TEAR LIPOCALIN MUTEINS BINDING IL-4 R ALPHA

Cross-reference to related applications

[0001] The present application claims priority to US provisional application.61/61/352,461 filed with the USPTO on June 8, 2010, the entire contents of which is incorporated herein by reference in its entirety.

Field of the invention

[0002] The present invention relates to muteins of human tear lipocalin that bind to IL-4 receptor alpha. The invention also relates to corresponding nucleic acid molecules encoding such a mutein and to a method for their generation. The invention further relates a method for producing such a mutein. Finally, the invention is directed to a pharmaceutical composition that includes such a lipocalin mutein as well as to various uses of the mutein.

Background

[0003] Proteins that selectively bind to selected targets by way of non-covalent interaction play a crucial role as reagents in biotechnology, medicine, bioanalytics as well as in the biological and life sciences in general. Antibodies, i.e. immunoglobulins, are a prominent example of this class of proteins. Despite the manifold needs for such proteins in conjunction with recognition, binding and/or separation of ligands/targets, almost exclusively immunoglobulins are currently used. The application of other proteins with defined ligand-binding characteristics, for example the lectins, has remained restricted to special cases.

[0004] Additional proteinaceous binding molecules that have antibody-like functions are the members of the lipocalin family, which have naturally evolved to bind ligands. Lipocalins occur in many organisms, including vertebrates, insects, plants and bacteria. The members of the lipocalin protein family (Pervaiz, S., & Brew, K. (1987) *FASEB J.* **1**,

209-214) are typically small, secreted proteins and have a single polypeptide chain. They are characterized by a range of different molecular-recognition properties: their ability to bind various, principally hydrophobic molecules (such as retinoids, fatty acids, cholesterols, prostaglandins, biliverdins, pheromones, tastants, and odorants), their binding to specific cell-surface receptors and their formation of macromolecular complexes. Although they have, in the past, been classified primarily as transport proteins, it is now clear that the lipocalins fulfill a variety of physiological functions. These include roles in retinol transport, olfaction, pheromone signaling, and the synthesis of prostaglandins. The lipocalins have also been implicated in the regulation of the immune response and the mediation of cell homoeostasis (reviewed, for example, in Flower, D.R. (1996) *Biochem. J.* **318**, 1-14 and Flower, D.R. et al. (2000) *Biochim. Biophys. Acta* **1482**, 9-24).

[0005] The lipocalins share unusually low levels of overall sequence conservation, often with sequence identities of less than 20%. In strong contrast, their overall folding pattern is highly conserved. The central part of the lipocalin structure consists of a single eight-stranded anti-parallel β -sheet closed back on itself to form a continuously hydrogen-bonded β-barrel. This β-barrel forms a central cavity. One end of the barrel is sterically blocked by the N-terminal peptide segment that runs across its bottom as well as three peptide loops connecting the β-strands. The other end of the β-barrel is open to the solvent and encompasses a target-binding site, which is formed by four flexible peptide loops. It is this diversity of the loops in the otherwise rigid lipocalin scaffold that gives rise to a variety of different binding modes each capable of accommodating targets of different size, shape, and chemical character (reviewed, e.g., in Flower, D.R. (1996), supra; Flower, D.R. et al. (2000), supra, or Skerra, A. (2000) *Biochim. Biophys. Acta* **1482**, 337-350).

[0006] International patent application WO 99/16873 discloses polypeptides of the lipocalin family with mutated amino acid positions in the region of the four peptide loops, which are arranged at the end of the cylindrical β-barrel structure encompassing the binding pocket, and which correspond to those segments in the linear polypeptide sequence that includes the amino acid positions 28 to 45, 58 to 69, 86 to 99, and 114 to 129 of the bilin-binding protein of *Pieris brassicae*. Members of the lipocalin familiy have been reported to be post-translationally modified, e.g. phosphorylation and glycosylation of tear lipocalin (e.g. You, J., et al. (2010) *Electrophoresis* **31**, 1853-1861). Nevertheless no post-translational modification is required for their molecular recognition properties.

[0007] International patent application WO 00/75308 discloses muteins of the bilin-

binding protein, which specifically bind digoxigenin, whereas the international patent applications WO 03/029463 and WO 03/029471 relate to muteins of the human neutrophil gelatinase-associated lipocalin (hNGAL) and apolipoprotein D, respectively. In order to further improve and fine tune ligand affinity, specificity as well as folding stability of a lipocalin variant various approaches using different members of the lipocalin family have been proposed (Skerra, A. (2001) *Rev. Mol. Biotechnol.* **74**, 257-275; Schlehuber, S., and Skerra, A. (2002) *Biophys. Chem.* **96**, 213-228), such as the replacement of additional amino acid residues. The PCT publication WO 2006/56464 discloses muteins of human neutrophile gelatinase- associated lipocalin with binding affinity for CTLA-4 in the low nanomolar range.

[0008] International patent application WO 2005/19256 discloses muteins of tear lipocalin with at least one binding site for different or the same target ligand and provides a method for the generation of such muteins of human tear lipocalin. According to this PCT application, certain amino acid stretches within the primary sequence of tear lipocalin, in particular the loop regions that include amino acids 7-14, 24-36, 41-49, 53-66, 69-77, 79-84, 87-98, and 103-110 of mature human tear lipocalin, are subjected to mutagenesis in order to generate muteins with binding affinities. The resulting muteins have binding affinities for the selected ligand (K_D) in the nanomolar range, in most cases >100 nM. International patent application WO 2008/015239 discloses muteins of tear lipocalin binding a given non-natural ligand, including the IL-4 receptor alpha. Binding affinities are in the nanomolar range, reaching as low as almost 1×10^{-10} M in surface plamon resonance experiments.

[0009] Human tear lipocalin (TLPC or Tlc), also termed lipocalin-1, tear pre-albumin or von Ebner gland protein, was originally described as a major protein of human tear fluid (approximately one third of the total protein content) but has also been identified in several other secretory tissues including prostate, adrenal gland, thymus, mammary gland, testis, nasal mucosa and tracheal mucosa as well as corticotrophs of the pituitary gland. Homologous proteins have been found in rhesus monkey, chimpanzee, rat, mouse, pig, hamster, cow, dog and horse. Tear lipocalin is an unusual lipocalin member in that it exhibits an unusually broad ligand specificity, when compared to other lipocalins, and in its high promiscuity for relative insoluble lipids (see Redl, B. (2000) *Biochim. Biophys. Acta* **1482**, 241-248). This feature of tear lipocalin has been attributed to the protein's function in inhibiting bacterial and fungal growth at the cornea. A remarkable number of lipophilic

compounds of different chemical classes such as fatty acids, fatty alcohols, phospholipids, glycolipids and cholesterol are endogenous ligands of this protein. Interestingly, in contrast to other lipocalins the strength of ligand (target) binding correlates with the length of the hydrocarbon tail both for alkyl amides and fatty acids. Thus, tear lipocalin binds most strongly the least soluble lipids (Glasgow, B.J. et al. (1995) *Curr. Eye Res.* **14**, 363-372; Gasymov, O.K. et al. (1999) *Biochim. Biophys. Acta* **1433**, 307-320). The 1.8-Å crystal structure of tear lipocalin revealed an unusually large cavity inside its β-barrel (Breustedt, D.A. et al. (2005) *J. Biol. Chem.* **280**, 1, 484-493).

[0010] Despite this progress it would be still desirable to have a human tear lipocalin mutein that has improved binding properties for IL-4 receptor alpha, in particular of higher binding affinity, simply for the reason to further improve the suitability of muteins of human tear lipocalin in diagnostic and therapeutic applications.

Summary of the invention

[0011] Accordingly, it is an object of the invention to provide a further human tear lipocalin mutein with high binding affinity for IL 4 receptor alpha.

[0012] This object is accomplished by a human tear lipocalin mutein with the features set out in the claims, in particular in claim 1.

[0013] In a first aspect the present invention provides a mutein of human tear lipocalin. The mutein binds to IL 4 receptor alpha. The mutein includes a mutated amino acid residue at any one or more of the sequence positions 27, 28, 30, 31, 33, 53, 57, 61, 64, 66, 80, 83, 104-106 and 108 of the linear polypeptide sequence of the mature human tear lipocalin. The mutein further includes a mutated amino acid residue at any two or more of the sequence positions 26, 32, 34, 55, 56, 58 and 63 of the linear polypeptide sequence of the mature human tear lipocalin. The amino acid sequence of the mutein of human tear lipocalin includes one of the following sets of amino acid combinations: (1) Ser 26, Glu 34, Leu 55, Lys 58, (2) Ser 26, Asn 34, Ala 55, Lys 58, (3) Ser 26, Val 34, (4) Pro 26, Ser 34 (5) Pro 26, Ala 55, (6) Leu 26, Trp 34, Ala 55, (7) Leu 26, Trp 34, Ile 58, (8) Asn 26, Asp 34, (9) Asn 26, Ala 55, (10) Tyr 26, His 34, Ala 55, (11) Tyr 26, His 34, Ala 58, (12) Lys 26, Arg 34, Ala 55, (13) Lys 26, Arg 34, Asn 58, (14) Glu 26, Gly 34, Ala 55 and (15) Glu 26, Gly 34, Leu 58.

[0014] The term "position" when used in accordance with the invention means the position of either an amino acid within an amino acid sequence depicted herein or the

position of a nucleotide within a nucleic acid sequence depicted herein. The term "corresponding" as used herein also includes that a position is not only determined by the number of the preceding nucleotides/amino acids. Accordingly, the position of a given amino acid in accordance with the invention which may be substituted may very due to deletion or addition of amino acids elsewhere in a (mutant or wild-type) lipocalin. Similarly, the position of a given nucleotide in accordance with the present invention which may be substituted may vary due to deletions or additional nucleotides elsewhere in a mutein or wild type lipocalin 5'-untranslated region (UTR) including the promoter and/or any other regulatory sequences or gene (including exons and introns).

[0015] Thus, under a "corresponding position" in accordance with the invention it is preferably to be understood that nucleotides/amino acids may differ in the indicated number but may still have similar neighboring nucleotides/amino acids. Said nucleotides/amino acids which may be exchanged, deleted or added are also comprised by the term "corresponding position".

[0016] Specifically, in order to determine whether a nucleotide residue or amino acid residue of the amino acid sequence of a lipocalin different from a Tlc lipocalin mutein of the invention corresponds to a certain position in the nucleotide sequence or the amino acid sequence of a Tlc lipocalin mutein as described, in particular any of SEQ ID NOs: 2-11or that having one or more amino acid substitutions at position 27, 28, 30, 31, 33, 53, 57, 61, 64, 66, 80, 83, 104-106 and 108 of the linear polypeptide sequence of Tlc (SEQ ID NO: 20), a skilled artisan can use means and methods well-known in the art, e.g., alignments, either manually or by using computer programs such as BLAST2.0, which stands for Basic Local Alignment Search Tool or ClustalW or any other suitable program which is suitable to generate sequence alignments. Accordingly, a lipocalin mutein of any of SEQ ID Nos: 2-11 or that having one or more amino acid substitutions at position 27, 28, 30, 31, 33, 53, 57, 61, 64, 66, 80, 83, 104-106 and 108 of the linear polypeptide sequence of Tlc (SEQ ID NO: 20) can serve as "subject sequence", while the amino acid sequence of a lipocalin different from Tlc serves as "query sequence".

[0017] In a second aspect the present invention provides a method of generating a mutein of human tear lipocalin. The mutein binds to IL-4 receptor alpha. The method includes subjecting a nucleic acid molecule encoding a human tear lipocalin to mutagenesis at any one or more of the amino acid sequence positions 27, 28, 30, 31, 33, 53, 57, 61, 64, 66, 80, 83, 104-106 and 108 of the linear polypeptide sequence of mature

human tear lipocalin. Further, the method includes subjecting the nucleic acid molecule encoding a human tear lipocalin to mutagenesis at any two or more of the amino acid sequence positions 26, 32, 34, 55, 56, 58 and 63 of the linear polypeptide sequence of the mature human tear lipocalin. At least one of the two or more of the amino acid sequence positions 26, 32, 34, 55, 56, 58 and 63 of the linear polypeptide sequence of the mature human tear lipocalin. As a result one or more nucleic acids encoding a mutein of human tear lipocalin are obtained. The amino acid sequence of the encoded mutein includes one of the following sets of amino acid combinations: (1) Ser 26, Glu 34, Leu 55, Lys 58, (2) Ser 26, Asn 34, Ala 55, Lys 58, (3) Ser 26, Val 34, (4) Pro 26, Ser 34, (5) Pro 26, Ala 55, (6) Leu 26, Trp 34, Ala 55, (7) Leu 26, Trp 34, Ile 58, (8) Asn 26, Asp 34, (9) Asn 26, Ala 55, (10) Tyr 26, His 34, Ala 55, (11) Tyr 26, His 34, Ala 58, (12) Lys 26, Arg 34, Ala 55, (13) Lys 26, Arg 34, Asn 58, (14) Glu 26, Gly 34, Ala 55, and (15) Glu 26, Gly 34, Leu 58. The method also includes expressing the one or more mutein encoding nucleic acid molecules thus obtained in an expression system. The method thereby includes obtaining one or more muteins. Further the method includes enriching the one or more muteins thus obtained that bind to IL 4 receptor alpha by means of selection and/or isolation.

[0018] In a third aspect the present invention provides a nucleic acid molecule. The nucleic acid molecule includes a nucleotide sequence that encodes a mutein according to the first aspect.

[0019] In a fourth aspect the present invention provides a host cell. The host cell contains a nucleic acid molecule according to the third aspect.

[0020] In a fifth aspect the present invention provides a pharmaceutical composition. The pharmaceutical composition includes a mutein of human tear lipocalin according to the first aspect. The pharmaceutical composition further includes a pharmaceutically acceptable excipient.

[0021] The invention will be better understood with reference to the detailed description when considered in conjunction with the non-limiting examples and the accompanying drawings.

Description of the Figures

[0022] **Figure 1** shows the polypeptide sequence of S191.4-B24, a mutein of human tear lipocalin possessing binding affinity for the IL-4 receptor alpha.

[0023] **Figure 2** shows the polypeptide sequences of exemplary muteins with high

affinity for IL-4 receptor alpha (SEQ ID NOs: 2-11).

[0024] **Figure 3** shows the inhibition of TF-1 cell proliferation by increased amounts of muteins of the invention in the presence of IL-4 (**A**) and IL-13 (**B**).

[0025] **Figure 4** depicts IC_{50} values of Fig. 3 and data of Biacore measurements of the binding of a human tear lipocalin muteins of the invention to IL-4 receptor alpha, such as human IL-4 receptor alpha.

Detailed description

[0026] The present invention provides muteins of human tear lipocalin that have a particularly high affinity to IL-4 receptor alpha. IL-4 receptor alpha as a target of a mutein of the present invention is typically a mammalian protein, such as a human protein. In vivo IL-4 receptor alpha can bind interleukin 4 and interleukin 13 to regulate IgE antibody production in B cells.

[0027] Binding affinities of muteins according to the invention have been found to be typically of a K_D below 0.1 nM and in some embodiments about 1 picomolar (pM) (cf. Fig. 4). The lipocalin muteins of the invention are accordingly able to bind IL-4 receptor alpha with detectable affinity, i.e. with a dissociation constant of at least 200 nM. In some embodiments a lipocalin mutein of the invention binds IL-4 receptor alpha with a dissociation constant for IL-4 receptor alpha of at least about 10 nM, about 1 nM, about 0.1 nM, about 10 pM or even less. The binding affinity of a mutein to a selected target, in the present case IL-4 receptor alpha, can be measured and thereby K_D values of a mutein-ligand complex be determine by a multitude of methods known to those skilled in the art. Such methods include, but are not limited to, fluorescence titration, competition ELISA, calorimetric methods, such as isothermal titration calorimetry (ITC), and surface plasmon resonance (BIAcore). Examples for such methods are detailed below (See e.g. Example 2).

[0028] Human interleukin-4 receptor alpha chain may have the amino acid sequence of SWISS PROT Data Bank Accession No. P24394 (SEQ ID NO: 18) or of fragments thereof. An illustrative example of a fragment of human interleukin-4 receptor alpha chain includes amino acids 26 to 232 of IL-4 receptor alpha. The amino acid sequence of the human IL-13 receptor alpha 1 is shown in SEQ ID NO: 19.

[0029] In general, the term “fragment”, as used herein with respect to protein ligands of the tear lipocalin muteins of the invention, relates to N-terminally and/or C-terminally

shortened protein or peptide ligands, which retain the capability of the full length ligand to be recognized and/or bound by a mutein according to the invention.

[0030] A human tear lipocalin mutein binding IL-4 receptor alpha may act as an IL-4 antagonist and/or IL-13 antagonist or as an inverse IL-4 agonist and/or inverse IL-13 agonist. An inverse agonist binds to the same binding-site as an agonist for a particular receptor and reverses constitutive activity of the respective receptor. So far IL-4 receptors have not been reported to possess intrinsic kinase activity, so that a mutein of the invention may typically act as an IL-4 antagonist and/or IL-13 antagonist. In one embodiment, the human tear lipocalin muteins act as antagonists of human IL-4 and/or human IL-13. In some embodiments the mutein is cross-reactive with the cynomolgus IL-4 IL-4 receptor alpha and as such acts as an antagonist of cynomolgus ligands such as IL-4 and/or IL-13. In some embodiments the mutein is cross-reactive with the marmoset IL-4 IL-4 receptor alpha and as such acts as an antagonist of marmoset ligands such as IL-4 and/or IL-13.

[0031] IL-4 receptor alpha can be taken to define a non-natural ligand of human tear lipocalin. The term “non-natural ligand” refers to a compound, which does not bind to native mature human tear lipocalin under physiological conditions. The term "human tear lipocalin" as used herein refers to the mature human tear lipocalin corresponding to the protein of the SWISS-PROT Data Bank Accession Number P31025. Mature human tear lipocalin does not include the N-terminal signal peptide that is included in the sequence of SWISS-PROT Accession Number P31025 (see Fig. 2).

[0032] The amino acid sequence of a mutein of the invention has a high sequence identity to mature human tear lipocalin when compared to sequence identities with other lipocalins (supra). In this general context the amino acid sequence of a mutein of the invention is at least substantially similar to the amino acid sequence of mature human tear lipocalin. A respective sequence of a mutein of the invention, being substantially similar to the sequences of mature human tear lipocalin, has in some embodiments at least 70 %, at least 75 %, at least 80 %, at least 82 %, at least 85 %, at least 87 %, at least 90% identity, including at least 95% identity to the sequence of mature human tear lipocalin, with the proviso that the altered position or sequence is retained.

[0033] By "identity" is meant a property of sequences that measures their similarity or relationship. Identity is measured by dividing the number of identical residues by the total number of residues and multiplying the product by 100. As two illustrative examples, the

mutein of the SEQ ID NO: 3 has a sequence identity of 83.3 % with the amino acid sequence of mature human tear lipocalin, and the mutein of the SEQ ID NO: 7 has an amino acid sequence identity of 82.0 % with mature human tear lipocalin.

[0034] "Gaps" are spaces in an alignment that are the result of additions or deletions of amino acids. Thus, two copies of exactly the same sequence have 100% identity, but sequences that are less highly conserved, and have deletions, additions, or replacements, may have a lower degree of identity. Those skilled in the art will recognize that several computer programs are available for determining sequence identity using standard parameters, for example Blast (Altschul, et al. (1997) Nucleic Acids Res. **25**, 3389-3402), Blast2 (Altschul, et al. (1990) J. Mol. Biol. **215**, 403-410), and Smith-Waterman (Smith, et al. (1981) J. Mol. Biol. **147**, 195-197).

[0035] The term “mutated” or “mutant” in reference to a nucleic acid or a polypeptide refers to the exchange, deletion, or insertion of one or more nucleotides or amino acids, respectively, compared to the naturally occurring nucleic acid or polypeptide. A mutein of the present invention includes at least three substitutions in comparison to the corresponding native human tear lipocalin.

[0036] In some embodiments a mutein according to the invention includes at least two amino acid substitutions, including 2, 3, 4, 5 or more amino acid substitutions of a native amino acid by an arginine residue. The substituted amino acid may in some embodiments be located at any of positions 27, 30, 57 and 83 with respect to the amino acid sequence of mature human tear lipocalin.

[0037] In some embodiments a mutein according to the invention includes an amino acid substitution of a native cysteine residue at positions 61 and/or 153 by a serine residue. In this context it is noted that it has been found that removal of the structural disulfide bond (on the level of a respective naïve nucleic acid library) of wild type tear lipocalin that is formed by the cystein residues 61 and 153 (cf. Breustedt, et al., 2005, supra) provides tear lipocalin muteins that are not only stably folded but in addition are also able to bind a given non-natural ligand with high affinity. Without wishing to be bound by theory, it is also believed that the elimination of the structural disulde bond provides the further advantage of allowing for the (spontaneous) generation or deliberate introduction of non-natural artifical disulfide bonds into muteins of the invention (see Examples), thereby increasing the stability of the muteins, for example. In some embodiments a mutein according to the invention includes an amino acid substitution of a native cysteine

residue at position 101 by a serine residue. Further, in some embodiments a mutein according to the invention includes an amino acid substitution of a native arginine residue at positions 111 by a proline residue. In some embodiments a mutein according to the invention includes an amino acid substitution of a native lysine residue at positions 114 by a tryptophan residue.

[0038] A mutein of human tear lipocalin according to the invention typically has one of an asparagine, a gutamic acid, a proline, a leucine, a lysine, a serine and a tyrosine at the position that corresponds to amino acid position 26 of mature human tear lipocalin. In some embodiments a mutein of the invention has a sequence in which amino acid position 34 is unchanged relative to mature human tear lipocalin, and where the sequence of the mutein includes the amino acid substitutions Arg 26 → Ser, Met 55 → Leu, Ser 58 → Lys. In some embodiments a mutein of the invention has a sequence that includes the amino acid substitutions Arg 26 → Pro and Glu 34 → Ser. In some embodiments a mutein of the invention has a sequence that includes the amino acid substitutions Arg 26 → Pro and Met 55 → Ala. In some embodiments a mutein of the invention has a sequence that includes the amino acid substitutions Arg 26 → Ser and Glu 34 → Val. In some embodiments a mutein of the invention has a sequence that includes the amino acid substitutions Arg 26 → Leu, Glu 34 → Trp and Met 55 → Ala. In some embodiments a mutein of the invention has a sequence that includes the amino acid substitutions Arg 26 → Leu, Glu 34 → Trp and Ser 58 → Ile. In some embodiments a mutein of the invention has a sequence that includes the amino acid substitutions Arg 26 → Ser, Glu 34 → Asn, Met 55 → Ala, and Ser 58 → Lys. In some embodiments a mutein of the invention has a sequence that includes the amino acid substitutions Arg 26 → Asn and Glu 34 → Asp. In some embodiments a mutein of the invention has a sequence that includes the amino acid substitutions Arg 26 → Asn and Met 55 → Ala. In some embodiments a mutein of the invention has a sequence that includes the amino acid substitutions Arg 26 → Tyr, Glu 34 → His and Met 55 → Ala. In some embodiments a mutein of the invention has a sequence that includes the amino acid substitutions Arg 26 → Tyr, Glu 34 → His and Ser 58 → Ala. In some embodiments a mutein of the invention has a sequence that includes the amino acid substitutions Arg 26 → Lys, Glu 34 → Arg and Met 55 → Ala. In some embodiments a mutein of the invention has a sequence that includes the amino acid substitutions Arg 26 → Lys, Glu 34 → Arg and Ser 58 → Asn. In some embodiments a mutein of the invention has a sequence that includes the amino acid substitutions Arg 26 → Glu, Glu 34 → Arg and Met 55 → Ala. In some embodiments a mutein of the invention has a sequence that

includes the amino acid substitutions Arg 26 → Glu, Glu 34 → Arg and Ser 58 → Leu.

[0039] In some embodiments a human tear lipocalin mutein of the invention that binds IL-4 receptor alpha has, when compared to the amino acid sequence of mature human tear lipocalin, a mutated amino acid residue at sequence position 58 or at sequence position 63. In some embodiments the sequence of the mutein of the invention is selected in such a way that, when compared to the amino acid sequence of mature human tear lipocalin, not at both amino acid positions 26 and 34 a serine is present.

[0040] The lipocalin mutein may further include with respect to the amino acid sequence of mature human tear lipocalin one or more, including at least two, at least three or at least four amino acid substitutions of native amino acid residues by cysteine residues at any of positions 26-28, 30-34, 53, 55-58, 61, 63, 64, 66, 80, 83, 104-106, and 108 of native mature human tear lipocalin. In some embodiments a mutein according to the invention includes an amino acid substitution of a native amino acid by a cysteine residue at positions 28 or 105 with respect to the amino acid sequence of mature human tear lipocalin. In some embodiments a mutein according to the invention includes an amino acid substitution of a native amino acid by a cysteine residue at positions 28 or 105 with respect to the amino acid sequence of mature human tear lipocalin.

[0041] In some embodiments a mutein according to the invention includes a mutated amino acid residue at any three or more, including 3, 4, 5, 6 or 7, of the sequence positions 26, 32, 34, 55, 56, 58 and 63 of the linear polypeptide sequence of the mature human tear lipocalin.

[0042] In some embodiments the lipocalin mutein of the invention has a mutated amino acid residue at position 26 that is one of asparagine, glutamine, proline, leucine, lysine, serine and tyrosine. In some embodiments the lipocalin mutein of the invention has a mutated amino acid residue at position 32 that is one of histidine, lysine, tyrosine and valine. In some embodiments the lipocalin mutein of the invention has a mutated amino acid residue at position 34 that is one of arginine, aspartic acid, asparagine, histidine, serine, tryptophan and valine. In some embodiments the lipocalin mutein of the invention has a mutated amino acid residue at position 55 that is one of alanine and leucine. In some embodiments the lipocalin mutein of the invention has a mutated amino acid residue at position 56 that is one of alanine, glutamine, histidine, methionine, leucine and lysine. In some embodiments the lipocalin mutein of the invention has a mutated amino acid residue at position 58 that is one of alanine, arginine, asparagine, histidine, isoleucine and lysine.

In some embodiments the lipocalin mutein of the invention has a mutated amino acid residue at position 63 that is one of glutamine, lysine, proline and serine.

[0043] In some embodiments a mutein according to the invention includes at least one of the substitutions Met 31 → Ala, Leu 33 → Tyr, Ser 61 → Trp, Asp 80 → Ser, Glu 104 → Leu, His 106 → Pro and Lys 108 → Gln. In some embodiments a mutein according to the invention includes two or more, such as 3, 4, 5, 6 or all of the substitutions Met 31 → Ala, Leu 33 → Tyr, Ser 61 → Trp, Asp 80 → Ser, Glu 104 → Leu, His 106 → Pro and Lys 108 → Gln. In some embodiments a mutein according to the invention includes a substitution Val 53 → Phe or Val 53 → Leu. The mutated amino acid residue may also include a substitution Val 64 → Tyr or Val 64 → Met. It may also include a substitution Ala 66 → Leu or Ala 66 → Asp.

[0044] In some embodiments a mutein of human tear lipocalin according to the invention includes a substituted amino acid of at least one or of both of the cysteine residues occurring at each of the sequences positions 61 and 153 by another amino acid and the mutation of at least three amino acid residue at any one of the sequence positions 26-28, 30-34, 53, 55-58, 63, 64, 66, 80, 83, 104-106, and 108 of the linear polypeptide sequence of mature human tear lipocalin. The positions 26-28 and 30-34 are included in the AB loop, the positions 53 and 55 are located at the very end of a beta-sheet and following positions 56-58 are included in the CD loop. Surprisingly, the positions 63, 64 and 66, are included within a beta-sheet (βD), and the position 80 is located in a α-helical region. Position 83 is a single loop-defining amino acid between this α-helical region and a beta-sheet (βF). The positions 104-106 and 108 are included in the GH loop in the binding site at the open end of the β-barrel structure of tear lipocalin. The definition of these regions is used herein in accordance with Flower (Flower, 1996, supra, Flower, et al., 2000, supra) and Breustedt et al. (2005, supra). Such a mutein may include at least 2, including 3, 4, 5, 6, 8, 10, 12, 14, 15, 16, 17 or 18 mutated amino acid residues at the sequence positions 26-34, 55-58, 63, 64, 80, 83, 104-106, and 108 of the linear polypeptide sequence of mature human tear lipocalin. In some embodiments the mutein includes the amino acid substitutions Cys 61 → Ala, Phe, Lys, Arg, Thr, Asn, Tyr, Met, Ser, Pro or Trp and Cys 153 → Ser or Ala. Such a substitution has proven useful to prevent the formation of the naturally occurring disulphide bridge linking Cys 61 and Cys 153, and thus to facilitate handling of the mutein. However, tear lipocalin muteins that binds IL-4 receptor alpha and that have the disulphide bridge formed between Cys 61 and Cys 153

are also part of the present invention

[0045] In some embodiments the mutein includes at least one amino acid substitution, which may be an additional amino acid substitution, selected from Arg 111 → Pro and Lys 114 → Trp. A mutein of the invention may further include the cysteine at position 101 of the sequence of native mature human tear lipocalin substituted by another amino acid. This substitution may, for example, be the mutation Cys 101 → Ser or Cys 101 → Thr.

[0046] As defined above, a mutein of the invention includes at least one amino acid substitution, which is located at a sequence position of the positions 27, 28, 30, 31, 33, 53, 57, 61, 64, 66, 80, 83, 104-106 and 108 of the linear polypeptide sequence of the mature human tear lipocalin. In some embodiments a mutein of the invention includes two or more, such as 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 1, 15 or 16 amino acid substitutions of these sequence positions of the mature human tear lipocalin. In one embodiment the mutein has a mutated amino acid residue at each of the sequence positions 27, 28, 30, 31, 33, 53, 57, 61, 64, 66, 80, 83, 104-106 and 108 of the linear polypeptide sequence of the mature human tear lipocalin.

[0047] In some embodiments the mutated amino acid residue at any one or more of the sequence positions 27, 28, 30, 31, 33, 53, 57, 61, 64, 66, 80, 83, 104-106 and 108 of the linear polypeptide sequence of the mature human tear lipocalin with respect to the amino acid sequence of mature human tear lipocalin includes one or more of the substitutions Met 31 → Ala, Leu 33 → Tyr, Ser 61 → Trp, Asp 80 → Ser, Glu 104 → Leu, His 106 → Pro and Lys 108 → Gln. In some embodiments a mutein of the invention includes two or more, such as 3, 4, 5, 6 or 7 amino acid substitutions of these sequence positions of the mature human tear lipocalin.

[0048] In the residual region, i.e. the region differing from sequence positions 26-28, 30-34, 53, 55-58, 63, 64, 66, 80, 83, 104-106, and 108, a lipocalin mutein of the invention may include the wild type (natural) amino acid sequence outside the mutated amino acid sequence positions. In some embodiments a lipocalin mutein according to the invention may also carry one or more amino acid mutations at a sequence position/ positions as long as such a mutation does, at least essentially not hamper or not interfere with the binding activity and the folding of the mutein. Such mutations can be accomplished very easily on DNA level using established standard methods. Illustrative examples of alterations of the amino acid sequence are insertions or deletions as well as amino acid

substitutions. Such substitutions may be conservative, i.e. an amino acid residue is replaced with an amino acid residue of chemically similar poperties, in particular with regard to polarity as well as size. Examples of conservative substitutions are the replacements among the members of the following groups: 1) alanine, serine, and threonine; 2) aspartic acid and glutamic acid; 3) asparagine and glutamine; 4) arginine and lysine; 5) isoleucine, leucine, methionine, and valine; and 6) phenylalanine, tyrosine, and tryptophan. On the other hand, it is also possible to introduce non-conservative alterations in the amino acid sequence. In addition, instead of replacing single amino acid residues, it is also possible to either insert or delete one or more continuous amino acids of the primary structure of tear lipocalin as long as these deletions or insertion result in a stable folded/functional mutein (see for example, the experimental section in which muteins with truncated N- and C-terminus are generated).

[0049] Such modifications of the amino acid sequence include directed mutagenesis of single amino acid positions in order to simplify sub-cloning of the mutated lipocalin gene or its parts by incorporating cleavage sites for certain restriction enzymes. In addition, these mutations can also be incorporated to further improve the affinity of a lipocalin mutein for a given target. Furthermore, mutations can be introduced in order to modulate certain characteristics of the mutein such as to improve folding stability, serum stability, protein resistance or water solubility or to reduce aggregation tendency, if necessary. For example, naturally occurring cysteine residues may be mutated to other amino acids to prevent disulphide bridge formation. It is also possible to deliberately mutate other amino acid sequence position to cysteine in order to introduce new reactive groups, for example for the conjugation to other compounds, such as polyethylene glycol (PEG), hydroxyethyl starch (HES), biotin, peptides or proteins, or for the formation of non-naturally occurring disulphide linkages. Exemplary possibilities of such a mutation to introduce a cysteine residue into the amino acid sequence of a human tear lipocalin mutein include the substitutions Thr 40→ Cys, Glu 73→ Cys, Arg 90→ Cys, Asp 95→ Cys, and Glu 131→ Cys. The generated thiol moiety at the side of any of the amino acid positions 40, 73, 90, 95 and/or 131 may be used to PEGylate or HESylate the mutein, for example, in order to increase the serum half-life of a respective tear lipocalin mutein.

[0050] The present invention also encompasses muteins as defined above, in which the first four N-terminal amino acid residues of the sequence of mature human tear lipocalin (His-His-Leu-Leu; positions 1-4) and/or the last two C-terminal amino acid

residues (Ser-Asp; positions 157-158) of the sequence of mature human tear lipocalin have been deleted (cf. also the Examples and the attached Sequence Listings). Another possible mutation of the wild type sequence is to change the amino acid sequence at sequence positions 5 to 7 (Ala Ser Asp) to Gly Gly Asp as described in PCT application WO 2005/019256.

[0051] In some embodiments a tear lipocalin mutein according to the invention has one or more, including 2, 3, 4, 5, 6, 7, 8, 9, 10, 11 or 12 of the following amino acid substitutions in comparison to mature human tear lipocalin: Glu 27 → Arg; Phe 28 → Cys; Glu 30 → Arg; Met 31 → Ala; Leu 33 → Tyr; Ile 57 → Arg; Ser 61→ Trp; Asp 80 → Ser; Lys 83 → Arg; Glu 104 → Leu; Leu 105 → Cys; His 106 → Pro; Lys 108 → Gln. In some embodiments the mutein includes all of these amino acid substitutions. In some embodiments the mutein further includes the set of amino acid substitutions Val 53 → Phe, Val 64 → Tyr, Ala 66 → Leu. In other embodiments the mutein further includes the set of amino acid substitutions Val 53 → Leu, Val 64 → Met, Ala 66 → Asp.

[0052] In some embodiments a tear lipocalin mutein according to the invention includes the combination of amino acid substitutions Arg 26 → Ser; Asn 32 → Tyr; Met 55 → Leu; Leu 56 → Gln; Ser 58 → Lys in comparison to mature human tear lipocalin. In some embodiments a tear lipocalin mutein according to the invention includes the combination of amino acid substitutions Arg 26 → Pro; Asn 32 → Tyr; Glu 34 → Ser; Met 55 → Ala; Leu 56 → Gln; Glu 63 → Lys in comparison to mature human tear lipocalin. In some embodiments a tear lipocalin mutein according to the invention includes the combination of amino acid substitutions Arg 26 → Leu; Asn 32 → Phe; Glu 34 → Trp; Met 55 → Ala; Ser 58 → Ile; Glu 63 → Ser in comparison to mature human tear lipocalin. In some embodiments a tear lipocalin mutein according to the invention includes the combination of amino acid substitutions Arg 26 → Ser; Asn 32 → Tyr; Glu 34 → Val; Met 55 → Ala; Leu 56 → Ala; Ser 58 → Ile; Glu 63 → Ser in comparison to mature human tear lipocalin. In some embodiments a tear lipocalin mutein according to the invention includes the combination of amino acid substitutions Arg 26 → Ser; Asn 32 → Val; Glu 34 → Asn; Met 55 → Ala; Leu 56 → Gln; Ser 58 → Lys; Glu 63 → Lys in comparison to mature human tear lipocalin. In some embodiments a tear lipocalin mutein according to the invention includes the combination of amino acid substitutions Arg 26 → Tyr; Asn 32 → Tyr; Glu 34 → His; Met 55 → Ala; Leu 56 → His; Ser 58 → Ala; Glu 63 → Lys in comparison to mature human tear lipocalin. In some embodiments a tear lipocalin

mutein according to the invention includes the combination of amino acid substitutions Arg 26 → Lys; Asn 32 → Tyr; Glu 34 → Arg; Met 55 → Ala; Leu 56 → Lys; Ser 58 → Asn; Glu 63 → Pro in comparison to mature human tear lipocalin. In some embodiments a tear lipocalin mutein according to the invention includes the combination of amino acid substitutions Arg 26 → Glu; Asn 32 → His; Glu 34 → Gly; Met 55 → Ala; Leu 56 → Met; Ser 58 → Leu; Glu 63 → Lys in comparison to mature human tear lipocalin.

[0053] In some embodiments a mutein of the invention includes with respect to the amino acid sequence of mature human tear lipocalin at least 6, 8, 10, 12, 14 or 16 amino acid substitutions selected from the group consisting of Arg 26 → Ser, Pro; Glu 27 → Arg; Phe 28 → Cys; Glu 30 → Arg; Met 31 → Ala; Asn 32 → Tyr, His; Leu 33 → Tyr; Glu 34 → Gly, Ser, Ala, Asp, Lys, Asn, Thr, Arg; Leu 56 → Gln; Ile 57 → Arg; Ser 58 → Ile, Ala, Arg, Val, Thr, Asn, Lys, Tyr, Leu, Met; Asp 80 → Ser; Lys 83 → Arg; Glu 104 → Leu; Leu 105 → Cys; His 106 → Pro; and Lys 108 → Gln.

[0054] Additionally, such a mutein may further include at least one amino acid substitution selected from the group consisting of Met 39 → Val; Thr 42 → Met, Ala; Thr 43 → Ile, Pro, Ala; Glu 45 → Lys, Gly; Asn 48 → Asp, His, Ser, Thr; Val 53 → Leu, Phe, Ile, Ala, Gly, Ser; Thr 54 → Ala, Leu; Met 55 → Leu, Ala, Ile, Val, Phe, Gly, Thr, Tyr; Glu 63 → Lys, Gln, Ala, Gly, Arg; Val 64 → Gly, Tyr, Met, Ser, Ala, Lys, Arg, Leu, Asn, His, Thr, Ile; Ala 66 → Ile, Leu, Val, Thr, Met; Glu 69 → Lys, Gly; Lys 70 → Arg, Gln, Glu; Thr 78 → Ala; Ile 89 → Val; Asp 95 → Asn, Ala, Gly; and Tyr 100 → His.

[0055] In one embodiment the human tear lipocalin mutein binding IL-4 receptor alpha includes the amino acid substitutions: Arg 26 → Ser, Glu 27 → Arg, Phe 28 → Cys, Glu 30 → Arg; Met 31 → Ala, Leu 33 → Tyr, Leu 56 → Gln, Ile 57 → Arg, Asp 80 → Ser, Lys 83 → Arg, Glu 104 → Leu, Leu 105 → Cys, His 106 → Pro, and Lys 108 → Gln.

[0056] In some embodiments the human tear lipocalin mutein binding IL-4 receptor alpha includes one of the following sets of amino acid substitutions:

(1) Arg 26 → Ser; Glu 27 → Arg; Phe 28 → Cys; Glu 30 → Arg; Met 31 → Ala; Asn 32 → Tyr; Leu 33 → Tyr; Glu 34 → Gly; Leu 56 → Gln; Ile 57 → Arg; Ser 58 → Ile; Asp 80 → Ser; Lys 83 → Arg; Glu 104 → Leu; Leu 105 → Cys; His 106 → Pro; Lys 108 → Gln;

(2) Arg 26 → Ser; Glu 27 → Arg; Phe 28 → Cys; Glu 30 → Arg; Met 31 → Ala; Asn 32 → Tyr; Leu 33 → Tyr; Glu 34 → Lys; Leu 56 → Gln; Ile 57 → Arg; Ser 58 → Asn; Asp 80 → Ser; Lys 83 → Arg; Glu 104 → Leu; Leu 105 → Cys; His 106 →

Pro; Lys 108 → Gln;

(3) Arg 26 → Ser; Glu 27 → Arg; Phe 28 → Cys, Glu 30 → Arg; Met 31 → Ala; Asn 32 → Tyr; Leu 33 → Tyr; Leu 56 → Gln; Ile 57 → Arg; Ser 58 → Arg; Asp 80 → Ser; Lys 83 → Arg; Glu 104 → Leu; Leu 105 → Cys; His 106 → Pro; Lys 108 → Gln;

(4) Arg 26 → Ser; Glu 27 → Arg; Phe 28 → Cys; Glu 30 → Arg; Met 31 → Ala; Asn 32 → Tyr; Leu 33 → Tyr; Glu 34 → Ser; Leu 56 → Gln; Ile 57 → Arg; Asp 80 → Ser; Lys 83 → Arg; Glu 104 → Leu; Leu 105 → Cys; His 106 → Pro; Lys 108 → Gln;

(5) Arg 26 → Ser; Glu 27 → Arg; Phe 28 → Cys; Glu 30 → Arg; Met 31 → Ala; Asn 32 → His; Leu 33 → Tyr; Glu 34 → Ser; Leu 56 → Gln; Ile 57 → Arg; Ser 58 → Ala; Asp 80 → Ser; Lys 83 → Arg; Glu 104 → Leu; Leu 105 → Cys; His 106 → Pro; Lys 108 → Gln;

(6) Arg 26 → Ser; Glu 27 → Arg; Phe 28 → Cys; Glu 30 → Arg; Met 31 → Ala; Asn 32 → Tyr; Leu 33 → Tyr; Glu 34 → Asp; Leu 56 → Gln; Ile 57 → Arg; Ser 58 → Lys; Asp 80 → Ser; Lys 83 → Arg; Glu 104 → Leu; Leu 105 → Cys; His 106 → Pro; Lys 108 → Gln; and

(7) Arg 26 → Ser;Glu 27 → Arg; Phe 28 → Cys; Glu 30 → Arg; Met 31 → Ala; Asn 32 → Tyr; Leu 33 → Tyr; Glu 34 → Gly; Leu 56 → Gln; Ile 57 → Arg; Asp 80 → Ser; Lys 83 → Arg; Glu 104 → Leu; Leu 105 → Cys; His 106 → Pro; Lys 108 → Gln.

[0057] The human tear lipocalin mutein of the invention may include, consist essentially of or consist of any one of the amino acid sequences set forth in SEQ ID NOs: 3-11 or a fragment or variant thereof. The term “fragment” as used herein in connection with the muteins of the invention relates to proteins or peptides derived from full-length mature human tear lipocalin that are N-terminally and/or C-terminally shortened, i.e. lacking at least one of the N-terminal and/or C-terminal amino acids. Such fragments may include at least 10, more such as 20 or 30 or more consecutive amino acids of the primary sequence of mature human tear lipocalin and are usually detectable in an immunoassay of mature human tear lipocalin.

[0058] The term “variant” as used in the present invention relates to derivatives of a protein or peptide that include modifications of the amino acid sequence, for example by substitution, deletion, insertion or chemical modification. Such modifications do in some

embodiments not reduce the functionality of the protein or peptide. Such variants include proteins, wherein one or more amino acids have been replaced by their respective D-stereoisomers or by amino acids other than the naturally occurring 20 amino acids, such as, for example, ornithine, hydroxyproline, citrulline, homoserine, hydroxylysine, norvaline. However, such substitutions may also be conservative, i.e. an amino acid residue is replaced with a chemically similar amino acid residue. Examples of conservative substitutions are the replacements among the members of the following groups: 1) alanine, serine, and threonine; 2) aspartic acid and glutamic acid; 3) asparagine and glutamine; 4) arginine and lysine; 5) isoleucine, leucine, methionine, and valine; and 6) phenylalanine, tyrosine, and tryptophan.

[0059] Such a mutein may include with respect to the amino acid sequence of mature human tear lipocalin at least 6, 8, 10, 12, 14 or 16 amino acid substitutions selected from the group consisting of Arg 26 → Ser; Glu 27 → Ile; Glu 30 → Ser; Met 31 → Gly; Asn 32 → Arg; Leu 33 → Ile; Glu 34 → Tyr; Leu 56 → Lys, Glu, Ala, Met; Ile 57 → Phe; Ser 58 → Arg; Asp 80 → Ser, Pro; Lys 83 → Glu, Gly; Glu 104 → Leu; Leu 105 → Ala; His 106 → Val; and Lys 108 → Thr and may further include at least one amino acid substitution selected from the group consisting of Leu 41→ Phe; Glu 63 → Lys; Val 64 → Met; Asp 72 → Gly; Lys 76 → Arg, Glu; Ile 88 → Val, Thr; Ile 89 → Thr; Arg 90 → Lys; Asp 95 → Gly; Phe 99 → Leu; and Gly 107 → Arg, Lys, Glu.

[0060] In one specific embodiment, such a mutein includes the amino acid substitutions: Arg 26 → Ser, Glu 27 → Ile, Glu 30 → Ser, Met 31 → Gly, Asn 32 → Arg, Leu 33 → Ile, Glu 34 → Tyr, Ile 57 → Phe, Ser 58 → Arg, Lys 83 → Glu, Glu 104 → Leu, Leu 105 → Ala, His 106 → Val, and Lys 108 → Thr.

[0061] A tear lipocalin mutein of the invention may exist as a monomeric protein. In some embodiments a lipocalin mutein according to the invention may be able to spontaneously dimerise or oligomerise. The use of lipocalin muteins that form stable monomers may be advantageous in some applications, e.g. because of faster diffusion and better tissue penetration. In other embodiments the use of a lipocalin mutein that spontaneously forms stable homodimers or multimers may be advantageous, since such multimers can provide (further) increased affinity and/or avidity to a given target. Furthermore, oligomeric forms of the lipocalin mutein may have slower dissociation rates or prolonged serum half-life. If dimerisation or multimerisation of muteins that form stable monomers is desired, this can for example be achieved by fusing respective

oligomerization domains such as jun-fos domains or leucin-zippers to muteins of the invention or by the use of “Duocalins” (see also below).

[0062] A lipocalin mutein according to the present invention can be obtained by means of mutagenesis of a naturally occurring form of human tear lipocalin. The term "mutagenesis" as used herein means that the experimental conditions are chosen such that the amino acid naturally occurring at a given sequence position of human tear lipocalin (Swiss-Prot data bank entry P31025) can be substituted by at least one amino acid that is not present at this specific position in the respective natural polypeptide sequence. The term "mutagenesis" also includes the (additional) modification of the length of sequence segments by deletion or insertion of one or more amino acids. Thus, it is within the scope of the invention that, for example, one amino acid at a chosen sequence position is replaced by a stretch of three random mutations, leading to an insertion of two amino acid residues compared to the length of the respective segment of the wild type protein. Such an insertion of deletion may be introduced independently from each other in any of the peptide segments that can be subjected to mutagenesis in the invention. In one exemplary embodiment of the invention, an insertion of several mutations may be introduced into the loop AB of the chosen lipocalin scaffold (cf. International Patent Application WO 2005/019256 which is incorporated by reference its entirety herein). The term "random mutagenesis" means that no predetermined single amino acid (mutation) is present at a certain sequence position but that at least two amino acids can be incorporated with a certain probability at a predefined sequence position during mutagenesis.

[0063] The coding sequence of human tear lipocalin (Redl, B. et al. (1992) *J. Biol. Chem.* **267**, 20282-20287) is used as a starting point for the mutagenesis of the peptide segments selected in the present invention. For the mutagenesis of the recited amino acid positions, the person skilled in the art has at his disposal the various established standard methods for site-directed mutagenesis. A commonly used technique is the introduction of mutations by means of PCR (polymerase chain reaction) using mixtures of synthetic oligonucleotides, which bear a degenerate base composition at the desired sequence positions. For example, use of the codon NNK or NNS (wherein N = adenine, guanine or cytosine or thymine; K = guanine or thymine; S = adenine or cytosine) allows incorporation of all 20 amino acids plus the amber stop codon during mutagenesis, whereas the codon VVS limits the number of possibly incorporated amino acids to 12,

since it excludes the amino acids Cys, Ile, Leu, Met, Phe, Trp, Tyr, Val from being incorporated into the selected position of the polypeptide sequence; use of the codon NMS (wherein M = adenine or cytosine), for example, restricts the number of possible amino acids to 11 at a selected sequence position since it excludes the amino acids Arg, Cys, Gly, Ile, Leu, Met, Phe, Trp, Val from being incorporated at a selected sequence position. In this respect it is noted that codons for other amino acids (than the regular 20 naturally occurring amino acids) such as selenocystein or pyrrolysine can also be incorporated into a nucleic acid of a mutein. It is also possible, as described by Wang, L., et al. (2001) *Science* **292**, 498-500, or Wang, L., and Schultz, P.G. (2002) *Chem. Comm.* **1**, 1-11, to use "artificial" codons such as UAG which are usually recognized as stop codons in order to insert other unusual amino acids, for example o-methyl-L-tyrosine or p-aminophenylalanine.

[0064] The use of nucleotide building blocks with reduced base pair specificity, as for example inosine, 8-oxo-2'deoxyguanosine or 6(2-deoxy-□-D-ribofuranosyl)-3,4-dihydro-8H-pyrimindo-1,2-oxazine-7-one (Zaccolo et al. (1996) *J. Mol. Biol.* **255**, 589-603), is another option for the introduction of mutations into a chosen sequence segment.

[0065] A further possibility is the so-called triplet-mutagenesis. This method uses mixtures of different nucleotide triplets, each of which codes for one amino acid, for incorporation into the coding sequence (Virnekäs B, et al., 1994 *Nucleic Acids Res* **22**, 5600-5607).

[0066] One possible strategy for introducing mutations in the selected regions of the respective polypeptides is based on the use of four oligonucleotides, each of which is partially derived from one of the corresponding sequence segments to be mutated. When synthesizing these oligonucleotides, a person skilled in the art can employ mixtures of nucleic acid building blocks for the synthesis of those nucleotide triplets which correspond to the amino acid positions to be mutated so that codons encoding all natural amino acids randomly arise, which at last results in the generation of a lipocalin peptide library. For example, the first oligonucleotide corresponds in its sequence - apart from the mutated positions - to the coding strand for the peptide segment to be mutated at the most N-terminal position of the lipocalin polypeptide. Accordingly, the second oligonucleotide corresponds to the non-coding strand for the second sequence segment following in the polypeptide sequence. The third oligonucleotide corresponds in turn to the coding strand for the corresponding third sequence segment. Finally, the fourth oligonucleotide

corresponds to the non-coding strand for the fourth sequence segment. A polymerase chain reaction can be performed with the respective first and second oligonucleotide and separately, if necessary, with the respective third and fourth oligonucleotide.

[0067] The amplification products of both of these reactions can be combined by various known methods into a single nucleic acid that includes the sequence from the first to the fourth sequence segments, in which mutations have been introduced at the selected positions. To this end, both of the products can for example be subjected to a new polymerase chain reaction using flanking oligonucleotides as well as one or more mediator nucleic acid molecules, which contribute the sequence between the second and the third sequence segment. In the choice of the number and arrangement within the sequence of the oligonucleotides used for the mutagenesis, the person skilled in the art has numerous alternatives at his disposal.

[0068] The nucleic acid molecules defined above can be connected by ligation with the missing 5'- and 3'-sequences of a nucleic acid encoding a lipocalin polypeptide and/or the vector, and can be cloned in a known host organism. A multitude of established procedures are available for ligation and cloning. For example, recognition sequences for restriction endonucleases also present in the sequence of the cloning vector can be engineered into the sequence of the synthetic oligonucleotides. Thus, after amplification of the respective PCR product and enzymatic cleavage the resulting fragment can be easily cloned using the corresponding recognition sequences.

[0069] Longer sequence segments within the gene coding for the protein selected for mutagenesis can also be subjected to random mutagenesis via known methods, for example by use of the polymerase chain reaction under conditions of increased error rate, by chemical mutagenesis or by using bacterial mutator strains. Such methods can also be used for further optimization of the target affinity or specificity of a lipocalin mutein. Mutations possibly occurring outside the segments of experimental mutagenesis are often tolerated or can even prove to be advantageous, for example if they contribute to an improved folding efficiency or folding stability of the lipocalin mutein.

[0070] In a method according to the invention a nucleic acid molecule encoding a human tear lipocalin is firstly subjected to mutagenesis at one or more of the amino acid sequence positions 27, 28, 30, 31, 33, 53, 57, 61, 64, 66, 80, 83, 104-106 and 108 of the linear polypeptide sequence of mature human tear lipocalin. Secondly the nucleic acid molecule encoding a human tear lipocalin is also subjected to mutagenesis at two or more of

the amino acid sequence positions 26, 32, 34, 55, 56, 58 and 63 of the linear polypeptide sequence of the mature human tear lipocalin. Of these latter amino acid sequence positions at least one position to be mutated is selected from amino acid sequence position 58 and amino acid sequence position 63.

[0071] In one embodiment of the invention, the method for the generation of a mutein of human tear lipocalin includes mutating at least 2, 3, 4, 5, 6, 8, 10, 12, 14, 15, 16, or 17 of the codons of any of the amino acid sequence positions 26-28, 30-34, 53, 55-58, 63, 64, 66, 80, 83, 104-106, and 108 of the linear polypeptide sequence of mature human tear lipocalin. In one embodiment all 22 of the codons of amino acid sequence positions 26, 27, 28, 30, 31, 32, 33, 34, 53, 55, 56, 57, 58, 63, 64, 66, 80, 83, 104, 105, 106, and 108 of the linear polypeptide sequence of mature human tear lipocalin are mutated.

[0072] In one embodiment of the afore-mentioned method, additionally at least 2, 3, 4, 5, 6, 8, 10, 12, 14, or 15 of the codons of any of the amino acid sequence positions 26-28, 30-34, 53, 55-58, 63, 64, 66, 80, 83, 104-106, and 108 of the linear polypeptide sequence of mature human tear lipocalin are mutated.

[0073] In a further embodiment of the invention, the methods according to the invention include the mutation of both of the codons encoding cysteine at positions 61 and 153 in the linear polypeptide sequence of mature human tear lipocalin. In one embodiment position 61 is mutated to encode an alanine, phenylalanine, lysine, arginine, threonin, asparagine, tyrosine, methionine, serine, proline or a tryptophane residue, to name only a few possibilities. In embodiments where position 153 is mutated, an amino acid such as a serine or alanine can be introduced at position 153.

[0074] In another embodiment of the invention as described herein, the codons encoding amino acid sequence positions 111 and/or 114 of the linear polypeptide sequence of mature human tear lipocalin are mutated to encode for example an arginine at position 111 and a tryptophane at position 114.

[0075] Another embodiment of the methods of the invention, involves mutagenesis of the codon encoding the cysteine at position 101 of the linear polypeptide sequence of mature human tear lipocalin so that this codon encodes any other amino acid. In one embodiment the mutated codon encoding position 101 encodes a serine. Accordingly, in some embodiments either two or all three of the cystein codons at position 61, 101 and 153 are replaced by a codon of another amino acid.

[0076] According to the method of the invention a mutein is obtained starting from a nucleic acid encoding human tear lipocalin. Such a nucleic acid is subjected to mutagenesis and introduced into a suitable bacterial or eukaryotic host organism by means of recombinant DNA technology. Obtaining a nucleic acid library of tear lipocalin can be carried out using any suitable technique that is known in the art for generating lipocalin muteins with antibody-like properties, i.e. muteins that have affinity towards a given target. Examples of such combinatorial methods are described in detail in the international patent applications WO 99/16873, WO 00/75308, WO 03/029471, WO 03/029462, WO 03/029463, WO 2005/019254, WO 2005/019255, WO 2005/019256, or WO 2006/56464 for instance. The content of each of these patent applications is incorporated by reference herein in its entirety. After expression of the nucleic acid sequences that were subjected to mutagenesis in an appropriate host, the clones carrying the genetic information for the plurality of respective lipocalin muteins, which bind a given target can be selected from the library obtained. Well known techniques can be employed for the selection of these clones, such as phage display (reviewed in Kay, B.K. et al. (1996) *supra*; Lowman, H.B. (1997) *supra* or Rodi, D.J., and Makowski, L. (1999) *supra*), colony screening (reviewed in Pini, A. et al. (2002) *Comb. Chem. High Throughput Screen.* **5**, 503-510), ribosome display (reviewed in Amstutz, P. et al. (2001) *Curr. Opin. Biotechnol.* **12**, 400-405) or mRNA display as reported in Wilson, D.S. et al. (2001) *Proc. Natl. Acad. Sci. USA* **98**, 3750-3755 or the methods specifically described in WO 99/16873, WO 00/75308, WO 03/029471, WO 03/029462, WO 03/029463, WO 2005/019254, WO 2005/019255, WO 2005/019256, or WO 2006/56464.

[0077] The nucleic acid molecule encoding the mutein is expressed using any suitable expression system. The obtained mutein or muteins is/are enriched by means of selection and/ or isolation. The selection may for example be carried out under competitive conditions. Competitive conditions as used herein means that selection of muteins encompasses at least one step in which the muteins and the given non-natural ligand of human tear lipocalin, i.e.the IL 4 receptor alpha, are brought in contact in the presence of an additional ligand, which competes with binding of the muteins to IL 4 receptor alpha. This additional ligand may be a physiological ligand of the target, e.g. IL 4, an excess of the target itself or any other non-physiological ligand of the target that binds at least an overlapping epitope to the epitope recognized by the muteins of the invention and thus interferes with target binding of the muteins. Alternatively, the additional ligand competes with binding of the muteins by complexing an epitope distinct from the binding

site of the muteins to the target by allosteric effects.

[0078] An embodiment of the phage display technique (reviewed in Kay, B.K. et al. (1996), *supra*; Lowman, H. B. (1997) *supra* or Rodi, D.J., & Makowski, L. (1999), *supra*) using temperent M13 phage is given as an example of a selection method that can be employed in the present invention. Another embodiment of the phage display technology that can be used for selection of muteins of the invention is the hyperphage phage technology as described by Broders et al. (Broders et al. (2003) "Hyperphage. Improving antibody presentation in phage display." *Methods Mol. Biol.* **205**:295-302). Other temperent phage such as f1 or lytic phage such as T7 may be employed as well. For the exemplary selection method, M13 phagemids are produced which allow the expression of the mutated lipocalin nucleic acid sequence as a fusion protein with a signal sequence at the N-terminus, such as the OmpA-signal sequence, and with the capsid protein pIII of the phage M13 or fragments thereof capable of being incorporated into the phage capsid at the C-terminus. The C-terminal fragment ΔpIII of the phage capsid protein that includes amino acids 217 to 406 of the wild type sequence is may be used to produce the fusion proteins. In one embodiment a C-terminal fragment of pIII is used, in which the cysteine residue at position 201 is missing or is replaced by another amino acid.

[0079] Accordingly, a further embodiment of the methods of the invention involves operably fusing a nucleic acid coding for the one or more muteins of human tear lipocalin and resulting from mutagenesis at the 3' end with a gene coding for the coat protein pIII of a filamentous bacteriophage of the M13-family or for a fragment of this coat protein, in order to select at least one mutein for the binding of a given ligand.

[0080] The fusion protein may include additional components such as an affinity tag, which allows the immobilization, detection and/or purification of the fusion protein or its parts. Furthermore, a stop codon can be located between the sequence regions encoding the lipocalin or its muteins and the phage capsid gene or fragments thereof, wherein the stop codon, such as an amber stop codon, is at least partially translated into an amino acid during translation in a suitable suppressor strain.

[0081] For example, the phasmid vector pTLPC27, now also called pTlc27 that is described here can be used for the preparation of a phagemid library encoding human tear lipocalin muteins. The inventive nucleic acid molecules coding for the tear lipocalin muteins are inserted into the vector using the two *Bst*XI restriction sites. After ligation a suitable host strain such as *E. coli* XL1-Blue is transformed with the resulting nucleic acid

mixture to yield a large number of independent clones. A respective vector can be generated for the preparation of a hyperphagemid library, if desired.

[0082] The resulting library is subsequently superinfected in liquid culture with an appropriate M13-helper phage or hyperphage in order to produce functional phagemids. The recombinant phagemid displays the lipocalin mutein on its surface as a fusion with the coat protein pIII or a fragment thereof, while the N-terminal signal sequence of the fusion protein is normally cleaved off. On the other hand, it also bears one or more copies of the native capsid protein pIII supplied by the helper phage and is thus capable of infecting a recipient, in general a bacterial strain carrying an F- or F'-plasmid. In case of hyperphage display, the hyperphagemids display the lipocalin muteins on their surface as a fusion with the infective coat protein pIII but no native capsid protein. During or after infection with helper phage or hyperphage, gene expression of the fusion protein between the lipocalin mutein and the capsid protein pIII can be induced, for example by addition of anhydrotetracycline. The induction conditions are chosen such that a substantial fraction of the phagemids obtained displays at least one lipocalin mutein on their surface. In case of hyperphage display induction conditions result in a population of hyperphagemids carrying between three and five fusion proteins consisting of the lipocalin mutein and the capsid protein pIII. Various methods are known for isolating the phagemids, such as precipitation with polyethylene glycol. Isolation typically occurs after an incubation period of 6-8 hours.

[0083] The isolated phasmids can then be subjected to selection by incubation with the desired target, wherein the target is presented in a form allowing at least temporary immobilization of those phagemids which carry muteins with the desired binding activity as fusion proteins in their coat. Among the various embodiments known to the person skilled in the art, the target can, for example, be conjugated with a carrier protein such as serum albumin and be bound via this carrier protein to a protein binding surface, for example polystyrene. Microtiter plates suitable for ELISA techniques or so-called "immuno-sticks" can for instance be used for such an immobilization of the target. Alternatively, conjugates of the target with other binding groups, such as biotin, can be used. The target can then be immobilized on a surface which selectively binds this group, for example microtiter plates or paramagnetic particles coated with streptavidin, neutravidin or avidin. If the target is fused to an Fc portion of an immunoglobulin, immobilization can also be achieved with surfaces, for example microtiter plates or paramagnetic particles, which are coated

with protein A or protein G.

[0084] Non-specific phagemid-binding sites present on the surfaces can be saturated with blocking solutions as they are known for ELISA methods. The phagemids are then typically brought into contact with the target immobilized on the surface in the presence of a physiological buffer. Unbound phagemids are removed by multiple washings. The phagemid particles remaining on the surface are then eluted. For elution, several methods are possible. For example, the phagemids can be eluted by addition of proteases or in the presence of acids, bases, detergents or chaotropic salts or under moderately denaturing conditions. One such method is the elution using buffers of pH 2.2, wherein the eluate is subsequently neutralized. Alternatively, a solution of the free target can be added in order to compete with the immobilzed target for binding to the phagemids or target-specific phagemids can be eluted by competition with immunoglobulins or natural liganding proteins which specifically bind to the target of interest.

[0085] Afterwards, *E. coli* cells are infected with the eluted phagemids. Alternatively, the nucleic acids can be extracted from the eluted phagemids and used for sequence analysis, amplification or transformation of cells in another manner. Starting from the *E. coli* clones obtained in this way, fresh phagemids or hyperphagemids are again produced by superinfection with M13 helper phages or hyperphage according to the method described above and the phagemids amplified in this way are once again subjected to a selection on the immobilized target. Multiple selection cycles are often necessary in order to obtain the phagemids with the muteins of the invention in sufficiently enriched form. The number of selection cycles is in some embodiments chosen in such a way that in the subsequent functional analysis at least 0.1 % of the clones studied produce muteins with detectable affinity for the given target. Depending on the size, i.e. the complexity of the library employed, 2 to 8 cycles are typically required to this end.

[0086] For the functional analysis of the selected muteins, an *E. coli* strain is infected with the phagemids obtained from the selection cycles and the corresponding double stranded phasmid DNA is isolated. Starting from this phasmid DNA, or also from the single-stranded DNA extracted from the phagemids, the nucleic acid sequences of the selected muteins of the invention can be determined by the methods known in the art and the amino acid sequence can be deduced therefrom. The mutated region or the sequence of the entire tear lipocalin mutein can be subcloned on another expression vector and expressed in a suitable host organism. For example, the vector pTLPC26 now also called

pTlc26 can be used for expression in *E. coli* strains such as *E.coli* TG1. A muteins of tear lipocalin thus produced can be purified by various biochemical methods. A tear lipocalin mutein produced, for example with pTlc26, may carry an affinity peptide, a so called affinity tag, for instance at its C-terminus and can therefore be purified by affinity chromatography. Examples of an affinity tag include, but are not limited to biotin, the Strep-tag, Strep-tag II (Schmidt et al., supra), oligohistidine, polyhistidine, an immunoglobulin domain, maltose-binding protein, glutathione-S-transferase (GST) or calmodulin binding peptide (CBP).

[0087] Some affinity tags are haptens, for example but not limited to, dinitrophenol and digoxigenin. Some affinity tags are epitope tags, such as the FLAG®-peptide (Asp-Tyr-Lys-Asp-Asp-Asp-Asp-Lys-Gly), the T7 epitope (Ala-Ser-Met-Thr-Gly-Gly-Gln-Gln-Met-Gly), maltose binding protein (MBP), the HSV epitope of the sequence Gln-Pro-Glu-Leu-Ala-Pro-Glu-Asp-Pro-Glu-Asp of herpes simplex virus glycoprotein D, the hemagglutinin (HA) epitope of the sequence Tyr-Pro-Tyr-Asp-Val-Pro-Asp-Tyr-Ala, the VSV-G epitope of the Vesicular Stomatitis viral glycoprotein (Cys-Tyr-The-Asp-Ile-Glu-Met-Asn-Arg-Leu-Lys), the E epitope tag of the sequence Gly-Ala-Pro-Val-Pro-Tyr-Pro-Asp-Pro-Leu-Glu-Pro-Arg, the E2 epitope tag of the sequence Gly-Val-Ser-Ser-Thr-Ser-Ser-Asp-Phe-Arg-Asp-Arg, the Tag-100 epitope tag of C-termini of mammalian MAPK/ERK kinases of the sequence Glu-Glu-Thr-Ala-Arg-Phe-Gln-Pro-Gly-Tyr-Arg-Ser, the S-tag of the sequence Lys-Glu-Thr-Ala-Ala-Ala-Lys-Phe-Glu-Arg-Gln-His-Met-Asp-Ser, the “myc” epitope of the transcription factor c-myc of the sequence Glu-Gln-Lys-Leu-Ile-Ser-Glu-Glu-Asp-Leu and the small V5 epitope present on the P and V proteins of the paramyxovirus of Simian Virus 5 (Gly-Lys-Pro-Ile-Pro-Asn-Pro-Leu-Leu-Gly-Leu-Asp-Ser-Thr). In addition, but generally not as a single tag, a solubility-enhancing tag such as NusA, thioredoxin (TRX), small ubiquitin-like modifier (SUMO), and ubiquitin (Ub) may be used. Haptens and epitope tags may be used in combination with a corresponding antibody or an antibody like proteinaceous molecule as binding partner. The S-peptide epitope of the sequence Lys-Glu-Thr-Ala-Ala-Ala-Lys-Phe-Glu-Arg-Gln-His-Met-Asp-Ser may be used as an epitope tag in connection with a respective antibody or in combination with the S-protein as a binding partner (Hackbarth, JS, et al., BioTechniques (2004) 37, 5, 835-839).

[0088] The selection can also be carried out by means of other methods. Many corresponding embodiments are known to the person skilled in the art or are described in

the literature. Moreover, a combination of methods can be applied. For example, clones selected or at least enriched by "phage display" can additionally be subjected to "colony screening". This procedure has the advantage that individual clones can directly be isolated with respect to the production of a tear lipocalin mutein with detectable binding affinity for a target.

[0089] In addition to the use of *E. coli* as host organism in the "phage display" technique or the "colony screening" method, other bacterial strains, yeast or also insect cells or mammalian cells can be used for this purpose. Further to the selection of a tear lipocalin mutein from a random library as described above, evolutive methods including limited mutagenesis can also be applied in order to optimize a mutein that already possesses some binding activity for the target with respect to affinity or specificity for the target after repeated screening cycles.

[0090] It is readily apparent to the skilled person that complex formation is dependent on many factors such as concentration of the binding partners, the presence of competitors, ionic strength of the buffer system etc. Selection and enrichment is generally performed under conditions allowing the isolation of lipocalin muteins having, in complex with the desired target, a dissociation constant of at least 200 nM. However, the washing and elution steps can be carried out under varying stringency. A selection with respect to the kinetic characteristics is possible as well. For example, the selection can be performed under conditions, which favor complex formation of the target with muteins that show a slow dissociation from the target, or in other words a low k_{off} rate. Alternatively, selection can be perfomed under conditions, which favour fast formation of the complex between the mutein and the target, or in other words a high k_{on} rate. As a further illustrative alternative, the screening can be performed under conditions that select for improved thermostability of the muteins (compared to either wild type tear lipocalin or a mutein that already has affinity towards a pre-selected target).

[0091] Once a mutein with affinity to a given target has been selected, it is additionally possible to subject such a mutein to another mutagenesis in order to subsequently select variants of even higher affinity or variants with improved properties such as higher thermostability, improved serum stability, thermodynamic stability, improved solubility, improved monomeric behavior, improved resistance against thermal denaturation, chemical denaturation, proteolysis, or detergents etc. This further mutagenesis, which in case of aiming at higher affinity can be considered as in vitro

"affinity maturation", can be achieved by site specific mutation based on rational design or a random mutation. Another possible approach for obtaining a higher affinity or improved properties is the use of error-prone PCR, which results in point mutations over a selected range of sequence positions of the lipocalin mutein. The error-prone PCR can be carried out in accordance with any known protocol such as the one described by Zaccolo et al. (1996) J. Mol. Biol. **255**, 589-603. Other methods of random mutagenesis that are suitable for such purposes include random insertion/deletion (RID) mutagenesis as described by Murakami, H et al. (2002) Nat.Biotechnol. **20**, 76-81 or nonhomologous random recombination (NRR) as described by Bittker, J. A et al. (2002) Nat. Biotechnol. **20**,1024-1029. If desired, affinity maturation can also be carried out according to the procedure described in WO 00/75308 or Schlehuber, S. et al., (2000) J. Mol. Biol. **297**, 1105-1120, where muteins of the bilin-binding protein having high affinity to digoxigenin were obtained.

[0092] A tear lipocalin mutein of the invention may be used for complex formation with IL 4 receptor alpha. The mutein may also be able to bind an immunogenic fragment of IL 4 receptor alpha. An immunogenic fragment of IL-4 recptor alpha is a fragment that has one or more epitopes, mimotopes or other antigenic determinants, and is thus capable of inducing an immune response or against which an antibody can be raised. The immunogenic fragment may include a single epitope or may have a plurality of epitopes. Since an antigen-presenting system, e.g. a carrier protein, may be used to provide the size required for recognition by an immune system, no particular size limitation applies to the immunogenic fragment. Hence, the immunogenic fragment may also be a "hapten", i.e. a fragment that need not be antigenic per se or may have low immunogenicity, in particular due to its small molecular weight and accordingly size. Typically an immunogenic fragment can, alone or when presented on a carrier, be bound by an immunoglobulin or by a T cell receptor (TCR) if presented by MHC molecules. An immunogenic fragment is typically, alone or when presented in the form of the antigen-presenting system, capable of inducing a humoral immune response and/or cellular immune response leading for instance to the activation of B- and/or T-lymphocytes.

[0093] The target of a mutein of the invention is the alpha chain of the interleukin-4 receptor, a transmembrane protein, which contains an extracellular domain of 207 amino acids. A secreted form of the extracellular domain exists, sIL-4R alpha, which is also known as CD124 and capable of blocking IL-4 activities. A mutein of the invention may be

able to bind sIL-4 receptor alpha as well as any portion of the extracellular domain of IL 4 receptor alpha.

[0094] In this context it is also noted that the complex formation between the respective mutein and its ligand is influenced by many different factors such as the concentrations of the respective binding partners, the presence of competitors, pH and the ionic strength of the buffer system used, and the experimental method used for determination of the dissociation constant K_D (for example fluorescence titration, competition ELISA or surface plasmon resonance, just to name a few) or even the mathematical algorithm which is used for evaluation of the experimental data.

[0095] Therefore, it is also clear to the skilled person that the K_D values (dissociation constant of the complex formed between the respective mutein and its ligand) given here may vary within a certain experimental range, depending on the method and experimental setup that is used for determining the affinity of a particular lipocalin mutein for a given ligand. This means, there may be a slight deviation in the measured K_D values or a tolerance range depending, for example, on whether the K_D value was determined by surface plasmon resonance (Biacore) or by competition ELISA.

[0096] Also included in the scope of the present invention are forms of the above muteins, in which the respective mutein has been altered or modified with respect to its potential immunogenicity.

[0097] Cytotoxic T-cells recognize peptide antigens on the cell surface of an antigen-presenting cell in association with a class I major histocompatibility complex (MHC) molecule. The ability of the peptides to bind to MHC molecules is allele specific and correlates with their immunogenicity. In order to reduce immunogenicity of a given protein, the ability to predict which peptides in a protein have the potential to bind to a given MHC molecule is of great value. Approaches that employ a computational threading approach to identify potential T-cell epitopes have been previously described to predict the binding of a given peptide sequence to MHC class I molecules (Altuvia et al. (1995) *J. Mol. Biol.* **249**, 244-250).

[0098] Such an approach may also be utilized to identify potential T-cell epitopes in the muteins of the invention and to make depending on its intended use a selection of a specific mutein on the basis of its predicted immunogenicity. It may be furthermore possible to subject peptide regions which have been predicted to contain T-cell epitopes to additional mutagenesis to reduce or eliminate these T-cell epitopes and thus minimize

immunogenicity. The removal of amphipathic epitopes from genetically engineered antibodies has been described (Mateo et al. (2000) *Hybridoma* **19,** 6, 463-471) and may be adapted to the muteins of the present invention.

[0099] The muteins thus obtained may possess a minimized immunogenicity, which is desirable for their use in therapeutic and diagnostic applications, such as those described below.

[00100] For some applications, it is also useful to employ the muteins of the invention in a labeled form. Accordingly, the invention is also directed to lipocalin muteins which are conjugated to a label selected from the group consisting of enzyme labels, radioactive labels, colored labels, fluorescent labels, chromogenic labels, luminescent labels, haptens, digoxigenin, biotin, metal complexes, metals, and colloidal gold. The mutein may also be conjugated to a low molecular weight organic compound. The term "low molecular weight organic compound" as used herein denotes a monomeric carbon-based compound, which may have aliphatic, alicyclic and/or aromatic moieties. In typical embodiments the low molecular weight organic compound is an organic compound that has a main chain of at least two carbon atoms, and in some embodiments not more than 7 or 12 rotatable carbon bonds. Such a compound has a molecular weight in the range from about 100 to about 2000 Dalton, such as from about 100 to about 1000 Dalton. It may optionally include one or two metal atoms.

[00101] In general, it is possible to label the lipocalin mutein with any appropriate chemical substance or enzyme, which directly or indirectly generates a detectable compound or signal in a chemical, physical, optical, or enzymatic reaction. An example for a physical reaction and at the same time optical reaction/marker is the emission of fluorescence upon irradiation or the emission of X-rays when using a radioactive label. Alkaline phosphatase, horseradish peroxidase or β-galactosidase are examples of enzyme labels (and at the same time optical labels) which catalyze the formation of chromogenic reaction products. In general, all labels commonly used for antibodies (except those exclusively used with the sugar moiety in the Fc part of immunoglobulins) can also be used for conjugation to the muteins of the present invention. The muteins of the invention may also be conjugated with any suitable therapeutically active agent, e.g., for the targeted delivery of such agents to a given cell, tissue or organ or for the selective targeting of cells, e.g., of tumor cells without affecting the surrounding normal cells. Examples of such therapeutically active agents include radionuclides, toxins, small

organic molecules, and therapeutic peptides (such as peptides acting as agonists/antagonists of a cell surface receptor or peptides competing for a protein binding site on a given cellular target). The lipocalin muteins of the invention may, however, also be conjugated with therapeutically active nucleic acids such as antisense nucleic acid molecules, small interfering RNAs, micro RNAs or ribozymes. Such conjugates can be produced by methods well known in the art.

[00102] In one embodiment, the muteins of the invention may also be coupled to a targeting moiety that targets a specific body region in order to deliver the inventive muteins to a desired region or area within the body. One example wherein such modification may be desirable is the crossing of the blood-brain-barrier. In order to cross the blood-brain barrier, the muteins of the invention may be coupled to moieties that facilitate the active transport across this barrier (see Gaillard PJ, et al., Diphtheria-toxin receptor-targeted brain drug delivery. *International Congress Series,* 2005 1277, 185-198 or Gaillard PJ, et al. Targeted delivery across the blood-brain barrier. *Expert Opin Drug Deliv.* 2005 2, 2, 299-309. Such moieties are for example available under the trade name 2B-Trans™ (to-BBB technologies BV, Leiden, NL).

[00103] As indicated above, a mutein of the invention may in some embodiments be conjugated to a moiety that extends the serum half-life of the mutein (in this regard see also PCT publication WO 2006/56464 where such conjugation strategies are described with references to muteins of human neutrophile gelatinase-associated lipocalin with binding affinity for CTLA-4). The moiety that extends the serum half-life may be a polyalkylene glycol molecule, hydroxyethyl starch, fatty acid molecules, such as palmitic acid (Vajo & Duckworth 2000, *Pharmacol. Rev*. **52**, 1-9), an Fc part of an immunoglobulin, a CH3 domain of an immunoglobulin, a CH4 domain of an immunoglobulin, albumin or a fragment thereof, an albumin binding peptide, or an albumin binding protein, transferrin to name only a few. The albumin binding protein may be a bacterial albumin binding protein, an antibody, an antibody fragment including domain antibodies (see US patent 6,696,245, for example), or a lipocalin mutein with binding activity for albumin. Accordingly, suitable conjugation partners for extending the half-life of a lipocalin mutein of the invention include albumin (Osborn, B.L. et al., 2002, *J. Pharmacol. Exp. Ther.* **303**, 540-548), or an albumin binding protein, for example, a bacterial albumin binding domain, such as the one of streptococcal protein G (König, T., & Skerra, A. (1998) *J. Immunol. Methods* **218**, 73-83). Other examples of albumin binding peptides that can be used as conjugation partner are,

for instance, those having a Cys-Xaa_1-Xaa_2-Xaa_3-Xaa_4-Cys consensus sequence, wherein Xaa_1 is Asp, Asn, Ser, Thr, or Trp; Xaa_2 is Asn, Gln, His, Ile, Leu, or Lys; Xaa_3 is Ala, Asp, Phe, Trp, or Tyr; and Xaa_4 is Asp, Gly, Leu, Phe, Ser, or Thr as described in US patent application 2003/0069395 or Dennis et al. (Dennis, M. S., Zhang, M., Meng, Y. G., Kadkhodayan, M., Kirchhofer, D., Combs, D. & Damico, L. A. (2002) *J Biol Chem* **277**, 35035-35043).

[00104] In other embodiments, albumin itself or a biological active fragment of albumin can be used as conjugation partner of a lipocalin mutein of the invention. The term "albumin" includes all mammal albumins such as human serum albumin or bovine serum albumin or rat albumine. The albumin or fragment thereof can be recombinantly produced as described in US patent 5,728,553 or European patent applications EP 0 330 451 and EP 0 361 991. Recombinant human albumin (Recombumin®) Novozymes Delta Ltd. (Nottingham, UK) can be conjugated or fused to a lipocalin mutein in order to extend the half-life of the mutein.

[00105] If the albumin-binding protein is an antibody fragment it may be a domain antibody. Domain Antibodies (dAbs) are engineered to allow precise control over biophysical properties and *in vivo* half-life to create the optimal safety and efficacy product profile. Domain Antibodies are for example commercially available from Domantis Ltd. (Cambridge, UK and MA, USA).

[00106] Using transferrin as a moiety to extend the serum half-life of the muteins of the invention, the muteins can be genetically fused to the N or C terminus, or both, of non-glycosylated transferrin. Non-glycosylated transferrin has a half-life of 14-17 days, and a transferrin fusion protein will similarly have an extended half-life. The transferrin carrier also provides high bioavailability, biodistribution and circulating stability. This technology is commercially available from BioRexis (BioRexis Pharmaceutical Corporation, PA, USA). Recombinant human transferrin (DeltaFerrin™) for use as a protein stabilizer/half-life extension partner is also commercially available from Novozymes Delta Ltd. (Nottingham, UK).

[00107] If an Fc part of an immunoglobulin is used for the purpose to prolong the serum half-life of the muteins of the invention, the *SynFusion*™ technology, commercially available from Syntonix Pharmaceuticals, Inc (MA, USA), may be used. The use of this Fc-fusion technology allows the creation of longer-acting biopharmaceuticals and may for example consist of two copies of the mutein linked to the Fc region of an antibody to

improve pharmacokinetics, solubility, and production efficiency.

[00108] Yet another alternative to prolong the half-life of a mutein of the invention is to fuse to the N-or C-terminus of a mutein of the invention long, unstructured, flexible glycine-rich sequences (for example poly-glycine with about 20 to 80 consecutive glycine residues). This approach disclosed in WO2007/038619, for example, has also been term "rPEG" (recombinant PEG).

[00109] If polyalkylene glycol is used as conjugation partner, the polyalkylene glycol can be substituted, unsubstituted, linear or branched. It can also be an activated polyalkylene derivative. Examples of suitable compounds are polyethylene glycol (PEG) molecules as described in WO 99/64016, in US Patent 6,177,074 or in US Patent 6,403,564 in relation to interferon, or as described for other proteins such as PEG-modified asparaginase, PEG-adenosine deaminase (PEG-ADA) or PEG-superoxide dismutase (see for example, Fuertges et al. (1990) The Clinical Efficacy of Poly(Ethylene Glycol)-Modified Proteins *J. Control. Release* **11**, 139-148). The molecular weight of such a polymer, such as polyethylene glycol, may range from about 300 to about 70.000 Dalton, including, for example, polyethylene glycol with a molecular weight of about 10.000, of about 20.000, of about 30.000 or of about 40.000 Dalton. Moreover, as e.g. described in US patents 6,500,930 or 6,620,413, carbohydrate oligo- and polymers such as starch or hydroxyethyl starch (HES) can be conjugated to a mutein of the invention for the purpose of serum half-life extension.

[00110] If one of the above moieties is conjugated to the human tear lipocalin mutein of the invention, conjugation to an amino acid side chain can be advantageous. Suitable amino acid side chains may occur naturally in the amino acid sequence of human tear lipocalin or may be introduced by mutagenesis. In case a suitable binding site is introduced via mutagenesis, one possibility is the replacement of an amino acid at the appropriate position by a cysteine residue. In one embodiment, such mutation includes at least one of Thr 40→ Cys, Glu 73→ Cys, Arg 90→ Cys, Asp 95→ Cys or Glu 131→ Cys substitution. The newly created cysteine residue at any of these positions can in the following be utilized to conjugate the mutein to moiety prolonging the serum half-life of the mutein, such as PEG or an activated derivative thereof.

[00111] In another embodiment, in order to provide suitable amino acid side chains for conjugating one of the above moieties to the muteins of the invention artificial amino acids may be introduced by mutagenesis. Generally, such artificial amino acids are designed to

be more reactive and thus to facilitate the conjugation to the desired moiety. One example of such an artifical amino acid that may be introduced via an artificial tRNA is para-acetyl-phenylalanine.

[00112] For several applications of the muteins disclosed herein it may be advantageous to use them in the form of fusion proteins. In some embodiments, the inventive human tear lipocalin mutein is fused at its N-terminus or its C-terminus to a protein, a protein domain or a peptide such as a signal sequence and/or an affinity tag.

[00113] For pharmaceutical applications a mutein of the invention may be fused to a fusion partner that extends the *in vivo* serum half-life of the mutein (see again PCT publication WO 2006/56464 where suitable fusion partner are described with references to muteins of human neutrophile gelatinase-associated lipocalin with binding affinity for CTLA-4). Similar to the conjugates described above, the fusion partner may be an Fc part of an immunoglobulin, a CH3 domain of an immunoglobulin, a CH4 domain of an immunoglubolin, albumin, an albumin binding peptide or an albumin binding protein, to name only a few. Again, the albumin binding protein may be a bacterial albumin binding protein or a lipocalin mutein with binding activity for albumin. Accordingly, suitable fusion partners for extending the half-life of a lipocalin mutein of the invention include albumin (Osborn, B.L. et al. (2002) *supra J. Pharmacol. Exp. Ther.* **303**, 540-548), or an albumin binding protein, for example, a bacterial albumin binding domain, such as the one of streptococcal protein G (König, T., & Skerra, A. (1998) *J. Immunol. Methods* **218**, 73-83). The albumin binding peptides described in Dennis et al, supra (2002) or US patent application 2003/0069395 having a Cys-Xaa_1-Xaa_2-Xaa_3-Xaa_4-Cys consensus sequence, wherein Xaa_1 is Asp, Asn, Ser, Thr, or Trp; Xaa_2 is Asn, Gln, His, Ile, Leu, or Lys; Xaa_3 is Ala, Asp, Phe, Trp, or Tyr; and Xaa_4 is Asp, Gly, Leu, Phe, Ser, or Thr can also be used as fusion partner. It is also possible to use albumin itself or a biological active fragment of albumin as fusion partner of a lipocalin mutein of the invention. The term “albumin” includes all mammal albumins such as human serum albumin or bovine serum albumin or rat serum albumin. The recombinant production of albumin or fragments thereof is well known in the art and for example described in US patent 5,728,553, European patent application EP 0 330 451 or EP 0 361 991.

[00114] The fusion partner may confer new characteristics to the inventive lipocalin mutein such as enzymatic activity or binding affinity for other molecules. Examples of suitable fusion proteins are alkaline phosphatase, horseradish peroxidase, gluthation-S-

transferase, the albumin-binding domain of protein G, protein A, antibody fragments, oligomerization domains, lipocalin muteins of same or different binding specificity (which results in the formation of "Duocalins", cf. Schlehuber, S., and Skerra, A. (2001), Duocalins, engineered ligand-binding proteins with dual specificity derived from the lipocalin fold.*Biol. Chem.* **382**, 1335-1342) or toxins.

[00115] In particular, it may be possible to fuse a lipocalin mutein of the invention with a separate enzyme active site such that both "components" of the resulting fusion protein together act on a given therapeutic target. The binding domain of the lipocalin mutein attaches to the disease-causing target, allowing the enzyme domain to abolish the biological function of the target.

[00116] Affinity tags such as the Strep-tag® or Strep-tag® II (Schmidt, T.G.M. et al. (1996) *J. Mol. Biol.* **255**, 753-766), the *myc*-tag, the FLAG-tag, the His_6-tag or the HA-tag or proteins such as glutathione-S-transferase also allow easy detection and/or purification of recombinant proteins are further examples of suitable fusion partners. Finally, proteins with chromogenic or fluorescent properties such as the green fluorescent protein (GFP) or the yellow fluorescent protein (YFP) are suitable fusion partners for a lipocalin mutein of the invention as well.

[00117] The term "fusion protein" as used herein also includes lipocalin muteins according to the invention containing a signal sequence. Signal sequences at the N-terminus of a polypeptide direct this polypeptide to a specific cellular compartment, for example the periplasm of *E. coli* or the endoplasmatic reticulum of eukaryotic cells. A large number of signal sequences is known in the art. An illustrative signal sequence for secretion a polypeptide into the periplasm of *E. coli* is the OmpA-signal sequence.

[00118] The present invention also relates to nucleic acid molecules (DNA and RNA) that include nucleotide sequences coding for muteins as described herein. Since the degeneracy of the genetic code permits substitutions of certain codons by other codons specifying the same amino acid, the invention is not limited to a specific nucleic acid molecule encoding a mutein of the invention but encompasses all nucleic acid molecules that include nucleotide sequences encoding a functional mutein.

[00119] Therefore, the present invention also includes a nucleic acid sequence encoding a mutein according to the invention that has a mutation at at least one codon of any of the amino acid sequence positions 26-34, 56-58, 80, 83, 104-106 and 108 of the

linear polypeptide sequence of native mature human tear lipocalin, wherein the codons encoding at least one of the cysteine residues at sequence positions 61 and 153 of the linear polypeptide sequence of the mature human tear lipocalin have been mutated to encode any other amino acid residue.

[00120] The invention as disclosed herein also includes nucleic acid molecules encoding tear lipocalin muteins, which include additional mutations outside the indicated sequence positions of experimental mutagenesis. Such mutations are often tolerated or can even prove to be advantageous, for example if they contribute to an improved folding efficiency, serum stability, thermal stability or ligand binding affinity of the mutein.

[00121] A nucleic acid molecule disclosed in this application may be "operably linked" to a regulatory sequence (or regulatory sequences) to allow expression of this nucleic acid molecule.

[00122] A nucleic acid molecule, such as DNA, is referred to as "capable of expressing a nucleic acid molecule" or capable "to allow expression of a nucleotide sequence" if it includes sequence elements which contain information regarding to transcriptional and/or translational regulation, and such sequences are "operably linked" to the nucleotide sequence encoding the polypeptide. An operable linkage is a linkage in which the regulatory sequence elements and the sequence to be expressed are connected in a way that enables gene expression. The precise nature of the regulatory regions necessary for gene expression may vary among species, but in general these regions include a promoter which, in prokaryotes, contains both the promoter *per se*, i.e. DNA elements directing the initiation of transcription, as well as DNA elements which, when transcribed into RNA, will signal the initiation of translation. Such promoter regions normally include 5' non-coding sequences involved in initiation of transcription and translation, such as the -35/-10 boxes and the Shine-Dalgarno element in prokaryotes or the TATA box, CAAT sequences, and 5'-capping elements in eukaryotes. These regions can also include enhancer or repressor elements as well as translated signal and leader sequences for targeting the native polypeptide to a specific compartment of a host cell.

[00123] In addition, the 3' non-coding sequences may contain regulatory elements involved in transcriptional termination, polyadenylation or the like. If, however, these termination sequences are not satisfactory functional in a particular host cell, then they may be substituted with signals functional in that cell.

[00124] Therefore, a nucleic acid molecule of the invention can include a regulatory

sequence, such as a promoter sequence. In some embodiments a nucleic acid molecule of the invention includes a promoter sequence and a transcriptional termination sequence. Suitable prokaryotic promoters are, for example, the *tet* promoter, the *lac*UV5 promoter or the T7 promoter. Examples of promoters useful for expression in eukaryotic cells are the SV40 promoter or the CMV promoter.

[00125] The nucleic acid molecules of the invention can also be part of a vector or any other kind of cloning vehicle, such as a plasmid, a phagemid, a phage, a baculovirus, a cosmid or an artificial chromosome.

[00126] In one embodiment, the nucleic acid molecule is included in a phasmid. A phasmid vector denotes a vector encoding the intergenic region of a temperent phage, such as M13 or f1, or a functional part thereof fused to the cDNA of interest. After superinfection of the bacterial host cells with such an phagemid vector and an appropriate helper phage (e.g. M13K07, VCS-M13 or R408) intact phage particles are produced, thereby enabling physical coupling of the encoded heterologous cDNA to its corresponding polypeptide displayed on the phage surface (see e.g. Lowman, H.B. (1997) *Annu. Rev. Biophys. Biomol. Struct.* **26,** 401-424, or Rodi, D.J., and Makowski, L. (1999) *Curr. Opin. Biotechnol.* **10,** 87-93).

[00127] Such cloning vehicles can include, aside from the regulatory sequences described above and a nucleic acid sequence encoding a lipocalin mutein of the invention, replication and control sequences derived from a species compatible with the host cell that is used for expression as well as selection markers conferring a selectable phenotype on transformed or transfected cells. Large numbers of suitable cloning vectors are known in the art, and are commercially available.

[00128] The DNA molecule encoding lipocalin muteins of the invention, and in particular a cloning vector containing the coding sequence of such a lipocalin mutein can be transformed into a host cell capable of expressing the gene. Transformation can be performed using standard techniques. Thus, the invention is also directed to a host cell containing a nucleic acid molecule as disclosed herein.

[00129] The transformed host cells are cultured under conditions suitable for expression of the nucleotide sequence encoding a fusion protein of the invention. Suitable host cells can be prokaryotic, such as *Escherichia coli* (*E. coli*) or *Bacillus subtilis*, or eukaryotic, such as *Saccharomyces cerevisiae*, *Pichia pastoris*, SF9 or High5 insect cells, immortalized mammalian cell lines (e.g. HeLa cells or CHO cells) or primary mammalian

cells.

[00130] The invention also relates to a method for the production of a mutein of the invention, wherein the mutein, a fragment of the mutein or a fusion protein of the mutein and another polypeptide is produced starting from the nucleic acid coding for the mutein by means of genetic engineering methods. The method can be carried out *in vivo*, the mutein can for example be produced in a bacterial or eucaryotic host organism and then isolated from this host organism or its culture. It is also possible to produce a protein in vitro, for example by use of an *in vitro* translation system.

[00131] When producing the mutein *in vivo* a nucleic acid encoding a mutein of the invention is introduced into a suitable bacterial or eukaryotic host organism by means of recombinant DNA technology (as already outlined above). For this purpose, the host cell is first transformed with a cloning vector that includes a nucleic acid molecule encoding a mutein of the invention using established standard methods. The host cell is then cultured under conditions, which allow expression of the heterologous DNA and thus the synthesis of the corresponding polypeptide. Subsequently, the polypeptide is recovered either from the cell or from the cultivation medium.

[00132] In some tear lipocalin muteins of the invention, the naturally occurring disulfide bond between Cys 61 and Cys 153 is removed. Accordingly, such muteins (or any other tear lipocalin mutein that does not include an intramolecular disulfide bond) can be produced in a cell compartment having a reducing redox milieu, for example, in the cytoplasma of Gram-negative bacteria. In case a lipocalin mutein of the invention includes intramolecular disulfide bonds, it may be desired to direct the nascent polypeptide to a cell compartment having an oxidizing redox milieu using an appropriate signal sequence. Such an oxidizing environment may be provided by the periplasm of Gram-negative bacteria such as *E. coli*, in the extracellular milieu of Gram-positive bacteria or in the lumen of the endoplasmatic reticulum of eukaryotic cells and usually favors the formation of structural disulfide bonds. It is, however, also possible to produce a mutein of the invention in the cytosol of a host cell, such as *E. coli*. In this case, the polypeptide can either be directly obtained in a soluble and folded state or recovered in form of inclusion bodies, followed by renaturation *in vitro*. A further option is the use of specific host strains having an oxidizing intracellular milieu, which may thus allow the formation of disulfide bonds in the cytosol (Venturi M, et al. (2002) *J. Mol. Biol.* **315**, 1-8).

[00133] However, a mutein of the invention may not necessarily be generated or

produced only by use of genetic engineering. Rather, a lipocalin mutein can also be obtained by chemical synthesis such as Merrifield solid phase polypeptide synthesis or by *in vitro* transcription and translation. It is for example possible that promising mutations are identified using molecular modeling and then to synthesize the wanted (designed) polypeptide *in vitro* and investigate the binding activity for a given target. Methods for the solid phase and/or solution phase synthesis of proteins are well known in the art (see e.g. Bruckdorfer, T. et al. (2004) *Curr. Pharm. Biotechnol.* **5**, 29-43).

[00134] In another embodiment, the muteins of the invention may be produced by *in vitro* transcription/translation employing well-established methods known to those skilled in the art.

[00135] The invention also relates to a pharmaceutical composition, which includes at least one inventive mutein of human tear lipocalin or a fusion protein or conjugate thereof and a pharmaceutically acceptable excipient.

[00136] The lipocalin muteins according to the invention can be administered via any parenteral or non-parenteral (enteral) route that is therapeutically effective for proteinaceous drugs. The muteins of the invention can be administered systemically or topically in formulations containing conventional non-toxic pharmaceutically acceptable excipients or carriers, additives and vehicles as desired.

[00137] In one embodiment of the present invention the pharmaceutical is administered parenterally to a mammal, and in particular to humans. The pharmaceutical composition may be an aqueous solution, an oil-in water emulsion or a water-in-oil emulsion.

[00138] In this regard it is noted that transdermal delivery technologies, e.g. iontophoresis, sonophoresis or microneedle-enhanced delivery, as described in Meidan VM and Michniak BB (2004) Am. J. Ther. 11, 4, 312-316, can also be used for transdermal delivery of the muteins described herein. The muteins of the invention can be administered systemically or topically in formulations containing a variety of conventional non-toxic pharmaceutically acceptable excipients or carriers, additives, and vehicles.

[00139] The dosage of the mutein applied may vary within wide limits to achieve the desired preventive effect or therapeutic response. It will, for instance, depend on the affinity of the compound for a chosen ligand as well as on the half-life of the complex between the mutein and the ligand *in vivo*. Further, the optimal dosage will depend on the

biodistribution of the mutein or its fusion protein or its conjugate, the mode of administration, the severity of the disease/disorder being treated as well as the medical condition of the patient. For example, when used in an ointment for topical applications, a high concentration of the tear lipocalin mutein can be used. However, if wanted, the mutein may also be given in a sustained release formulation, for example liposomal dispersions or hydrogel-based polymer microspheres, like PolyActive™ or OctoDEX™ (cf. Bos et al., Business Briefing: Pharmatech 2003, 1-6). Other sustained release formulations available are for example PLGA based polymers (PR pharmaceuticals), PLA-PEG based hydrogels (Medincell) and PEA based polymers (Medivas).

[00140] Accordingly, the muteins of the present invention can be formulated into compositions using pharmaceutically acceptable ingredients as well as established methods of preparation. The pharmaceutical composition may also contain additives, such as, for example, fillers, binders, wetting agents, glidants, stabilizers, preservatives, emulsifiers, and furthermore solvents or solubilizers or agents for achieving a depot effect. The latter is that fusion proteins may be incorporated into slow or sustained release or targeted delivery systems, such as liposomes and microcapsules.

[00141] The formulations can be sterilized by numerous means, including filtration through a bacteria-retaining filter, or by incorporating sterilizing agents in the form of sterile solid compositions which can be dissolved or dispersed in sterile water or other sterile medium just prior to use.

[00142] A lipocalin mutein described herein can be administered to an organism, including a human patient per se, or in a pharmaceutical composition where it may include or be mixed with pharmaceutically active ingredients or suitable carriers or excipient(s). Techniques for formulation and administration of a respective lipocalin mutein composition resemble or are identical to those of low molecular weight compounds well established in the art. Exemplary routes include, but are not limited to, oral, transdermal, and parenteral delivery.

[00143] A composition that includes a lipocalin mutein of the invention may for instance be applied onto the skin or onto a wound. In some embodiments one may administer a lipocalin mutein or a respective composition in a local rather than systemic manner, for example, via injection.

[00144] Pharmaceutical compositions that include a lipocalin mutein of the present invention may be manufactured in a manner that is itself known, e. g., by means of

conventional mixing, dissolving, granulating, dragee-making, levigating, emulsifying, encapsulating, entrapping or lyophilizing processes. A pharmaceutical composition for use in accordance with the present invention thus may be formulated in conventional manner using one or more physiologically acceptable carriers including excipients and auxiliaries that facilitate processing of the hydrogel and/or peptide/peptoid into preparations that can be used pharmaceutically. Proper formulation is dependent upon the route of administration chosen.

[00145] For injection, the lipocalin mutein or a respective composition may be formulated in aqueous solutions, for instance in physiologically compatible buffers such as Hanks's solution, Ringer's solution, or physiological saline buffer. For transmucosal administration, penetrants appropriate to the barrier to be permeated are used in the formulation. Such penetrants are generally known in the art.

[00146] For oral administration, the lipocalin mutein or a respective composition can be formulated readily by combining them with pharmaceutically acceptable carriers well known in the art. Such carriers enable the lipocalin mutein or a respective composition, as well as a pharmaceutically active compound where present, to be formulated as tablets, pills, dragees, capsules, liquids, gels, syrups, slurries, suspensions and the like, for oral ingestion by a patient to be treated. Pharmaceutical preparations for oral use can be obtained by adding a solid excipient, optionally grinding a resulting mixture, and processing the mixture of granules, after adding suitable auxiliaries, if desired, to obtain tablets or dragee cores. Suitable excipients are, in particular, fillers such as sugars, including lactose, sucrose, mannitol, or sorbitol; cellulose preparations such as, for example, maize starch, wheat starch, rice starch, potato starch, gelatine, gum tragacanth, methyl cellulose, hydroxypropylmethyl-cellulose, sodium carboxymethylcellulose, and/or polyvinylpyrrolidone (PVP). If desired, disintegrating agents may be added, such as the cross-linked polyvinyl pyrrolidone, agar, or alginic acid or a salt thereof such as sodium alginate.

[00147] Dragee cores are provided with suitable coatings. For this purpose, concentrated sugar solutions may be used, which may optionally contain gum arabic, talc, polyvinyl pyrrolidone, carbopol gel, polyethylene glycol, and/or titanium dioxide, lacquer solutions, and suitable organic solvents or solvent mixtures. Dyestuffs or pigments may be added to the tablets or dragee coatings for identification or to characterize different combinations of active compound doses.

[00148] Pharmaceutical preparations that can be used orally include push-fit capsules made of gelatine, as well as soft, sealed capsules made of gelatine and a plasticizer, such as glycerol or sorbitol. The push-fit capsules can contain the active ingredients in admixture with filler such as lactose, binders such as starches, and/or lubricants such as talc or magnesium stearate and, optionally, stabilizers. In soft capsules, the peptides/peptoids may be suspended in suitable liquids, such as fatty oils, liquid paraffin, or liquid polyethylene glycols. In addition, stabilizers may be added. All formulations for oral administration should be in dosages suitable for such administration.

[00149] The lipocalin mutein may be formulated for parenteral administration by injection, e.g., by intramuscular injections or bolus injection or continuous infusion. Formulations for injection may be presented in unit dosage form, e. g., in ampules or in multi-dose containers, with an added preservative. The respective compositions may take such forms as suspensions, solutions or emulsions in oily or aqueous vehicles, and may contain formulatory agents such as suspending, stabilizing and/or dispersing agents.

[00150] The subject in need of such a treatment may be a mammal, such as a human, a dog, a mouse, a rat, a pig, an ape such as cymologous to name only a few illustrative examples. A mutein of the present invention can be use to treat any disease or disorder that involves IL-4 receptor alpha, in the development of such disease or disorder can be displayed to the expression product of a nucleic acid library of the present invention or displayed to otherwise obtained muteins of tear lipocalin.

[00151] In this context it is noted that a variety of tumor cells express a greater number of high affinity IL-4 receptors than normal cells. Such cells include solid human tumor such as melanoma, breast cancer, ovarian carcinoma, mesothelioma, glioblastoma, astrocytoma, renal cell carcinoma, head and neck carcinoma, AIDS associated Kaposi‘s sarcoma = AIDS KS, hormone dependent and independent prostate carcinoma cells, and primary cultures from prostate tumors, for example (cf. Garland, L, et al., (2005) Journal of Immunotherapy 28, 4, 376-381, Rand, RW, et al. Clinical Cancer Research. (2000) 6, 2157-2165; Husain SR, et al. (1999) Nature Medicine 5, 817-822; Puri RK, et al. Cancer Research (1996) 56, 5631-5637; Debinski W, et al, or Husain SR, et al. Cancer Research (1998) 58, 3649-3653, Kawakami K, et al. (2000) Cancer Research, 60, 2981-2987; or Strome SE, et al. Clinical Cancer Research (2002) 8, 281-286, for example. Specific examples of cells with documents overexpression of IL-4 receptors include, but are not limited to, Burkitt lymphoma cell line Jijoye (B-cell lymphom), prostate carcinoma (LNCaP,

DU145), head and neck carcinoma (SCC, KCCT873), Pranceatic cancer (PANC-1 cell line), SCC-25: 13.000 (+/-500) h head and neck cancer cell line (ATCC). IL4R alpha chain plays a major role in IL4-internalization. Accordingly, when fused or conjugated to a toxin, the tear lipocalin muteins binding to IL-4 Receptor alpha chain can therefore also be used for the treatment of tumors (cancer). Examples of suitable toxins include Pseudomonas exotoxin, pertussis-toxin, diphtheria toxin, ricin, saporin, pseudomonas exotoxin, calicheamicin or a derivative thereof, a taxoid, a maytansinoid, a tubulysin and a dolastatin analogue. Examples of dolastatin analogues include, but are not limited to, auristatin E, monomethylauristatin E, auristatin PYE and auristatin PHE.

[00152] For the treatment of cancer, it is also possible to conjugate muteins binding to IL-4 Receptor alpha chain to a cystostatic agent. Examples of such cystostatic agents include Cisplatin, Carboplatin, Oxaliplatin, 5-Fluorouracil, Taxotere (Docetaxel), Paclitaxel, Anthracycline (Doxorubicin), Methotrexate, Vinblastin, Vincristine, Vindesine, Vinorelbine, Dacarbazine, Cyclophosphamide, Etoposide, Adriamycine, Camptotecine, Combretatastin A-4 related compounds, sulfonamides, oxadiazolines, benzo[b]thiophenessynthetic spiroketal pyrans, monotetrahydrofuran compounds, curacin and curacin derivatives, methoxyestradiol derivatives and Leucovorin.

[00153] As is evident from the above disclosure, a mutein of the present invention or a fusion protein or a conjugate thereof can be employed in many applications. In general, such a mutein can be used in all applications antibodies are used, except those with specifically rely on the glycosylation of the Fc part.

[00154] Therefore, in another aspect of the invention, the invented muteins of human tear lipocalin are used for the detection of a given non-natural ligand of human tear lipocalin. Such use may include the steps of contacting the mutein with a sample suspected of containing the given ligand under suitable conditions, thereby allowing formation of a complex between the mutein and the given ligand, and detecting the complexed mutein by a suitable signal.

[00155] The detectable signal can be caused by a label, as explained above, or by a change of physical properties due to the binding, i.e. the complex formation, itself. One example is plasmon surface resonance, the value of which is changed during binding of binding partners from which one is immobilized on a surface such as a gold foil.

[00156] The muteins of human tear lipocalin disclosed herein may also be used for the separation of a given non-natural ligand of human tear lipocalin. Such use may include the

steps of contacting the mutein with a sample supposed to contain said ligand under suitable conditions, thereby allowing formation of a complex between the mutein and the given ligand, and separating the mutein/ligand complex from the sample.

[00157] In both the use of the mutein for the detection of a given non-natural ligand as well as the separation of a given ligand, the mutein and/or the target may be immobilized on a suitable solid phase.

[00158] The human tear lipocalin muteins of the invention may also be used to target a compound to a preselected site. For such a purpose the mutein is contacted with the compound of interest in order to allow complex formation. Then the complex that includes the mutein and the compound of interest are delivered to the preselected site. This use is in particular suitable, but not restricted to, for delivering a drug (selectively) to a preselected site in an organism, such as an infected body part, tissue or organ which is supposed to be treated with the drug. Besides formation of a complex between mutein and compound of interest, the mutein can also be reacted with the given compound to yield a conjugate of mutein and compound. Similar to the above complex, such a conjugate may be suitable to deliver the compound to the preselected target site. Such a conjugate of mutein and compound may also include a linker that covalently links mutein and compound to each other. Optionally, such a linker is stable in the bloodstream but is cleavable in a cellular environment.

[00159] The muteins disclosed herein and its derivatives can thus be used in many fields similar to antibodies or fragments thereof. In addition to their use for binding to a support, allowing the target of a given mutein or a conjugate or a fusion protein of this target to be immobilized or separated, the muteins can be used for labeling with an enzyme, an antibody, a radioactive substance or any other group having biochemical activity or defined binding characteristics. By doing so, their respective targets or conjugates or fusion proteins thereof can be detected or brought in contact with them. For example, muteins of the invention can serve to detect chemical structures by means of established analytical methods (e.g. ELISA or Western Blot) or by microscopy or immunosensorics. Here, the detection signal can either be generated directly by use of a suitable mutein conjugate or fusion protein or indirectly by immunochemical detection of the bound mutein via an antibody.

[00160] Numerous possible applications for the inventive muteins also exist in medicine. In addition to their use in diagnostics and drug delivery, a mutant polypeptide of

the invention, which binds, for example, tissue- or tumor-specific cellular surface molecules can be generated. Such a mutein may, for example, be employed in conjugated form or as a fusion protein for "tumor imaging" or directly for cancer therapy. Accordingly, the present invention provides a diagnostic composition comprising an inventive mutein and means for diagnosis such as excipients, buffers, labels (which can be used to label the muteins).

[00161] Thus, the present invention also involves the use of the human tear lipocalin muteins of the invention for complex formation with a given non-natural ligand.

[00162] Another related use of a mutein described herein is target validation, i.e. the analysis whether a polypeptide assumed to be involved in the development or progress of a disease or disorder is indeed somehow causative of that disease or disorder. This use for validating a protein as a pharmacological drug target takes advantage of the ability of a mutein of the present invention to specifically recognize a surface area of a protein in its native conformation, i.e. to bind to a native epitope. In this respect, it is to be noted that this ability has been reported only for a limited number of recombinant antibodies. However, the use of an inventive mutein for validation of a drug target is not limited to the detection of proteins as targets, but also includes the detection of protein domains, peptides, nucleic acid molecules, organic molecules or metal complexes.

Exemplary Embodiment of the Invention

[00163] **Fig. 1** shows the primary structure of a previously disclosed human tear lipocalin mutein (S191.4-B24) that exhibits binding affinity for IL-4 receptor alpha. The first 21 residues (underlined) constitute the signal sequence, which is cleaved upon periplasmic expression. The N-terminal T7-tag (italic) and the C-terminal Streptag-II (bold) are part of the characterized protein. Fig. 1 also shows that 4 N-terminal amino acid residues (His1 His2 Leu3 Ala4) as well as the two last C-terminal amino acid residues (Ser157 and Asp158) of the wild type tear lipocalin are deleted in this mutein.

[00164] **Figure 2** shows the polypeptide sequences of exemplary muteins with high affinity for IL-4 receptor alpha (SEQ ID Nos: 2-11). Numbers indicated by 'SwissProt P31025' show the corresponding amino acid position numbering of the unprocessed precursor sequence of the entry of the SwissProt database. Wt TLc26 essentially corresponds to the sequence of wild type tear lipocalin in the vector pTLc26. However, wt TLc26 does not include a disulfide bond, since the cystein residues at positions 61 and 153 of the mature protein are replaced by serine residues. Likewise, a serine residue at

positions 101 of the mature protein is replaced by a serine residue. In addition, at position 111 of the mature protein an arginine residue has been replaced by a proline and at position 114 of the mature protein a lysine residue has been replaced by a tryptophan. Furthermore, the two C-terminal amino acids included in the sequence of SwissProt entry P31025 are not included in the sequence. AB4004 is a randomization library disclosed in international patent application WO 2008/015239, in which the mutated positions are indicated in bold. Randomized sequences are identical to J14 with the exception of position 53 and 55. M3-B24(PSM) indicates hot spots from PSM B24.

[00165] **Figure 3** shows the results of TF-1 cell proliferation assays. TF-1 cells were incubated for 1 hour at 37 °C with the indicated muteins of the invention (S276.2 K04, S308.5 F08, S308.5 N01, S308.5 L20, S308.5 L04, S308.5 N20, S308.3 O10, S191.4 B24 [SEQ ID Nos: 2-11]) in a dilution series before addition of 0.8 ng/ml IL-4 (**A**) or 12 ng/ml IL-13 (**B**) for 72 h. Proliferation was measured by ^{3}H-thymidine incorporation.

[00166] **Figure 4** depicts IC_{50} values from Fig. 3 and results of Biacore measurements of the muteins of human tear lipocalin with affinity for IL-4 receptor alpha. Approximately 400 RU of IL-4 receptor alpha-Fc was captured on a CM-5 chip, which had previously been coated with an anti human-Fc monoclonal antibody. Subsequently, mutein in a single concentration of 25 nM was passed over the flowcell and changes in resonance units recorded. Reference signals from a flow cell that was equally treated apart from not having any IL-4 receptor alpha-Fc was subtracted and the resulting data fitted to a 1:1 Langmuir model using the BIAevaluation software. Due to the slow dissociation kinetics of the double referencing was used by subtracting the signals from a flow cell that was equally treated apart from not having any IL-4 receptor alpha-Fc and subtracting the signal from an experiment where only sample buffer was injected. The resulting data was fitted to a 1:1 Langmuir model with mass-transport limitation using the BIAevaluation software.

[00167] Unless otherwise indicated, established methods in the art of recombinant gene technology were used.

Examples

Example 1: Affinity maturation of the mutein S191.4-B24 using a site-directed random approach

[00168] A library of variants based on the mutein S191.4-B24 (SEQ ID NO:2) that is

described in PCT application WO 2008/015239 was designed by randomization of the positions 26, 32, 34, 55, 56, 58 and 63 to allow for all 20 amino acids on these positions. The library was constructed essentially as described in Example 1 of WO 2008/015239.

[00169] Phagemid selection was carried out as described in Example 2 of WO 2008/015239 using limited target concentration (0.5 nM and 0.1 nM of IL-4 receptor alpha, Peprotech) combined with extended washing times together with a competitive monoclonal antibody against IL-4 receptor alpha (MAB230, R&D Systems; 1 hour washing) or short incubation times (10 minutes), respectively. Three or four rounds of selection were performed.

[00170] Preparative production of IL-4 receptor alpha-specific muteins was carried out using *E. coli* K12 strain JM83 harbouring the respective mutein encoded on the expression vector pTLPC10 (SEQ ID No: 1) or, where larger amounts of protein were needed,*E. coli* strain W3110 harbouring the respective expression vector as described in WO 2008/015239.

Example 2: Affinity measurement using Biacore

[00171] Affinity measurements were performed essentially as described in Example 9 of WO 2006/56464 with the modifications that approximately 400 RU of IL-4 receptor alpha-Fc (R&D Systems) was immobilized (instead of 2000 RU of human CTLA-4 or murine CTLA-4-Fc used as target in WO 2006/56464) and 100 µl of mutein was injected at a concentration of 25 nM (instead of 40 µl sample purified lipocalin muteins at concentrations of 5 - 0.3 µM as used in WO 2006/56464).

Example 3: TF-1 cell proliferation assays

[00172] IL-4 and IL-13-stimulated TF-1 cell proliferation assays were performed essentially as described in Lefort et al. (Lefort S., et al., (1995) *FEBS Lett.* **366,** 2-3, 122-126) and as described in Example 10 of PCT application WO 2008/015239. TF-1 cells were incubated for 1 hour at 37 °C with the indicated muteins muteins of the invention (S276.2 K04, S308.5 F08, S308.5 N01, S308.5 L20, S308.5 L04, S308.5 N20, S308.3 O10, S191.4 B24 [SEQ ID Nos: 3-11]) in a dilution series before addition of 0.8 ng/ml IL-4 (a) or 12 ng/ml IL-13 (b) for 72 h. Proliferation was measured by ^{3}H-thymidine incorporation. The results from a TF-1 proliferation assay is depicted in Figure and shows that the high affinity variants S276.2 K04, S308.5 F08, S308.5 N01, S308.5 L20, S308.5 L04, S308.5 N20, and S308.3O10 are potent antagonists of IL-4 as well as IL-13 induced

signalling and proliferation.

[00173] One skilled in the art would readily appreciate that the present invention is well adapted to carry out the objects and obtain the ends and advantages mentioned, as well as those inherent therein. Further, it will be readily apparent to one skilled in the art that varying substitutions and modifications may be made to the invention disclosed herein without departing from the scope and spirit of the invention. The compositions, methods, procedures, treatments, molecules and specific compounds described herein are presently representative of certain embodiments are exemplary and are not intended as limitations on the scope of the invention. Changes therein and other uses will occur to those skilled in the art which are encompassed within the spirit of the invention are defined by the scope of the claims. The listing or discussion of a previously published document in this specification should not necessarily be taken as an acknowledgement that the document is part of the state of the art or is common general knowledge.

[00174] The invention illustratively described herein may suitably be practiced in the absence of any element or elements, limitation or limitations, not specifically disclosed herein. Thus, for example, the terms “comprising”, “including,” containing”, etc. shall be read expansively and without limitation. Additionally, the terms and expressions employed herein have been used as terms of description and not of limitation, and there is no intention in the use of such terms and expressions of excluding any equivalents of the features shown and described or portions thereof, but it is recognized that various modifications are possible within the scope of the invention claimed. Thus, it should be understood that although the present invention has been specifically disclosed by exemplary embodiments and optional features, modification and variation of the inventions embodied therein herein disclosed may be resorted to by those skilled in the art, and that such modifications and variations are considered to be within the scope of this invention.

[00175] The invention has been described broadly and generically herein. Each of the narrower species and subgeneric groupings falling within the generic disclosure also form part of the invention. This includes the generic description of the invention with a proviso or negative limitation removing any subject matter from the genus, regardless of whether or not the excised material is specifically recited herein.

[00176] Other embodiments are within the following claims. In addition, where features

or aspects of the invention are described in terms of Markush groups, those skilled in the art will recognize that the invention is also thereby described in terms of any individual member or subgroup of members of the Markush group.

Claims

1. A mutein of human tear lipocalin having binding affinity to IL-4 receptor alpha, characterised in at least one substitution at a position corresponding to the amino acid position of mature human tear lipocalin selected from the group consisting of: Pro, Leu, Asn, Tyr, Lys or Glu at position 26; Lys at position 32; Trp, Asn, Val, His or Arg at position 34; Leu, Ala, His, Lys or Met at position 56; Leu at position 58; or Lys, Ser, Gln or Pro at position 63, wherein the mutein has a stronger binding affinity than S191.4-B24 (SEQ ID NO:2).

2. The mutein of human tear lipocalin according to claim 1, wherein the mutein comprises:

(i) a mutated amino acid residue at any one or more of the sequence positions 27, 28, 30, 31, 33, 53, 57, 61, 64, 66, 80, 83, 104 -106 and 108 of the linear polypeptide sequence of the mature human tear lipocalin,

(ii) a mutated amino acid residue at any 2 or more of the sequence positions 26, 32, 34, 55, 56, 58 and 63 of the linear polypeptide sequence of the mature human tear lipocalin, wherein the amino acid sequence of the mutein comprises one of the following sets of amino acid combinations:

(1) Ser 26, Glu 34, Leu 55, Lys 58,
(2) Ser 26, Asn 34, Ala 55, Lys 58,
(3) Ser 26, Val 34,
(4) Pro 26, Ser 34,
(5) Pro 26, Ala 55,
(6) Leu 26, Trp 34, Ala 55,
(7) Leu 26, Trp 34, Ile 58,
(8) Asn 26, Asp 34,
(9) Asn 26, Ala 55,
(10) Tyr 26, His 34, Ala 55,
(11) Tyr 26, His 34, Ala 58,
(12) Lys 26, Arg 34, Ala 55,
(13) Lys 26, Arg 34, Asn 58,
(14) Glu 26, Gly 34, Ala 55,
(15) Glu 26, Gly 34, Leu 58.

3. The mutein of claim 1 or 2, further comprising at least one of the following: Ala 31, Tyr 33, Trp 61, Ser 80, Leu 104, Pro 106 and Gln 108.

4. The mutein of any one of claims 1-3, further comprising at least one of the following: Phe 53, Leu 53, Tyr 64, Met 64, Leu 66 and Asp 66.

5. The mutein of any one of claims 1-4, further comprising at least one of the following: Arg 27, Cys 28, Arg 30, Ala 31, Tyr 33, Arg 57, Trp 61, Ser 80, Arg 83, Leu 104, Cys 105, Pro 106 and Gln 108.

6. The mutein of any one of claims 1-5, further comprising one of the following sets of amino acid substitutions:

(1) Phe 53, Tyr 64, Leu 66

(2) Leu 53, Met 64, Asp 66.

7. The mutein of any one of claims 1 or 2, comprising one of the following sets of amino acid substitutions:

Ser 26; Tyr 32; Leu 55; Gln 56; Lys 58;
Pro 26; Tyr 32; Ser 34; Ala 55; Gln 56; Lys 63;
Leu 26; Phe 32; Trp 34; Ala 55; Ile 58; Ser 63;
Ser 26; Tyr 32; Val 34; Ala 55; Ala 56; Ile 58; Ser 63;
Ser 26; Val 32; Asn 34; Ala 55; Gln 56; Lys 58; Lys 63;
Asn 26; Lys 32; Asp 34; Ala 55; His 56; Arg 58; Gln 63;
Tyr 26; Tyr 32; His 34; Ala 55; His 56; Ala 58; Lys 63;
Lys 26; Tyr 32; Arg 34; Ala 55; Lys 56; Asn 58; Pro 63; and
Glu 26; His 32; Gly 34; Ala 55; Met 56; Leu 58; Lys 63.

8. The mutein of any one of claims 1 or 2 as set forth in any one of SEQ ID NOs: 3-11 or a fragment thereof.

9. A method of generating a mutein of human tear lipocalin, wherein the mutein binds to IL 4 receptor alpha and wherein the mutein has a stronger binding affinity than S191.4-B24 (SEQ ID NO:2), comprising:

(a) subjecting a nucleic acid molecule encoding a human tear lipocalin to mutagenesis at

(i) any one or more of the amino acid sequence positions 27, 28, 30, 31, 33, 53, 57, 61, 64, 66, 80, 83, 104 -106 and 108 of the linear polypeptide sequence of mature human tear lipocalin,

(ii) any two or more of the amino acid sequence positions 26, 32, 34, 55, 56, 58 and 63 of the linear polypeptide sequence of the mature human tear lipocalin,

thereby obtaining one or more nucleic acids encoding a mutein of human tear lipocalin, wherein

if mutated at position 26, the mutation comprises Ser, Pro, Leu, Asn, Tyr, Lys or Glu;

if mutated at position 32, the mutation comprises Tyr, Val, Lys or His;

if mutated at position 34, the mutation comprises Glu, Ser, Trp, Asn, Val, Asp, His, Arg or Gly;

if mutated at position 55, the mutation comprises Leu or Ala;

if mutated at position 56, the mutation comprises Gln, Leu, Ala, His, Lys or Met;

if mutated at position 58, the mutation comprises Lys, Ser, Ile, Arg, Ala, Asn or Leu;

if mutated at position 63, the mutation comprises Glu, Lys, Ser, Gln or Pro,

(b) expressing the one or more mutein nucleic acid molecule(s) obtained in (a) in an expression system, thereby obtaining one or more mutein(s), and

(c) enriching the one or more mutein(s) obtained in (b) and selecting muteins that bind to IL 4 receptor alpha with a dissociation constant of less than 10 nM by means of selection and/or isolation.

10. The method according to claim 9, wherein step (c) further comprises:

(ci) providing IL 4 receptor alpha or an immunogenic fragment thereof,

(cii) contacting the one or more muteins with the IL 4 receptor alpha or the immunogenic fragment thereof, thereby allowing the formation of a complex between the IL 4 receptor alpha or the immunogenic fragment thereof and a mutein having binding affinity.

11. The method of claims 9 or 10, wherein the selection in step (c) is carried out under competitive conditions.

12. The method according to any one of claims 9 to 11, wherein the mutein comprises one of the following sets of amino acids:

(1) Ser 26, Glu 34, Leu 55, Lys 58,

(2) Ser 26, Asn 34, Ala 55, Lys 58,

(3) Ser 26, Val 34,

(4) Pro 26, Ser 34,

(5) Pro 26, Ala 55,

(6) Leu 26, Trp 34, Ala 55,

(7) Leu 26, Trp 34, Ile 58,

(8) Asn 26, Asp 34,

(9) Asn 26, Ala 55,
(10) Tyr 26, His 34, Ala 55,
(11) Tyr 26, His 34, Ala 58,
(12) Lys 26, Arg 34, Ala 55,
(13) Lys 26, Arg 34, Asn 58,
(14) Glu 26, Gly 34, Ala 55,
(15) Glu 26, Gly 34, Leu 58.

13. A mutein obtainable or obtained according to any one of claims 9 to 12.

14. A nucleic acid molecule comprising a nucleotide sequence encoding a mutein of any one of claims 1 to 8 and 13.

15. A host cell containing a nucleic acid molecule of claim 14.

16. A pharmaceutical composition comprising a mutein of human tear lipocalin as defined in any one of claims 1 to 8 and 13 and a pharmaceutically acceptable excipient.

17. A diagnostic composition comprising a mutein of human tear lipocalin as defined in any one of claims 1 to 8 and 13 and means for diagnosis.