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(54) **FLUOROGENIC SUBSTRATE FOR ADAMTS13**

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**C12Q 1/00** (2006.01)  
**A61K 38/00** (2006.01)

(52) **U.S. Cl.**  
USPC ..... **435/4**; 530/324; 530/300

(58) **Field of Classification Search**  
None  
See application file for complete search history.

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(57) **ABSTRACT**

Disclosed are fluorogenic substrates for measuring ADAMTS13 activity or ADAMTS13 inhibitor activity. Substrates can comprise an oligopeptide which can consist of up to 80 amino acids of sequence of von Willebrand Factor (VWF). The oligopeptide can include modifications of sequence of VWF, including an amino-terminal glycine, a scissile Y-M peptide, and a cysteine substitution located from 1 to 12 amino acids from the scissile Y-M in the carboxy terminal direction. A substrate can further comprise a fluorophore and a fluorescence quencher bound to the oligopeptide on opposite sides of the scissile Y-M peptide, wherein the fluorescence quencher is not identical to the fluorophore. An oligopeptide can be encoded by a nucleic acid sequence which can also encode a His tag. An oligopeptide can be expressed in a cell or microorganism. Also disclosed are methods of using a fluorogenic substrate to measure ADAMTS13 activity or ADAMTS13 inhibitor activity.

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FIG. 1

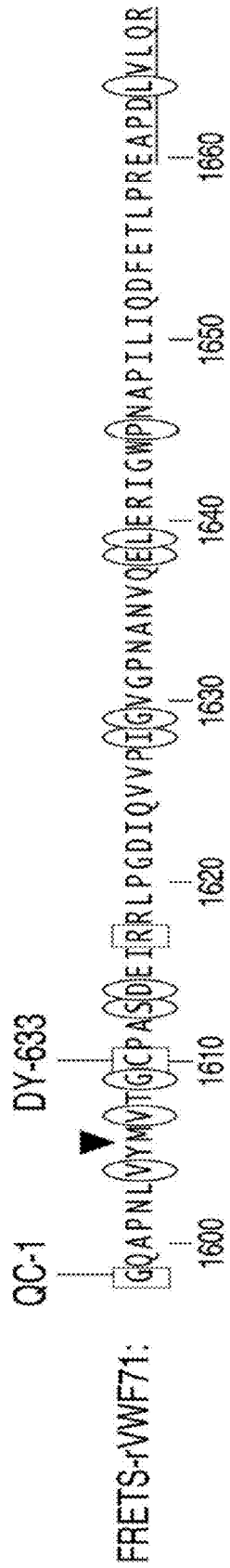


FIG. 2

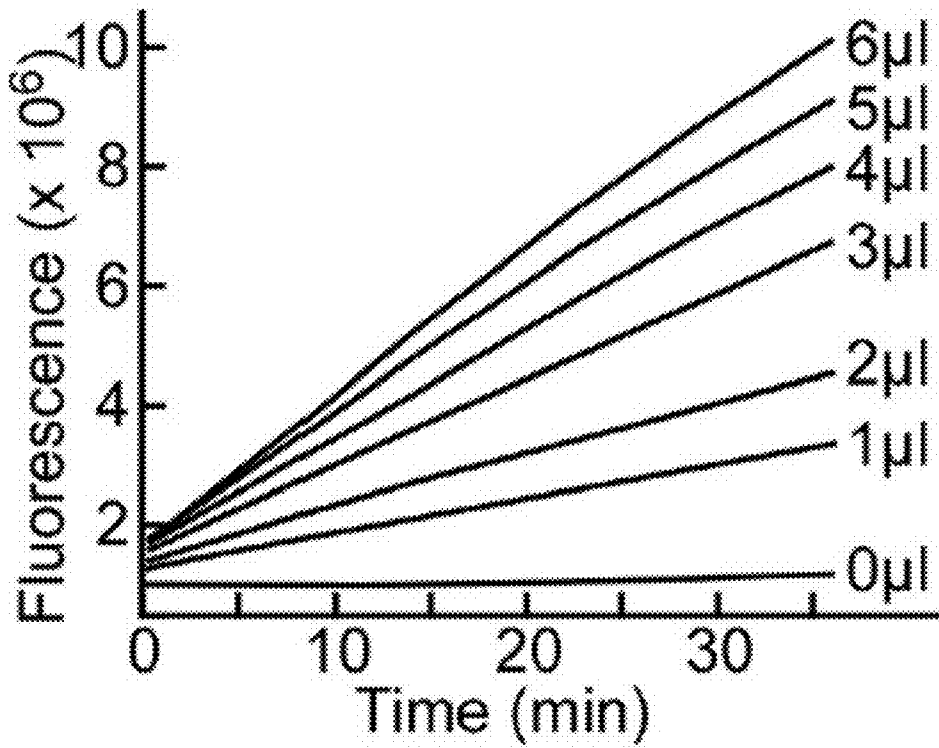


FIG. 3

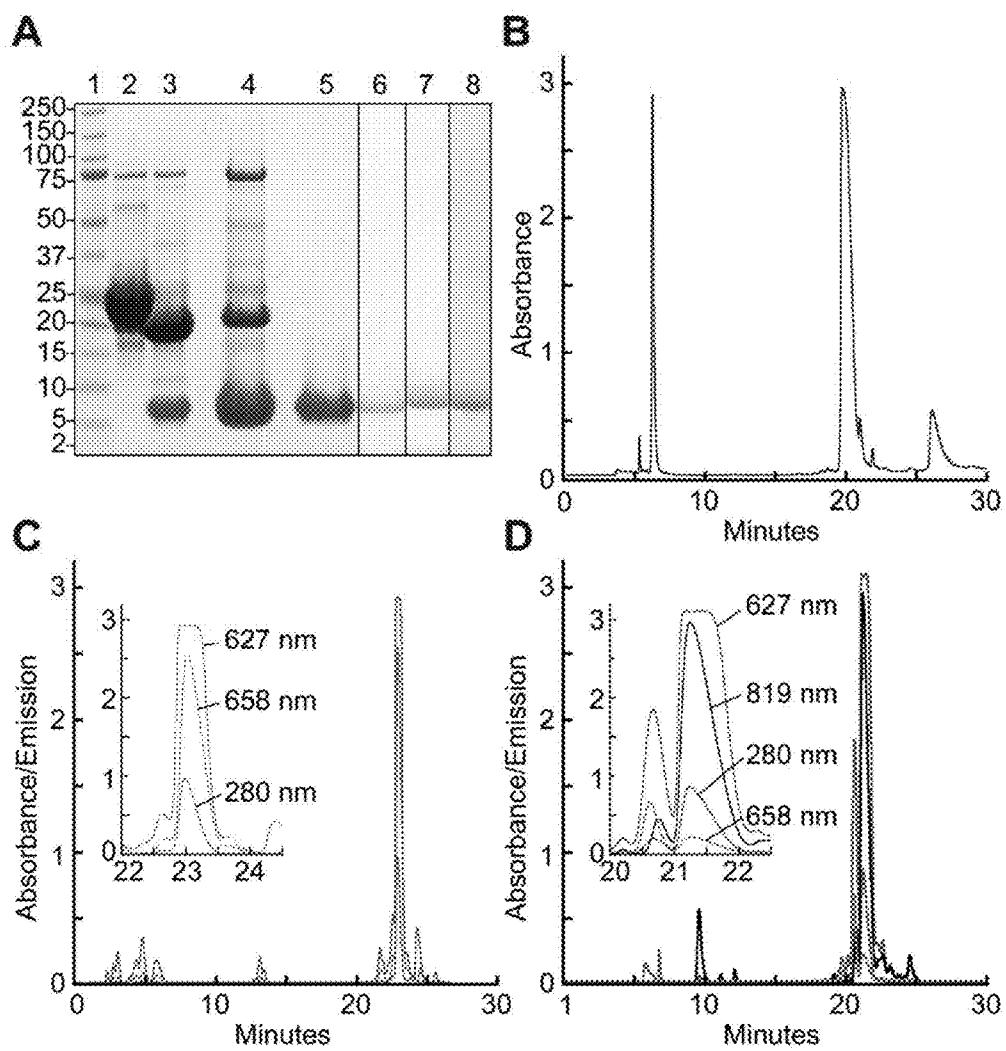


FIG. 4

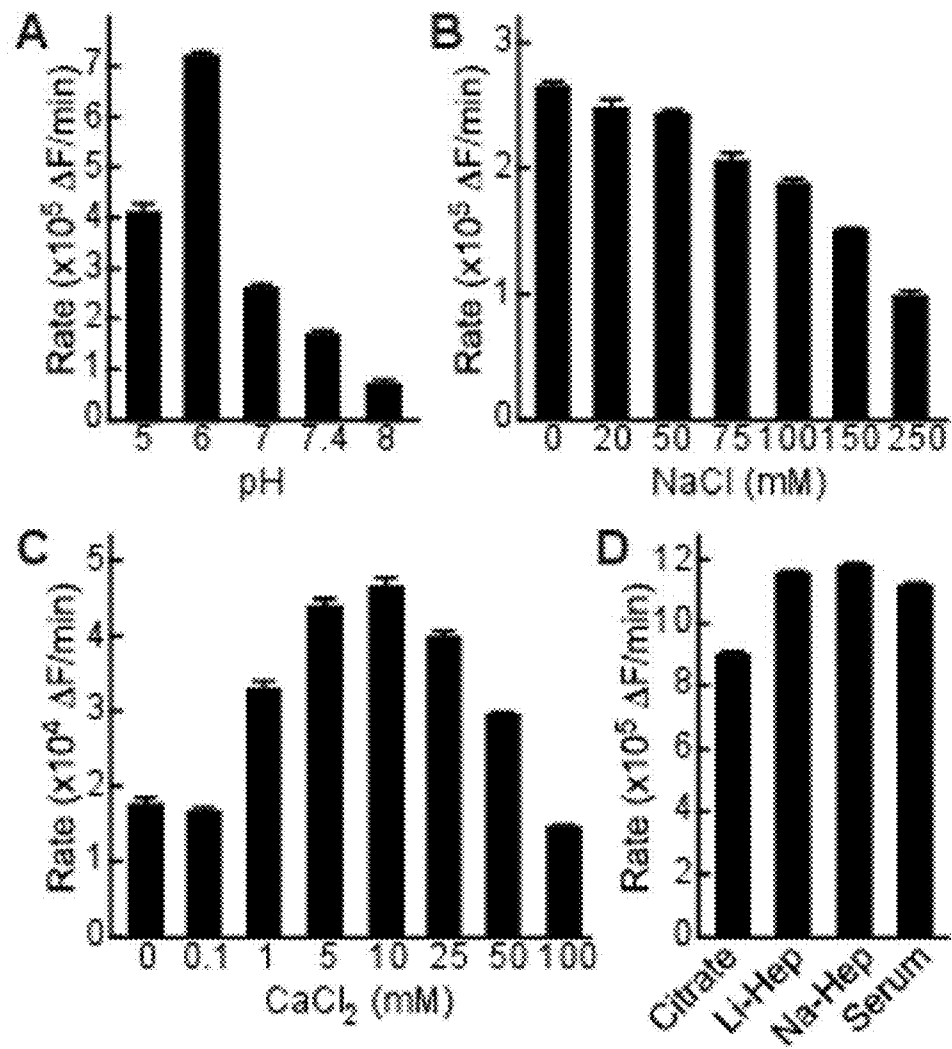


FIG. 5

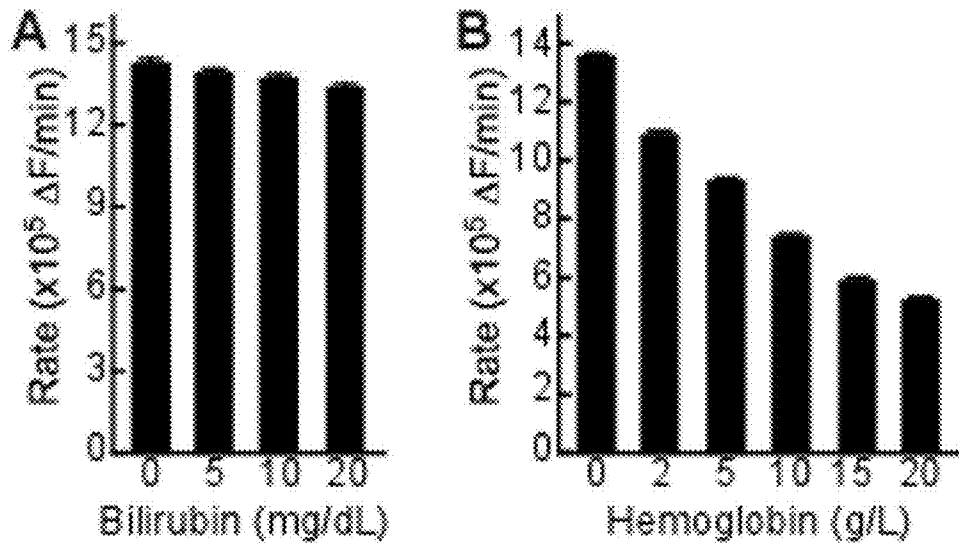


FIG. 6

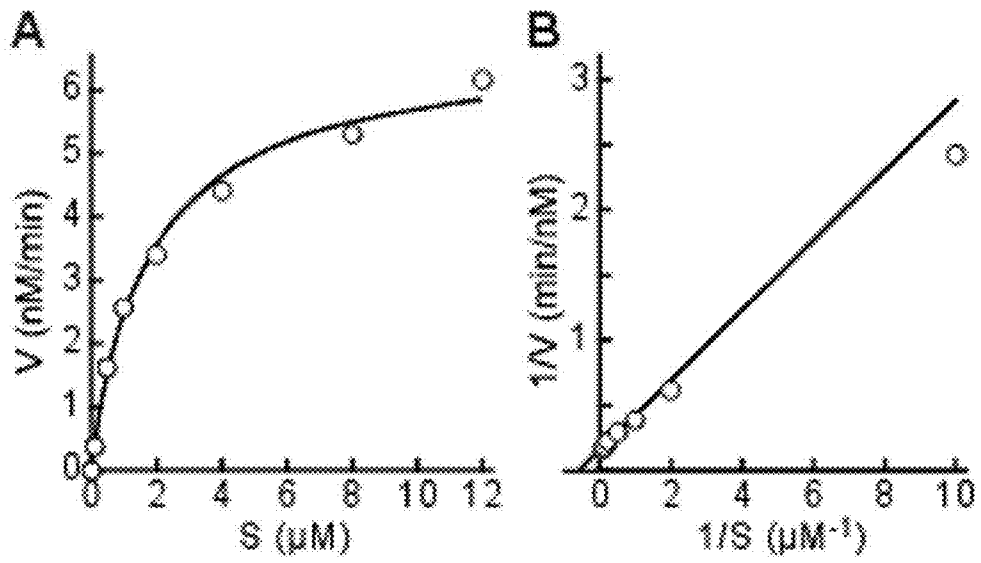




FIG. 7

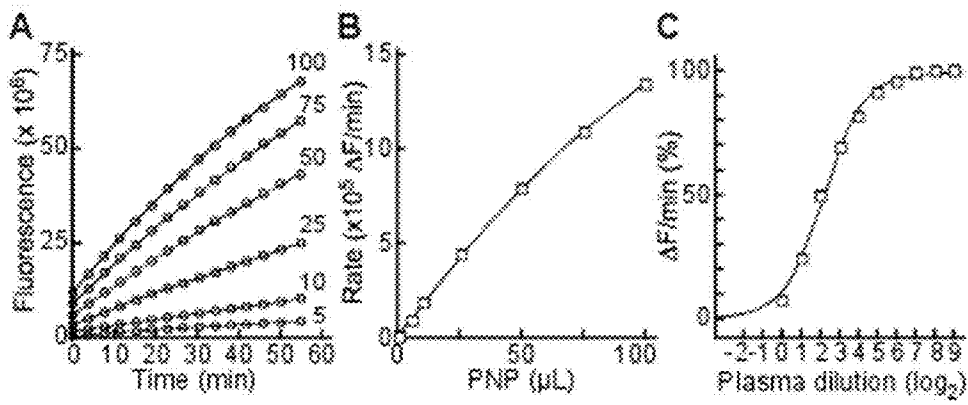
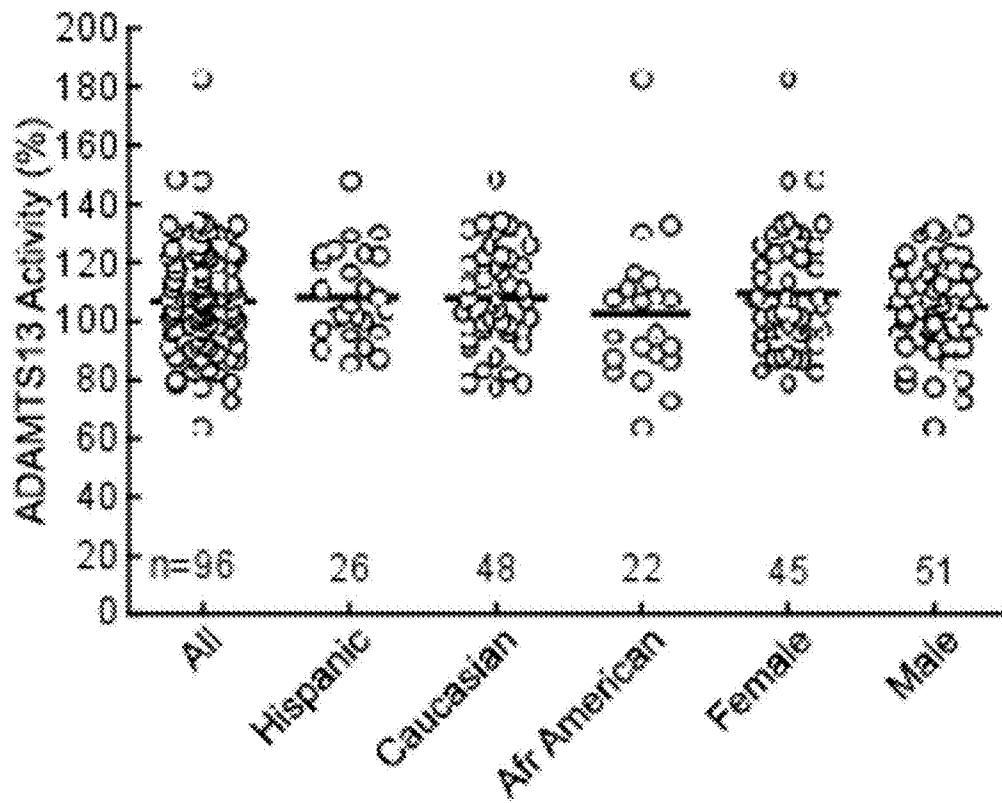


FIG. 8



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## FLUOROGENIC SUBSTRATE FOR ADAMTS13

### CROSS-REFERENCE TO RELATED APPLICATIONS

This application claims priority to U.S. Provisional Application Ser. No. 61/508,782 filed on Jul. 18, 2011, which is incorporated herein by reference in its entirety.

### STATEMENT REGARDING FEDERALLY SPONSORED RESEARCH OR DEVELOPMENT

This invention was made with government support under Grants R01 HL072917-08 and R01 HL089746-04 awarded by the National Institutes of Health. The Government has certain rights in the invention.

### INCORPORATION-BY-REFERENCE OF MATERIAL SUBMITTED IN COMPUTER READABLE FORM

The Sequence Listing, which is a part of the present disclosure, includes a computer readable form. The subject matter of the Sequence Listing is incorporated herein by reference in its entirety. The information recorded in computer readable form is identical to the written sequence listing.

### INTRODUCTION

ADAMTS13 is a metalloprotease that cleaves von Willebrand Factor (VWF) and inhibits the growth of platelet thrombi. ADAMTS13 is a normal component of blood. In some disease states, such as, for example idiopathic thrombotic thrombocytopenic purpura (idiopathic TTP), levels of ADAMTS13 levels can be below normal.

Several studies suggest that ADAMTS13 activity and inhibitor assays can provide useful prognostic information. Measurement of ADAMTS13 activity in blood, plasma or serum can be useful, for example to identify patients with severe ADAMTS13 deficiency (Tsai, H.-M.; *Blood*. 87:4235-4244, 1996; Downes, K. A., et al., *J. Clin. Apheresis*. 19:86-89, 2004), or with confounding potential causes of thrombotic microangiopathy such as stem cell (Adachi, T., et al., *Int. J. Hematol.* 83:415-419, 2006) or solid organ transplants (Pham, P. T., et al., *Transplantation*. 74:1077-1080, 2002; Mal, H., et al., *Transplantation*. 81:1628-1632, 2006), systemic lupus erythematosus (Ahmed, S., et al., *Am J. Med.* 116:786-787, 2004; Rick, M. E., et al., *Am. J. Hematol.* 75:96-100, 2004; Fujisaki, K., et al., *Clin Nephrol.* 64:305-310, 2005; Coppo, P., et al., *Medicine*. 83:233-244, 2004), preeclampsia or HELLP syndrome (Lattuada, A et al., *Haematologica*. 88:1029-1034, 2003; George, J. N., *Curr. Opin. Hematol.* 10:339-344, 2003). In such situations, ADAMTS13 testing can facilitate diagnosis and expedite treatment.

However, the clinical use of ADAMTS13 data requires assays that are rapid, robust, and feasible for most hospital laboratories. Available assays for ADAMTS13 activity can be technically challenging, slow to perform, insensitive to inhibitors, and difficult to automate (Furlan, M., et al., *Blood*. 87:4223-4234, 1996; Gerritsen, H. E., et al., *Thromb Haemost.* 82:1386-1369, 1999).

FRETS-VWF73 is a fluorogenic ADAMTS13 substrate that corresponds to VWF Asp<sup>1596</sup>-Arg<sup>1668</sup> (73 residues), replacing Gln<sup>1599</sup> with 2,3-diaminopropionic acid (A2pr) linked to N-methyl anthranilate (Nma), and Asn<sup>1610</sup> with A2pr-2,4-dinitrophenyl (Dnp) (Kokame, K., et al., *Br. J. Haematol.* 129:93-100, 2005).

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In FRETS-VWF73, Gln<sup>1599</sup> is replaced with A2pr (Nma), and Asn<sup>1610</sup> is replaced with A2pr (Dnp). Nma absorbs at 340 nm and emits at 440 nm; nearby Dnp quenches this fluorescence. When the Tyr<sup>1605</sup>-Met<sup>1606</sup> bond is cleaved, quenching is relieved, producing a fluorescence signal proportional to product concentration. FRETS-VWF73 assays require <1 hour and have been adapted to measure inhibitors. Results with FRETS-VWF73 and multimeric VWF substrates are generally congruent (Kokame, K., et al., *Br. J. Haematol.* 129:93-100, 2005; Kremer Hovinga, J. A., et al., *J. Thromb. Haemost.* 4:1146-1148, 2006; Groot, E., et al., *J. Thromb. Haemost.* 4:698-699, 2006; Tripodi, A., et al., *J. Thromb. Haemost.* 6:1534-1541, 2008; Peyvandi, F., et al., *J. Thromb. Haemost.* 8:631-640, 2010).

Despite these attributes, FRETS-VWF73 has several limitations. For example, it is chemically synthesized, and expensive (~\$10/data point). Furthermore, assay conditions are not physiological (pH 6.0, low ionic strength) (Kokame, K., et al., *Br. J. Haematol.* 129:93-100, 2005). Nma absorption at 340 nm and emission at 450 nm make assays susceptible to interference from autofluorescence, absorbance and quenching by plasma proteins such as hemoglobin and bilirubin (Meyer, S. C., et al., *J. Thromb. Haemost.* 5:866-867, 2007). To avoid such problems, in these methods plasma is diluted  $\geq 1:20$  in ADAMTS13 assays using FRETS-VWF73 substrate. This dilution limits assay sensitivity to ~3% of normal ADAMTS13 levels, prevents the detection of some inhibitors, and can overestimate ADAMTS13 activity for patients with low titer inhibitors (Kremer Hovinga, J. A., et al., *Blood*. 115:1500-1511, 2010).

Current assays of ADAMTS13 involve dilutions of  $\geq 1:20$  and cannot detect inhibitors with low affinity and rapid dissociation kinetics. Alternatively, ELISA methods are used that measure antibody bound to immobilized ADAMTS13 and results are reported as " $\mu\text{g}$  antibody/mL," analogous to assays for anti-PF4 antibodies. However, this approach cannot distinguish inhibitory from noninhibitory antibodies that may have no biological significance.

United States Patent Application 20100240050 of Bhatia, S. N., et al., entitled "Methods and Products For In Vivo Enzyme Profiling," discloses a "pro-diagnostic reagent" comprising "a carrier domain linked to an enzyme susceptible domain which is linked to a signature molecule." A "signature molecule" can be a peptide, nucleic acid, small molecule, fluorophore/quencher, carbohydrate, or particle, such as a peptide comprising a fluorophore and a quencher. However, the carrier domain comprises a particle, for example, a micro-particle or a nanoparticle. Although this publication lists ADAMTS13 and VWF as an enzyme and substrate, none of the peptide sequences disclosed in this application appear to derive from VWF or could be used as a target substrate for ADAMTS13.

U.S. Pat. Nos. 7,468,258 and 7,927,864 to Owen disclose a peptide consisting of a sequence from VWF which further comprises self-quenching fluorophores. This fluorogenic ADAMTS13 substrate relies on the autoquenching properties of the disclosed paired fluorophores as an alternative to fluorophore-quencher combinations.

However, there is an unmet need for an ADAMTS13 substrate that can be used with minimal sample dilution.

### SUMMARY

The present inventors have developed fluorogenic substrates which can be used to measure ADAMTS13 activity as well as ADAMTS13 inhibitor activity. An ADAMTS13 substrate of the present teachings comprises an oligopeptide. In

some embodiments, an oligopeptide of the present teachings can be encoded in a vector and expressed in a cell or a microorganism such as *E. coli*. An ADAMTS13 substrate of the present teachings can further comprise a fluorophore and a fluorescence quencher.

In some embodiments, a fluorogenic ADAMTS13 substrate of the present teachings comprises an oligopeptide consisting of no more than 80 amino acids of sequence of von Willebrand Factor (VWF). In various embodiments, the oligopeptide can comprise a scissile tyrosine-methionine (Y-M) peptide, a cysteine substitution located from 1 to 12 amino acids from the scissile Y-M in the carboxy terminal direction; a C-terminal segment; a fluorophore; and a fluorescence quencher.

In some configurations, an oligopeptide of an ADAMTS13 substrate can consist of no more than 73 amino acids. In some configurations, an oligopeptide of an ADAMTS13 substrate can consist of no more than 72 amino acids. In some configurations, an oligopeptide of an ADAMTS13 substrate can consist of no more than 71 amino acids.

In some configurations, the cysteine substitution can be situated 3 amino acids from the scissile Y-M peptide. In some configurations, the cysteine substitution can be a N1610C substitution of a VWF sequence.

In some configurations, an oligopeptide can comprise a substitution of lysine K 1617 for an amino acid that does not comprise a primary amine on its side chain that can react with amine-reactive reagents (such as a succinimidyl ester or an isothiocyanate). In some configurations, an oligopeptide can comprise a K1617R substitution.

In some configurations, an oligopeptide can comprise a substitution of glutamic acid E1798 of VWF with glycine, so that the oligopeptide comprises a glycine at its amino terminal.

In some configurations, a probe can comprise an oligopeptide consisting of a sequence set forth as SEQ ID NO: 1:

DREQAPNLVYVMTGCPASDEIKRLPGDIQVVPIGVGPANVQELERIGW  
PNAPILIQDFETLPREAPDLVLQR.

In some configurations, a probe can comprise an oligopeptide consisting of a sequence set forth as SEQ ID NO: 2:

GQAPNLVYVMTGCPASDEIRRLPGDIQVVPIGVGPANVQELERIGWPN  
APILIQDFETLPREAPDLVLQR.

In various aspects, an ADAMTS13 substrate of the present teachings can comprise a fluorophore and a fluorescence quencher situated on opposite sides of a scissile Y-M peptide. In various configurations, a fluorophore can have an absorption maximum of >550 nm or >630 nm.

In various aspects, a fluorophore can have an emission maximum of >600 nm or >650 nm.

In some aspects, a fluorophore can comprise at least one sulfate.

In some aspects, a fluorophore can be an Alexa Fluor® 594 maleimide (Life Technologies Corporation, Carlsbad, Calif.) or a DyLight® 633 maleimide (Thermo Fisher Scientific, Rockford, Ill.).

In various aspects, a quencher can have an absorption maximum of >550 nm or >630 nm.

In some aspects, a quencher can comprise at least one sulfate.

In some aspects, a quencher can be a QSY21-succinimidyl ester or an IRDye QC-1 N-hydroxy succinimidyl ester.

In some aspects, a probe can be soluble in water at >50 μM. In some aspects, a probe can be soluble in water at >200 μM.

In some embodiments, the present teachings include a vector comprising a nucleic acid sequence encoding an oligopeptide of an ADAMTS13 substrate described herein.

In some aspects, a nucleic acid sequence can further encode an N-terminal His tag.

In some aspects, a nucleic acid sequence can further comprise a sequence encoding thioredoxin.

In some aspects, a nucleic acid sequence can further comprise a sequence encoding an oligopeptide further comprises a sequence encoding a Tobacco Etch Virus (TEV) protease cleavage site.

In some aspects, a vector can be a plasmid or a virus, such as a bacteriophage.

In some embodiments, the present teachings include methods of determining presence, absence or quantity of ADAMTS13 activity in a sample.

In some embodiments, the inventors disclose assays for detecting and quantifying activity of inhibitors of ADAMTS13. In some configurations, an assay of the present teachings can be substantially more sensitive than alternative existing assays for detecting inhibitors of ADAMTS13.

In various configurations, methods of the present teachings can comprise: forming a mixture comprising a sample and a probe as described herein; and measuring fluorescence at one or more time points after forming the mixture.

In some configurations, the sample can be a blood sample, a serum sample, or a plasma sample.

In some configurations, the sample can be undiluted or diluted less than 20-fold in an assay of ADAMTS13 activity or ADAMTS13 inhibitor activity. In some configurations, the sample can be diluted less than 2-fold. In some configurations, the sample can be heparinized plasma. In some configurations, the sample can be citrated plasma. In some configurations, the sample can be serum. In some configurations, the sample can be plasma anticoagulated with any protease inhibitor that does not interfere with ADAMTS13 activity.

In some aspects, an ADAMTS13 assay or ADAMTS13 inhibitor assay using a probe of the present teachings can be insensitive to >20 mg/dL (250 μM) conjugated bilirubin.

In some configurations, sensitivity of assays for inhibitors of ADAMTS13 activity can be increased by using recombinant ADAMTS13 (including concentrated recombinant ADAMTS13) as the enzyme source instead of pooled normal plasma.

In some embodiments, the present teachings include methods of producing a probe described herein.

In some configurations, these methods can comprise expressing, in a cell, an oligopeptide encoded by a vector described herein. Alternatively, an oligopeptide can be generated by chemical means such as, for example but not limited to, the solid-phase method of Merrifield, R., *J. Am. Chem. Soc.* 85: 2149-2154, 1963. In various configurations, an oligopeptide can be digested with His-tagged TEV protease to yield an oligopeptide comprising VWF sequence comprising a cysteine as described herein. In various configurations, an oligopeptide comprising a cysteine can be reacted with a fluorophore comprising a maleimide. In various configurations, an oligopeptide can comprise a primary amine such as the N-terminal primary amine which can be reacted with a fluorescence quencher comprising a succinimidyl ester or other amine-reactive moiety such as, without limitation, an isothiocyanate.

Alternatively, an oligopeptide can be generated by chemical means such as, for example but not limited to, the solid-

phase method of Merrifield, R., J. Am. Chem. Soc. 85: 2149-2154, 1963. In various configurations, an oligopeptide can be digested with His-tagged TEV protease to yield an oligopeptide comprising VWF sequence comprising a cysteine as described herein. In various configurations, an oligopeptide comprising a cysteine can be reacted with a fluorescence quencher comprising a maleimide. In various configurations, an oligopeptide can comprise a primary amine such as the N-terminal primary amine which can be reacted with a fluorophore comprising a succinimidyl ester or other amine-reactive moiety such as, without limitation, an isothiocyanate. The present disclosure includes the following aspects, without limitation:

1. A probe comprising:

an oligopeptide consisting of no more than 80 amino acids of sequence of von Willebrand Factor (VWF), said oligopeptide comprising a scissile Y-M peptide, a cysteine substitution located from 1 to 12 amino acids from the scissile Y-M in the carboxy terminal direction, and a C-terminal segment;

a fluorophore; and

a fluorescence quencher,

wherein the fluorophore and the fluorescence quencher are bound to the oligopeptide on opposite sides of the scissile Y-M peptide.

2. A probe in accordance with aspect 1, wherein the oligopeptide consists of no more than 73 amino acids.
3. A probe in accordance with aspect 1, wherein the oligopeptide consists of no more than 71 amino acids.
4. A probe in accordance with aspect 1, wherein the cysteine substitution is situated 3 amino acids from the scissile Y-M peptide.
5. A probe in accordance with aspect 1, wherein the cysteine substitution is a N1610C substitution of a VWF sequence.
6. A probe in accordance with aspect 1, wherein the oligopeptide comprises a substitution of lysine K1617 for an amino acid that does not comprise a primary amine on its side chain.
7. A probe in accordance with aspect 1, comprising a K1617R substitution.
8. A probe in accordance with aspect 1, wherein the oligopeptide consists of the sequence set forth as SEQ ID NO: 2,

GOAPNLVYVMVTGCPASDEIRRLPGDGIQVVPVIGVGPANVQELERIGWPN

APILIQDFETLPREAPDLVLQR.

9. A probe in accordance with aspect 1, wherein the fluorophore has an absorption maximum >550 nm.
10. A probe in accordance with aspect 1, wherein the fluorophore has an emission maximum >600 nm.
11. A probe in accordance with aspect 1, wherein the fluorophore has an absorption maximum >630 nm.
12. A probe in accordance with aspect 1, wherein the fluorophore has an emission maximum >650 nm.
13. A probe in accordance with aspect 1, wherein the fluorophore comprises at least one sulfate.
14. A probe in accordance with aspect 1, wherein the fluorophore is Alexa Fluor 594 maleimide.
15. A probe in accordance with aspect 1, wherein the fluorophore is DyLight 633 maleimide.
16. A probe in accordance with aspect 1, wherein the quencher has an absorption maximum >550 nm.
17. A probe in accordance with aspect 1, wherein the quencher has an absorption maximum >630 nm.
18. A probe in accordance with aspect 1, wherein the quencher comprises at least one sulfate.

19. A probe in accordance with aspect 1, wherein the quencher is QSY21-succinimidyl ester.
20. A probe in accordance with aspect 1, wherein the quencher is IRDye QC-1 N-hydroxy succinimidyl ester.
21. A probe in accordance with aspect 1, wherein the quencher is attached to the oligopeptide at the amino terminal of the oligopeptide.
22. A probe in accordance with aspect 1, wherein the fluorophore is attached to the oligopeptide at the cysteine.
23. A probe in accordance with aspect 1, wherein the probe is soluble in water at >50  $\mu$ M.
24. A vector comprising a nucleic acid sequence encoding an oligopeptide of aspect 1.
25. A vector in accordance with aspect 24, further comprising a nucleic acid sequence encoding an N-terminal His tag.
26. A vector in accordance with aspect 24, wherein the nucleic acid sequence encoding an oligopeptide further comprises a sequence encoding thioredoxin.
27. A vector in accordance with aspect 24, wherein the nucleic acid sequence encoding an oligopeptide further comprises a sequence encoding a Tobacco Etch Virus (TEV) protease cleavage site.
28. A vector in accordance with aspect 24, wherein the vector is a plasmid.
29. A method of determining presence, absence or quantity of ADAMTS13 activity in a sample, comprising: forming a mixture comprising a sample and a probe of aspect 1; and measuring fluorescence at one or more time points after forming the mixture, wherein the sample is diluted less than 20-fold.
30. A method in accordance with aspect 29, wherein the sample is a serum sample.
31. A method in accordance with aspect 30, wherein the serum sample is an undiluted serum sample.
32. A method in accordance with aspect 30, wherein the serum sample is a concentrated serum sample.
33. A method in accordance with aspect 29, wherein the sample is a plasma sample.
34. A method in accordance with aspect 33, wherein the plasma sample is an undiluted plasma sample.
35. A method in accordance with aspect 33, wherein the plasma sample is a concentrated plasma sample.
36. A method of determining presence, absence or quantity of ADAMTS13 inhibitor activity in a sample, comprising: forming a mixture comprising a sample, a source of ADAMTS13, and a probe of aspect 1; and measuring fluorescence at one or more time points after forming the mixture.
37. A method in accordance with aspect 36, further comprising inactivating ADAMTS13 activity endogenous to the sample prior to forming the mixture.
38. A method in accordance with aspect 37, wherein the inactivating ADAMTS13 activity comprises heating the sample.
39. A method in accordance with aspect 38, wherein the heating the sample comprises heating the sample to about 56° C. for about 30 min.
40. A method in accordance with aspect 36, wherein the source of ADAMTS13 is selected from the group consisting of normal plasma, recombinant ADAMTS13, and a combination thereof.
41. A method in accordance with aspect 36, wherein the sample is a serum sample.
42. A method in accordance with aspect 41, wherein the serum sample is an undiluted serum sample.

43. A method in accordance with aspect 41, wherein the serum sample is a concentrated serum sample.
44. A method in accordance with aspect 36, wherein the sample is a plasma sample.
45. A method in accordance with aspect 44, wherein the plasma sample is an undiluted plasma sample.
46. A method in accordance with aspect 44, wherein the plasma sample is a concentrated plasma sample.
47. A method of producing a probe of aspect 1, comprising:

expressing in a cell, an oligopeptide encoded by the vector of aspect 24;

digesting the oligopeptide with His-tagged TEV protease to yield an oligopeptide comprising VWF sequence comprising a cysteine of aspect 5; and

reacting the cysteine with a fluorophore comprising a maleimide, and reacting a primary amine of the oligopeptide such as the N-terminal primary amine with a fluorescence quencher comprising a succinimidyl ester.

#### BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 illustrates an optimized fluorogenic ADAMTS13 substrate set forth as SEQ ID No: 2.

FIG. 2 illustrates FRETs-rVWF71 assays.

FIG. 3 illustrates preparation of FRETs-rVWF71.

FIG. 4 illustrates ADAMTS13 activity toward FRETs-rVWF71.

FIG. 5 illustrates effect of bilirubin and hemoglobin on FRETs-rVWF71.

FIG. 6 illustrates kinetics of FRETs-rVWF71 cleavage by plasma ADAMTS13.

FIG. 7 illustrates plasma concentration dependence of FRETs-rVWF71 assays.

FIG. 8 illustrates plasma ADAMTS13 activity of healthy controls.

#### DETAILED DESCRIPTION

The present inventors have developed fluorogenic substrates which can be used to measure ADAMTS13 activity. An ADAMTS13 substrate of the present teachings comprises an oligopeptide which can be encoded in a vector and expressed in a cell or a microorganism such as *E. coli*. ADAMTS13 substrate of the present teachings can further comprise a fluorophore and fluorescence quencher.

The present inventors have shown that the substrate, FRETs-rVWF71, can be cleaved by ADAMTS13 and that it can be used as an ADAMTS13 substrate without significant interference in undiluted serum or plasma. Contact between a probe of the present teachings and ADAMTS13 can lead to cleavage of the probe which can result in a positive signal, i.e.,

an increase in fluorescence. In various configurations, the increase in fluorescence can be quantified and can serve as a measure of ADAMTS13 activity comprised by a sample. Furthermore, the quantifiable fluorescence signal can be used to measure ADAMTS13 inhibitor activity in a sample.

In various configurations, FRETs-rVWF71 allows ADAMTS13 activity and ADAMTS13 inhibitor activity assays to be performed in undiluted plasma or serum. In various configurations of the present teachings, an assay using FRETs-rVWF71 can be approximately 8-fold more sensitive than an assay using FRETs-VWF73, when using equal concentrations of plasma as the enzyme source. In various configurations, FRETs-rVWF71 can be used in undiluted plasma, which can increase sensitivity another 30-fold relative to FRETs-VWF73, or 240-fold overall. Because the substrate can be used in undiluted plasma, it can be used to detect ADAMTS13 inhibitors that have not been possible to detect with FRETs-VWF73.

In some embodiments, the inventors disclose assays for detecting and quantifying activity of inhibitors of ADAMTS13. In some configurations, an assay of the present teachings can be substantially more sensitive than alternative assays for detecting inhibitors of ADAMTS13. In some configurations, an assay of the present teachings can detect inhibitor activity at <0.5 "Bethesda-like Units" (BU) per mL. For example, a sample from a subject with TTP and undetectable ADAMTS13 activity, and in which a FRETs-VWF73-based assay reported an inhibitor titer  $\leq 1$  U/ml. The same sample tested with a FRETs-rVWF71-based assay of the present teachings revealed an inhibitor titer of 6 U/ml.

In some configurations, sensitivity to inhibitors can be increased by using concentrated recombinant ADAMTS13 as the enzyme source instead of pooled normal plasma.

In some embodiments, an ADAMTS13 substrate of the present teachings can be a probe comprising an oligopeptide consisting of no more than 80 amino acids of sequence of von Willebrand Factor (VWF), wherein said oligopeptide comprises a scissile Y-M peptide, a cysteine substitution located from 1 to 12 amino acids from the scissile Y-M in the carboxy terminal direction, and a C-terminal segment; a fluorophore; and a fluorescence quencher. In some configurations, an oligopeptide of an ADAMTS13 substrate can consist of no more than 73 amino acids. In some configurations, an oligopeptide of an ADAMTS13 substrate can consist of no more than 71 amino acids.

In some configurations, the cysteine substitution can be situated 3 amino acids from the scissile Y-M peptide. In some configurations, the cysteine substitution can be a N1610C substitution of a VWF sequence, wherein the entire sequence of human VWF is:

(Genbank GenBank: AAB59458.1).

(SEQ ID NO: 3)

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1  miparfagvl lalalilpgt lcaegtrgrs starcslfgs dfvntfdgsm ysfagycsyl
61  laggcqkrxf siigdfqngk rvslsvylge ffdihlfvng tvtgqgdrvs mpyaskglyl
121 eteagyykls geaygfvari dsgsnfqvll sdryfnktcg lcgfnfiae ddfmtqegt1
181 tsdpydfans walssgeqwc eraspsssc nissgemqkg lweqcqlks tsvfarchpl
241 vdpepfvalc ektlceagg lecacpalle yartcaqegm vlygwtghsa cspvcpagme

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301 yrqcvspcar tcqslhinem cqercvdgcs cpegqlldeg lcvestecpc vhsgekryppg  
 361 tslsrucentc icrnswicws neecpgeclv tgqshfksfd nryftfsgic qyllardecd  
 421 hsfisivietv qcaddrdavr trsvtvlrpg lhnslvklkh gagvamdgqd iqlpllkgd  
 481 rightvtasv rlsygedlqm dwdgrgrllv klspvyagkt cglcgnyngn qgddfltpsg  
 541 laeprvedfg nawklhgdcq dlqkqhsdpc alnprmrfrs eeacavltsp tfeachrav  
 601 plpylrncry dvscscsdgre clcgalasya aacagrgrvr awrepgrcel ncpkgqvylq  
 661 cgtpcnltrc slsyypeecn eaclegcfcp pglymdergd cvpkaqcpcy ydgeifqped  
 721 ifsdhhtmcy cedgfmhctm sgvpqslldp avlssplshr skrslscrpp mvklvcpadn  
 781 laeaglectk tcqnydlecm smgcvsgclc ppgmvrhenr cvalercpcf hqgkeyapge  
 841 tvkigcntcv crdrkwnctd hvcdatcsti gmahyltfdg lkylfpgecq yvlvqdydcs  
 901 npgtfrilvg nkgcshpsvk ckkrvtilve ggeielfdge vnvkrpmkde thfevvesgr  
 961 yiilllgkal svvwrhlsci svvlkqtyqe kvcglcgnfd gignndltss nlqveedpvd  
 1021 fgnskwvssq cadtrkvpld sspatchnni mkqtmvdssc riltsdvfdq cnklvdpepy  
 1081 ldvcydtcs cesigdcacf cdtiaayahv caqhkgvvtw rtaticpqsc eernlrengy  
 1141 ecewrynsca pacqvctqhp eplacpvqcv egchahcppg kildellqtc vdpedcpvce  
 1201 vagrrfasgk kvtlnpdpe hcqichcdv nltceacqep gglvvpptda pvspttlyve  
 1261 disepplhdf ycsrllldlv lldgssrlse aefevlkafv vdmmerlris qkwrvravve  
 1321 yhdgshayig lkdrkrpsel rriassqvka gsqvastsev lkytlfqifs kidrpeasri  
 1381 allmasqep qrmsrnfvr yvqglkkkvi vipvgigpha nlkqirlied qapenkafvl  
 1441 ssvdeleqqr deivsyldcl apeappptlp phmaqvtvqp gllgvstlqp krnsmlvda  
 1501 fvlegsdkig eadfnrskef meeviqrmdv gqdsihvtvl qsymvtvey pfseaqsqgd  
 1561 ilqrvreiry gqgnrtntgl alrylsdhsf lvsqgdrepa pnlvymvtgn pasdeikrlp  
 1621 gdiqvvpigv gpnanvqele rigwvnapil iqdfetlpre apdlvlqrcc sgegqlqptl  
 1681 spapdcsql dvillldgss sfpasyfdem ksfakafisk anigprltqv svlqygsitt  
 1741 idvpwvvpve kahllslvdv mqreggpsi gdalgfavry ltsemhgap gaskavvilv  
 1801 tdvsvdsda aadaarsrv tvfpigigdr ydaaqlrila gpagsnvvk lqriedlptm  
 1861 vtlgnsflhk lcsqfvricm dedgnekprg dvwtlpdqch tvtcqpdqgt llkshrvncd  
 1921 rglrpscpcns qspvkeetc gcrwtcpcvc tgsstrhivt fdggnfkltg scsyvlfqnk  
 1981 eqdlevilhn gacspgarqg cmksievkhs alsvelhscm evtvngrlvs vpyvggnmev  
 2041 nvygaimhev rfnhlghift ftpqnnfql qlspktfask tyglcgicde ngandfmlrd  
 2101 gtvttldwktl vqewtvqrpg qtcqpilleq clvpdsshcq vlllplfaec hkvlapatfy  
 2161 aicqqdschq eqvceviasy ahlcrtnqvc vdwrtpdfca mscppslvyn hcehgpcprhc  
 2221 dgnvsscghd psegcfcpd kvmlgscvp eeactqcige dgvdhgflea wvpdhqpcqi  
 2281 ctclsrgrkvn cttqpcptak aptcglceva rlrqndqcc peyecvcdpv scdlppvphc  
 2341 erglqptltn pgecrpnftc acrkeckrv sppsccphrl ptlrktqccd eyecacncvn  
 2401 stvscplgyl astatndcgc tttclpdkv cvhrstiyvp gqfweegcdv ctctdmedav  
 2461 mglrvaqcsq kpcedscrsg ftyvlhegec cgrclpsace vvtgsprgds qsswksvgsq  
 2521 waspenpcli necvrveev fiqqrnvscp qlevpvcpsg fqlscktsac cpscrerme  
 2581 acmlngtvig pkgtmidvc ttcrcmvqvg visgfklectr kttcnpcplg ykeenntgec  
 2641 cgrclptact iqlrggqimt lkrdetlqdg cdthfckvne rgeyfwekrv tgcppfdehk

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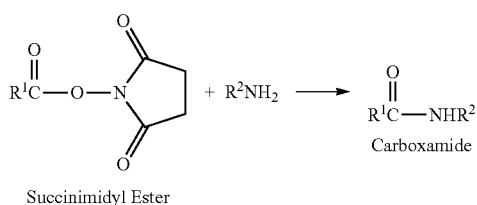
2701 claeqgkink ipgtccdtce epeenditar lqyvkvgsck sevevdihyc qgkaskamy

2761 sidindvqdq cscscptrte pmqvalhctn gsvvyhevl n ameckcsprk csk

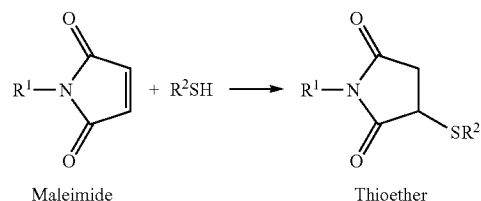
In some aspects, an oligopeptide of the present teachings can comprise, prior to derivatization, a thiol (—SH) such as a cysteine, for example a N1610C substitution (FIG. 1). In various configurations, a thiol such as that of a cysteine can be derivatized with a thiol-reactive moiety such as a maleimide moiety of a fluorophore or a fluorescence quencher. In some aspects, prior to derivatization, an oligopeptide of the present teachings can have a single thiol; in various configurations, this thiol can be available for derivatization.

In some aspects, an oligopeptide of the present teachings can comprise, prior to derivatization, at least one amine such as an amino terminal primary amine, or an amino acid comprising a primary amine such as a lysine. An amine of an oligopeptide can be available for derivatization with an amine-reactive moiety such as an N-hydroxysuccinimide (NHS) moiety of a fluorophore or a fluorescence quencher. In some aspects, prior to derivatization, an oligopeptide can have a single primary amine that is available for derivatization. In some aspects, an oligopeptide of the present teaching can include a substitution of a lysine of wild type VWF, such as a lysine K1617 for an amino acid that does not comprise a primary amine on its side chain, such as a K1617R substitution (FIG. 1).

In some configurations, an oligopeptide can comprise a substitution of E1598G, which can provide an amino terminal end after release from a precursor. These mutations can result in an oligopeptide having one thiol and one primary amine, which can be available for addition of one fluorophore and one quencher to the oligopeptide (FIG. 1). In various embodiments, these reactive sites can be located on opposite sides of a protease cleavage target site such as the Y-M peptide of von Willebrand Factor. In various configurations, the succinimidyl ester and the maleimide can react through the reactions below, wherein R<sup>1</sup> can be a fluorophore or fluorescence quencher and R<sup>2</sup> can be an oligopeptide.



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Accordingly, as used herein, oligopeptides reacted with such reactive precursors may not include a reactive moiety such as a succinimide in a final product. In various configurations of the present teachings, prior to cleavage of an oligopeptide comprising both a fluorophore and a quencher, fluorescence is quenched. When the Tyr1605-Met1606 bond is cleaved, quenching is relieved, thereby resulting in a fluorescence signal. In various configurations, the fluorescence signal can be proportional to product concentration. In various configurations, a fluorescence signal can be detected by means well known to skilled artisans, such as but not limited to fluorescence spectrometry.

In some configurations, a probe can comprise an oligopeptide consisting of a sequence set forth as SEQ ID NO: 1:

DEREQPNLVYVMTGNPASDEIKRLPGDIQVVP IGVGPNANVQELERIGW  
PNAPILIQDFETLPREAPDLVLQR.

In some configurations, a probe can comprise an oligopeptide consisting of a sequence set forth as SEQ ID NO: 2 and as FRETs-rVWF71 (FIG. 1):

GQAPNLVYVMTGCPASDEIRRLPGDIQVVP IGVGPNANVQELERIGWPN  
APILIQDFETLPREAPDLVLQR.

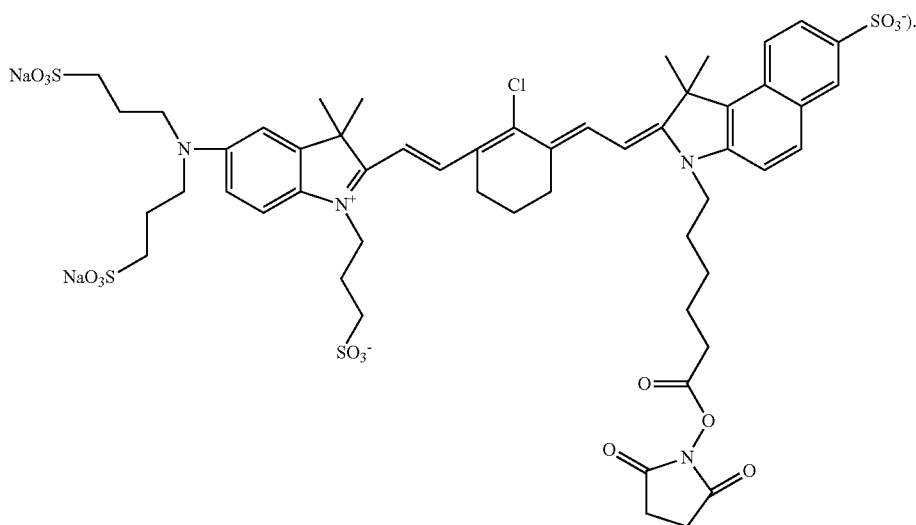
In various aspects, a fluorophore can have an absorption maximum of >550 nm or >630 nm. In various aspects, a fluorophore can have an emission maximum of >600 nm or >650 nm. In some aspects, a fluorophore, prior to reaction with an oligopeptide, can be, without limitation, an Alexa Fluor® 594 maleimide (Life Technologies Corp. Carlsbad, Calif.) (abs 590 nm, em 617 nm, ε 96,000) or a DyLight® 633 maleimide (Thermo, abs 638 nm, em 658 nm, ε 170,000).

In various aspects, a quencher can have an absorption maximum of >550 nm or >630 nm. In some aspects, a quencher, prior to reaction with an oligopeptide, can be, without limitation, a QSY21-succinimidyl ester (abs 661 nm, ε 90,000) or an IRDye QC-1 N-hydroxy succinimidyl ester (LI-COR, Lincoln, Nebr.; abs 737 nm, ε 96,000,



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In some aspects, a substrate of the present teachings can comprise a fluorophore or quencher that comprises at least one sulfate. In some aspects, a substrate of the present teachings can be soluble in water at concentrations  $>50 \mu\text{M}$ , or greater than that of FRETTS-VWF73.

In some embodiments, the present teachings include a vector comprising a nucleic acid sequence encoding an oligopeptide of an ADAMTS13 substrate described herein. In some aspects, a nucleic acid sequence can further encode a His tag such as an N-terminal His tag. In some aspects, a nucleic acid sequence can further comprise a sequence encoding thioredoxin. In some aspects, a nucleic acid sequence can further comprise a sequence encoding a Tobacco Etch Virus (TEV) protease cleavage site. In some aspects, a vector can be a plasmid or a virus, such as a bacteriophage.

In some embodiments, the present teachings include methods of determining presence, absence or quantity of ADAMTS13 activity in a sample. In various configurations, such methods can comprise: forming a mixture comprising a sample and a probe as described herein and measuring fluorescence at one or more time points after forming the mixture.

In some embodiments, the present teachings include methods of determining presence, absence or quantity of ADAMTS13 inhibitor activity in a sample. In various configurations, such methods can comprise: forming a mixture comprising a sample and a probe as described herein and measuring fluorescence at one or more time points after forming the mixture. Such methods can further comprise including in the mixture ADAMTS13 which can be obtained from a source such as, without limitation, ADAMTS13 produced in a microorganism using recombinant methods, or ADAMTS13 present in serum or plasma, such as normal pooled serum or plasma.

In some embodiments, the present teachings include methods of producing a probe described herein. In some configurations, these methods can comprise expressing, in a cell, an oligopeptide encoded by the vector of described herein, digesting the oligopeptide with His-tagged TEV protease to yield an oligopeptide comprising VWF sequence comprising a cysteine as described herein and reacting the cysteine with a fluorophore comprising a maleimide, and reacting a primary amine of the oligopeptide such as the N-terminal primary amine with a fluorescence quencher comprising a succinimidyl ester.

In some embodiments, the present teachings include methods of producing a probe described herein. In some configurations, these methods can comprise expressing, in a cell, an oligopeptide encoded by the vector of described herein, digesting the oligopeptide with His-tagged TEV protease to yield an oligopeptide comprising VWF sequence comprising a cysteine as described herein and reacting the cysteine with a fluorescence quencher comprising a maleimide, and reacting a primary amine of the oligopeptide such as the N-terminal primary amine with a fluorophore comprising a succinimidyl ester or another amine-reactive moiety such as an isothiocyanate.

## EXAMPLES

The following Examples are intended to be illustrative of various aspects of the present teachings and are not intended to be limiting of any aspect. While some of examples may include conclusions about the way the invention may function, the inventors do not intend to be bound by those conclusions, but put them forth only as possible explanations. Unless indicated by use of past tense, presentation of an example does not imply that an experiment or procedure was, or was not, conducted, or that results were, or were not, actually obtained.

The methods and compositions described herein utilize laboratory techniques well known to skilled artisans, and can be found in laboratory manuals such as Sambrook, J., et al., *Molecular Cloning: A Laboratory Manual*, 3rd ed. Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N.Y., 2001; Spector, D. L. et al., *Cells: A Laboratory Manual*, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N.Y., 1998; Ausubel, F. M., et al., ed., *Current Protocols in Molecular Biology*, Wiley Interscience, 2003; Nagy, A., et al., *Manipulating the Mouse Embryo: A Laboratory Manual* (Third Edition), Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N.Y., 2003; Hedrickson et al., *Organic Chemistry* 3rd edition, McGraw Hill, New York, 1970; Carruthers, W., and Coldham, I., *Modern Methods of Organic Synthesis* (4th Edition), Cambridge University Press, Cambridge, U.K., 2004; Graham Solomons T. W., et al., *Organic Chemistry* 9th edition, Wiley, John & Sons, Incorporated, 2007.

As used herein, the singular forms "a", "an" and "the" are intended to include the plural forms as well, unless the context indicates otherwise. Methods of administration of pharmaceuticals and dosage regimes can be determined according

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to standard principles of pharmacology well known skilled artisans, using methods provided by standard reference texts such as Remington: the Science and Practice of Pharmacy (Alfonso R. Gennaro ed. 19th ed. 1995); Hardman, J. G., et al., Goodman & Gilman's The Pharmacological Basis of Therapeutics, Ninth Edition, McGraw-Hill, 1996; and Rowe, R. C., et al., Handbook of Pharmaceutical Excipients, Fourth Edition, Pharmaceutical Press, 2003.

These and all other publications cited in this disclosure are incorporated herein by reference, each in its entirety.

## Example 1

This example illustrates a substrate oligopeptide.

The inventors designed a chimeric construct in plasmid pET-32 Xa/LIC (Novagen, Billerica, Mass.) encoding an N-terminal His-tag, thioredoxin, Tobacco Etch Virus (TEV) protease cleavage site, a Gly residue, and VWF (Gln<sup>1599</sup>-Arg<sup>1668</sup>). Two mutations were made in the VWF segment: N1610C introduced a thiol group for alkylation, and K1617R removed an amino group that competes with the peptide N-terminus for chemical modification. This oligopeptide was expressed in IPTG-induced BL21 cells, purified on Ni-NTA-agarose, and digested with His-tagged TEV protease (made by the present inventors). TEV protease and thioredoxin were removed on Ni-NTA-agarose, and the C-terminal peptide (FIG. 1) was purified by C18 reverse phase HPLC. >100 mg of this peptide was prepared, sufficient for thousands of assays.

In FIG. 1, the scissile Y-M bend is indicated (triangle) and the C-terminal segment of VWF domain A2 which can promote efficient cleavage by ADAMTS13 consists of the carboxy terminal sequence EAPDLVLQR (SEQ ID NO: 6) (underlined), Residues mutated in von Willebrand Disease type 2A, associated with increased proteolysis, are circled. FRETs-rVWF71 has the mutations E1598G, N1610C, and K1617R (boxed). Cys1610 is modified with DyLight 633 and the N-terminus is modified with IRDye QC-1.

## Example 2

This example illustrates fluorescence donors and quenchers.

The present inventors have tested several combinations of donor and quencher groups chosen for lack of interference from blood proteins. For example, modification at Cys<sup>1610</sup> with Alexa Fluor 594 maleimide (abs 590 nm, em 617 nm,  $\epsilon$  96,000) and at N-terminal Gly with QSY21-succinimidyl ester (abs 661 nm,  $\epsilon$  90,000) gave superior fluorescence quenching and sensitivity compared to FRETs-VWF73, but this substrate had poor solubility. In one embodiment, a substrate of the present teachings is modified at Cys<sup>1610</sup> with DyLight 633-maleimide (Thermo, abs 638 nm, em 658 nm,  $\epsilon$  170,000), and at N-terminal Gly with IRDye QC-1 N-hydroxysuccinimidyl ester (LI-COR, abs 737 nm,  $\epsilon$  96,000). These dyes incorporate sulfate groups and have markedly increased water solubility. After RP-HPLC purification the doubly-labeled substrate (FRETs-rVWF71) is soluble at >50  $\mu$ M. These dyes absorb/emit in the near-infrared, which does not overlap with the spectrum of blood proteins, hemoglobin, or bilirubin.

## Example 3

This example illustrates ADAMTS13 activity assays.

Preliminary studies show that paired samples of citrated plasma and serum have the same ADAMTS13 activity, which is stable for extended times at 4° C. or -20° C. These findings

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are consistent with published data (Gerritsen, H. E., et al., Blood. 98:1654-1661, 2001; Furlan, M., et al., Blood. 87:4223-4234, 1996).

In these experiments, the FRETs-rVWF71 substrate was utilized in a microtiter format (FIG. 2), which is compatible with fluorescence plate readers that are supported by laboratory equipment suppliers and available in many clinical hemostasis laboratories, such as the BIO-TEK FLx800 Fluorometer from DiaPharma (West Chester, Ohio) or Technoclone (Vienna). Reactions were performed in 200  $\mu$ L 5 mM Bis-Tris, pH 6, 25 mM CaCl<sub>2</sub>, at room temperature with added plasma as shown (FIG. 2). Fluorescence was monitored in a Victor<sup>2</sup> V microplate reader (Perkin Elmer Life Sciences, Boston, Mass.) with 638 nm excitation filter and 658 nm emission filter.

Using 1  $\mu$ M substrate with varying amounts of plasma or serum and optimal filters for each substrate,  $\Delta F$ /min for VWF-rFRETs71 is 8-fold greater compared to VWF-FRETs73. In addition, assays with VWF-rFRETs71 are linear with time and linear with enzyme up to >95% serum or plasma added, with no significant background fluorescence or interference from blood constituents. The combination of stronger fluorescence signal and compatibility with undiluted plasma make FRETs-rVWF71 ~250-fold more sensitive than FRETs-VWF73.

## Example 4

This example illustrates ADAMTS13 inhibitor assays.

In this prophetic example, activity of ADAMTS13 inhibitors is examined. In these experiments, antibody source can be patient serum, optionally heat-treated to inactivate endogenous ADAMTS13 (56° C., 30 min) (Zheng, X. L., et al., Blood. 103:4043-4049, 2004). The ADAMTS13 source is normal pooled serum (or plasma), or recombinant ADAMTS13 to further increase sensitivity. In these assays, a fixed amount of ADAMTS13 (5  $\mu$ L) and variable amounts of inhibitor (90  $\mu$ L serum and serial dilutions) are mixed and preincubated (15 min). FRETs-rVWF71 is added (5  $\mu$ L) and rates of product generation are analyzed by standard methods to yield an inhibitor titer in "Bethesda-like" units. The assay permits the detection of inhibitors at <<0.5 "BU" per mL.

## Example 5

This prophetic example illustrates an inhibitor assay based on FRETs-rVWF71 cleavage in minimally diluted Li<sup>+</sup>-heparin plasma.

We have designed an inhibitor assay based on FRETs-rVWF71 cleavage in minimally diluted Li<sup>+</sup>-heparin plasma. The inhibitor source is patient plasma, optionally heat-treated to inactivate endogenous ADAMTS13 (56° C., 30 min). Equal amounts of pooled normal plasma and inhibitor (and serial dilutions) are mixed and preincubated. Samples (100  $\mu$ L) are assayed with FRETs-rVWF71 (200  $\mu$ L final volume) and rates of product generation are analyzed to yield an inhibitor titer in "Bethesda-like" units.

## Example 6

This example illustrates ADAMTS13 activity assays.

In these experiments (FIG. 7A), reactions were performed in 200  $\mu$ L 50 mM HEPES, pH 7.4, 150 mM NaCl, 10 mM CaCl<sub>2</sub>, 100  $\mu$ g/ml BSA, 0.05% Tween 20, at 30° C. with added plasma, in a Perkin Elmer Victor2V plate reader with 635/15 nm excitation and 660/10 nm emission filters. FIG. 7B shows initial slope vs. PNP standard curve.

After RP-HPLC purification, the doubly-labeled substrate "FRETs-rVWF71" is soluble at >200  $\mu$ M. In addition, assays with FRETs-rVWF71 were insensitive to >20 mg/dL (250

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$\mu\text{M}$ ) conjugated bilirubin, which is incompatible with FRET5-VWF73 (Meyer, S. C., et al., *J. Thromb. Haemost.* 5: 866-867, 2007).

In these experiments, samples were analyzed from 100 healthy controls. The mean ADAMTS13 activity was 1.11 U/mL with SD 0.21 U/dL (referenced to local pooled normal plasma). Intra-assay and inter-assay coefficients of variation were <3%. Results obtained with FRET5-rVWF71 and FRET5-VWF73 were congruent, with a coefficient of variation of 3.8% for healthy controls assayed by both methods.

#### Example 7

This example illustrates preparation of fluorogenic FRET5-rVWF71 substrate.

In these experiments, a precursor substrate peptide was expressed from a plasmid that encodes thioredoxin, a His-tag, a TEV cleavage site, and VWF Gln<sup>1599</sup>-Arg<sup>1668</sup>. The mutation K1617R introduced a unique Cys, and the mutation K1617R removed a primary amine that otherwise would have competed with subsequent chemical modification of the N-terminus. The expressed chimeric protein was readily soluble and was purified by chromatography on Ni-NTA agarose (FIG. 3) with a yield of >50 mg/L of LB culture medium. Removal of thioredoxin by cleavage with His-tagged TEV protease gave the product rVWF71, which has an extra Gly before VWF residues Gln<sup>1599</sup>-Arg<sup>1668</sup>. Peptide rVWF71 was purified by Ni-NTA chromatography to remove His-tagged thioredoxin and TEV protease, followed by reverse-phase HPLC (FIG. 3) to yield >10 mg rVWF71/L of LB culture medium.

FIG. 3A shows polyacrylamide 4-12% gel electrophoresis of intermediates and purified FRET5-rVWF71. Lanes correspond to: 1, protein markers; 2, thioredoxin-VWF71 fusion protein eluted from Ni-NTA agarose; 3, after cleavage by TEV protease; 4, unbound products after rechromatography on Ni-NTA agarose; 5, after purification by HPLC; 6, rVWF71 peptide modified with DyLight 633 and purified by HPLC; 7, purified FRET5-rVWF71; 8, purified FRET5-rVWF71. Except for lanes 6 and 7, which are unstained, gels were stained with Simply Blue SafeStain.

FIG. 3B shows purification of rVWF71 peptide. In these experiments, thioredoxin-VWF71 fusion protein was digested with TEV protease, chromatographed on Ni-NTA agarose to remove His-tagged TEV protease and thioredoxin, and purified by HPLC on a C18 column in 0.1% TFA developed with a 20-90% acetonitrile gradient. The eluate was monitored for absorbance at 280 nm. The peak at 20 min corresponds to panel A, lane 5.

FIG. 3C shows purification of singly modified rVWF71. In these experiments, DyLight 633-rVWF71 was purified by HPLC on a C18 column in 50 mM TEAA, pH 6.0, developed with a 20-45% acetonitrile gradient. The eluate was monitored for absorbance at 280 nm and 627 nm, and for emission at 658 nm after excitation at 635 nm. The inset shows an expanded view of the peak at 23 min with traces labeled by wavelength.

FIG. 3D shows purification of FRET5-rVWF71. In these experiments, after reaction with IRDye QC-1 N-hydroxy-succinimidyl ester, FRET5-rVWF71 was purified by HPLC as described for DyLight 633-rVWF71 (panel C) with additional monitoring of absorbance at 819 nm. The inset shows an expanded view of the peak at 21 min with traces labeled by wavelength.

In these experiments, DyLight 633 maleimide (Thermo Scientific, Waltham, Mass.) 1 mg/100  $\mu\text{L}$  in dimethyl sulfoxide (DMSO) was added dropwise with stirring to peptide VWF71 (10 mg) in  $\leq 2$  mL of 100 mM sodium phosphate, pH 7.1, in the dark, and stirred overnight at room temperature. The product DyLight 633-rVWF71 was purified by HPLC on

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C18 using the TEAA/acetonitrile solvent system as described for the purification of VWF71. DyLight 633-rVWF71 was lyophilized and desalted on a small column of PD-10 in  $\leq 2$  mL of 100 mM sodium phosphate, pH 7.9. IRDye QC-1 N-hydroxysuccinimide ester 0.5 mg in 100  $\mu\text{L}$  DMSO was added dropwise with stirring and the solution was stirred overnight at room temperature protected from light. The product FRET5-rVWF71 was purified by reverse phase HPLC, lyophilized, dissolved in  $\leq 0.5$  mL of deionized water and applied onto a column (7 $\times$ 230 mm) of Amberlite IR120 sodium form. FRET5-rVWF71 product was eluted with deionized water and lyophilized or concentrated by ultrafiltration to  $\geq 250$   $\mu\text{M}$ . The concentration was verified by amino acid analysis (The Protein Chemistry Laboratory, Texas A&M University). Working stocks stored at  $-20^\circ\text{C}$ . were thawed and refrozen repeatedly over several months without any change in stability or chemical properties.

The rVWF71 was modified with DyLight 633 maleimide in  $\sim 70\%$  yield and purified by HPLC (FIG. 3). The rVWF71 had been reduced with 10 mM dithiothreitol before purification by HPLC and buffer exchange to maximize the efficiency of alkylation by DyLight 633 maleimide. The N-terminal Gly of DyLight 633-rVWF71 was modified with IRDye QC-1 N-hydroxy-succinimidyl ester in  $\sim 90\%$  yield and purified by HPLC. The final FRET5-rVWF71 product was converted into the sodium salt, which was stable and soluble at >250  $\mu\text{M}$ .

Absorbance maxima for FRET5-rVWF71 were observed at 627 nm and 819 nm, consistent with the presence of both dyes. Upon excitation at 635 nm, the fluorescence emission at 660 nm of uncleaved FRET5-rVWF71 and fully cleaved FRET5-rVWF71 showed little dependence on pH, varying <7% between pH 5 and pH 10.

#### Example 8

This example illustrates the optimization of FRET-rVW71 cleavage by plasma ADAMTS13.

In these experiments, matched samples of serum, and plasma anticoagulated with sodium citrate, Li<sup>+</sup>-heparin and Na<sup>+</sup>-heparin were obtained from volunteer healthy donors with informed consent according to a human studies protocol approved by the Washington University Institutional Review Board.

Assays were performed at 30 $^\circ\text{C}$ . Samples (100  $\mu\text{L}$ ) of plasma or serum, diluted as necessary in assay buffer, were pipetted in duplicate in 96 well white microplates (Optiplate-96, PerkinElmer, Waltham, Mass.). Reaction was initiated by addition of 100  $\mu\text{L}$  substrate in assay buffer. Cleavage of FRET5-rVWF71 was detected as an increase in fluorescence compared to control reactions without added enzyme at 2 min intervals using a Victor<sup>2</sup>V Multilabel Counter (PerkinElmer, Waltham, Mass.) or Synergy H1 Hybrid Multi-Mode Microplate Reader (Biotek, Winooski, Vt.) equipped with 635 $\pm$ 10 nm excitation and 660 $\pm$ 10 nm emission filters. Initial velocities were determined by fitting progress curves to the polynomial  $\Delta F = A + Bt + Ct^2$ , where  $\Delta F$  is the change in fluorescence,  $t$  is time,  $A$  is the y-intercept,  $B$  is the initial velocity (slope), and the term  $Ct^2$  accounts for any decline in velocity due to substrate consumption or photobleaching.

ADAMTS13 assays with the FRET5-VWF73 substrate (Kokame, K., et al. *Br. J. Haematol.* 129:93-100, 2005) (Peptide International, Louisville, Ky.) were performed according to the manufacturer's instructions. Hemoglobin was prepared by hypotonic lysis of red blood cells from a voluntary donor and quantified with Drabkin's reagent (Sigma, St. Louis, Mo.).

Optimal conditions for cleavage of FRET5-rVWF71 were similar to those reported for ADAMTS13 cleavage of other substrates (Kokame, K., et al. *Br. J. Haematol.* 129:93-100,

2005; Tsai, H. M., et al. *Blood*. 87:4235-44, 1996; Anderson, P. J., et al. *J. Biol. Chem.* 281:850-7, 2006). Activity was maximal at pH 6, very low ionic strength, and 5-10 mM CaCl<sub>2</sub> (FIG. 4). Under conditions of physiological ionic strength (150 mM NaCl) and pH 7.4, the rate of reaction was decreased ~50%.

To optimize compatibility with minimally diluted plasma samples, the standard assay buffer was made 50 mM HEPES, pH 7.4, 150 mM NaCl, and 10 mM CaCl<sub>2</sub>, supplemented with 0.05% Tween-20 and 400 µg/mL bovine serum albumin. Under these conditions, assays of serum or plasma anticoagulated with citrate, Li<sup>+</sup>-heparin or Na<sup>+</sup>-heparin gave comparable results. Values for citrated plasma were decreased as expected from dilution by the citrate anticoagulant solution (FIG. 4). Li<sup>+</sup>-heparin plasma is commonly used for clinical chemistry assays and therefore was selected for ADAMTS13 assay development.

Bilirubin interferes with ADAMTS13 assays that use the FRETTS-VWF73 substrate because bilirubin absorbs light at the same wavelengths as the chromophores in the substrate (Hovinga, J. A. K., et al. *J. Thromb. Haemost.* 5:866-7, 2007). However, the spectrum of bilirubin does not overlap with DyLight 633 and IRDye QC-1, and bilirubin ≤20 mg/dL did not inhibit ADAMTS13 activity assays with FRETTS-rVWF71 (FIG. 5). Hemoglobin absorbs at 550 nm and also interferes with FRETTS-VWF73 assays. In addition, hemoglobin directly inhibits ADAMTS13 regardless of the assay method (Studt, J. D., et al. *Blood*. 105:542-4, 2005). As expected, hemoglobin ≤20 g/L did not affect the detection of FRETTS-rVWF71 cleavage products, but did inhibit ADAMTS13 activity with an IC<sub>50</sub> of 10-15 g/L (FIG. 5).

#### Example 9

This example illustrates the kinetics of FRETTS-rVWF71 cleavage.

In these experiments, FRETTS-rVWF71 cleavage by plasma ADAMTS13 (50 µL) was assessed in 200 µL reactions containing 50 mM HEPES, pH 7.4, 10 mM CaCl<sub>2</sub>, 150 mM NaCl, and 0.05% Tween-20, and varying concentrations of FRETTS-rVWF71. The concentration of ADAMTS13 in PNP is ~1.03 µg/ml (Feys, H. B., et al. *J. Thromb. Haemost.* 4:955-62, 2006), or ~6 nM for a molecular mass of 170 kDa. For each concentration of substrate, the relationship between product fluorescence and concentration was determined directly by cleaving FRETTS-rVWF71 to completion with excess recombinant ADAMTS13 MDTCS (Feys, H. B., et al. *J. Thromb. Haemost.* 7:2088-95, 2009) and measuring the fluorescence. Control assays contained plasma that was completely deficient in ADAMTS13. The initial velocities (nM/min) as a function of substrate concentration were fitted to the Michealis-Menten equation by nonlinear regression analysis (Prism, GraphPad).

Plasma ADAMTS13 cleaved FRETTS-rVWF71 with a  $K_m$  of 1.8 µM and a  $k_{cat}$  of 6.8 min<sup>-1</sup> at 30° C. (FIG. 6). For comparison, ADAMTS13 cleaves FRETTS-VWF73 in the same buffer with a  $K_m$  of 3.2 µM and a  $k_{cat}$  of 58 min<sup>-1</sup> at 37° C. (Anderson, P. J., et al. *J. Biol. Chem.* 281:850-7, 2006). Therefore, the larger dyes of FRETTS-rVWF71 do not impair substrate binding to ADAMTS13 but decrease the rate of catalysis ~8-fold.

#### Example 10

This example demonstrates FRETTS-rVWF71 assay performance.

Progress curves for cleavage of FRETTS-rVWF71 by plasma ADAMTS13 were approximately linear with time for

at least 60 minutes (FIG. 7A). To account for small time-dependent decreases in reaction rate with high concentrations of enzyme, initial rates were obtained by fitting to a second order polynomial. Reaction rate increased approximately linearly with the volume of added plasma. For maximum sensitivity reactions were performed with 100 µL plasma in a reaction volume of 200 µL. Standard calibration curves were constructed with PNP (FIG. 7B).

Results with FRETTS-rVWF71 and FRETTS-VWF73 assayed under standard conditions (Kokame, K., et al. *Br. J. Haematol.* 129:93-100, 2005) correlated well with an inter-assay CV of 3.8%. Using 1 µM substrate and 5 µL PNP, the change in fluorescence (ΔF/min) for FRETTS-rVWF71 (50 mM HEPES, pH 7.4, 150 mM NaCl) was 8-fold greater than for FRETTS-VWF73 (5 mM Bis-Tris, pH 6).

The FRETTS-rVWF71 assay for ADAMTS13 activity has been adapted to measure autoantibody inhibitors of ADAMTS13 in a manner analogous to the measurement of factor VIII inhibitors in "Bethesda-like" units.

Plasma samples with ADAMTS13 protease activity <10% were tested for the presence of an ADAMTS13 inhibitor. Plasma samples were serially diluted two-fold with assay buffer (50 mM HEPES, pH 7.4, 150 mM NaCl, 10 mM CaCl<sub>2</sub>, and 0.05% Tween-20) to yield 8 dilutions. In duplicate, PNP (50 µL) was mixed with an equal volume (50 µL) of undiluted test plasma and each serially diluted sample. Control conditions included PNP mixed with an equal volume of cleavage buffer. The microplate containing the samples was sealed with adhesive film (Sealplate, Excel Scientific, Victorville, Calif.) and incubated at 37° C. for 1 h. Reaction was initiated by addition of 100 µL of assay buffer containing 2 µM FRETTS-rVWF71. Fluorescence emission was monitored and initial reaction rates were determined.

The inhibitor titer was determined by fitting the initial reaction rates to a four parameter logistic model or sigmoidal dose response equation by nonlinear regression (Prism):

$$\text{Rate}(D) = \text{Max} + (\text{Max} - \text{Min}) / (1 + 10^{((\log T - \log D) * H)})$$

Where Rate(D) is the initial rate with test plasma at dilution D, Max is the maximum rate in the absence of test plasma, Min is the rate with no active enzyme (fixed at 0), H is the Hill slope, and T is the inhibitor titer.

For selected patient samples, ADAMTS13 activity and inhibitor titer were measured with FRETTS-VWF73 at the BloodCenter of Wisconsin reference laboratory.

A fixed amount of ADAMTS13 in PNP (100 µL) was preincubated with serially diluted plasma samples and assayed for ADAMTS13 activity. Reaction rates were analyzed according to a model for competitive sigmoidal dose-response inhibition to obtain the dilution of plasma at which ADAMTS13 activity is decreased by 50%, and the inverse of the dilution is the inhibitor titer (FIG. 7C). This assay design allows the detection of inhibitors with a titer <0.5 U/ml.

#### Example 11

This example illustrates ADAMTS13 assays in healthy donors and patients with TTP.

In these experiments, frozen Li<sup>+</sup>-heparin plasma samples were obtained from 100 healthy controls (Biological Specialty Corp., Colmar, Pa.) deidentified except for demographic information on gender, age, and ethnicity. Pooled normal Li<sup>+</sup>-heparin plasma (PNP) for assay standardization was prepared from at least 35 donors. For each healthy control, 25 µL and 100 µL plasma samples were assayed in duplicate for ADAMTS13 activity. Up to 20 healthy controls were assayed per 96 well microplate. For cross-validation, ≥3

samples were randomly selected from each assayed batch and reanalyzed with the subsequent batch. A standard curve was constructed from duplicate assays of PNP 5  $\mu$ L, 25  $\mu$ L, 50  $\mu$ L, 75  $\mu$ L and 100  $\mu$ L. ADAMST13 activity of randomly selected controls samples was determined by the FRETTS-VWF73 method.

Li<sup>+</sup>-heparin plasma was obtained from individuals suspected to have TPP with their informed consent according to a human studies protocol approved by the Washington University Institutional Review Board. Anticoagulated blood was centrifuged at 2000 $\times$ g for 10 min at 10° C., and the blood cells debris were discarded. The supernatant was assayed immediately and the remainder stored in aliquots at -80° C.

ADAMST13 activity was assayed in Li<sup>+</sup>-heparin plasmas from 96 healthy controls (FIG. 8). Using a PNP standard prepared from 35 donors, the mean ADAMTS13 activity was 107 $\pm$ 18% (SD). Intra-assay and inter-assay coefficients of variation were less <2%. Mean ADAMTS13 activity was not significantly different based on gender or ethnicity (FIG. 8). There was no significant relationship between ADAMTS13 activity and age.

ADAMTS13 assays using both FRETTS-rVWF71 and FRETTS-VWF73 substrates were performed on samples from several patients with idiopathic TTP (Table 1). Both substrates gave consistent results for ADAMTS13 activity, although FRETTS-rVWF71 assays were 20-fold more sensitive. Three patients did not have detectable ADAMTS13 inhibitors by either assay. For the remaining patients, inhibitor assays with FRETTS-rVWF71 gave titers 1.4-fold to >5-fold higher than inhibitor assays with FRETTS-VWF73. This difference in sensitivity for inhibitors reflects the 1:20 dilution of plasma required for assays with FRETTS-VWF73.

TABLE 1

Comparison of ADAMTS13 assays for patients with TTP				
Patient	ADAMTS13 activity (%)		ADAMTS13 Inhibitor (units/ml)	
	FRETTS-rVWF71	FRETTS-VWF73	FRETTS-rVWF71	FRETTS-VWF73
UPN 323	7.6	<5	3.8	1.0
UPN 330	1.8	<5	4.5	1.8
UPN 333	2.2	<5	1.9	0.9
UPN 334	0.2 <sup>a</sup>	<5 <sup>b</sup>	23 <sup>a</sup>	3.6 <sup>b</sup>
UPN 335	10	<5	<0.4	<0.4
UPN 336	1.8	<5	3.1	0.6
UPN 337	1.2	<5	<0.4	<0.4
UPN 339	30	14	<0.4	<0.4
UPN 340	1.9	<5	<0.4	1.8
UPN 342	9.5	7	2.7	1.9
UPN 346	3.3	<5	ND	ND

<sup>a</sup>Sample of Sep. 5, 2011.

<sup>b</sup>Sample of Sep. 13, 2011.

ND, not done.

### Example 12

This example illustrates construction of a vector of the present teachings.

To construct a plasmid comprising a sequence encoding amino acid residues Gln1599-Arg1668 (VWF70) of von Willebrand factor (VWF), DNA sequence of VWF was amplified from pSVHvWF1 (Matsushita, T., and Sadler, J. E. J. Biol. Chem. 270:13406-14, 1995) using primers with Ligation Independent Cloning overhangs (Aslanidis, C. and de Jong, P. J. Nucleic. Acids. Res. 18:6069-74, 1990):

Forward,  
(SEQ ID NO: 4)  
GGTAATGAGGGTCGCGAGAACCTTTATTTCAGGGCCAGGCGCCC

Reverse,  
(SEQ ID NO: 5)  
AGAGGAGAGTTAGAGCCTCACCTCTGCAGCACCAGGTC

The forward primer encodes a tobacco etch virus (TEV) protease cleavage site and the reverse primer introduces a stop codon. The PCR product was purified and ligated into pET-32 Xa/LIC (Novagen, Billerica, Mass.) to yield a plasmid that encodes thioredoxin, a His-tag, a TEV cleavage site, a Gly residue, and VWF Gln1599-Arg1668. The mutations D1610C and K1617R were introduced using a site-directed mutagenesis kit (Stratagene, Santa Clara, Calif.) to yield plasmid pET32XaTEVvWF70. The sequence was confirmed by ABI BigDye V3.1 terminator cycle sequencing.

### Example 13

This example illustrates recombinant peptide preparation.

In these experiments, plasmid pET32XaTEVvWF70 was transformed into *E. coli* BL21 (DE3). Single colonies were inoculated in 5 mL of LB medium containing 50  $\mu$ g/mL ampicillin and grown for 5 h at 37° C. with shaking at 250 rpm. While maintaining the antibiotic and growth conditions, the 5 mL culture was transferred into 50 mL of LB for 1 h, after which the 50 mL culture was transferred into 1 L of LB. When the culture reached optical density at 600 nm of ~0.6-0.7, isopropyl  $\beta$ -D-1-thiogalactopyranoside (IPTG) was added to 1 mM and incubation was continued for 4 hours. The cell pellet was obtained by centrifugation at 8,000 $\times$ g for 15 min at 4° C. and stored at -80° C. if not processed immediately.

The cell pellet was thawed in water at room temperature for 10 min and resuspended in 10 mL/g pellet of B-PER Protein Extraction Reagent (Pierce), 10  $\mu$ L/mL benzonase (Sigma, St. Louis, Mo.) 10  $\mu$ L/mL, Halt Protease Inhibitor Cocktail (Pierce) and 1 mM phenylmethylsulfonyl fluoride. The suspension was incubated for 15 min at room temperature with gentle shaking (Orbitron Rotator I, Boeckel Scientific). The supernatant was recovered by centrifugation at 20,000 $\times$ g for 15 min at 4° C., diluted with an equal volume of His-binding buffer (20 mM sodium phosphate, pH 7.4, 500 mM NaCl, and 10 mM imidazole), and applied to a 5 mL column of nickel-nitrilotriacetic acid (Ni-NTA) agarose beads (Agarose Beads Technologies, Spain) equilibrated with His-binding buffer. The Ni-NTA column was washed with 50 mL of His-binding buffer and bound protein was eluted with 300 mM imidazole in His-binding buffer. The eluted protein was concentrated to 10 mL by ultrafiltration and dialyzed in a Slide-A-Lyzer cassette (MWCO 7,000, Pierce) overnight with two buffer changes against TEV protease cleavage buffer (50 mM Tris-HCl, pH 8.0, 1 mM dithiothreitol, and 0.5 mM EDTA).

The protein concentration was determined by the BCA method (BioRad, Hercules, Calif.) and the affinity tags cleaved by adding 1 mg of His-tagged TEV protease (Kapust, R. B. and Waugh, D. S. Protein. Sci. 8:1668-74, 1999) per 10 mg of protein and incubation for 16-20 h at room temperature. The solution was dialyzed overnight against His-binding buffer in a Slide-A-Lyzer cassette (MWCO 3,000), and cleaved affinity tags and TEV protease were removed by adsorption on Ni-NTA agarose. Dithiothreitol was added to 10 mM and the unbound product was further purified by HPLC on a C18 column (300  $\text{Å}$ , 5  $\mu$ m, 150 $\times$ 10 mm, GraceVydac) equilibrated with buffer A (50 mM triethylammonium

acetate (TEAA), pH 6.0) and developed at 2 mL/min with 3 min of 35% buffer B (60% acetonitrile/40% 50 mM TEAA, pH 6.0) followed by a linear gradient of 35%-75% buffer B for 35 min. Alternatively, the column was equilibrated with 0.1% trifluoroacetic acid (TFA) and developed at 2 mL/min with 20% buffer C (0.092% TFA, 90% acetonitrile) for 5 min,

followed by a linear gradient of 20% to 90% buffer C for 35 min. Fractions containing pure VWF71 peptide were lyophilized using a Speed Vac concentrator (Savant), dissolved in a minimum volume of 100 mM sodium phosphate, pH 7.1, and desalted on a small column of PD-10 in the same buffer.

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 35 40 45  
 Ser Met Tyr Ser Phe Ala Gly Tyr Cys Ser Tyr Leu Leu Ala Gly Gly  
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 Cys Gln Lys Arg Ser Phe Ser Ile Ile Gly Asp Phe Gln Asn Gly Lys  
 65 70 75 80



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Arg	Val	Glu	Asp	Phe	Gly	Asn	Ala	Trp	Lys	Leu	His	Gly	Asp	Cys	Gln
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Asp	Leu	Gln	Lys	Gln	His	Ser	Asp	Pro	Cys	Ala	Leu	Asn	Pro	Arg	Met
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Thr	Arg	Phe	Ser	Glu	Glu	Ala	Cys	Ala	Val	Leu	Thr	Ser	Pro	Thr	Phe
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His	Leu	Ser	Ile	Ser	Val	Val	Leu	Lys	Gln	Thr	Tyr	Gln	Glu	Lys	Val
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	1235					1240						1245			
Asp	Ala	Pro	Val	Ser	Pro	Thr	Thr	Leu	Tyr	Val	Glu	Asp	Ile	Ser	
	1250					1255						1260			
Glu	Pro	Pro	Leu	His	Asp	Phe	Tyr	Cys	Ser	Arg	Leu	Leu	Asp	Leu	
	1265					1270						1275			
Val	Phe	Leu	Leu	Asp	Gly	Ser	Ser	Arg	Leu	Ser	Glu	Ala	Glu	Phe	
	1280					1285						1290			
Glu	Val	Leu	Lys	Ala	Phe	Val	Val	Asp	Met	Met	Glu	Arg	Leu	Arg	
	1295					1300						1305			
Ile	Ser	Gln	Lys	Trp	Val	Arg	Val	Ala	Val	Val	Glu	Tyr	His	Asp	
	1310					1315						1320			
Gly	Ser	His	Ala	Tyr	Ile	Gly	Leu	Lys	Asp	Arg	Lys	Arg	Pro	Ser	
	1325					1330						1335			



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Pro Trp Asn Val Val Pro	Glu Lys Ala His Leu	Leu Ser Leu Val
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Asp Val Met Gln Arg Glu	Gly Gly Pro Ser Gln	Ile Gly Asp Ala
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Leu Gly Phe Ala Val Arg	Tyr Leu Thr Ser Glu	Met His Gly Ala
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Arg Pro Gly Ala Ser Lys	Ala Val Val Ile Leu	Val Thr Asp Val
1790	1795	1800
Ser Val Asp Ser Val Asp	Ala Ala Ala Asp Ala	Ala Arg Ser Asn
1805	1810	1815
Arg Val Thr Val Phe Pro	Ile Gly Ile Gly Asp	Arg Tyr Asp Ala
1820	1825	1830
Ala Gln Leu Arg Ile Leu	Ala Gly Pro Ala Gly	Asp Ser Asn Val
1835	1840	1845
Val Lys Leu Gln Arg Ile	Glu Asp Leu Pro Thr	Met Val Thr Leu
1850	1855	1860
Gly Asn Ser Phe Leu His	Lys Leu Cys Ser Gly	Phe Val Arg Ile
1865	1870	1875
Cys Met Asp Glu Asp Gly	Asn Glu Lys Arg Pro	Gly Asp Val Trp
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Thr Leu Pro Asp Gln Cys	His Thr Val Thr Cys	Gln Pro Asp Gly
1895	1900	1905
Gln Thr Leu Leu Lys Ser	His Arg Val Asn Cys	Asp Arg Gly Leu
1910	1915	1920
Arg Pro Ser Cys Pro Asn	Ser Gln Ser Pro Val	Lys Val Glu Glu
1925	1930	1935
Thr Cys Gly Cys Arg Trp	Thr Cys Pro Cys Val	Cys Thr Gly Ser
1940	1945	1950
Ser Thr Arg His Ile Val	Thr Phe Asp Gly Gln	Asn Phe Lys Leu
1955	1960	1965
Thr Gly Ser Cys Ser Tyr	Val Leu Phe Gln Asn	Lys Glu Gln Asp
1970	1975	1980
Leu Glu Val Ile Leu His	Asn Gly Ala Cys Ser	Pro Gly Ala Arg
1985	1990	1995
Gln Gly Cys Met Lys Ser	Ile Glu Val Lys His	Ser Ala Leu Ser
2000	2005	2010
Val Glu Leu His Ser Asp	Met Glu Val Thr Val	Asn Gly Arg Leu
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Val Ser Val Pro Tyr Val	Gly Gly Asn Met Glu	Val Asn Val Tyr
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Gly Ala Ile Met His Glu	Val Arg Phe Asn His	Leu Gly His Ile
2045	2050	2055
Phe Thr Phe Thr Pro Gln	Asn Asn Glu Phe Gln	Leu Gln Leu Ser
2060	2065	2070
Pro Lys Thr Phe Ala Ser	Lys Thr Tyr Gly Leu	Cys Gly Ile Cys
2075	2080	2085
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Thr Thr Asp Trp Lys Thr	Leu Val Gln Glu Trp	Thr Val Gln Arg
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2120	2125	2130

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Gln Gln	Asp Ser Cys His Gln	Glu Gln Val Cys Glu	Val Ile Ala
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Ser Tyr	Ala His Leu Cys Arg	Thr Asn Gly Val Cys	Val Asp Trp
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Arg Thr	Pro Asp Phe Cys Ala	Met Ser Cys Pro Pro	Ser Leu Val
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2210	2215		2220
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Pro Asp	Lys Val Met Leu Glu	Gly Ser Cys Val Pro	Glu Glu Ala
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Cys Thr	Gln Cys Ile Gly Glu	Asp Gly Val Gln His	Gln Phe Leu
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Glu Ala	Trp Val Pro Asp His	Gln Pro Cys Gln Ile	Cys Thr Cys
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Ala Lys	Ala Pro Thr Cys Gly	Leu Cys Glu Val Ala	Arg Leu Arg
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Gln Asn	Ala Asp Gln Cys Cys	Pro Glu Tyr Glu Cys	Val Cys Asp
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Pro Val	Ser Cys Asp Leu Pro	Pro Val Pro His Cys	Glu Arg Gly
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Leu Gln	Pro Thr Leu Thr Asn	Pro Gly Glu Cys Arg	Pro Asn Phe
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2360	2365		2370
Ser Cys	Pro Pro His Arg Leu	Pro Thr Leu Arg Lys	Thr Gln Cys
2375	2380		2385
Cys Asp	Glu Tyr Glu Cys Ala	Cys Asn Cys Val Asn	Ser Thr Val
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Ser Cys	Pro Leu Gly Tyr Leu	Ala Ser Thr Ala Thr	Asn Asp Cys
2405	2410		2415
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Arg Ser	Thr Ile Tyr Pro Val	Gly Gln Phe Trp Glu	Glu Gly Cys
2435	2440		2445
Asp Val	Cys Thr Cys Thr Asp	Met Glu Asp Ala Val	Met Gly Leu
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Arg Val	Ala Gln Cys Ser Gln	Lys Pro Cys Glu Asp	Ser Cys Arg
2465	2470		2475
Ser Gly	Phe Thr Tyr Val Leu	His Glu Gly Glu Cys	Cys Gly Arg
2480	2485		2490
Cys Leu	Pro Ser Ala Cys Glu	Val Val Thr Gly Ser	Pro Arg Gly
2495	2500		2505
Asp Ser	Gln Ser Ser Trp Lys	Ser Val Gly Ser Gln	Trp Ala Ser
2510	2515		2520
Pro Glu	Asn Pro Cys Leu Ile	Asn Glu Cys Val Arg	Val Lys Glu
2525	2530		2535

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Glu Val Phe Ile Gln Gln Arg Asn Val Ser Cys Pro Gln Leu Glu  
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 Val Pro Val Cys Pro Ser Gly Phe Gln Leu Ser Cys Lys Thr Ser  
 2555 2560 2565  
 Ala Cys Cys Pro Ser Cys Arg Cys Glu Arg Met Glu Ala Cys Met  
 2570 2575 2580  
 Leu Asn Gly Thr Val Ile Gly Pro Gly Lys Thr Val Met Ile Asp  
 2585 2590 2595  
 Val Cys Thr Thr Cys Arg Cys Met Val Gln Val Gly Val Ile Ser  
 2600 2605 2610  
 Gly Phe Lys Leu Glu Cys Arg Lys Thr Thr Cys Asn Pro Cys Pro  
 2615 2620 2625  
 Leu Gly Tyr Lys Glu Glu Asn Asn Thr Gly Glu Cys Cys Gly Arg  
 2630 2635 2640  
 Cys Leu Pro Thr Ala Cys Thr Ile Gln Leu Arg Gly Gly Gln Ile  
 2645 2650 2655  
 Met Thr Leu Lys Arg Asp Glu Thr Leu Gln Asp Gly Cys Asp Thr  
 2660 2665 2670  
 His Phe Cys Lys Val Asn Glu Arg Gly Glu Tyr Phe Trp Glu Lys  
 2675 2680 2685  
 Arg Val Thr Gly Cys Pro Pro Phe Asp Glu His Lys Cys Leu Ala  
 2690 2695 2700  
 Glu Gly Gly Lys Ile Met Lys Ile Pro Gly Thr Cys Cys Asp Thr  
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 Cys Glu Glu Pro Glu Cys Asn Asp Ile Thr Ala Arg Leu Gln Tyr  
 2720 2725 2730  
 Val Lys Val Gly Ser Cys Lys Ser Glu Val Glu Val Asp Ile His  
 2735 2740 2745  
 Tyr Cys Gln Gly Lys Cys Ala Ser Lys Ala Met Tyr Ser Ile Asp  
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 Ile Asn Asp Val Gln Asp Gln Cys Ser Cys Cys Ser Pro Thr Arg  
 2765 2770 2775  
 Thr Glu Pro Met Gln Val Ala Leu His Cys Thr Asn Gly Ser Val  
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2810

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45

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38

<210> SEQ ID NO 6  
 <211> LENGTH: 9

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&lt;212&gt; TYPE: PRT

&lt;213&gt; ORGANISM: Homo sapiens

&lt;400&gt; SEQUENCE: 6

Glu Ala Pro Asp Leu Val Leu Gln Arg

1

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What is claimed is:

1. A probe comprising:

an oligopeptide consisting of no more than 80 amino acids of sequence of von Willebrand Factor (VWF), said oligopeptide comprising a scissile Y1605-M1606 peptide of a VWF sequence set forth as SEQ ID NO: 3, a cysteine substitution located from 1 to 12 amino acids from the scissile Y-M in the carboxy terminal direction, and a C-terminal segment;

a fluorophore: and

a fluorescence quencher,

wherein the fluorophore and the fluorescence quencher are bound to the oligopeptide on opposite sides of the scissile Y-M peptide and wherein the oligopeptide comprises a substitution of lysine K1617 of a VWF sequence set forth as SEQ ID NO: 3 with an amino acid that does not comprise a primary amine on its side chain.

2. A probe in accordance with claim 1, wherein the cysteine substitution is a substitution of an amino acid located at least 3 amino acids from the scissile Y-M peptide.

3. A probe in accordance with claim 1, wherein the cysteine substitution is a N1610C substitution of a VWF sequence set forth as SEQ ID NO: 3.

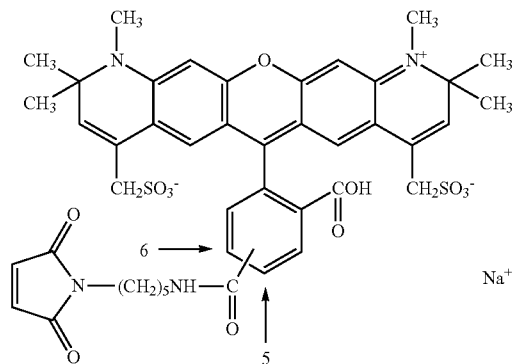
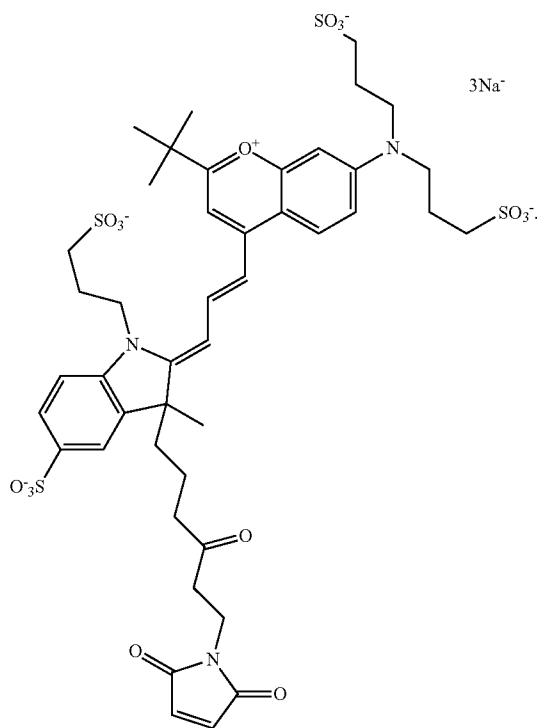
4. A probe in accordance with claim 1 wherein the oligopeptide consists of the sequence set forth as SEQ ID NO: 2.

5. A probe in accordance with claim 1, wherein the fluorophore has an absorption maximum >550 nm.

6. A probe in accordance with claim 1, wherein the fluorophore has an emission maximum >600 nm.

7. A probe in accordance with claim 1 wherein the fluorophore is selected from the group consisting of Alexa Fluor®594 maleimide (fluorophore)

and DyLight®633 maleimide (fluorophore)



8. A probe in accordance with claim 1 wherein the quencher has an absorption maximum >550 nm.

9. A probe in accordance with claim 1, wherein at least one of the fluorophore and the quencher comprises at least one sulfate.

10. A probe in accordance with claim 1, wherein the quencher is selected from the group consisting of QSY® (fluorescence quencher) 21- succinimidyl ester and IRDye® (infrared dye) QC-1 N-hydroxy succinimidyl ester.

11. A probe in accordance with claim 1, wherein the probe is soluble in water at >50 μM.

12. A method of determining presence, absence or quantity of ADAMTS13 activity in a sample, comprising: forming a mixture comprising a sample and a probe of claim 1; and measuring fluorescence at one or more time points after forming the mixture, wherein the sample is diluted less than 20-fold.

13. A method in accordance with claim 12, wherein the sample is selected from the group consisting of a serum sample, an undiluted serum sample, a concentrated serum sample, a plasma sample, an undiluted plasma sample and a concentrated plasma sample.

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14. A method of determining presence, absence or quantity of ADAMTS13 inhibitor activity in a sample, comprising; forming a mixture comprising a sample, as source of ADAMTS13, and a probe of claim 1; and measuring fluorescence at one or more time points after forming the mixture. 5

15. A method in accordance with claim 14, further comprising inactivating ADAMTS13 activity endogenous to the sample prior to forming the mixture.

16. A method in accordance with claim 14, wherein the source of ADAMTS13 is selected from the group consisting of normal plasma, recombinant ADAMTS13 and a combination thereof. 10

17. A method in accordance with claim 14, wherein the sample is selected from the group consisting of a serum sample, an undiluted serum sample, a concentrated serum sample, a plasma sample, an undiluted plasma sample and a concentrated plasma sample. 15

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18. A probe in accordance with claim 1, wherein the C-terminal segment consists of sequence EAPDLVLQR (SEQ ID NO: 6).

19. A probe comprising:  
 an oligopeptide consisting of no more than 80 amino acids of sequence of von Willebrand Factor (VWF), said oligopeptide comprising a scissile Y-M peptide, a cysteine substitution located from 1 to 12 amino acids from the scissile Y-M in the carboxy terminal direction, and a C-terminal segment;  
 as fluorophore; and  
 as fluorescence quencher,  
 wherein the fluorophore and the fluorescence quencher are bound to the oligopeptide on opposite sides of the scissile Y-M peptide and wherein the oligopeptide consists of the sequence set forth as SEQ ID NO: 2.

\* \* \* \* \*