISRVGSTTYYADSVKG	RFTISRDNAKNSLYLQMNSLRAEDTAVYYCAR
26H3H	IIYPYDSDTRYSPSFQG	QVTISADKSINTAYLQWSSLKASDTAMFYCAS
19G6H	YISSSGSTMYYADSVKG	RFTISRVNAKNSLYLQMNSLRAEDTAVYYCAR
22B2H	IIDPNDSDTRYSPSFQG	QVTISADKSIHTAYLQWSSLKASDTAMYYCAT
24A2H	YISSSGSAIYYADSVKG	RFTISRDNPKNSLYLQMNSLRAEDTAVYYCAR
26E9H	YISSSGSTSYCADSVKG	RFTISRDNAKNSLYLQMNSLRAEDTAVYYCAR
19E6H	AISGSGGSTYYADSVKG	RFTISRDNSKNTLYLQMNSLRAEDTAVYYCAK
22F5H	YISSTGSTLYYADSVKG	RFTISRDNAKNSLYLQMDSLRADDAAVYYCTR
25C9H	YISSSGSAIHYADSVKG	RFTISRDNAKNSLYLQMNSLRAEDTAVYYCAR
26C10H	YISSSGSAIHYADSVKG	RFTISRDNAKNSLYLQMDSLRAEDTAVFYCAR
17C8H	YISNSGSAMYYADSVKG	RFTISRDNARNSLYLQMNSLRAEDTAVYYCAR
1A10H (a)	GFDPAEGKIISAQKFQD	RVTMTRDTSTSTVYMELSSLRSEDTAVYYCAR
27G8H (b)	NIKQDGSEKYYVDSVKG	RFTISRDNAKNSLYLQMNSLRAGDTAVYYCAR

Table 2 (cont'd)

Heavy	CDR3	FR4
1A10H	LDFSSWFDP	WGQGTLVTVSS
11E7H1	DYSSGWFYFDY	WGRGTLVTVSS
11E7H2	EHWNYAFDI	WGQGTMVTVSS
2F12H	ESSSAWFDP	WGQGTLVTVSS
14E4H	DRSSAWDEAFDI	WGQGTMVTVSS
3A5H	LDFSSWFDP	WGQGTLVTVSS
10D7H	EFSSGWSYFDY	WGQGTLVTVSS
27D8H	DYSSGWYYFDY	WGQGTLVTVSS
18A11H	GSSSSWFDP	WGQGTLVTVSS
20D7H	EHSSGYWYFDL	WGRGALVTVSS
23H6H	EYSSGWYYFDY	WGRGTLVTVSS
26G2H	EYSSGWAYFDY	WGQGTLVTVSS
27G8H	EGGYDWNYADYYGMDV	WGQGTTVTVSS
26C7H	DYSSGWYYFDY	WGQGTLVTVSS
26H3H	HRLWLGEFPGPLNI	WGQGTMVTVSS
19G6H	DRSSGLVSFDY	WGQGTLVTVSS
22B2H	HRLWLGTLPGGFYI	WGQGTMVTVSS
24A2H	DFSSGYYYFDY	WGHGTLVTVSS
26E9H	DYSSGWFYFDY	WGQGTLVTVSS
19E6H	APYSSSWALGLGMDV	WGQGTTVTVSS
22F5H	EYSSGWFFFDY	WGQGTLVTVSS
25C9H	EYSSGWAYFDY	WGQGTLVTVSS
26C10H	DHSSGYWYFDL	WGRGTLVTVSS
17C8H	EYSSGWFFFES	WGQGTLVTVSS
1A10H (a)	LDFSSWFDP	WGQGTLVTVSS
27G8H (b)	EGGYDWNYADYYGMDV	WGQGTTVTVSS

The amino acid sequences of the antibodies were further analyzed for similarities. The kappa light chains were grouped into three groups, and a consensus sequence was developed for each group. There were three antibodies with lambda light chains, none of which bore enough similarity to each other to form a group of related sequences from which a consensus sequence could be developed. Two of the variants developed varied in the lambda light chain. The heavy chains were grouped into four groups with a single heavy chain categorized into a fifth group, and a consensus sequence was developed for groups 1 through 4. These results are shown in Table 3(a) and 3(b) below; consensus sequences are shown in the Sequence Listing. The numbers in parentheses indicate the SEQ ID NO in the Sequence Listing.

Table 3(a): Grouping of antibodies by kappa light chain, with corresponding heavy chain

Kappa Group 1 (10 members)	Heavy chain group (H1 – H5)	Kappa Group 2 (9 members)	Heavy chain group (H1 – H5)	Kappa Group 3 (4 members)	Heavy chain group (H1 – H5)
20D7K (10)	H1 (38)	11E7K2 (3)	H1 (31)	22B2K (16)	H4 (45)
11E7K1 (2)	H1 (30)	10D7K(7)	H1 (35)	19G6K (15)	H1 (44)
26C10K (20)	H1 (51)	3A5K(6)	H2 (34)	26H3K (14)	H4 (43)
23H6K (11)	H1 (39)	1A10K (1)	H2 (29)	26H3K(c) (27)	H4 (43)
26C7K (13)	H1(42)	25C9K (22)	H1 (50)		
24A2K (17)	H1 (46)	26G2K (24)	H1 (40)		
27D8K (8)	H1 (36)	18A11K (9)	H2 (37)		

22F5 (19)	H1 (49)	2F12K (4)	H2 (32)		
26E9K (18)	H1 (47)	1A10K(d) (28)	H2 (53)		
17C8K (21)	H1 (52)				

Table 3(b): *Grouping of antibodies by lambda light chain, with corresponding heavy chain*

Lambda chains (3 antibodies, no consensus)	Heavy chain group (H1 – H5)
14E4 (5)	H1 (33)
27G8 (12)	H3 (41)
27G8(a) (25)	H3 (54)
27G8(b) (26)	H3 (54)
19E6 (23)	H5 (48)

CDR boundaries within the consensus sequences (which may vary, as discussed previously) were as follows: Kappa Group 1 CDR1 24-35, CDR2 51-57, CDR3 90-99; Kappa Group 2 CDR1 24-34, CDR2 51-56, CDR3 89-97; Kappa Group 3 CDR1 24-34, CDR2 50-56, CDR3 89-97; Heavy Chain Group 1 CDR1 31-35, CDR2 50-66, CDR3 99-110; Heavy Chain Group 2 CDR1 31-35, CDR2 50-66, CDR3 99-107; Heavy Chain Group 3 CDR1 31-35, CDR2 50-66, CDR3 99-114; and Heavy Chain Group 4 CDR1 31-35, CDR2 50-66, CDR3 99-114.

EXAMPLE 3: FUNCTIONAL ASSAYS

This example describes various assays that were used to characterize the antibodies.

HUT78 adhesion assay.

Coated plates (for example, Costar® 3368 96-well plates; Corning Incorporated Life Sciences, Lowell MA) are prepared by coating 96-well plates overnight at 4°C with 20 microG/mL MAdCAM-1 (or a similar concentration of human IgG1 as a coating control) diluted in phosphate buffer pH9.0. The coating is removed and the plates are blocked with 100microL of 3% BSA/PBS, incubated for 1hr or more at room temp. The plates are washed three times with Hank's balanced salt solution (HBSS).

HUT78 cells (a human T cell lymphoma cell line that exhibits the features of a mature T cell line with inducer/helper phenotype; ATCC TIB 161), grown to confluency, are pelleted and washed 3X in HBSS, then resuspended in HBSS at appropriate concentration to yield ~30,000 cells in 50microL.

Antibodies to be tested are diluted to twice the final concentration, and then titrated 1:4 in calcium-free, magnesium free HBSS containing 1%BSA with 1mM Mn^{2+}. Fifty microL of antibody titration or control is added to each well of a VEE bottom plate, followed by 50 microL of HUT78 cells. The cells and antibodies are incubated at 4°C for 30minutes, then added to the coated plates and incubated at 37°C for 40 minutes. Cells on coated plates are washed three times in room temperature HBSS, by flicking HBSS off between washes. The adherent cells are freeze-thawed at -20 C followed by the addition of 100 microL of CyQuant® dye/lysis buffer (a buffer used in a fluorescence-based cell quantification assays useful in high-throughput screening Molecular Probes®, Life Technologies Corporation, Carlsbad, CA. The fluorescent signal from each well is quantitated at 485 nm excitation and 530 nm emission, for example using a Tecan GENiosPro, a multi-label microplate reader (Tecan Group Ltd. Männedorf, Switzerland).

Human CD4+ cell adhesion assay

Plates are coated with human MAdCAM-1-Fc or human IgG (3 microG/ml in 20 mM phosphate buffer, pH 9.0, 130 mM NaCl), 100 microL/well, at 4 ^{0}C overnight then blocked with 200microL/well blocking reagent (3% bovine serum albumen in PBS) at room temperature for at least two hours. Plates are then washed three times with adhesion buffer (30mM HEPES, pH 7.4, 120mM NaCl, 1mM $MnCl_2$, 10 g/ml Human IgG).

Serial dilutions of antibodies to be tested are prepared, and added to the plate (35microL/well); isolated $CD4^+$ T cells (250,000 cells/35microL/well) are added and the plates are incubated at 4 ^{0}C for 2 hrs. After washing three times with adhesion buffer, plates are frozen at -20 ^{0}C overnight. Detection reagent (100 microL. well CyQUANT® reagent; Life Technologies Corporation, Carlsbad, CA) is added and plates are incubated at 37 ^{0}C for 45 minutes. Results are determined by reading fluorescence at 485nm excitation and 530nm emission.

EC50 in binding human CD4+CD45RA- memory T cells

Human peripheral blood mononuclear cells (PBMC; fresh or frozen and thawed, for example in phosphate buffered saline with 2%FBS) are washed and resuspended in HEPES buffer (30mM HEPES+140nM NaCl) with 1%BSA, with or without 1mM $MnCl_2$ (depending on the experiment; Mn^{2+} is necessary for MAdCAM-1 binding) and plated into 96 well plates (10^6 cells/well). Cells are incubated with 10 microG/ml human IgG for 30 minutes on ice to block nonspecific binding. Cells are then incubated with serial dilutions of biotinylated anti-alpha4beta7 antibodies in 96 well plates for one hour on ice, followed by addition of 1:100 dilution of streptavidin-phycoerythrin (PE; Jackson ImmunoResearch Laboratories Inc., West Grove, PA), 4 microL CD3-Pacific Blue, CD4-PerCP-Cy5.5 and CD45RA-fluorescein isothiocyanate (FITC) (BD Biosciences, San Jose CA) for a final volume of 100 microL, and incubated for another hour on ice. Cells were washed twice with HEPES buffer (with or without $MnCl_2$, correspondingly) and then fixed in 200microL HEPES buffer plus 0.5% paraformaldehyde (again, with or without $MnCl_2$, correspondingly). The percentage of positive alpha4beta7 antibody binding CD4+CD45RA- memory T cells is determined using a fluorescence activated cell sorter (FACS), for example, a BD™ LSR II benchtop flow cytometer (BD Biosciences, San Jose CA). EC50 is defined as the concentration of alpha4beta7 antibody at which 50% of the alpha4beta7 sites on CD4CD45RA- memory cells are bound by the alpha4beta7 antibody.

IC50 in blocking MAdCAM-1-Fc binding to human CD4+CD45RA- memory T cells.

PBMC (fresh or frozen as described previously) are washed and resuspended in HEPES buffer (30mM HEPES+140nM NaCl) with 1%BSA and 1mM MnCl. to a final concentration of 10^7 cells/ml. Cells are blocked as described previously; after blocking, cells are incubated with a serial dilution of anti-alpha4beta7 antibody (or appropriate control) in 96 well plates for 30 minutes on ice, and then with 0.3 microG/ml biotinylated MAdCAM-1-Fc protein for another one hour.

After two washes in HEPES buffer with 1mM MnCl, cells are treated with 1:100 dilution of streptavidin-PE, 4 microL CD3-Pacific Blue, CD4-PerCP-Cy5.5 and CD45RA-FITC as described previously, in a final volume of 100 microL. After one hour incubation on ice, cells are washed twice with HEPES buffer with 1mM MnCl and then fixed in 200 microL buffer plus 0.5% paraformaldehyde. The percentage of positive MAdCAM-1-Fc binding CD4+CD45RA- memory T

cells is determined by fluorescence activated cell sorter (FACS) analysis, as described previously. IC50 is defined as the concentration of alpha4beta7 antibody at which MAdCAM-1-Fc binding to alpha4beta7 on CD4CD45RA- memory cells is inhibited by 50%.

Alpha4beta7 induction by retinoic acid on activated T cells

Isolated human PBMC are activated by anti CD3 (plate bound, 5 microG/ml), human IL-2 (20 ng/ml) in the presence or absence of retinoic acid (1000 nM) for 7 days. The activated cells are washed twice with staining buffer (PBS plus 0.5%BSA and 1mM MnCl) and incubated with 100 microG/ml human Ig for 30 minutes to block non-specific binding. The cells are first incubated in a serial dilution of anti-alpha4beta7 antibodies for 30 minutes on ice, and then stained with 1 microG/ml biotinylated MAdCAM-1-Fc for another 30 minutes. After twice washing with staining buffer, cells are stained with Streptavidin-PE (1:1000) for 30 minutes. The cells are analyzed by fluorescence activated cell sorting, for example, with a FACSCalibur™ (BD Biosciences, San Jose CA). Cells prepared in this manner may be used for additional experiments such as competition assays.

Competition Assays

Alpha4beta7 antibodies were also examined for their ability to compete with other anti-alpha4beta7 and/or beta7 antibodies in binding to alpha4 beta7 expressing cells by fluorometric microvolume assay technology or FMAT, substantially as described by Fiscella, et al., *Nature Biotechnology* 21:302-307; 2003. Briefly, cells expressing high levels of alpha4beta7 are prepared, for example by transient co- transfection of cells with nucleic acids encoding alpha4 and nucleic acids expressing beta7. Stable cell lines are prepared in a similar manner, using cells and protocols appropriate for stable transfection. Transfected cells are screened, for example, by FACS, using antibodies to alpha4, antibodies to beta7, and/or ligand (i.e., MAdCAM-1, for example, a MAdCAM-1-Fc fusion protein). Cells may undergo several cycles of sorting and selection to yield clonal cell lines with reproducible, elevated levels of alpha4beta7 expression.

Binding to S250N Mutant

Antibodies were also evaluated for their ability to recognize the S250N point mutant in the beta7 chain, which is known to be critical for ACT-1 binding (J Immunol. 159:1497, 1997). 293 cells transiently co-expressing alpha4beta7 having the S250N mutation in the beta chain (ref) are prepared in a similar manner as that previously described for preparation of cells expressing high levels of alpha4beta7.

Briefly a total $1x10^6$ transfected cells for profile are collected using cell-dissociation solution and spun down at 1000 rpm for 5 min. The cells are then blocked with 0.5 ml blocking buffer (1% goat serum/PBS) for 30min to 1hr at 4°C with shaking. For MAdCAM-1-Fc staining, cells are incubated for 1hr at 4°C with shaking in Mn^{2+} Buffer (1mM $MnCl_2$ in 30mM HEPES + 1% goat serum). Cells are then spun down at 1000 rpm/5min, and 0.5ml of fresh blocking buffer with along with 10microG/ml of primary antibody is added, followed by incubation for 30min to 1hr at 4°C with shaking. After two washes with 4 ml of cold PBS (each wash), secondary antibody (i.e., goat-anti IgG-Phycoerythrin-conjugated antibody; Southern Biotech, 1:250 diluted or 0.1 microG /10^6 cells) in 0.5ml

blocking buffer is added and cells are incubated for 20-30min at 4°C. Cells are washed one final time with 4 mls of cold PBS, then resuspended in 0.5 ml of FACS buffer for profile.

Binding to Single Nucleotide Polymorphisms (SNPs)

For SNP analysis of the α4 subunit, α4 gene exons 1-28 from 90 individuals (180 haploid genomes) representing different ethnic groups were amplified by polymerase chain reaction (PCR) and subsequently sequenced. Three candidate SNPs in the coding region of the α4 gene were identified, and one of the three resulted in an amino acid change (Arg878Gln). Similarly for the β7 subunit SNP analysis, the coding region of β7 gene exons 2-15 from 90 individuals (180 haploid genomes) representing different ethnic groups were PCR amplified and subsequently sequenced. Three SNPs were identified, and two of them resulted in amino acid changes. The in-house SNP analysis data were compared with information in the NCBI database (NCBI: National Center for Biotechnology Information a division of the National Library of Medicine (NLM) at the National Institutes of Health (NIH)). Only the A/G mutation resulting in Gln878Arg in the α4 subunit occurs at high frequency - 20% or 30% in both in-house SNP and the public database respectively. The other SNPs occur at low frequency. This information is summarized in Table 4 below.

Table 4: Frequency of SNPs in human beta7 and alpha4

	SNP	Alternative allele frequency – NCBI database	Alternative allele frequency – in-house analysis	Location
beta7	E97V	NA	A(0.989)/T(0.019)	extracellular
	R213S	C(0.975)/A(0.25)	No observation	extracellular
	G611E	NA	No observation	extracellular
	G629S	NA	A(0.989)/T(0.019)	extracellular
	H672Y	NA	No observation	extracellular
alpha4	V824A	T(0.972)/C(0.028)	No observation	extracellular
	Q878R	A(0.648)/G(0.352)	A(0.783)/G(0.217)	extracellular
	R1007S	NA	No observation	intracellular

Point mutant constructs representing amino acid altering SNPs (a4b7(E97V); a4b7(R213S); a4b7(G629S; a4(V824A)b7; a4(Q878R)b7) in the extracellular domains of both alpha4 and beta7 were generated. Each point mutant construct was transfected to 293 cells along with the wild-type partner expression construct. Transfected 293 cells were first stained with 1 microG/ml of human IgG or anti-alpha4beta7 antibody, washed with PBS, and the stained with phycoerytherin conjugated secondary antibody goat-anti human IgG. After washing with PBS, cells were analyzed by fluorescence activated cell sorting, for example, with a FACSCalibur™ (BD Biosciences, San Jose CA). Fluorescence staining intensity (geometric mean) for each antibody staining is indicated in Table 5 below.

Table 5: Binding to SNPs

	wt	E97V	R213S	G629S	V824A	Q878R
IgG	8	7	7	7	7	7
1A10	122	135	31	80	102	70

3A5	124	135	31	80	110	71
2F12	129	134	37	80	112	73
18A11	92	105	30	63	82	56
22B2	97	108	58	65	85	58
26H3	93	106	49	62	82	55
27G8	102	116	59	68	88	58
26G2	99	113	38	64	86	58
17C8	93	96	49	58	74	51
19G6	94	108	46	59	67	46
25C9	96	106	33	59	77	51

These results indicated that all of the antibodies tested bound to the known SNPs of alpha4beta7.

The activities of various alpha4beta7heterodimer specific antibodies in several different assays are compared in Table 6 below.

Table 6: Characterization of antibodies to alpha4beta7

Antibody	HUT78 Adhesion IC50 (ng/ml)	MAdCAM-1 Competition IC50 (ng/mL)	CD4+CD45RA- Cell Binding EC50 (ng/mL)	a4b7(S250N) Binding
1A10	6.1	6.2	4.9	-
3A5	7.5	6.2	5.6	-
2F12	11.4	4.6	3.3	-
18A11	7.4	7.3	4.7	-
22B2	3.7	23.2	5.1	-
26H3	8.9	14.1	9.3	-
27G8	14.9	8.7	6.3	-
26G2	6.9	99.6	32.6	+
17C8	6.8	31.1	22.9	+
19G6	12.2	103.3	32.9	+
25C9	13.7	77.6	NA	+

Soler et al. reported the binding specificity of a humanized anti-alpha4beta7 antibody known as vedolizumab (J Pharmacol Exp Ther 330:864; 2009). This antibody was reported to have an EC50 on memory CD4+ T lymphocytes of 0.042 microgr/ml (42 ng/ml). Vedolizumab also inhibited the binding of soluble MAdCAM-1 to alpha4beta7hi memory T cells with an IC50 of 0.034 microgr/ml (34 ng/ml). In contrast, many of the antibodies shown in Table 6 have an EC50 on memory T cells (i.e., CD4+CD45RA- cells) of less than 10 ng/ml, and all of them have an EC50 of less than 35 ng/ml (all also have an EC50 of greater than 0.1 ng/ml in this assay). Additionally, several of the antibodies shown in Table 6 demonstrated an IC50 in a MAdCAM competition assay of less than 10 ng/ml, and many demonstrated an IC50 of less than 30 ng/ml (all also exhibited an IC50 of greater than 0.1 ng/ml in this assay). Although Soler et al. made no mention of the ability of vedolizumab to bind an S250N mutant of alpha4beta7, the murine antibody ACT-1 from which vedolizumab is derived is known to be unable to bind an S250N mutant (Tidswell et al., J Immunol 159:1497; 1997), and according to Soler et

al., vedolizumab and ACT-1 exhibit the same antigen specificity. Thus, vedolizumab also does not bind the S250N mutant, in contrast to several of the antibodies shown in Table 6.

EXAMPLE 4: ADDITIONAL ANALYSIS

Several representative antibodies with different properties in the afore-mentioned functional assays were chosen for additional analysis as described below.

Binding affinity to human a4b7

To measure cell binding affinity of human anti-alpha4beta7antibodies, a Kinetic Exclusion Assay which measures binding events in solution phase can be used to calculate the equilibrium dissociation constant, K_d. KinExA® Technology (Sapidyne Instruments, Boise, ID) was used, substantially as described previously by Xie et. al. *J. Imm, Methods* 304:1 (2005) and Rathanaswami et. al. *Anal. Biochem.* 373:52 (2008). Briefly, HUT78 cells expressing human alpha4beta7 were titrated 1 in 3 from ~50^6 cells/mL to ~400 cells/mL and then equilibrated with a final concentration of either 2 or 30 pM of mAb 2F12 or 18A11 and 30 or 500 pM for 17C8 for 18 hours at 4^0C. The free antibody remaining in the supernatant at equilibrium was measured by KinExA® technology by passing the supernatant over PMMA beads pre-coated with goat anti-human Fc and detected with goat anti-human (H+L) Cy5 (substantially as described by Rathanaswami et al. *Biochem Biophys Research Commun*:1004 (2005). The equilibrium dissociation constant (K_d) is obtained using KinExA® software by "n-curve analysis" which fits all of the given curves to a single K_d value simultaneously (Rathanswami et al. 2005 and Xie et al., *supra*).; results are shown below in Table 7.

Table 7: Binding affinity of antibodies

Antibody	Kd (pM)	Ratio (low Ab)	% error	Kd low	Kd high
2F12	4.56	0.44	3.80	1.94	11.12
18A11	0.90	2.22	4.40	0.23	2.29
17C8	29.36	1.02	3.58	12.09	74.53

These antibodies demonstrated a Kd in a KinExA® assay greater than 0.05pM, but less than 80 pM, less than 15 pM, or less than 5 pM.

PK/PD Characteristics

A single-dose pharmacokinetic (PK) and pharmacodynamic (PD) study of three fully human anti-alpha4beta7 antibodies in male cynomologous monkeys was conducted following intravenous (IV; 5 mg/kg) or subcutaneous (SC; 0.5 or 5 mg/kg) administration. Similar initial PK exposure (C_0; concentration at time zero) and distribution within the central circulation after 5 mg/kg IV was observed. After SC administration, both C_{max} (maximum concentration in serum) and AUC (area under the concentration-time curve) exhibited dose-proportionality within the 0.5-5 mg/kg SC dose range for all three antibodies. The absolute bioavailability after SC for the three tested antibodies ranged from 44 to 68%.

Free alpha4beta7 on T cells before and after antibody treatment were quantified by PE-conjugated anti-alpha4beta7 antibody 27G8. The background level was controlled by staining with PE-

conjugated anti-alpha4beta7 antibody 27G8 in the presence of 10mg/ml antibody being tested before and after antibody treatment. Fractional saturation was determined by percentage of free alpha4beta7 sites in comparison to the pre-treatment for each antibody. CD4-PerCP, CD99-APC and CD28-FITC were used to distinguish naive, central memory and effector memory cells. Fractional saturation of alpha4beta7 on naive T cells was shown because of the variability and limited assay sensitivity in the cynomologous memory cell population. For 18A11 treatment, all three dosage groups remained saturated from day 1 to day 14. At day 29, all three groups lost coverage. In both 2F12 and 17C8 treatment, loss of target coverage was observed on day 14 at the low dosage (0.5mg/Kg) group.

In addition to free alpha4beta7 detection, target saturation was also determined by staining with PE-conjugated anti-human antibody A35 in the absence or presence of 10mg/ml antibody. Total alpha4beta7 sites were estimated with pre-incubation of samples with 10mg/ml anti-alpha4beta7 antibody followed by staining with anti-human antibody. Target saturation was determined by the percentage of total alpha4beta7 sites occupied by the anti-alpha4beta7 antibodies for each sample. The three fully human anti-alpha4beta7 antibodies demonstrated saturation of alpha4beta7 that was maintained at mean of 81 to 100% within 14 days after 5 mg/kg IV.

PK/PD modeling was conducted on serum anti-alpha4beta7 antibody concentrations and corresponding alpha4beta7 receptor saturation data using a direct E_{max} model. The model estimated PD parameters are E_{max} (maximum alpha4beta7 receptor saturation) of 92%, EC_{50} (anti-alpha4beta7 antibody concentration at which 50% of the E_{max} was reached) of 52 ng/mL, and E_0 (initial alpha4beta7 receptor saturation) of 18%. All three antibodies exhibited potent in vivo PD effects on saturating alpha4beta7 receptors with average $t_{1/2}$ of ~3-5 days in cynomolgus monkeys.

CLAIMS :

1. An isolated, alpha4beta7 heterodimer specific antigen binding protein having a heavy chain variable region comprising CDR1, CDR2 and CDR3 from SEQ ID NO:37, and a light chain variable region comprising CDR1, CDR2 and CDR3 from SEQ ID NO:9.

2. The heterodimer specific antigen binding protein of claim 1, wherein the light chain variable region is at least 90% identical to SEQ ID NO:9 by containing substitutions in the framework regions set forth in SEQ ID NO:56, and the heavy chain variable region is at least 90% identical to SEQ ID NO:37 by containing substitutions in the framework regions set forth in SEQ ID NO:59.

3. The heterodimer specific antigen binding protein of claim 2, wherein the light chain variable region comprises SEQ ID NO:9, and the heavy chain variable region is selected from the group consisting of a polypeptide that comprises SEQ ID NO:37 and a polypeptide that comprises SEQ ID NO:37 wherein the N-terminal amino acid is converted to pyroglutamic acid.

4. The isolated, alpha4beta7 heterodimer specific antigen binding protein of claim 1, which further comprises a light chain constant region and a heavy chain constant region.

5. The isolated, alpha4beta7 heterodimer specific antigen binding protein of claim 4 wherein the light chain constant region is a kappa-type light chain constant region; and the heavy chain constant region is selected from the group consisting of:

a') a constant region from an IgD antibody;
b') a constant region from an IgE antibody;
c') a constant region from an IgM antibody;
d') a constant region from an IgG1 antibody;
e') a constant region from an IgG2 antibody;
f') a constant region from an IgG3 antibody;
g') a constant region from an IgG4 antibody; and
h') a constant region from an IgG4 antibody having at least one mutation in a hinge region that alleviates a tendency to form intra-H chain disulfide bond.

6. The isolated, alpha4beta7 heterodimer specific antigen binding protein of claim 4 wherein the light chain constant region is selected from the group consisting of:

a) a polypeptide comprising SEQ ID NO:70;

b) a polypeptide at least 90% identical to SEQ ID NO:70;

c) a polypeptide of a) which incorporates a post-translational modification selected from the group consisting of conversion of glutamine to pyroglutamic acid, deamidation of an amino acid, isomerization of an amino acid, glycation of an amino acid, oxidation of an amino acid, N-linked glycosylation, and O-linked glycosylation;

d) a polypeptide having an amino acid sequence as set forth in SEQ ID NO:70 from which one, two, three, four or five N-terminal amino acids have been removed;

e) a polypeptide having an amino acid sequence as set forth in SEQ ID NO:70 from which one, two, three, four or five C-terminal amino acids have been removed; and

f) combinations of the above; and the heavy chain constant region is selected from the group consisting of:

a') a polypeptide comprising SEQ ID NO:72;

b') a polypeptide at least 90% identical to SEQ ID NO:72;

c') a polypeptide of a') which incorporates a post-translational modification selected from the group consisting of conversion of glutamine to pyroglutamic acid, deamidation of an amino acid, isomerization of an amino acid, glycation of an amino acid, oxidation of an amino acid, N-linked glycosylation, and O-linked glycosylation;

d') a polypeptide having an amino acid sequence as set forth in SEQ ID NO:72 from which one, two, three, four or five N-terminal amino acids have been removed;

e') a polypeptide having an amino acid sequence as set forth in SEQ ID NO:72 from which one, two, three, four or five C-terminal amino acids have been removed; and

f')combinations of the above.

7. The isolated, alpha4beta7 heterodimer specific antigen binding protein of claim 2 which further comprises a light chain constant region and a heavy chain constant region.

8. The isolated, alpha4beta7 heterodimer specific antigen binding protein of claim 7 wherein the light chain constant region is a kappa-type light chain constant region; and the heavy chain constant region is selected from the group consisting of:

a') a constant region from an IgD antibody;

b') a constant region from an IgE antibody;

c') a constant region from an IgM antibody;

d') a constant region from an IgG1 antibody;

e') a constant region from an IgG2 antibody;

f') a constant region from an IgG3 antibody;

g') a constant region from an IgG4 antibody; and

h') a constant region from an IgG4 antibody having at least one mutation in a hinge region that alleviates a tendency to form intra-H chain disulfide bond.

9. The isolated, alpha4beta7 heterodimer specific antigen binding protein of claim 7 wherein the light chain constant region is selected from the group consisting of:

a) a polypeptide comprising SEQ ID NO:70;

b) a polypeptide at least 90% identical to SEQ ID NO:70;

c) a polypeptide of a) which incorporates a post-translational modification selected from the group consisting of conversion of glutamine to pyroglutamic acid, deamidation of an amino acid, isomerization of an amino acid, glycation of an amino acid, oxidation of an amino acid, N-linked glycosylation, and O-linked glycosylation; and

d) a polypeptide having an amino acid sequence as set forth in SEQ ID NO:70 from which one, two, three, four or five N-terminal amino acids have been removed;

e) a polypeptide having an amino acid sequence as set forth in SEQ ID NO:70 from which one, two, three, four or five C-terminal amino acids have been removed; and

f) combinations of the above; and the heavy chain constant region is selected from the group consisting of:

a') a polypeptide comprising SEQ ID NO:72;

b') a polypeptide at least 90% identical to SEQ ID NO:72;

c') a polypeptide of a') which incorporates a post-translational modification selected from the group consisting of conversion of glutamine to pyroglutamic acid, deamidation of an amino acid, isomerization of an amino acid, glycation of an amino acid, oxidation of an amino acid, N-linked glycosylation, and O-linked glycosylation; and

d') a polypeptide having an amino acid sequence as set forth in SEQ ID NO:72 from which one, two, three, four or five N-terminal amino acids have been removed;

e') a polypeptide having an amino acid sequence as set forth in SEQ ID NO:72 from which one, two, three, four or five C-terminal amino acids have been removed; and

f')combinations of the above.

10. The isolated, alpha4beta7 heterodimer specific antigen binding protein of claim 3, which further comprises a light chain constant region and a heavy chain constant region.

11. The isolated, alpha4beta7 heterodimer specific antigen binding protein of claim 10 wherein the light chain constant region is a kappa-type light chain constant region; and the heavy chain constant region is selected from the group consisting

of:

a') a constant region from an IgD antibody;
b') a constant region from an IgE antibody;
c') a constant region from an IgM antibody;
d') a constant region from an IgG1 antibody;
e') a constant region from an IgG2 antibody;
f') a constant region from an IgG3 antibody;
g') a constant region from an IgG4 antibody; and
h') a constant region from an IgG4 antibody having at least one mutation in a hinge region that alleviates a tendency to form intra-H chain disulfide bond.

12. The isolated, alpha4beta7 heterodimer specific antigen binding protein of claim 10 wherein the light chain constant region is selected from the group consisting of:

a) a polypeptide comprising SEQ ID NO:70;
b) a polypeptide at least 90% identical to SEQ ID NO:70;
c) a polypeptide of a) which incorporates a post-translational modification selected from the group consisting of conversion of glutamine to pyroglutamic acid, deamidation of an amino acid, isomerization of an amino acid, glycation of an amino acid, oxidation of an amino acid, N-linked glycosylation, and O-linked glycosylation;
d) a polypeptide having an amino acid sequence as set forth in SEQ ID NO:70 from which one, two, three, four or five N-terminal amino acids have been removed;
e) a polypeptide having an amino acid sequence as set forth in SEQ ID NO:70 from which one, two, three, four or five C-terminal amino acids have been removed; and
f) combinations of the above; and the heavy chain constant region is selected from the group consisting of:

a') a polypeptide comprising SEQ ID NO:72;
b') a polypeptide at least 90% identical to SEQ ID NO:72;
c') a polypeptide of a') which incorporates a post-translational modification selected from the group consisting of conversion of glutamine to pyroglutamic acid, deamidation of an amino acid, isomerization of an amino acid, glycation of an amino acid, oxidation of an amino acid, N-linked glycosylation, and O-linked glycosylation; and
d') a polypeptide having an amino acid sequence as set forth in SEQ NO:72 from which one, two, three, four or five N-terminal amino acids have been removed;
e') a polypeptide having an amino acid sequence as set forth in SEQ ID NO:72 from which one, two, three, four or five C-terminal amino acids have been removed; and
f')combinations of the above.

13. An isolated, alpha4beta7 heterodimer specific antigen binding protein having a heavy chain variable region and a light chain variable region, wherein the heavy chain variable region is encoded by a nucleic acid that encodes SEQ ID NO:37, and the light chain variable region is encoded by a nucleic acid that encodes SEQ ID NO:9.

14. The isolated, alpha4beta7 heterodimer specific antigen binding protein of claim 13 wherein the nucleic acid that encodes the heavy chain variable region comprises SEQ ID NO:64, and the nucleic acid that encodes the light chain variable region comprises SEQ ID NO:63.

15. The isolated, alpha4beta7 heterodimer specific antigen binding protein of claim 13, which further comprises a light chain constant region and a heavy china constant region, wherein the light chain constant region is encoded by a nucleic acid that encodes SEQ ID NO:70 and the heavy chain constant region is encoded by a nucleic acid that encodes SEQ ID NO:72.

16. The isolated, alpha4beta7 heterodimer specific antigen binding protein of claim 15 wherein the light chain constant region is encoded by a nucleic acid that comprises SEQ ID NO:69, and the heavy chain constant region is encoded by a nucleic acid that comprises SEQ ID NO:71.

17. The isolated, alpha4beta7 heterodimer specific antigen binding protein of claim 14, which further comprises a light chain constant region and a heavy china constant region, wherein the light chain constant region is encoded by a nucleic acid that encodes SEQ ID NO:70 and the heavy chain constant region is encoded by a nucleic acid that encodes SEQ ID NO:72.

18. The isolated, alpha4beta7 heterodimer specific antigen binding protein of claim 17 wherein the light chain constant region is encoded by a nucleic acid that comprises SEQ NO:69, and the heavy chain constant region is encoded by a nucleic acid that comprises SEQ ID NO:71.

19. A composition for treating inflammatory bowel disease, pancreatitis, insulin-dependent diabetes mellitus, mastitis, cholecystitis, cholangitis, pericholangitis, chronic bronchitis, chronic sinusitis, asthma, or graft versus host disease, comprising the protein of claim 1 and a physiologically acceptable diluent, excipient or carrier.

20. A composition for treating inflammatory bowel disease, pancreatitis, insulin-dependent diabetes mellitus, mastitis, cholecystitis, cholangitis, pericholangitis, chronic bronchitis, chronic sinusitis, asthma, or graft versus host disease, comprising the protein of claim 2 and a physiologically acceptable diluent, excipient or carrier.

21. A composition for treating inflammatory bowel disease, pancreatitis, insulin-dependent diabetes mellitus, mastitis, cholecystitis, cholangitis, pericholangitis, chronic bronchitis, chronic sinusitis, asthma, or graft versus host disease, comprising the protein of claim 3 and a physiologically acceptable diluent, excipient or carrier.

22. A composition for treating inflammatory bowel disease, pancreatitis, insulin-dependent diabetes mellitus, mastitis, cholecystitis, cholangitis, pericholangitis, chronic bronchitis, chronic sinusitis, asthma, or graft versus host disease, comprising the protein of claim 4 and a physiologically acceptable diluent, excipient or carrier.

23. A composition for treating inflammatory bowel disease, pancreatitis, insulin-dependent diabetes mellitus, mastitis, cholecystitis, cholangitis, pericholangitis, chronic bronchitis, chronic sinusitis, asthma, or graft versus host disease, comprising protein of claim 5 and a physiologically acceptable diluent, excipient or carrier.

24. A composition for treating inflammatory bowel disease, pancreatitis, insulin-dependent diabetes mellitus, mastitis, cholecystitis, cholangitis, pericholangitis, chronic bronchitis, chronic sinusitis, asthma, or graft versus host disease, comprising the protein of claim 6 and a physiologically acceptable diluent, excipient or carrier.

25. A composition for treating inflammatory bowel disease, pancreatitis, insulin-dependent diabetes mellitus, mastitis, cholecystitis, cholangitis, pericholangitis, chronic bronchitis, chronic sinusitis, asthma, or graft versus host disease, comprising the protein of claim 7 and a physiologically acceptable diluent, excipient or carrier.

26. A composition for treating inflammatory bowel disease, pancreatitis, insulin-dependent diabetes mellitus, mastitis, cholecystitis, cholangitis, pericholangitis, chronic bronchitis, chronic sinusitis, asthma, or graft versus host disease, comprising the protein of claim 8 and a physiologically acceptable diluent, excipient or carrier.

27. A composition for treating inflammatory bowel disease, pancreatitis, insulin-dependent diabetes mellitus, mastitis, cholecystitis, cholangitis, pericholangitis, chronic bronchitis, chronic sinusitis, asthma, or graft versus host disease, comprising the protein of Claim 9 and a physiologically acceptable diluent, excipient or carrier.

28. A composition for treating inflammatory bowel disease, pancreatitis, insulin-dependent diabetes mellitus, mastitis, cholecystitis, cholangitis, pericholangitis, chronic bronchitis, chronic sinusitis, asthma, or graft versus host disease, comprising the protein of claim 10 and a physiologically acceptable diluent, excipient or carrier.

29. A composition for treating inflammatory bowel disease, pancreatitis, insulin-dependent diabetes mellitus, mastitis, cholecystitis, cholangitis, pericholangitis,

chronic bronchitis, chronic sinusitis, asthma, or graft versus host disease, comprising the protein of claim 11 and a physiologically acceptable diluent, excipient or carrier.

30. A composition for treating inflammatory bowel disease, pancreatitis, insulin-dependent diabetes mellitus, mastitis, cholecystitis, cholangitis, pericholangitis, chronic bronchitis, chronic sinusitis, asthma, or graft versus host disease, comprising the protein of claim 12 and a physiologically acceptable diluent, excipient or carrier.

31. A composition for treating inflammatory bowel disease, pancreatitis, insulin-dependent diabetes mellitus, mastitis, cholecystitis, cholangitis, pericholangitis, chronic bronchitis, chronic sinusitis, asthma, or graft versus host disease, comprising the protein of claim 13 and a physiologically acceptable diluent, excipient or carrier.

32. A composition for treating inflammatory bowel disease, pancreatitis, insulin-dependent diabetes mellitus, mastitis, cholecystitis, cholangitis, pericholangitis, chronic bronchitis, chronic sinusitis, asthma, or graft versus host disease, comprising the protein of claim 14 and a physiologically acceptable diluent, excipient or carrier.

33. A composition for treating inflammatory bowel disease, pancreatitis, insulin-dependent diabetes mellitus, mastitis, cholecystitis, cholangitis, pericholangitis, chronic bronchitis, chronic sinusitis, asthma, or graft versus host disease, comprising the protein of claim 15 and a physiologically acceptable diluent, excipient or carrier.

34. A composition for treating inflammatory bowel disease, pancreatitis, insulin-dependent diabetes mellitus, mastitis, cholecystitis, cholangitis, pericholangitis, chronic bronchitis, chronic sinusitis, asthma, or graft versus host disease, comprising the protein of claim 16 and a physiologically acceptable diluent, excipient or carrier.

35. A composition for treating inflammatory bowel disease, pancreatitis, insulin-dependent diabetes mellitus, mastitis, cholecystitis, cholangitis, pericholangitis, chronic bronchitis, chronic sinusitis, asthma, or graft versus host disease, comprising the protein of claim 17 and a physiologically acceptable diluent, excipient or carrier.

36. A composition for treating inflammatory bowel disease, pancreatitis, insulin-dependent diabetes mellitus, mastitis, cholecystitis, cholangitis, pericholangitis, chronic bronchitis, chronic sinusitis, asthma, or graft versus host disease, comprising the protein of claim 18 and a physiologically acceptable diluent, excipient or carrier.

37. An isolated nucleic acid that encodes the antigen binding protein of any one of Claims 1 through 18.

38. A vector comprising the nucleic acid of Claim 37.

39. An isolated host cell transfected or transformed with the vector of Claim 38.

40. A method for the production of an antigen binding protein comprising culturing the host cell of Claim 39 to express the protein and recovering the protein from the culture medium.