

CAUSTIC STABLE CHROMATOGRAPHY LIGANDS

Related Applications

[0001] The present application claims the benefit of priority of U.S. Provisional Patent Application No. 61/203,664, filing date December 24, 2008, the entire content of which is incorporated herein in its entirety.

Field of the Invention

[0002] The present invention relates to chromatography ligands having improved caustic stability, *e.g.*, ligands based on immunoglobulin-binding proteins such as, *Staphylococcal* protein A, as well as chromatography matrices comprising such ligands.

Background

[0003] Ligands used in affinity chromatography typically confer a high selectivity for the target molecule, thereby resulting in high yield and fast and economical purification of target molecules. *Staphylococcal* protein A (SpA) based reagents and chromatography matrices have found a widespread use in the field of affinity chromatography for capture and purification of antibodies as well as in antibody detection methods due to its ability to bind IgG without significantly affecting the affinity of the immunoglobulin for antigen.

[0004] Accordingly, various reagents and media comprising protein A-ligands have been developed and are commercially available including, for example, ProSep®-vA High Capacity, ProSep® vA Ultra and ProSep® UltraPlus (Millipore) and Protein A Sepharose™, MabSelect™, MabSelect Xtra™ and MabSelect SuRe® (GE Healthcare) and Poros MabCapture A™ (Applied Biosystems).

[0005] In order to maintain selectivity and binding capacity of the chromatography ligands including, *e.g.*, resins including SpA based chromatography ligands, the ligand bound resins, referred to as chromatography matrices, have to be cleaned and are typically cleaned under alkaline conditions, *e.g.*, with sodium hydroxide. For example, a standard process which is used for cleaning and restoring the matrix is a cleaning-in-place (CIP) alkaline protocol, which typically involves

treatment of the matrix with 1M NaOH, pH 14. However, such harsh treatment is often undesirable, especially, where the ligand is a protein or a protein-based molecule.

Summary of the Invention

[0006] The present invention provides alkaline-stable SpA-based chromatography ligands which, for example, are capable of withstanding repeated cleaning-in-place (CIP) cycles. More specifically, ligands according to the invention are able to withstand conventional alkaline cleaning for a prolonged period of time, which renders the ligands attractive candidates, especially for cost-effective large-scale purification of immunoglobulins.

[0007] In one aspect of the present invention, an alkaline-stable chromatography ligand comprises two or more domains of SpA. For example, in some embodiments according to this aspect, an alkaline-stable chromatography ligand is provided, which comprises two or more B domains or two or more Z domains of Staphylococcus protein A (SpA), or a functional fragment or variant thereof, where the two or more B domains or two or more Z domains are attached to a chromatography resin at more than one site on the resin.

[0008] In some embodiments according to this aspect, the ligand comprises three or more B domains or three or more Z domains of SpA, or a functional fragment or variant thereof, where the three or more B domains or three or more Z domains are attached to a chromatography resin at more than one site on the resin. In some other embodiments, the ligand comprises four or more B domains or four or more Z domains of SpA, where the four or more B domains or four or more Z domains are attached to a chromatography resin at more than one site on the resin. In yet other embodiments, the ligand comprises five or more B domains or five or more Z domains of SpA; or six or more B domains or six or more Z domains of SpA; or seven or more B domains or seven or more Z domains of SpA, where the five or more B domains, or five or more Z domains, or six or more B domains, or six or more Z domains, or seven or more B domains or seven or more Z domains are attached to a chromatography resin at more than one site on the resin.

[0009] In another aspect according to the present invention, an alkaline-stable chromatography ligand comprises one or more isolated E, D, A, B, C or Z domains of *Staphylococcus* protein A, where the one or more isolated domains comprise one or more

amino acid residues at position n+1 mutated to any naturally occurring amino acid except cysteine (C), serine (S), alanine (A), glycine (G), asparagine (N), or glutamine (Q). In some embodiments, n represents the asparagine residue at position 23 of an isolated SpA domain. Exemplary ligands having a mutation at position 24 of an isolated SpA domain where n represents an asparagine are shown in Table I.

Table I

SpA domains including modifications	Designation
E, D, A, B, C or Z domain-glutamic acid 24 or aspartic acid 24 replaced with methionine	E-D24M D-E24M A-E24M B-E24M C-E24M Z-E24M
E, D, A, B, C or Z domain-glutamic acid 24 or aspartic acid 24 replaced with isoleucine	E-D24I D-E24I A-E24I B-E24I C-E24I Z-E24I
E, D, A, B, C or Z domain-glutamic acid 24 or aspartic acid 24 replaced with phenylalanine	E-D24F D-E24F A-E24F B-E24F C-E24F Z-E24F
E, D, A, B, C or Z domain-glutamic acid 24 or aspartic acid 24 replaced with threonine	E-D24T D-E24T A-E24T B-E24T C-E24T Z-E24T
E, D, A, B, C or Z domain-glutamic acid 24 or aspartic acid 24 replaced with proline	E-D24P D-E24P A-E24P B-E24P C-E24P Z-E24P
E, D, A, B, C or Z domain-glutamic acid 24 or aspartic acid 24 replaced with tryptophan	E-D24W D-E24W A-E24W B-E24W

	C-E24W Z-E24W
E, D, A, B, C or Z domain-glutamic acid 24 or aspartic acid 24 replaced with arginine	E-D24R D-E24R A-E24R B-E24R C-E24R Z-E24R

E, D, A, B, C or Z domain-glutamic acid 24 or aspartic acid 24 replaced with valine	E-D24V D-E24V A-E24V B-E24V C-E24V Z-E24V
E, D, A, B, C or Z domain-glutamic acid 24 or aspartic acid 24 replaced with leucine	E-D24L D-E24L A-E24L B-E24L C-E24L Z-E24L
E, D, A, B, C or Z domain-glutamic acid 24 or aspartic acid 24 replaced with tyrosine	E-D24Y D-E24Y A-E24Y B-E24Y C-E24Y Z-E24Y
E, D, A, B, C or Z domain-glutamic acid 24 or aspartic acid 24 replaced with histidine	E-D24H D-E24H A-E24H B-E24H C-E24H Z-E24H
E, D, A, B, C or Z domain-glutamic acid 24 or aspartic acid 24 replaced with lysine	E-D24K D-E24K A-E24K B-E24K C-E24K Z-E24K
E, D, A, B, C or Z domain-glutamic acid 24 or aspartic acid 24 replaced with glutamic acid	D-E24D A-E24D B-E24D C-E24D Z-E24D

[0010] The single letter codes for the naturally occurring amino acids as well as the corresponding three letter codons encoding for each amino acid are depicted in Table II. In general, due to the degeneracy of the codon, more than one three letter codon can encode for the same amino acid.

Table II

	T	C	A	G
T	TTT Phe (F) TTC " TTA Leu (L) TTG "	TCT Ser (S) TCC " TCA " TCG "	TAT Tyr (Y) TAC TAA Stop TAG Stop	TGT Cys (C) TGC TGA Stop TGG Trp (W)
C	CTT Leu (L) CTC " CTA " CTG "	CCT Pro (P) CCC " CCA " CCG "	CAT His (H) CAC " CAA Gln (Q) CAG "	CGT Arg (R) CGC " CGA " CGG "
A	ATT Ile (I) ATC " ATA " ATG Met (M)	ACT Thr (T) ACC " ACA " ACG "	AAT Asn (N) AAC " AAA Lys (K) AAG "	AGT Ser (S) AGC " AGA Arg (R) AGG "
G	GTT Val (V) GTC " GTA " GTG "	GCT Ala (A) GCC " GCA " GCG "	GAT Asp (D) GAC " GAA Glu (E) GAG "	GGT Gly (G) GGC " GGA " GGG "

[0011] Also encompassed by the present invention is a chromatography matrix comprising a ligand according to one or more aspects of the invention coupled to a solid support such as, *e.g.*, at least one insoluble carrier.

[0012] Additionally, provided herein are methods of using the ligands described herein. Accordingly, a method of affinity purifying one or more target molecules (*e.g.*, immunoglobulins) from a sample is provided, where the method comprising the steps of: (a) providing a sample comprising one or more target molecules (*e.g.*, immunoglobulins); (b) contacting the sample with a matrix according to the invention under conditions such that the one or more target molecules (*e.g.*, immunoglobulins) bind to the matrix; and (c)

recovering the one or more bound target molecules (*e.g.*, immunoglobulins) by eluting under suitable conditions such as, for example, a suitable pH.

[0013] In some embodiments, an alkaline-stable chromatography ligand according to the present invention retains at least 95% of its binding capacity after 5 hours, or after 10 hours, after 15 hours, or after 20 hours, or after 25 hours, or after 30 hours of incubation in 0.5 M NaOH.

[0014] The immunoglobulins which are capable of being bound by the various ligands described herein include, *e.g.*, IgG, IgA and IgM, or any fusion protein comprising antibody and any fragment of antibody.

[0015] Also provided herein are nucleic acid molecules encoding the various ligands described herein, as well as host cells including such nucleic acid molecules. In some embodiments, a host cell is a prokaryotic cell. In other embodiments, a host cell is a eukaryotic cell.

[0016] Additionally, the present invention encompasses a library of polypeptides comprising one or more ligands described herein, and functional fragments and variants thereof. In yet another embodiment, the present invention provides a library of nucleic acid molecules encoding one or more ligands encompassed by the present invention or encoding functional fragments and variants thereof.

[0017] In some embodiments, the present invention provides SpA based ligands which exhibit altered (increased or decreased) binding to a Fab portion of an immunoglobulin compared to the previously known SpA ligands, while retaining the ability to bind the Fc portion of the immunoglobulin. In one embodiment, an SpA based ligand according to the present invention exhibits decreased binding to a Fab portion of an immunoglobulin compared to wild type SpA. In an exemplary embodiment, an alkaline-stable chromatography ligand according to the present invention further includes the amino acid glycine at position 29 replaced with an alanine. In another embodiment, an alkaline stable chromatography ligand according to the present invention further includes the glycine at position 29 replaced with an amino acid other than alanine or tryptophan.

Brief Description of the Drawings

[0018] FIG. 1 depicts the nucleic acid sequences for the wild type (wt) IgG binding domains of SpA, represented by SEQ ID NOs:1-5. SEQ ID NO:1 represents

the nucleic acid sequence for the wt E domain; SEQ ID NO: 2 represents the nucleic acid sequence for the wt D domain; SEQ ID NO: 3 represents the nucleic acid sequence for the wt A domain; SEQ ID NO: 4 represents the nucleic acid sequence for the wt B domain; and SEQ ID NO: 5 represents the nucleic acid sequence for the wt C domain.

[0019] FIG. 2 depicts the nucleic acid sequence for the Z domain of SpA, represented by SEQ ID NO: 6.

[0020] FIG. 3 depicts the amino acid sequence alignments of the wild type (wt) IgG binding domains of SpA (E, D, A, B and C). SEQ ID NO: 7 represents the amino acid sequence of the wt E domain; SEQ ID NO: 8 represents the amino acid sequence of the wt D domain; SEQ ID NO: 9 represents the amino acid sequence of the wt A domain; SEQ ID NO: 10 represents the amino acid sequence of the wt B domain; and SEQ ID NO: 11 represents the amino acid sequence of the wt C domain.

[0021] FIG. 4 depicts the amino acid sequence of the Z domain, represented by SEQ ID NO: 12.

[0022] FIG. 5 depicts a schematic of the plasmid pJ56:8620.

[0023] FIG. 6 depicts the amino acid sequences for the his-tagged wt B domain of SpA as well as the various his-tagged n+1 mutants. SEQ ID NO: 13 represents the amino acid sequence for the his-tagged wt B domain of SpA and SEQ ID NOs: 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25 and 26 represent the amino acid sequences for the n+1 B domain mutants E24M, E24I, E24F, E24T, E24P, E24W, E24R, E24V, E24L, E24Y, E24H, E24K and E24D, respectively.

[0024] FIG. 7 depicts the nucleic acid sequences for the his-tagged wt B domain of SpA as well as nucleic acid sequences encoding the various his-tagged n+1 mutants. SEQ ID NO: 27 represents the nucleic acid sequence encoding the his-tagged wt B domain of SpA and SEQ ID NOs: 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38, 39 and 40 represent the nucleic acid sequences encoding the n+1 B domain mutants E24M, E24I, E24F, E24T, E24P, E24W, E24R, E24V, E24L, E24Y, E24H, E24K and E24D, respectively.

[0025] FIG. 8 depicts a bar graph summarizing the results of an exemplary experiment to assay for residual IgG binding in the various his-tagged n+1 B domain mutants, where n represents the asparagine at position 23, using the high-throughput ELISA based assay described herein. The X-axis represents various n+1 B domain mutants having a mutation at position 24 and the Y-axis represents the percent IgG

binding remaining after a six hour exposure to 1N NaOH. As shown in the graph, n+1 B domain mutants having amino acids which contain bulky side chains at position 24 exhibit increased caustic stability relative to the wt B domain.

[0026] FIG. 9 depicts the results of an exemplary experiment to assay for the percent retained IgG binding capacity of the various SpA domain trimers (*i.e.*, EEE, DDD, AAA, BBB, CCC and ZZZ) subsequent to their multipoint attachment to an agarose resin. nPrA refers to the wt SpA. The X-axis represents the number of cycles of caustic exposure, where each cycle consists of a 15 minute exposure to 0.5 N NaOH. The Y-axis represents the retained IgG binding capacity of the attached SpA ligands. As shown in the graph in FIG. 9, the ligands including C or Z domain trimers are roughly equivalent in their caustic stability and are more caustic stable than the ligands including B domain trimers, which are more caustic stable than the ligands including A domain trimers, which are more caustic stable than the ligands including E or D domain trimers.

[0027] FIG. 10 depicts the results of an exemplary experiment to assay for the percent retained IgG binding capacity of attached ligands according to the invention, containing three (B3), or four (B4), or five (B5) B domains, attached via multipoint attachment to an agarose resin. Also, attached ligands containing five B domains or seven B domains and additionally including a mutation to reduce Fab binding (G29A) are used, referred to as B5-NF and B7-NF, respectively. nPrA refers to the wt SpA. The X-axis represents the number of cycles of caustic exposure, where each cycle consists of a 15 minute exposure to 0.5 N NaOH. The Y-axis represents the retained IgG binding capacity of the attached SpA ligands. As shown in the graph in FIG. 10, the level or extent of caustic stability is directly proportional to the number of B domains in the ligands and is not altered by the G29A mutation to decrease Fab binding.

Detailed Description of the Invention

[0028] The present invention provides SpA based alkaline-stable chromatography ligands, and in particular, ligands based on one or more domains of SpA. Previously described exemplary SpA based alkaline-stable chromatography ligands include, for example, those described in International PCT patent application no. WO2008/039141, which discusses alkaline-stable chromatography ligands based on the C domain of SpA which are capable of binding the Fab portions of antibodies

and are coupled to an insoluble carrier at a single site using a terminal coupling group; and those described in U.S. Patent No. 6,831,161, which discusses SpA based alkaline based chromatography ligands where one or more asparagine amino acid residues have modified.

[0029] In order that the present disclosure may be more readily understood, certain terms are first defined. Additional definitions are set forth throughout the detailed description.

I. Definitions

[0030] As used herein, the term "SpA" or "protein A of *Staphylococcus aureus*," refers to a 42Kda multi-domain protein isolated from the bacterium *Staphylococcus aureus*. SpA is bound to the bacterial cell wall via its carboxy-terminal cell wall binding region, referred to as the X domain. At the amino-terminal region, it includes five immunoglobulin-binding domains, referred to as E, D, A, B, and C (Sjodhal, *Eur J Biochem.* Sep 78(2):471-90 (1977); Uhlen *et al.*, *J Biol Chem.* Feb 259(3):1695-702 (1984). Each of these domains contains approximately 58 amino acid residues, and they share 65-90% amino acid sequence identity. The Z domain of SpA is an engineered analogue of the B domain of SpA and includes an alanine instead of a glycine residue at position 29 (Nilsson, *et al.*, *Protein engineering*, Vol. 1, No. 2, 107-113, 1987.). Each of the E, D, A, B and C domains of SpA possess distinct Ig-binding sites. One site is for Fc γ (the constant region of IgG class of Ig) and the other is for the Fab portion of certain Ig molecules (the portion of the Ig that is responsible for antigen recognition). It has been reported that each of the domains contains a Fab binding site. The non-Ig binding portion of SpA is located at the C-terminus and is designated the X region or X-domain.

[0031] The cloning of the gene encoding SpA is described in U.S. Patent No. 5,151,350, the entire contents of which are incorporated by reference herein in their entirety.

[0032] The present invention provides SpA-based alkaline-stable chromatography ligands. In some aspects according to the present invention, an alkaline stable chromatography ligand comprises two or more, or three or more, or four or more, or five or more, or six or more, or seven or more of isolated wt B or Z

domain of SpA. In other aspects according to the present invention, an alkaline stable chromatography ligand comprises one or more isolated E, D, A, B, C or Z domains of SpA, where the one or more isolated domains comprise one or more amino acid residues at position n+1 mutated to an amino acid selected from the group consisting of tryptophan, arginine, threonine, isoleucine, valine and methionine, wherein n represents an asparagine.

[0033] In a particular embodiment, the present invention provides a chromatography ligand comprising two or more B domains of SpA, attached to a chromatography resin at more than one site on the resin. In another embodiment, the present invention provides a chromatography ligand comprising two or more Z domains of SpA, attached to a chromatography resin at more than one site on the resin. In yet another embodiment, a chromatography ligand comprises two or more C domains of SpA, attached to a chromatography resin at more than one site on the resin.

[0034] Also encompassed by the present invention are amino acid variants of SpA, which may differ from the parent amino acid sequence from which they are derived, in the substitution, deletion and/or insertion of one or more amino acids anywhere within the parent amino acid sequence and are alkaline stable. In some embodiments, amino acid sequence variants will possess at least about 70%, or at least about 80%, or at least about 85%, or at least about 90%, or at least about 95%, or at least about 96%, or at least about 97%, or at least about 98% identity with the parent sequence (*i.e.* wt SpA domains or Z domain), where such variants are alkaline stable. In a particular embodiment, variants of SpA further include the glycine amino acid residue at position 29 replaced by an amino acid residue other than alanine or tryptophan, while retaining its alkaline stability.

[0035] The term "functional variant" of a protein means herein a variant protein, where the function, in relation to the invention defined as alkaline stability, is essentially retained. Functional variants include, and are not limited to, SpA variants including more than domain of SpA, *e.g.*, dimers, trimers, multimers of various domains of SpA and SpA variants having a deletion, substitution and/or addition of one or more amino acids in one or more wild type domains of SpA, while retaining the alkaline stability, as defined herein.

[0036] The term "parental molecule" is used herein for the corresponding protein in the form before it is modified according to the invention or a mutation according to the invention has been introduced.

[0037] The term "sequence identity" means that two nucleotide or amino acid sequences, when optimally aligned, such as by the programs GAP or BESTFIT using default gap weights, share at least 70% sequence identity, or at least 80% sequence identity, or at least 85% sequence identity, or at least 90% sequence identity, or at least 95% sequence identity or more. For sequence comparison, typically one sequence acts as a reference sequence (*e.g.*, parent sequence), to which test sequences are compared. When using a sequence comparison algorithm, test and reference sequences are input into a computer, subsequence coordinates are designated, if necessary, and sequence algorithm program parameters are designated. The sequence comparison algorithm then calculates the percent sequence identity for the test sequence(s) relative to the reference sequence, based on the designated program parameters.

[0038] Optimal alignment of sequences for comparison can be conducted, *e.g.*, by the local homology algorithm of Smith & Waterman, *Adv. Appl. Math.* 2:482 (1981), by the homology alignment algorithm of Needleman & Wunsch, *J. Mol. Biol.* 48:443 (1970), by the search for similarity method of Pearson & Lipman, *Proc. Nat'l. Acad. Sci. USA* 85:2444 (1988), by computerized implementations of these algorithms (GAP, BESTFIT, FASTA, and TFASTA in the Wisconsin Genetics Software Package, Genetics Computer Group, 575 Science Dr., Madison, Wis.), or by visual inspection (see generally Ausubel *et al.*, *Current Protocols in Molecular Biology*). One example of algorithm that is suitable for determining percent sequence identity and sequence similarity is the BLAST algorithm, which is described in Altschul *et al.*, *J. Mol. Biol.* 215:403 (1990). Software for performing BLAST analyses is publicly available through the National Center for Biotechnology Information (publicly accessible through the National Institutes of Health NCBI internet server). Typically, default program parameters can be used to perform the sequence comparison, although customized parameters can also be used. For amino acid sequences, the BLASTP program uses as defaults a wordlength (W) of 3, an expectation (E) of 10, and the BLOSUM62 scoring matrix (see Henikoff & Henikoff, *Proc. Natl. Acad. Sci. USA* 89:10915 (1989)).

[0039] As used interchangeably herein, the terms “E domain,” “E domain of SpA,” and “E domain of *Staphylococcus* protein A,” refer to the polypeptide whose amino acid sequence is set forth in SEQ ID NO:7 or that encoded by, *e.g.*, the nucleotide sequence set forth in SEQ ID NO: 1. The “E domain” is a 51 amino acid polypeptide that folds into a three-helix bundle structure. It is capable of binding Fc via residues on the surface of helices 1 and 2, or to Fab via residues on the surface of helices 2 and 3. In some embodiments, an E domain according to the invention is at least 70% identical, or at least 80% identical, or at least 90% identical or at least 95% or more identical in sequence to the amino acid sequence set forth in SEQ ID NO:7.

[0040] As used interchangeably herein, the terms “D domain,” “D domain of SpA,” and “D domain of *Staphylococcus* protein A,” refer to the polypeptide whose amino acid sequence is set forth in SEQ ID NO: 8 or that encoded by *e.g.*, the nucleotide sequence set forth in SEQ ID NO: 2. The “D domain” is a 61 amino acid polypeptide that folds into a three-helix bundle structure. It is capable of Fc binding via residues on the surface of helices 1 and 2, or to Fab via residues on the surface of helices 2 and 3. In some embodiments, a D domain according to the invention is at least 70% identical, or at least 80% identical, or at least 90% identical or at least 95% or more identical in sequence to the amino acid sequence set forth in SEQ ID NO: 8.

[0041] As used interchangeably herein, the terms “A domain,” “A domain of SpA,” and “A domain of *Staphylococcus* protein A,” refer to the polypeptide whose amino acid sequence is set forth in SEQ ID NO: 3 or that encoded by, *e.g.*, the nucleotide sequence set forth in SEQ ID NO: 9. The “A domain” is a 58 amino acid polypeptide that folds into a three-helix bundle structure. It is capable of Fc binding via residues on the surface of helices 1 and 2, or to Fab via residues on the surface of helices 2 and 3. In some embodiments, an A domain according to the invention is at least 70% identical, or at least 80% identical, or at least 90% identical or at least 95% or more identical in sequence to the amino acid sequence set forth in SEQ ID NO: 3.

[0042] As used interchangeably herein, the terms “B domain,” “B domain of SpA,” and “B domain of *Staphylococcus* protein A,” refer to the polypeptide whose amino acid sequence is set forth in SEQ ID NO: 10 or that encoded by, *e.g.*, the nucleotide sequence set forth in SEQ ID NO: 4. The “B domain” is a 58 amino acid polypeptide that folds into a three-helix bundle structure. It is capable of Fc binding via residues on the surface of helices 1 and 2, or to Fab via residues on the surface of helices 2 and 3. In some embodiments, a B domain according to the invention is at

least 70% identical, or at least 80% identical, or at least 90% identical or at least 95% or more identical in sequence to the amino acid sequence set forth in SEQ ID NO: 10.

[0043] As used interchangeably herein, the terms “C domain,” “C domain of SpA,” and “C domain of *Staphylococcus* protein A,” refer to the polypeptide whose amino acid sequence is set forth in SEQ ID NO: 11 or that encoded by, *e.g.*, the nucleotide sequence set forth in SEQ ID NO: 5. The “C domain” is a 58 amino acid polypeptide that folds into a three-helix bundle structure. It is capable of Fc binding via residues on the surface of helices 1 and 2, or to Fab via residues on the surface of helices 2 and 3. In some embodiments, a C domain according to the invention is at least 70% identical, or at least 80% identical, or at least 90% identical or at least 95% or more identical in sequence to the amino acid sequence set forth in SEQ ID NO: 11.

[0044] As used interchangeably herein, the terms “Z domain,” “Z domain of SpA” and “Z domain of protein A,” refer to the three helix, 59 amino acid polypeptide that is a variant of the B domain of protein A. The amino acid sequence of the Z domain is set forth in SEQ ID NO: 12. An exemplary Z domain is described in Nilsson *et al.*, *Protein Engng.*, 1:107-113 (1997), the entire contents of which are incorporated by reference herein.

[0045] The term “alkaline-stable,” “alkaline stability,” “caustic stable” or “caustic stability,” as used herein, generally refers to the ability of a chromatography ligand according to the present invention, either alone or when immobilized onto a chromatography resin, to withstand repeated cleaning-in-place (CIP) cycles using alkaline wash without losing its binding capacity. In general, it is assumed that a resin, by itself, onto which a ligand according to the invention is immobilized, contributes to less than a 5% change in stability after having been soaked in 0.5 M NaOH for up to 30 hours. For example, in some embodiments, chromatography ligands according to the invention are able to withstand conventional alkaline cleaning for a prolonged period of time, which renders the ligands attractive candidates, especially for cost-effective large-scale purification of immunoglobulins. In some embodiments, a ligand according to the present invention exhibits an improved chemical stability in an alkaline environment, which may be defined, for example, as that having an increased pH value such as above about 10, or up to about 13 or 14. Alternatively, the alkaline environment can be defined by the concentration of a base, *e.g.*, about 1.0 M NaOH, or about 0.7 M NaOH, or about 0.5 M NaOH. In one embodiment, alkaline stability refers to the ability of an alkaline-stable chromatography ligand according to the present invention to retain at least 80%, or at least 85%, or at least

90%, or at least 95% of its binding capacity after 5 hours, or after 10 hours, after 15 hours, or after 20 hours, or after 25 hours, or after 30 hours of incubation in 0.5 M NaOH. In another embodiment, alkaline stability refers to a decrease in the binding capacity of the ligand by less than 70%, or less than 60%, or less than 50%, or less than 30% even after treatment with 0.5 M NaOH for 5 hours or 7.5 hours or 10 hours or 15 hours or 20 hours or 25 hours or 30 hours.

[0046] In some embodiments, SpA based chromatography ligands according to the present invention exhibit an increased or improved alkaline stability as compared to wild type SpA.

[0047] Alkaline stability can be readily measured by one of ordinary skill in the art using routine experimentation and/or as described herein.

[0048] The term "chromatography," as used herein, refers to a dynamic separation technique which separates the analyte of interest (*e.g.*, an immunoglobulin) from other molecules in the mixture and allows it to be isolated. Typically, in a chromatography method, a mobile phase (liquid or gas) transports a sample containing the analyte of interest across or through a stationary phase (normally solid) medium. Differences in partition or affinity to the stationary phase separate different analytes while mobile phase carries the different analytes out at different time.

[0049] The term "affinity chromatography," as used herein, refers to a mode of chromatography where the analyte to be separated is isolated by its interaction with a molecule (*e.g.*, an alkaline stable chromatography ligand) which specifically interacts with the analyte. In one embodiment, affinity chromatography involves the addition of a sample containing a target analyte (*e.g.*, an immunoglobulin) to a solid support which carries on it an alkaline stable chromatography ligand, as described herein.

[0050] The term "protein A affinity chromatography," as used herein, refers to the separation or isolation of substances using protein A or SpA ligands, such as those described herein, where the SpA or protein A ligand is immobilized, *e.g.*, on a solid support. Examples of protein A affinity chromatography media/resin known in the art include those having the protein A immobilized onto a controlled pore glass backbone, *e.g.*, PROSEP A™ and PROSEP vA™ media/resin (Millipore); those having protein A immobilized onto a polystyrene solid phase, *e.g.*, the POROS 50A™ and Poros MabCapture A™ media/resin (Applied Biosystems, Inc.); and those having

protein A immobilized on an agarose solid support, *e.g.*, rPROTEIN A SEPHAROSE FAST FLOW™ or MABSELECT™ columns (Amersham Biosciences).

[0051] The term “immunoglobulin,” “Ig” or “antibody” (used interchangeably herein) refers to a protein having a basic four-polypeptide chain structure consisting of two heavy and two light chains, said chains being stabilized, for example, by interchain disulfide bonds, which has the ability to specifically bind antigen. The term “single-chain immunoglobulin” or “single-chain antibody” (used interchangeably herein) refers to a protein having a two-polypeptide chain structure consisting of a heavy and a light chain, said chains being stabilized, for example, by interchain peptide linkers, which has the ability to specifically bind antigen. The term “domain” refers to a globular region of a heavy or light chain polypeptide comprising peptide loops (*e.g.*, comprising 3 to 4 peptide loops) stabilized, for example, by β -pleated sheet and/or intrachain disulfide bond. Domains are further referred to herein as “constant” or “variable”, based on the relative lack of sequence variation within the domains of various class members in the case of a “constant” domain, or the significant variation within the domains of various class members in the case of a “variable” domain. Antibody or polypeptide “domains” are often referred to interchangeably in the art as antibody or polypeptide “regions”. The “constant” domains of antibody light chains are referred to interchangeably as “light chain constant regions”, “light chain constant domains”, “CL” regions or “CL” domains. The “constant” domains of antibody heavy chains are referred to interchangeably as “heavy chain constant regions”, “heavy chain constant domains”, “CH” regions or “CH” domains. The “variable” domains of antibody light chains are referred to interchangeably as “light chain variable regions”, “light chain variable domains”, “VL” regions or “VL” domains. The “variable” domains of antibody heavy chains are referred to interchangeably as “heavy chain variable regions”, “heavy chain variable domains”, “VH” regions or “VH” domains.

[0052] Immunoglobulins or antibodies may be monoclonal or polyclonal and may exist in monomeric or polymeric form, for example, IgM antibodies which exist in pentameric form and/or IgA antibodies which exist in monomeric, dimeric or multimeric form. The term “fragment” refers to a part or portion of an antibody or antibody chain comprising fewer amino acid residues than an intact or complete antibody or antibody chain. Fragments can be obtained *via* chemical or enzymatic

treatment of an intact or complete antibody or antibody chain. Fragments can also be obtained by recombinant means. Exemplary fragments include Fab, Fab', F(ab')₂, Fc and/or Fv fragments.

[0053] The term "antigen-binding fragment" refers to a polypeptide portion of an immunoglobulin or antibody that binds an antigen or competes with intact antibody (*i.e.*, with the intact antibody from which they were derived) for antigen binding (*i.e.*, specific binding). Binding fragments can be produced by recombinant DNA techniques, or by enzymatic or chemical cleavage of intact immunoglobulins. Binding fragments include Fab, Fab', F(ab')₂, Fv, single chains, and single-chain antibodies.

[0054] Also encompassed are fusion proteins including an antibody or fragment thereof as a part of the fusion protein.

[0055] The terms "polynucleotide" and "nucleic acid molecule," used interchangeably herein, refer to polymeric forms of nucleotides of any length, either ribonucleotides or deoxyribonucleotides. These terms include a single-, double- or triple-stranded DNA, genomic DNA, cDNA, RNA, DNA-RNA hybrid, or a polymer comprising purine and pyrimidine bases, or other natural, chemically or biochemically modified, non-natural or derivatized nucleotide bases. The backbone of the polynucleotide can comprise sugars and phosphate groups (as may typically be found in RNA or DNA), or modified or substituted sugar or phosphate groups. In addition, a double-stranded polynucleotide can be obtained from the single stranded polynucleotide product of chemical synthesis either by synthesizing the complementary strand and annealing the strands under appropriate conditions, or by synthesizing the complementary strand *de novo* using a DNA polymerase with an appropriate primer. A nucleic acid molecule can take many different forms, *e.g.*, a gene or gene fragment, one or more exons, one or more introns, mRNA, cDNA, recombinant polynucleotides, branched polynucleotides, plasmids, vectors, isolated DNA of any sequence, isolated RNA of any sequence, nucleic acid probes, and primers. A polynucleotide may comprise modified nucleotides, such as methylated nucleotides and nucleotide analogs, uracyl, other sugars and linking groups such as fluororibose and thioate, and nucleotide branches. As used herein, "DNA" or "nucleotide sequence" includes not only bases A, T, C, and G, but also includes any of their analogs or modified forms of these bases, such as methylated nucleotides, internucleotide modifications such as uncharged linkages and thioates, use of sugar

analogs, and modified and/or alternative backbone structures, such as polyamides. In a particular embodiment, a nucleic acid molecule comprises a nucleotide sequence encoding a variant of SpA.

[0056] The term “Fc-binding,” “binds to an Fc portion” or “binding to an Fc portion” refers to the ability of an alkaline stable chromatography ligand described herein, to bind to the crystallizable part (Fc) of an antibody. In some embodiments, ligand according to the present invention binds an Fc portion of an antibody (*e.g.*, human IgG1, IgG2 or IgG4) with an affinity of at least 10^{-7} M, or at least 10^{-8} M, or at least 10^{-9} M.

[0057] As used herein, the term “Fab binding” or “binding to a Fab portion” refers to the ability of an alkaline stable chromatography ligand described herein, to bind to a Fab region of an antibody or an immunoglobulin molecule. The term “reduced binding to a Fab portion” refers to any decrease in binding to a Fab (or F(ab)₂) portion of an immunoglobulin molecule by an SpA based ligand according to the present invention relative to the wt SpA, where the ligand further includes a mutation in one or more amino acids. In an exemplary embodiment, a ligand according to the present invention further includes the glycine residue at position 29 replaced with an alanine. In another embodiment, a ligand according to the present invention further includes the glycine at position 29 replaced with an amino acid other than alanine or tryptophan. In one embodiment, binding to a Fab portion of an immunoglobulin molecule is undetectable using conventional techniques in the art and those described herein. Binding to an immunoglobulin molecule can be detected using well known techniques including those described herein and including but not limited to, for example, affinity chromatography and Surface Plasmon Resonance Analysis. In some embodiments, an immunoglobulin binding protein encompassed by the present invention binds an immunoglobulin molecule with an affinity of at least 10^{-10} M.

II. Generation of SpA Based Molecules for Use as Chromatography Ligands

[0058] The SpA based chromatography ligands encompassed by the present invention can be made using any suitable methods known in the art.

[0059] For example, as an initial step, standard genetic engineering techniques, *e.g.*, those described in the laboratory manual entitled Molecular Cloning

by Sambrook, Fritsch and Maniatis, may be used for the generation of nucleic acids which express the SpA ligand molecules described herein.

[0060] In some embodiments, a nucleic acid molecule encoding one or more domains of SpA or portions thereof can be cloned into a suitable vector for expression in an appropriate host cell. Suitable expression vectors are well known in the art and typically include the necessary elements for the transcription and translation of the variant SpA coding sequence.

[0061] SpA molecules described herein may also be synthesized chemically from amino acid precursors for fragments using methods well known in the art, including solid phase peptide synthetic methods such as the Boc (tert-butyloxycarbonyl) or Fmoc (9-fluorenylmethyloxy carbonyl) approaches (see, *e.g.*, U.S. Pat. Nos. 6,060,596; 4,879,378; 5,198,531; 5,240,680).

[0062] Expression of SpA molecules described herein can be accomplished in cells from eukaryotic hosts such as yeasts, insects or mammals, or in prokaryotic host cells, *e.g.*, bacteria such as *E. coli*.

[0063] In some embodiments, SpA molecules or fragments and variants thereof may be expressed on the surface of a bacteriophage such that each phage contains a DNA sequence that codes for an individual SpA molecule displayed on the phage surface. The affinity of the SpA molecule for an immunoglobulin can be readily assayed for using standard techniques in the art and those described herein, *e.g.*, ELISA and Biacore™ 2000 standard set-up (Biacore AB, Uppsala Sweden). It is desirable that the binding affinity of an SpA molecule of the present invention to an immunoglobulin is at least comparable with that of the parent molecule. Furthermore, it is desirable that the alkaline stability of the SpA molecule is generally improved over that of the parent molecule.

III. Assaying for Alkaline Stability of the SpA Molecules

[0064] Subsequent to the generation and purification of a suitable SpA ligand molecule, as described herein, the alkaline stability of the molecule can be assayed using standard techniques in the art and those described herein. For example, the alkaline stability of an SpA molecule according to the invention can be assayed using routine treatment with NaOH at a concentration of about 0.5M, *e.g.*, as described in the experimental part below.

[0065] In some embodiments, alkaline stable SpA molecules exhibit an “increased” or “improved” alkaline stability, meaning that the molecules are stable under alkaline conditions for an extended period of time relative to wild type SpA. Previously, it has been reported that SpA molecules based on the wild type C domain of SpA or having a mutation of one or more asparagine residues provides an improved chemical stability and hence a decreased degradation rate in environments wherein the pH is above about 10, such as up to about 13 or 14.

[0066] The present invention is based on the surprising and unexpected discovery of novel SpA molecules which exhibit increased alkaline stability even when they are based on domains other than the C domain or have mutations in amino acids other than asparagines. For example, the present invention provides B or Z domain based alkaline stable SpA molecules and SpA molecules which have a mutation in an amino acid at position n+1, where n represents an asparagine (*e.g.*, asparagine at position 23).

[0067] In some embodiments, subsequent to the generation of the SpA ligands according to the present invention, alkaline stability of the ligands is evaluated using a novel high throughput immunological assay, described in more detail in the Examples *infra*. The assay is based on the assumption that degradation of SpA in response to extended caustic exposure is reflected as a loss or reduction in IgG binding. Briefly, soluble SpA based ligands are treated for about 6 hours with either water or 1.0M NaOH. Hydrophobic interactions are used to attach microgram quantities of neutralized candidate ligands to a solid support in the form of an ELISA plate, *e.g.*, a 96 well plate. IgG binding is then assessed for each candidate ligand before and after exposure to 1.0M NaOH. Enhanced caustic stability is indicated when the amount of residual IgG binding to a ligand following caustic exposure exceeds that of wild type SpA or parental SpA from which it is derived.

IV. Supports Used for the Preparation of Chromatography Matrices

[0068] In some embodiments, alkaline stable SpA ligands encompassed by the present invention are attached to a support, *e.g.*, a solid support or a soluble support, to generate a chromatography matrix suitable for the separation of biomolecules such as, *e.g.*, immunoglobulins.

[0069] In some embodiments, a ligand according to the present invention is attached to a solid support. Without wishing to be bound by theory, it is contemplated

that any suitable solid support may be used for the attachment of a ligand according to the invention. For example, solid support matrices include, but are not limited to, controlled pore glass, silica, zirconium oxide, agarose, polymethacrylate, polyacrylate, polyacrylamide, and polystyrene.

[0070] It is contemplated that any porous material that contributes to less than a 5% change in alkaline stability of the attached ligand after soaking for about 30 hours in 0.5 M NaOH may be used as a solid support.

[0071] A porous material used as a solid support may be comprised of a hydrophilic compound, a hydrophobic compound, an oleophobic compound, an oleophilic compound or any combination thereof. The porous material may be comprised of a polymer or a copolymer. Examples of suitable porous materials, include, but are not limited to polyether sulfone, polyamide, *e.g.*, nylon, polysaccharides such as, for example, agarose and cellulose, polyacrylate, polymethacrylate, polyacrylamide, polymethacrylamide, polytetrafluoroethylene, polysulfone, polyester, polyvinylidene fluoride, polypropylene, polyethylene, polycarbonate, a fluorocarbon, *e.g.* poly (tetrafluoroethylene-co-perfluoro(alkyl vinyl ether)), glass, silica, zirconia, titania, ceramic, and metal.

[0072] The porous material may be comprised of an organic or inorganic molecules or a combination of organic and inorganic molecules and may be comprised of one or more functional groups, *e.g.*, a hydroxyl group, a thiol group, an amino group, a carbonyl group, or a carboxylic acid group, suitable for reacting, *e.g.*, forming covalent bonds for further chemical modification in order to covalently bond to a protein. In another embodiment, the porous material may not possess a functional group but can be coated with a layer of material that bears functional groups such as, an hydroxyl group, a thiol group, an amino acid group, a carbonyl group, or a carboxylic acid group.

[0073] In some embodiments, a conventional affinity separation matrix is used, *e.g.*, of organic nature and based on polymers that expose a hydrophilic surface to the aqueous media used, *i.e.* expose hydroxy (--OH), carboxy (--COOH), carbonyl (--CHO, or RCO-R'), carboxamido (--CONH₂, possibly in N-substituted forms), amino (--NH₂, possibly in substituted form), oligo- or polyethylenoxy groups on their external and, if present, also on internal surfaces. In one embodiment, the polymers may, for instance, be based on polysaccharides, such as dextran, starch, cellulose, pullulan, agarose *etc.*, which advantageously have been cross-linked, for instance with

bisepoxides, epihalohydrins, 1,2,3-trihalo substituted lower hydrocarbons, to provide a suitable porosity and rigidity. In another embodiment, the solid support comprises porous agarose beads. The various supports used in the present invention can be readily prepared according to standard methods known in the art, such as, for example, inverse suspension gelation described, *e.g.*, in Hjerten, *Biochim Biophys Acta* 79(2), 393-398 (1964). Alternatively, the base matrices can be commercially available products, such as Sepharose™ FastFlow (GE Healthcare, Uppsala, Sweden). In some embodiments, especially advantageous for large-scale separations, the support is adapted to increase its rigidity, and hence renders the matrix more suitable for high flow rates.

[0074] Alternatively, the solid support can be based on synthetic polymers, such as polyvinyl alcohol, polyhydroxyalkyl acrylates, polyhydroxyalkyl methacrylates, polyacrylamides, polymethacrylamides *etc.* In case of hydrophobic polymers, such as matrices based on divinyl and monovinyl-substituted benzenes, the surface of the matrix is often hydrophilised to expose hydrophilic groups as defined above to a surrounding aqueous liquid. Such polymers can be easily produced according to standard methods, see *e.g.*, Arshady, *Chimica e L'Industria* 70(9), 70-75 (1988). Alternatively, a commercially available product, such as Source™ (GE Healthcare, Uppsala, Sweden) and Poros (Applied BioSystems, Foster City, CA) may be used.

[0075] In yet other embodiments, the solid support comprises a support of inorganic nature, *e.g.* silica, zirconium oxide *etc.* The surface of inorganic matrices is often modified to include suitable reactive groups for further reaction to SpA and its variants. Examples include CM Zirconia (Ciphergen–BioSeptra (CergyPontoise, France) and CPG® (Millipore).

[0076] In some embodiments, the polymers may, for instance, be based on zirconia or silica or controlled pore glass, which may be modified to either contain reactive groups and/or sustain caustic soaking, to be coupled to ligands.

[0077] Exemplary solid support formats include, but are not limited to, a bead, a gel, a membrane, a cassette, a column, a chip, a slide, a plate or a monolith.

[0078] With respect to the format of a matrix, in one embodiment, it is in the form of a porous monolith. In an alternative embodiment, the matrix is in beaded or particle form that can be porous or non-porous. Matrices in beaded or particle form can be used as a packed bed or in a suspended form. Suspended forms include those

known as expanded beds and pure suspensions, in which the particles or beads are free to move. In case of monoliths, packed bed and expanded beds, the separation procedure commonly follows conventional chromatography with a concentration gradient. In case of pure suspension, batch-wise mode will be used. Also, solid support in forms such as a surface, a chip, a capillary, or a filter may be used.

[0079] The matrix could also be in the form of membrane in a cartridge. The membrane could be in flat sheet, spiral, or hollow fiber format.

[0080] In another embodiment, a ligand according to the present invention is attached to a soluble support, *e.g.*, a soluble polymer. Exemplary soluble supports include, but are not limited to, a bio-polymer such as, *e.g.*, a protein or a nucleic acid. In some embodiments, biotin may be used as a soluble polymer, *e.g.*, as described in US Patent Publication No. 20080108053. For example, biotin may be bound to a ligand, *e.g.*, an SpA based caustic stable ligand according to the present invention, which subsequent to being bound to the ligand can be used for isolating a protein of interest, *e.g.*, an antibody or fragment thereof, *e.g.*, present in a crude mixture and the protein of interest can be isolated or separated *via* precipitation of the biotin-ligand-protein polymer complex in either a reversible or irreversible fashion. The polymer may also be a synthetic soluble polymer, such as, for example, including but not limited, to a polymer containing negatively charged groups (carboxylic or sulfonic), positively charged groups (quarternary amine, tertiary amine, secondary or primary groups), hydrophobic groups (phenyl or butyl groups), hydrophilic groups (hydroxyl, or amino groups) or a combination of the above. Exemplary synthetic soluble polymers can be found in International PCT Publication No. WO2008091740 and U.S. Publication No. US20080255027, the entire teachings of each of which are incorporated by reference herein. These polymers, upon specific physical changes in one or more conditions such as pH, conductivity or temperature, can be used to purify the protein of interest *via* precipitation in either a reversible or an irreversible fashion. Synthetic soluble polymers may be used alone or may be coupled with a caustic stable ligand according to the present invention and used for capture/purification of a protein of interest such as, *e.g.*, an antibody or a fragment thereof, *via* precipitation in either a reversible or an irreversible fashion.

V. Methods for Attaching a Ligand to a Support

[0081] Any suitable technique may be used for attaching a ligand to a support, *e.g.*, a solid support including those well known in the art and described herein. For example, in some embodiments, the ligand may be attached to a support via conventional coupling techniques utilizing, *e.g.* amino and/or carboxy groups present in the ligand. For example, bisepoxides, epichlorohydrin, CNBr, N-hydroxysuccinimide (NHS) etc. are well-known coupling reagents. In some embodiments, a spacer is introduced between the support and the ligand, which improves the availability of the ligand and facilitates the chemical coupling of the ligand to the support. Alternatively, the ligand may be attached to the support by non-covalent bonding, such as physical adsorption or biospecific adsorption.

[0082] In various embodiments encompassed by the present invention, the ligand is attached to a solid support such as, for example, a chromatography resin at more than one site, thereby resulting in a chromatography matrix.

[0083] Attachment of an alkaline stable SpA based chromatography ligand to a solid support can be achieved via many different ways known, most of which are well known in the art, as well as those described herein. See, *e.g.*, Hermanson *et al.*, *Immobilized Affinity Ligand Techniques*, Academic Press, pp. 51-136 (1992).

[0084] For example, protein ligands can be coupled to a solid support via active groups on either the surface of the solid support or the protein ligand, such as, for example, hydroxyl, thiol, epoxide, amino, carbonyl, epoxide, or carboxylic acid group. Attachment can be achieved using known chemistries including, but not limited to, use of cyanogen bromide (CNBr), N-hydroxyl succinimide ester, epoxy (bisoxirane) activation, and reductive amination.

[0085] For example, thiol directed protein coupling has been described in the literature. See, *e.g.*, Ljungquist, *et al.* Eur. J. Biochem. Vol 186, pp. 558-561 (1989). This technique has been previously applied for coupling SpA to a solid support. Since wild type SpA does not contain thiol groups, the attachment is achieved by recombinantly inserting a thiol containing cysteine at the C-terminus of SpA. See, *e.g.*, U.S. Patent No. 6,399,750. Several commercial products such as MabSelect™, MabSelect™ Xtra and MabSelect™ SuRe are produced via this mechanism. It has been reported that this terminal cysteine only reacts with the epoxide group on the solid surface, thereby resulting in single point attachment of the SpA to the solid support. See, *e.g.*, Process Scale Bioseparations for the Biopharmaceutical Industry, CRC Press, 2006, page 473.

[0086] In case of the present invention, in certain embodiments, SpA based chromatography ligands comprising two or more domains of SpA are attached to a solid support at more than one site via non-discriminate, multipoint attachment. In general, SpA contains abundant free amino groups from numerous lysines in each domain. The attachment of an SpA domain to more than one site on a solid support, *e.g.*, a chromatography resin with epoxide or aldehyde group, can be achieved by reacting the amino group of lysine on SpA, via epoxide ring-opening or reductive amination, respectively. In certain embodiments, multipoint attachment can be achieved by the reaction of one or more naturally occurring amino acids on SpA having free hydroxyl groups, such as, for example, serine and tyrosine, with a support containing an epoxide group via a ring-opening reaction. Alternatively, multipoint attachment can be achieved, for example, by the reaction of naturally occurring amino acids on SpA having free carboxylic acid groups, such as, for example, aspartic acid and glutamic acid, with a support containing amino groups via, for example, *N,N'*-carbonyldiimidazole. Multipoint attachment of the ligand to support can also be achieved by a combination of all the above mechanisms.

[0087] To achieve caustic stability using the multimers of B and Z domains, this invention excludes the single cysteine mutation that leads to single point attachment.

[0088] SpA based chromatography ligands may also be attached to a solid support via an associative mechanism. For example, an associative group may interact with a ligand of interest non-covalently via ionic, hydrophobic or a combination of interactions, thereby to attach ligand of interest onto the solid surface.. This facilitates the high efficiency coupling of ligand to the solid matrix, for example, as described in US Patent Publication No. 20070207500A1, thereby resulting in ligand density higher than that without the associative groups. Associative groups suitable for use in the invention include charged species such as ionic species, and uncharged species such as hydrophobic species. The associative group may modify the solid support, *e.g.* by covalently binding directly with the solid support. Suitable examples of ionic species may include quaternary amines, tertiary amines, secondary amines, primary amines, a sulfonic group, carboxylic acid, or any combination thereof. Suitable examples of hydrophobic species may include a phenyl group, a butyl group, a propyl group, or any combination thereof. It is also contemplated that mixed mode species may be used. The associative group may also interact with the

protein ligand. Thus the interaction between the associative group and the protein ligand may be comprised of a mixture of interactions, *e.g.* ionic and hydrophobic species.

[0089] The associative group may be covalently coupled to the solid support by reacting a functional group on the solid support with a functional group on the associative group. Suitable functional groups include, but are not limited to amines, hydroxyl, sulfhydryl, carboxyl, imine, aldehyde, ketone, alkene, alkyne, azo, nitrile, epoxide, cyanogens and activated carboxylic acid groups. As an example, agarose beads contain hydroxyl groups which may be reacted with the epoxide functionality of a positively charged associative group, such as glycidyl trimethylammonium chloride. A skilled artisan will appreciate that a plurality of associative groups may be coupled to the solid support provided that at least one bifunctional associative group is used. Thus associative groups may be coupled in tandem to the solid support or they may be individually coupled directly to the solid support.

[0090] In some embodiments, the present invention provides associative groups and/ or protein ligands which may be coupled to a solid support via an intervening linker. The linker may comprise at least one functional group coupled to a linking moiety. The linking moiety may comprise any molecule capable of being coupled to a functional group. For example, the linking moiety may include any of an alkyl, an alkenyl, or an alkynyl group. The linking moiety may comprise a carbon chain ranging from 1 to 30 carbon atoms. In some embodiments the linker may be comprised of more than 30 carbon atoms. The linking moiety may comprise at least one hetero-atom such as nitrogen, oxygen and sulfur. The linking moiety may be comprised of a branched chain, an unbranched chain or a cyclic chain. The linking moiety may be substituted with two or more functional groups.

[0091] Choosing the appropriate buffer conditions for coupling a protein ligand to a solid support is well within the capability of the skilled artisan. Suitable buffers include any non-amine containing buffer such as carbonate, bicarbonate, phosphate and acetate buffers. When associative chemistry is used, salt concentration of the buffer will depend on the associative group used. For example, the salt concentration may be in the range of 5nM- 100 mM. Where a charged species is used, the salt concentration may be at least 5nM but less than 0.1M, at least 5nM but less than 0.01M, at least 5nM but less than 0.001M. In certain embodiments, the salt concentration may be 0.01M. Where a hydrophobic species is used a high salt

concentration is usually desirable. Thus the salt concentration may be greater than 0.001 M, greater than 0.01 M, or greater than 0.1 M.

[0092] In some embodiments, when associative chemistry is used, the reaction is performed at a temperature ranging from 0°C to 99°C. In certain embodiments the reaction method is practiced at a temperature less than 60°C, less than 40°C, less than 20°C, or less than 10°C. In some embodiments the method of the invention is practiced at a temperature of about 4°C. In other embodiments the method of the invention is practiced at a temperature of 20°C.

VI. Methods for Assaying for Alkaline Stability of the Attached Ligands

[0093] The increased alkaline stability of the ligands subsequent to their attachment to a support can be assayed by using well known techniques in the art and those described herein. For example, in some embodiments, alkaline stability of multimers of SpA domains attached to a resin, *e.g.*, B and Z domains of SpA attached to a chromatography resin, as described herein, can be confirmed by treatment of the resin with 0.5 M NaOH. It is to be understood that an increased stability means that the initial IgG binding capacity is retained during a longer period of time than what can be achieved by the wild type SpA molecule. For example, in case of the present invention, after 100 cycles, each including a 15 min treatment with 0.5 N NaOH, the percentage of retained capacity of the SpA ligands, *e.g.*, those comprising multiple B or Z domains, is at least 1.5 times more, 2.0 times more, 2.5 times more, or 3 times more than that of wild type SpA. In one embodiment, the alkaline stability of the bound ligand, as assayed by the retention of IgG binding capacity over time, is measured as follows. The binding capacity, referred to as Qd 50%, is measured by obtaining the volume of IgG loaded to a UV_{280nm} of 50% of the initial IgG concentration. Qd 50% of the initial virgin resin packed in a column is measured first. The resin is then exposed to about 10 cycles of 15 min exposure of 0.5 N NaOH at 0.8 ml/min. Qd 50% is measured again. This process is repeated until the resin is exposed to a total of about 100 cycles of 0.5 N NaOH. Qd 50% is measured one last time and the results from resins made from different ligands are compared with the wildtype SpA.

[0094] In another assay, caustic or alkaline stability of the resins is measured by static soaking of resins of interest. By soaking a measured amount of

resin in 0.5N NaOH for 25 hrs with gentle rotation and measuring IgG binding capacity before and after the NaOH soaking, the alkaline stability by way of retention of binding capacity of the resin can be determined.

VII. Methods of Purifying a Target Molecule Using a Chromatography Ligand of the Invention

[0095] In some embodiments, the present invention provides a method of purifying a target molecule from a mixture using the alkaline stable chromatography ligands described herein. The target molecule may be any molecule which is recognized by an alkaline stable chromatography ligand provided herein, where the ligand is coupled to a solid support. Examples of target molecules include immunoglobulins. The immunoglobulins may be polyclonal antibodies or a monoclonal antibody or a functional fragment thereof. Functional fragments include any fragment of an immunoglobulin comprising a variable region that still binds specifically to its antigen while at the same time retaining its ability to specifically bind to a protein ligand coupled to a solid support.

[0096] In some embodiments, a method of isolating a target molecule of interest using an alkaline stable chromatography ligand described herein includes the steps of: (a) contacting a solid support including an attached SpA based alkaline stable chromatography ligand with a mixture comprising a molecule of interest under conditions such that the target molecule specifically binds to the ligand; and (b) altering the conditions such that the target molecule is no longer bound to the ligand, thereby isolating the target molecule.

[0097] In some embodiments, the altering step includes altering the pH, such the target molecule is no longer bound to the ligand. In a particular embodiment, the pH is altered in a manner such that it is more acidic than the pH conditions in step (a). For example, in one embodiment, step (a) may be performed at a neutral pH, or a pH ranging from about 6 to about 8 and step (b) may be performed at an acidic pH, *e.g.*, a pH ranging from about 1 to about 5.

[0098] In another embodiment, step (b) comprises altering the salt concentration of the buffer in use, such that the target molecule is no longer bound to the ligand. For example, in one embodiment, a high salt concentration, *e.g.*, > 0.1 M, may be used in step (a) and a lower salt concentration, *e.g.*, <0.1M may be used in step (b). Conversely, in some embodiments, a low salt concentration, *e.g.*, < 0.1M

may be used in step (a) and a high salt concentration may be used in step (b). In still other embodiments both the pH and the salt concentration of the buffer may be altered between step (a) and step (b).

[0099] One skilled in the art can readily determine the conditions suitable for binding a target molecule to a ligand, and thereby alter the conditions to disrupt the binding of the molecule to the ligand.

[00100] This invention is further illustrated by the following examples which should not be construed as limiting. The contents of all references, patents and published patent applications cited throughout this application, as well as the Figures, are incorporated herein by reference.

Examples

Example 1: Generation of an SpA B Domain Variant Having an n+1 Mutation, where n Represents an Asparagine

[00101] In an exemplary experiment, a synthetic gene encoding the “B domain” of protein A is obtained from DNA 2.0 (Menlo Park, CA). The 5’ end of the gene includes a codon for an initiating methionine as well as six histidine codons at the 3’ end of the gene. The gene is provided in vector pJ56:8620 from DNA 2.0. The parent vector pJ56 confers resistance to ampicillin, kanamycin, chloramphenicol and gentamicin. Appropriate restriction enzyme sites for the subsequent cloning of the gene into expression vectors are introduced into the gene at both 5’ and 3’ ends. A plasmid map of the vector is shown in Figure 5.

[00102] Saturation mutagenesis is subsequently used to mutate glutamic acid at position 24 to all other naturally occurring amino acids except cysteine (C), serine (S), alanine (A), glycine (g), asparagine (N), and glutamine (Q) by PCR-based methods using the Phusion High-Fidelity DNA polymerase (New England Biolabs, Ipswich, MA). Primers are purchased from IDT DNA (Coralville, IA), as 100 μ M solution in Tris EDTA buffer. The mutagenic primers have the sequence CTGCCGAACCTGAACNN SGAACAACGCAACGG (SEQ ID NO: 41) where NNS represents the three bases encoding the amino acid at position 24. PCR is performed in 50 μ L reactions containing dNTPs (0.2 mM each), 125 ng of each primer, 50 ng of template plasmid, and 1 U of Phusion enzyme. PCR is carried out according to the scheme outlined in Table III.

Table III

Cycle description	Temperature	Time	# of cycles
Initial denaturation	95 °C	30 seconds	1 cycle
Denaturation	95 °C	30 seconds	18 cycles
Annealing	55 °C	60 seconds	
Extension	68 °C	6 minutes	

[00103] PCR reactions are treated with the restriction enzyme *DpnI* (New England Biolabs, Ipswich, MA) to reduce wild type background. To each 50 µL PCR reaction, about 1 µL of *DpnI* enzyme is added and the samples are incubated for about one hour at 37 °C.

[00104] *E. coli* NEB5α competent cells (New England Biolabs, Ipswich, MA) are transformed with 2 µL of the *DpnI*-treated PCR reaction. Cells are thawed on ice, and 2 µL of the PCR reaction is added to 25 µL of cells. Following about a 30 minute incubation on ice, cells are heat shocked for 30 seconds at about 42 °C. Cells are allowed to recover for about 5 minutes on ice, and then 125 µL of SOC media (New England BioLabs) is added. Cells are incubated for about one hour at 37 °C, and then 100 µL are plated on LB plates (Northeast Laboratory Services, Winslow, ME) containing 100 µg/mL ampicillin and grown overnight at about 37 °C. Positive clones are identified by testing for the expression of the proteins in total cell lysates using SDS PAGE.

In order to obtain purified DNA, individual colonies are picked for overnight culture in LB containing 100 µg/mL ampicillin. DNA is purified using spin mini-prep kits from Qiagen (Valencia, CA).

[00105] Mini-prepped DNA is sequenced to confirm the identity of each clone (MWG Biotech, Huntsville AL). The resulting plasmid is used to transform *E. coli* NEB5α competent cells as described above.

[00106] Following the identification of positive clones, 35 mls overnight cultures are grown in Terrific Broth with 100 µg/ml ampicillin and cells are pelleted by centrifugation at 13,500 g for 10 minutes. Pellets are resuspended in 10 mls of 20 mM imidazole in PBS (phosphate buffered saline), lysed by sonication and centrifuged at 13,500 g for 30 minutes to pellet insoluble debris. Lysates are subsequently applied to 750 µl of Ni-NTA resin that is pre-equilibrated with 10 column volumes of 20 mM imidazole in PBS. After washing with 20 column

volumes of 20 mM imidazole in PBS, samples are eluted from the resin with 200 mM imidazole in PBS. Purified protein is dialyzed overnight against PBS using Pierce Slide-A-Lyzer 3.5 MWCO dialysis cassettes. Following dialysis, total protein quantitation is accomplished using the Pierce MicroBCA assay, and samples are stored at -30 °C.

[00107] The amino acid and nucleic acid sequences for the wt his-tagged B domain and the various n+1 mutants are depicted in Figures 6 and 7.

Example 2: Assaying the Expressed Proteins for Alkaline Stability

[00108] Affinity purified wildtype and mutant SpA his-tagged constructs described in Example 1 are diluted with MilliQ water or with NaOH to a final concentration of 1N and incubated for six hours at room temperature. Biorad Micro Biospin 6 gel filtration columns are used to neutralize and buffer exchange 50 µl of each sample into PBS. Total protein quantitation is accomplished using the Pierce MicroBCA assay and samples are diluted to 10 µg/ml in PBS for loading onto the ELISA plate. Approximately 200 µl (2 µg) of treated PrA is adsorbed to the wells of an ELISA plate for 2-24 hours at 37 °C. Plates are blocked in Pierce Superblock Blocking Buffer in PBS (Superblock-PBS) for about 2 hours at room temperature. Approximately 200 µl of Sigma human gamma globulin diluted to 0.05 mg/ml in Superblock-PBS (10 µg) is added to each well of the plate and binding is allowed to proceed for about 1 hour at room temperature. Following three washes with PBS containing about 0.05% Tween20 (PBS-T), plates are incubated with 200 µl of a 1:10,000 dilution of a chicken IgY-HRP conjugate raised against human IgG for about 1 hour at room temperature. After the final three washes with PBS-T, plates are developed with 100 µl of Pierce 1-Step Slow TMB ELISA for 30 minutes at room temperature. The reaction is stopped by the addition of 100 µl of 1N HCL and primary IgG binding is quantitated as absorbance read at 450 nm. All samples are assayed in triplicate and data is analyzed as the change in primary IgG binding before and after caustic treatment.

[00109] The result of an exemplary experiment assaying for the alkaline stability of the various n+1 B domain mutants is summarized in the bar graph in Figure 8. The various constructs are depicted on the X-axis and the corresponding percent of IgG binding remaining after six hours of exposure to 1M NaOH is plotted

on the Y-axis As depicted in the bar graph in Figure 8, several of the n+1 mutants having a bulky amino acid at position 24 of the B domain exhibit enhanced caustic stability.

[00110] In another experiment, SpA ligands having two or more B domains or two or more Z domains of SpA are assayed for alkaline stability using the assay described above. In yet another experiment, SpA ligands having three B domains (B3), four B domains (B4), five B domains (B5), six B domains (B6) or seven B domains (B7) are assayed for alkaline stability using the assay described above.

[00111] In an exemplary experiment, the percent of residual IgG binding capacity of the ligands B3, B4 and B5, as assayed using the ELISA based method described herein is as follows. The B3 ligand retains about 79% of its residual IgG binding capacity following a six hour exposure to 1M NaOH; the B4 ligand retains about 86% of its residual IgG binding capacity following a six hour exposure to 1M NaOH; and the B5 ligand retains about 83% of its residual IgG binding capacity following a six hour exposure to 1M NaOH. The wt SpA ligand (nPrA) only retains about 49% of its residual IgG binding capacity following a six hour exposure to 1M NaOH.

Example 3: Attachment of wildtype SpA and SpA variants to a solid support.

[00112] Subsequent to the generation of the various SpA variants and identification of those which are caustic stable using one or more assays known in the art and those described in Example 2, the SpA variants are attached to a support, *e.g.*, a solid support. In an exemplary experiment, agarose beads (Sepharose 4B) (GE Healthcare, Piscataway NJ) are crosslinked using epichlorohydrin according to a previously described method (Porath and Fornstedt, *J. Chromatography*, 51:479 (1979)). The agarose beads are reacted with positively charged associative groups, *e.g.*, cations, according to the following method: 50 mL of beads are added to 40g of 75% wt glycidyl trimethylammonium chloride (GTMAC), 10 mL Milli-Q® water (Millipore Corp., Billerica, MA) and 1.67 g 50 % wt sodium hydroxide. The reaction is shaken vigorously (> 100 rpm) on a rotary shaker overnight at room temperature. The beads are then filtered and washed with three 100 mL volumes of Milli-Q® water (Millipore Corp, Billerica, MA).

[00113] The beads (50 mL, filtered cake) are added to a jar containing 15 mLs of 4.6M NaOH. The mixture is slurried and then 19.5 mL of butanediol diglycidylether (BUDGE) is added. This mixture is shaken at 35 °C for about 2 hours. The beads are then washed with 750 mL of Milli-Q® water (Millipore Corp, Billerica, MA) and equilibrated with 250 mL of 10 mM NaHCO₃.

[00114] Immediately following the BUDGE activation step, 10 mL of the filtered bead cake is added to 10mL solution of 10 mM NaHCO₃ containing a 15g/L concentration of wild type SpA or an SpA ligand according to the present invention. The mixture is capped in a glass vial and the vial is rotated in a hybridizer at 37 °C for about 2 hours. After two hours, the beads are washed with 30mLs of Milli-Q® water (Millipore Corp, Billerica, MA). The filtered bead cake (10mL) is added to a jar containing a 10 mL solution comprised of 1 mL of thioglycerol and 9 mL of a buffer solution with 0.2 M NaHCO₃ and 0.5 M NaCl. The mixture is slurried and rotated overnight at room temperature. The beads are then washed with 30 mL of the following buffers: 0.1 M Tris Buffer (pH 8), 0.1 M Tris Buffer with 0.15 M NaCl (pH 8), 50 mM Acetic Acid (pH 4.5), PBS (pH 7.4) with 0.002 % sodium azide.

[00115] Resin samples coupled with different SpA variants are labeled as follows: nPrA for wild type SpA; Z3 for SpA ligand containing three Z domains; E3 for SpA ligand containing three E domains; D3 for SpA ligand containing three D domains; A3 for SpA ligand containing three A domains, C3 for SpA ligand containing three C domains; B3 for SpA ligand containing three B domains; B4 for SpA ligand containing four B domains; B5 for SpA ligand containing 5 B domains; B5NF and B7NF containing 5 and 7 B domains, respectively, and additionally containing a G29A mutation at position 29. Following attachment to a resin, the various SpA variants are assayed for alkaline stability.

Example 4: Resin IgG binding capacity (Qd 50%) tests before and after caustic cycling

[00116] In an exemplary experiment, the various SpA constructs according to the invention are assayed for the IgG binding capacity following attachment to a support. In one exemplary experiment, a standard method for testing resin/media dynamic capacity using commercial polyclonal IgG is used. Briefly, resin coupled with an SpA ligand according to the present invention is packed into an Omnifit column (6.6 mm x 70 mm) in PBS, pH 7.4 and the flow rate is set at 300 cm/hr. The

packed column is equilibrated with PBS for 10 column volumes (CVs). Polyclonal IgG (Sigma-Aldrich, 2 mg/mL in PBS, pH 7.4) is loaded onto the column until UV_{280nm} reaches more than 50% of the initial IgG concentration. After washing with equilibration buffer, IgG is eluted with 0.1 M citric acid, pH 3.0. After each run, the media is sanitized using 6M Guanidine hydrochloride. Qd 50% is calculated based on the amount of IgG loaded when UV_{280nm} reaches 50% of the initial IgG concentration.

[00117] After the initial dynamic capacity measurement, the media is exposed to 10 cycles of 15 min, 0.5 N NaOH (flow rate 100 cm/hr) followed by another IgG dynamic capacity measurement. The media is subsequently contacted with another 10 cycles of 15 min, 0.5N NaOH exposure, followed by another dynamic binding capacity measurement. Dynamic capacity measurement is carried out after each sample is exposed to 100 cycles of 15 min 0.5 N NaOH.

[00118] In an exemplary experiment, the results of which are depicted in Figure 9, trimers of E, D, A, B, C, and Z domains containing natural B domain linkers are attached to an agarose resin and their caustic stability is measured by dynamic binding capacity over 100 cycles of caustic exposure, each cycle including a 15 minute exposure to 0.5 N NaOH. As summarized in the graph in Figure 9, the number of cycles are plotted on the X-axis and the IgG binding capacity is plotted on the Y-axis. The C and the Z domain trimers exhibit approximately the same level of caustic stability. The B domain trimers are more caustic stable than the A domain trimers, which are more caustic stable than the E and the D domain trimers. Therefore, the order of caustic stability can be summarized as follows C and Z>B>A>E and D.

[00119] In another exemplary experiment, agarose resins coupled with B domain variants: BBB (B3), BBBB (B4), BBBBB (B5), BBBBB-NF (B5-NF), and BBBBBBB-NF (B7-NF), are compared with wt SpA (nPrA) using the assay described above to assay for IgG dynamic binding capacity over 100 cycles of caustic exposure, each cycle including a 15 minute exposure to 0.5 N NaOH. As summarized in the graph in Figure 10, the extent of caustic stability is directly proportional to the number of domains and further that, the G29A mutation to reduce Fab binding does not affect the caustic stability.

[00120] The specification is most thoroughly understood in light of the teachings of the references cited within the specification which are hereby incorporated by reference. The embodiments within the specification provide an illustration of embodiments in this invention and should not be construed to limit its scope. The skilled artisan readily recognizes that many other embodiments are encompassed by this invention. All publications and inventions are incorporated by reference in their entirety. To the extent that the material incorporated by reference contradicts or is inconsistent with the present specification, the present specification will supercede any such material. The citation of any references herein is not an admission that such references are prior art to the present invention.

[00121] Unless otherwise indicated, all numbers expressing quantities of ingredients, cell culture, treatment conditions, and so forth used in the specification, including claims, are to be understood as being modified in all instances by the term "about." Accordingly, unless otherwise indicated to the contrary, the numerical parameters are approximations and may vary depending upon the desired properties sought to be obtained by the present invention. Unless otherwise indicated, the term "at least" preceding a series of elements is to be understood to refer to every element in the series. Those skilled in the art will recognize, or be able to ascertain using no more than routine experimentation, many equivalents to the specific embodiments of the invention described herein. Such equivalents are intended to be encompassed by the following claims.

[00122] Many modifications and variations of this invention can be made without departing from its spirit and scope, as will be apparent to those skilled in the art. The specific embodiments described herein are offered by way of example only and are not meant to be limiting in any way. It is intended that the specification and examples be considered as exemplary only, with a true scope and spirit of the invention being indicated by the following claims.

CLAIMS

What is claimed is:

1. An alkaline-stable chromatography ligand comprising two or more B domains or two or more Z domains of Staphylococcus protein A (SpA), or a functional fragment or variant thereof.
2. An alkaline-stable chromatography ligand comprising two or more B domains or two or more Z domains of Staphylococcus protein A (SpA), or a functional fragment or variant thereof, wherein the two or more B domains or two or more Z domains are attached to a chromatography resin at more than one site on the resin.
3. The alkaline-stable chromatography ligand of claim 1 or claim 2, wherein the ligand comprises three or more B domains or three or more Z domains of SpA.
4. The alkaline-stable chromatography ligand of claim 1 or claim 2, wherein the ligand comprises four or more B domains or four or more Z domains of SpA.
5. The alkaline-stable chromatography ligand of claim 1 or claim 2, wherein the ligand comprises five or more B domains or five or more Z domains of SpA.
6. The alkaline-stable chromatography ligand of claim 1 or claim 2, wherein the ligand comprises six or more B domains or six or more Z domains of SpA.
7. The alkaline-stable chromatography ligand of claim 1 or claim 2, wherein the ligand comprises seven or more B domains or seven or more Z domains of SpA.
8. The alkaline-stable chromatography ligand of any of the preceding claims, further comprising at least one amino acid mutation to alter Fab binding.
9. The alkaline-stable chromatography ligand of claim 8, wherein the mutation comprises substitution of glycine at position 29 with an alanine.
10. A chromatography matrix comprising a ligand according to any one of claims 1-9.
11. A method of affinity purifying one or more immunoglobulins from a sample, the method comprising the steps of:
 - (a) providing a sample comprising one or more immunoglobulins;
 - (b) contacting the sample with the matrix of claim 10 under conditions such that the one or more immunoglobulins bind to the matrix; and
 - (c) recovering the one or more bound immunoglobulins by eluting under suitable conditions.
12. The ligand according to any one of claims 1-9, wherein the ligand retains at least 95% of its binding capacity after 5 hours, or after 10 hours, or after 15 hours, or after 20 hours incubation in 0.5 M NaOH.

13. The ligand according to any one of claims 1-9, wherein the ligand is capable of binding at least 1.5 times, or 2 times, or 3 times, or more of IgG as compared to wtSpA.
14. An alkaline-stable chromatography ligand comprising one or more isolated E, D, A, B, C or Z domains of *Staphylococcus* protein A, wherein the one or more isolated domains comprise one or more amino acid residues at position n+1 mutated to any naturally occurring amino acid other than cysteine (C), serine (S), alanine (A), glycine (G), asparagine (N), or glutamine (Q), wherein n represents an asparagine.
15. An alkaline-stable chromatography ligand comprising one or more isolated E, D, A, B, C or Z domains of *Staphylococcus* protein A, wherein the one or more isolated domains comprise one or more amino acid residues at position n+1 mutated to an amino acid selected from the group consisting of arginine (R), aspartic acid (D), glutamic acid (E), histidine (H), isoleucine (I), leucine (L), lysine (K), methionine (M), phenylalanine (F), proline (P), threonine (T), tryptophan (W), tyrosine (Y) and valine (V), wherein n represents an asparagine.
16. The ligand of claim 14 or 15, wherein n is asparagine at position 23.
17. A caustic stable affinity chromatography matrix comprising three or more, or four or more, or five or more, or six or more, or seven or more B or Z domains of SpA attached to a solid support via multipoint attachment.
18. A method for identification of an SpA based chromatography ligand which is caustic stable, the method comprising the steps of:
- (a) treating an SpA based ligand with caustic;
 - (b) adding the caustic treated SpA ligand to a multiwell plate;
 - (c) contacting the SpA based ligand with a saturating amount of a first immunoglobulin, such that the first immunoglobulin binds to the ligand;
 - (d) contacting the bound first immunoglobulin with a molecule capable of binding the first immunoglobulin, wherein the molecule is conjugated with a detectable agent; and
 - (e) detecting the amount of a signal from the agent,
- wherein the amount of signal is proportional to the caustic stability of the ligand, thereby to identify the SpA based ligand as being caustic stable.
19. The method of claim 18, wherein the molecule is a second immunoglobulin.