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(57) Abstract:



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**LIPID BILAYER-INTEGRATED SPP1 CONNECTOR PROTEIN NANOPORE AND
SPP1 CONNECTOR PROTEIN VARIANTS FOR USE AS LIPID BILAYER-
INTEGRATED NANOPORE**

by

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RELATED APPLICATIONS

[0001] This application claims priority from U.S. Provisional Application Serial No. 62/104,375 filed January 16, 2015, the entire disclosure of which is incorporated herein by this reference.

GOVERNMENT INTEREST

[0002] This invention was made with government support under EB012135 and EB003730 awarded by the National Institutes of Health. The government has certain rights in the invention.

TECHNICAL FIELD

[0003] The presently-disclosed subject matter relates to an engineered SPP1 viral DNA-packaging motor connector protein that can be incorporated into a lipid membrane to form an electroconductive aperture.

BACKGROUND

[0004] Highly sensitive detection and characterization of minute quantities of chemicals and biochemicals represent desirable goals of modern analytical technologies. Robust molecular sensing devices would find uses in a wide range of biomedical, industrial,

environmental, forensic, security and other contexts, for example, in the detection and identification of pathogens and chemicals at extremely low concentrations for disease diagnosis and environmental monitoring, in high throughput DNA sequencing and other genomics applications, and elsewhere.

[0005] Analytical methodologies have been described that employ intermolecular affinity binding interactions, typically non-covalent in nature, to detect binding or “capture” of an analyte of interest by a specific affinity ligand, for instance, including detection of bacterial, viral, parasitic or other microbial pathogens or pathogen-associated antigens, and detection of antibodies, cancer markers, and other analytes (e.g., Kittigul et al., *Am J Trop Med Hyg.* 1998 Sep;59(3):352-6; Cordiano et al., *J Immunol Methods.* 1995 Jan 13;178(1):121-30; Olson et al., *J Immunol Methods.* 1990 Nov 6;134(1):71-9; Nerurkar et al., *J Clin Microbiol.* 1984 Jul;20(1):109-14; Jia et al., *J Virol Methods.* 2009 Oct;161(1):38-43; He et al., *Clin Vaccine Immunol.* 2007 May;14(5):617-23; Xu et al., *J Clin Microbiol.* 2006 Aug;44(8):2872-8; Che et al., *J Clin Microbiol.* 2004 Jun;42(6):2629-35; Hunt et al., Brown et al., *Am J Trop Med Hyg.* 2001 Sep;65(3):208-13; Loa et al., *Avian Dis.* 2000 Jul-Sep;44(3):498-506; Lubenko et al., *Transfus Med.* 2000 Sep;10(3):213-8; Chanteau et al., *Int J Tuberc Lung Dis.* 2000 Apr;4(4):377-83; Brinker et al., *J Clin Microbiol.* 1998 Apr;36(4):1064-9; Vyse et al., *J Virol Methods.* 1997 Jan;63(1-2):93-101; Peterson et al., *J Clin Microbiol.* 1997 Jan;35(1):208-12; Lairmore et al., *AIDS Res Hum Retroviruses.* 1993 Jun;9(6):565-71; Heller et al., *Vet Microbiol.* 1993 Oct;37(1-2):127-33; van Loon et al., *Epidemiol Infect.* 1992 Feb;108(1):165-74; Wolf-Rogers et al., *J Immunol Methods.* 1990 Oct 19;133(2):191-8; Barsoum et al., *Exp Parasitol.* 1990 Jul;71(1):107-13; Hierholzer et al., *J Clin Microbiol.* 1989 Jun;27(6):1243-9; Hurley et al., *J Immunoassay.* 1986;7(4):309-36; Wolff et al., *Cancer Res.* 53:2560-65 (1993); see generally, e.g., Harlow and Lane, *Antibodies: A Laboratory Manual*, Cold Spring Harbor Laboratory, 1988; Weir, D.M., *Handbook of Experimental Immunology*, 1986, Blackwell Scientific, Boston, MA).

[0006] Beyond detection of the presence of an analyte following its involvement in an affinity binding interaction, sophisticated technologies are emerging that permit characterization of the analyte, often by comparing a single- or multiparameter physicochemical profile of the analyte to type- characteristic profiles generated using one or more known reference standards, and hence referred to as “fingerprinting” techniques (e.g., Li et al., *Rapid Commun Mass Spectrom.* 2009 23(22):3533-3542; Ali et al., *J Agric Food Chem.* 2009; Leski et al., *Appl Environ Microbiol.* Sept. 18, 2009; Weinkopff et al., *J Parasitol.* June 18, 2009; Song et al., *Proteomics.* 2009 9(11):3090-9; Ortea et al., *J Agric*

Food Chem. 2009 57(13):5665-72; Amini, Pharmeur Sci Notes. 2009(1):11-6; Shi et al., Biol Pharm Bull. 2009 32(1):142-6; Sun et al., J Chromatogr A. 2009 1216(5):830-6; Yin et al., Phytopathology. 2003 93(8):1006-13; Roy et al., Clin Cancer Res. 2008 14(20):6610-7; Pei et al., Zhongguo Zhong Yao Za Zhi. 2008 33(14):1662-8; Arthur, Methods Mol Med. 2008, 141:257-70; Zhao et al., Se Pu 2008 26(1):43-9; Woo et al., Anal Chem. 2008 80(7):2419-25; Damodaran et al., Genomics Proteomics Bioinformatics. 2007 5(3-4):152-7; Fellstrom et al., J Microbiol Methods. 2008 72(2):133-40; Song et al., Conf Proc IEEE Eng Med Biol Soc. 2006 1:4556-9; De Vuyst et al., Int J Food Microbiol. 2008 125(1):79-90).

[0007] The use of transmembrane channels has been demonstrated in stochastic analyte detection (Bayley et al., 2001 Nature 13:225-230), an electrochemical approach relying on the real-time observation of individual binding events between single substrate molecules and a receptor, as evidenced by altered (e.g., decreased or increased in a statistically significant manner) electrical conductance by the channel (receptor) as a result of substrate (analyte) binding. A wide range of processes, such as the transport of DNA, RNA, pharmaceutical agents, peptides, proteins, and polymers, have been studied by such approaches, for example, using electrophysiological measurements of individual membrane channels (Thieffry et al., 1988 EMBO J 7:1449; Hinnah et al., 2002 Biophys J 83:899; Alcaayaga et al., 1992 FEBS Lett. 311:246-50; Benz et al., 1986 J Bacteriol 165:978; Movileanu et al., 2000 Nat. Biotechnol. 18:1091).

[0008] For instance, the transient blockade of ionic current through the *Staphylococcus aureus* alpha-hemolysin (α -HL) channel, a bacterial transmembrane pore-forming protein, has been used to measure the length of single-stranded DNA or RNA (Kasianowicz et al. Proc. Natl. Acad. Sci. USA 93, 13770-13773 (1996)). Subsequently, DNA hairpin molecules have been used to decelerate the DNA translocation rate through the alpha-hemolysin (α -HL) pore, to demonstrate the ability of a transmembrane ion channel to discriminate between single nucleotide polymorphisms (Vercoutere et al., 2001 Nat. Biotechnol. 19:248). Detection of base pair stacking and strand orientation within the pore have also been investigated (Vercoutere et al., 2003 Nucl. Ac. Res. 31:1311; Howorka et al., 2001 Nat. Biotechnol. 19:636; deGuzman et al., 2006 Nucl. Ac. Res. 34:6425). The channel of α -HL with a covalently attached adapter molecule has been shown to discriminate the nucleotides A, T, G, and C (Clarke et al., 2009 Nat. Nanotechnol. 4:265).

[0009] The α -HL channel has also been reported to provide some ability to translocate small peptides (See e.g., Mohammad, et al., "Protein Sensing with engineered protein nanopores," Methods Mol Biol. 2012, 870:21-37; Jetha, et al., "Nanopore Analysis of Wild-

Type and Mutant Prion Protein (PrP^c): Single Molecule Discrimination and PrP^c Kinetics,” PLOS ONE, 2013, 8(2); Mereuta, et al., “Slowing down single-molecule trafficking through a protein nanopore reveals intermediates for peptide translocation,” Scientific Reports, 2014, 4:3885; Rosen, et al., “Single-molecule site-specific detection of protein phosphorylation with a nanopore,” Nat Biotechnol, 2014 Feb, 32(2):179-81); Movileanu, “Watching single proteins using engineered nanopores,” Protein Pept Lett. 2014 March, 21(3):235-46). However, at only about 1.2 – 1.5 nm, only a limited number of peptides are capable of being translocated through the α -HL channel and there is not an ability to differentiate between different peptides.

[0010] Other protein channels that have been investigated include alamethicin for detection of polyethylene glycol (Bezrukov, 2000 J Membr Biol. 174:1-13), and the reengineered MspA protein from *M. smegmatis* for translocation of ssDNA (Butler et al., 2008 Proc. Nat. Acad. Sci. USA 105:20647). Most studies involving nucleic acid transport through nanopores have focused on α -HL. However, the limiting lumen diameter of α -HL (about 1.2 - 1.5 nm) and other channels has restricted their DNA and RNA applications to translocation of single-stranded nucleic acid (Song, 1996 Science 274:1859). A similar limitation was also reported for the MspA nanopore (Butler et al., 2008).

[0011] In a small number of other membrane pore systems, evidence of double-stranded DNA (dsDNA) transport across the membrane has been presented (Szabo et al., 2002 Cell Physiol Biochem 12:127; Mobasher et al. 2002 Eur J Biophys 31:389; Carneiro et al., 2003 Biochim Biophys Acta 1612:144), but these systems are not robust and represent poor candidates for widespread use such as biomedical applications, due to their undesirable voltage gating properties and the associated signal fluctuation. For this reason, their potential is considered limited and researchers have switched instead to fabricating synthetic metal or silicon nanopores for potential use in DNA sequencing (Smeets et al, 2006 Nano Lett 6:89; Wang et al., 2001 Nat. Biotechnol. 19:622; Iqbal et al., 2007 Nat. Nano 2:243). Such synthetic nanopores, however, suffer from shortcomings due to difficulties in reliably producing replicated structures having consistent properties from batch to batch, and also lack versatility with regard to the ability to engineer modifications to pore structures and/or to serve as substrates for modification by a wide range of chemical conjugation. As a result, the search for superior alternatives to currently available protein nanopores is still ongoing.

[0012] There is a need for improved compositions and methods that would provide a versatile membrane conductive channel platform for sensitively detecting and characterizing a wide range of analytes, having a lumen capable of accommodating dsDNA and

polypeptides, that can be reliably and reproducibly assembled, that is not susceptible to voltage gating under working conditions, and that can be readily modified to feature a wide variety of specific affinity receptors for use in the detection and characterization of different analytes. The presently disclosed invention embodiments fulfill such a need, and offer other related advantages.

SUMMARY

[0013] The presently-disclosed subject matter meets some or all of the above-identified needs, as will become evident to those of ordinary skill in the art after a study of information provided in this document.

[0014] This Summary describes several embodiments of the presently-disclosed subject matter, and in many cases lists variations and permutations of these embodiments. This Summary is merely exemplary of the numerous and varied embodiments. Mention of one or more representative features of a given embodiment is likewise exemplary. Such an embodiment can typically exist with or without the feature(s) mentioned; likewise, those features can be applied to other embodiments of the presently-disclosed subject matter, whether listed in this Summary or not. To avoid excessive repetition, this Summary does not list or suggest all possible combinations of such features.

[0015] The presently disclosed subject matter includes a nucleic acid molecule encoding an SPP1 connector polypeptide variant. In some embodiments, the SPP1 connector polypeptide variant comprises an amino acid sequence having at least 95% sequence identity to the sequence shown in SEQ ID NO:4, SEQ ID NO:6, SEQ ID NO:8, or SEQ ID NO:10. In some embodiments, the SPP1 connector polypeptide variant comprises an amino acid sequence having at least 95% sequence identity to the sequence of SEQ ID NO:4, SEQ ID NO:6, SEQ ID NO:8, or SEQ ID NO:10. In some embodiments, the SPP1 connector polypeptide variant comprises an amino acid sequence having at least 99% sequence identity to the sequence of in SEQ ID NO:4, SEQ ID NO:6, SEQ ID NO:8, or SEQ ID NO:10. In some embodiments, the SPP1 connector polypeptide variant comprises the amino acid sequence of SEQ ID NO:4, SEQ ID NO:6, SEQ ID NO:8, or SEQ ID NO:10. In some embodiments, the nucleic acid molecule encoding an SPP1 connector polypeptide variant includes the sequence of SEQ ID NO:3, SEQ ID NO:5, SEQ ID NO:7, or SEQ ID NO:9.

[0016] The presently disclosed subject matter further includes a SPP1 connector polypeptide variant, wherein the polypeptide comprises an amino acid sequence having at least 95% sequence identity to the sequence of SEQ ID NO:4, SEQ ID NO:6, SEQ ID NO:8,

or SEQ ID NO:10. at least 95% sequence identity to the sequence shown in SEQ ID NO:4, SEQ ID NO:6, SEQ ID NO:8, or SEQ ID NO:10. In some embodiments, the SPP1 connector polypeptide variant comprises an amino acid sequence having at least 95% sequence identity to the sequence of SEQ ID NO:4, SEQ ID NO:6, SEQ ID NO:8, or SEQ ID NO:10. In some embodiments, the SPP1 connector polypeptide variant comprises an amino acid sequence having at least 99% sequence identity to the sequence of in SEQ ID NO:4, SEQ ID NO:6, SEQ ID NO:8, or SEQ ID NO:10. In some embodiments, the SPP1 connector polypeptide variant comprises the amino acid sequence of SEQ ID NO:4, SEQ ID NO:6, SEQ ID NO:8, or SEQ ID NO:10.

[0017] The presently disclosed subject matter further includes a SPP1 connector polypeptide variant, wherein the polypeptide comprises an amino acid sequence encoded by a nucleic acid molecule having the sequence of SEQ ID NO:3, SEQ ID NO:5, SEQ ID NO:7, or SEQ ID NO:9.

[0018] The presently disclosed subject matter further includes a conductive channel-containing membrane, which includes (a) a membrane layer, and (b) a SPP1 connector polypeptide variant that is incorporated into the membrane layer to form an aperture through which conductance can occur when an electrical potential is applied across the membrane.

[0019] In some embodiments the SPP1 connector polypeptide variant of the conductive channel-containing membrane is selected from (i) a polypeptide comprising an amino acid sequence having at least 95% sequence identity to the sequence of SEQ ID NO:4, SEQ ID NO:6, SEQ ID NO:8, or SEQ ID NO:10; (ii) a polypeptide comprising an amino acid sequence having at least 99% sequence identity to the sequence of SEQ ID NO:4, SEQ ID NO:6, SEQ ID NO:8, or SEQ ID NO:10; (iii) a polypeptide comprising the amino acid sequence of SEQ ID NO:4, SEQ ID NO:6, SEQ ID NO:8, or SEQ ID NO:10; and (iv) a polypeptide comprising an amino acid sequence encoded by a nucleic acid molecule having the sequence of SEQ ID NO:3, SEQ ID NO:5, SEQ ID NO:7, or SEQ ID NO:9.

[0020] In some embodiments of the conductive channel-containing membrane the SPP1 connector polypeptide variant comprises a detectable label. In some embodiments of the conductive channel-containing membrane, the detectable label is selected from the group consisting of a colorimetric indicator, a GCMS tag compound, a fluorescent indicator, a luminescent indicator, a phosphorescent indicator, a radiometric indicator, a dye, an enzyme, a substrate of an enzyme, an energy transfer molecule, a quantum dot, a metal particle and an affinity label.

[0021] In some embodiments of the conductive channel-containing membrane, the membrane layer comprises a lipid layer. In some embodiments, the lipid layer comprises amphipathic lipids. In some embodiments, the amphipathic lipids comprise phospholipids and the lipid layer comprises a lipid bilayer. In some embodiments, the lipid layer is selected from the group consisting of a planar membrane layer and a liposome. In some embodiments, the liposome is selected from the group consisting of a multilamellar liposome and a unilamellar liposome. In some embodiments, the incorporated SPP1 connector polypeptide variant is mobile in the membrane layer.

[0022] In some embodiments, the conductive channel-containing membrane is capable of translocating a single stranded nucleic acid molecule, a double-stranded nucleic acid molecule, and/or a polypeptide through the aperture when the electrical potential is applied, wherein said conductive channel-containing membrane is capable of polypeptide detection, identification, sequencing, and discrimination.

[0023] The presently-disclosed subject matter further includes a method of sensing a molecule using a conductive channel-containing membrane as disclosed herein. In some embodiments, the molecule is a polypeptide. In some embodiments, the molecule is a nucleic acid molecule. In some embodiments, the nucleic acid molecule is a double-stranded nucleic acid molecule.

[0024] The presently-disclosed subject matter further includes a method of DNA sequencing using a conductive channel-containing membrane as disclosed herein.

BRIEF DESCRIPTION OF THE DRAWINGS

[0025] The novel features of the invention are set forth with particularity in the appended claims. A better understanding of the features and advantages of the present invention will be obtained by reference to the following detailed description that sets forth illustrative embodiments, in which the principles of the invention are used, and the accompanying drawings of which:

[0026] **Fig. 1:** Structure of the channel of SPP1 DNA packaging motor. (a) Side view and (b) top view from PDB file 2JES showing hydrophilic (red), hydrophobic (blue) and neutral (white) amino acids; and dimensions of the channel (PDB: 2JES). (c) Coomassie-blue stained SDS PAGE showing the purified SPP1 channel subunits gp6.

[0027] **Fig. 2:** Electrophysiological properties of Membrane-embedded SPP1 connector. (a) Current trace showing the insertion of SPP1 connector into the planar membrane with a characteristic step size of ~200 pA at 50 mV. (b) Conductance distribution based on 104

insertion events. (c) Current-Voltage trace acquired from $-50 \rightarrow +50$ mV. Buffer: 1 M KCl, 5 mM HEPES, pH 8.

[0028] Fig. 3: Peptide translocation through SPP1 connector. (a) Current trace showing a burst of current blockage events with characteristic current amplitude and dwell time (insert) indicating the translocation of TAT peptides. (b) Rate of peptide translocation as a function of peptide concentration ($n = 3$). (c) Histogram of current blockage percentage from 285 events. (d) Dwell time of peptide translocation events fitted with a single exponential function. Applied voltage: 50 mV; Buffer: 1 M KCl, 5 mM HEPES, pH 8.

[0029] Fig. 4: Determining the conformation states of TAT peptide. Current trace (left), current blockage distribution (middle) and conformation (right) for (a) Oxidized dimer state of TAT peptide; (b) reduced monomer state of TAT peptide; and (c) Cy3-conjugated TAT monomer. Applied voltage: 50 mV; Buffer: 1 M KCl, 5 mM HEPES, pH 8. Total number of events: 858 in A; 367 in B and 1128 in C.

[0030] Fig. 5: Single molecule fluorescent images validating TAT peptide translocation. (a) The upper row is the image showing the detection of Cy3-labeled TAT peptide from the fractions collected from patch clamp at 0, 20, 40 and 60 mins. The lower row shows the background resulting from the self-permeation of the TAT peptide which is the HIV membrane penetrating peptide. Excitation λ : 532 nm; laser power: 5 mW; 60 $\times$ objective (N.A. = 1.4, oil immersion); Exposure time: 500 ms. (b) Quantitative analysis showing the increase in Cy3-TAT peptide signal in presence of SPP1 connector compared to control without connector. The error bar is based on counting of the spots from different image areas. Applied voltage: 50 mV; Buffer: 1 M KCl, 5 mM HEPES, pH 8.

[0031] Fig. 6: Real-time assessment of the conformation states of TAT peptide. (a) Continuous current trace showing transition of oxidized dimer states to reduced monomer states after addition of reducing agent TCEP. (b) Quantitative analysis showing the fraction ($\square$) of dimer and monomer states as a function of reaction time. (c) Current blockage *versus* dwell time distribution over the course of reaction time. Applied voltage: 50 mV; Buffer: 1 M KCl, 5 mM HEPES, pH 8. (d) Quantitative analysis showing the reaction quotient Q_r as a function of reaction time.

[0032] Fig. 7: Assembly of SPP1 gp6 subunits into 12 subunit connector complexes as shown by 15-35% glycerol gradient ultracentrifugation. BSA was used for size control.

BRIEF DESCRIPTION OF THE SEQUENCE LISTING

[0033] SEQ ID NO: 1 is the nucleic acid sequence of a nucleotide encoding a wild type SPP1 connector channel polypeptide.

[0034] SEQ ID NO: 2 is the amino acid sequence of a wild type SPP1 connector channel polypeptide.

[0035] SEQ ID NO: 3 is the nucleic acid sequence of a nucleotide encoding an exemplary SPP1 connector channel polypeptide variant.

[0036] SEQ ID NO: 4 is the amino acid sequence of the exemplary SPP1 connector channel polypeptide variant encoded by the nucleotide of SEQ ID NO: 3.

[0037] SEQ ID NO: 5 is the nucleic acid sequence of a nucleotide encoding another exemplary SPP1 connector channel polypeptide variant.

[0038] SEQ ID NO: 6 is the amino acid sequence of the exemplary SPP1 connector channel polypeptide variant encoded by the nucleotide of SEQ ID NO: 5.

[0039] SEQ ID NO: 7 is the nucleic acid sequence of a nucleotide encoding another exemplary SPP1 connector channel polypeptide variant.

[0040] SEQ ID NO: 8 is the amino acid sequence of the exemplary SPP1 connector channel polypeptide variant encoded by the nucleotide of SEQ ID NO: 7.

[0041] SEQ ID NO: 9 is the nucleic acid sequence of a nucleotide encoding another exemplary SPP1 connector channel polypeptide variant.

[0042] SEQ ID NO: 10 is the amino acid sequence of the exemplary SPP1 connector channel polypeptide variant encoded by the nucleotide of SEQ ID NO: 9.

DESCRIPTION OF EXEMPLARY EMBODIMENTS

[0043] The details of one or more embodiments of the presently-disclosed subject matter are set forth in this document. Modifications to embodiments described in this document, and other embodiments, will be evident to those of ordinary skill in the art after a study of the information provided in this document. The information provided in this document, and particularly the specific details of the described exemplary embodiments, is provided primarily for clearness of understanding and no unnecessary limitations are to be understood therefrom. In case of conflict, the specification of this document, including definitions, will control.

[0044] The presently-disclosed subject matter includes, in some embodiments, a nucleic acid molecule encoding an SPP1 connector polypeptide variant. The SPP1 connector polypeptide variant can comprises an amino acid sequence having at least 90, 91, 92, 93, 94, 95, 96, 97, or 99% sequence identity to a sequence shown in SEQ ID NO:4, SEQ ID NO:6, SEQ ID NO:8, or SEQ ID NO:10. In some embodiments, the nucleic acid molecule encodes a SPP1 connector polypeptide variant comprising an amino acid sequence as set forth in SEQ ID NO:4, SEQ ID NO:6, SEQ ID NO:8, or SEQ ID NO:10. In some embodiments, the nucleic acid molecule has the sequence shown in SEQ ID NO:3, SEQ ID NO:5, SEQ ID NO:7, or SEQ ID NO:9.

[0045] The presently-disclosed subject matter further includes, in some embodiments, a SPP1 connector polypeptide variant comprising an amino acid sequence having at least 90, 91, 92, 93, 94, 95, 96, 97, or 99% sequence identity to a sequence shown in SEQ ID NO:4, SEQ ID NO:6, SEQ ID NO:8, or SEQ ID NO:10. In some embodiments, the SPP1 connector polypeptide variant comprises an amino acid sequences as set forth in SEQ ID NO:4, SEQ ID NO:6, SEQ ID NO:8, or SEQ ID NO:10. In some embodiments, the SPP1 connector polypeptide variant comprises an amino acid sequence encoded by a nucleic acid molecule having the sequence shown in SEQ ID NO:3, SEQ ID NO:5, SEQ ID NO:7, or SEQ ID NO:9.

[0046] A nucleic acid or polypeptide sequence can be compared to another sequence and described in terms of its percent sequence identity. In calculating percent sequence identity, two sequences are aligned and the number of identical matches of nucleotides or amino acid residues between the two sequences is determined. The number of identical matches is divided by the length of the aligned region (i.e., the number of aligned nucleotides or amino acid residues) and multiplied by 100 to arrive at a percent sequence identity value. It will be appreciated that the length of the aligned region can be a portion of one or both sequences up

to the full-length size of the shortest sequence. It will be appreciated that a single sequence can align differently with other sequences and hence, can have different percent sequence identity values over each aligned region. It is noted that the percent identity value is usually rounded to the nearest integer.

[0047] The alignment of two or more sequences to determine percent sequence identity is performed using the algorithm described by Altschul et al. (1997, Nucleic Acids Res., 25:3389-3402) as incorporated into BLAST (basic local alignment search tool) programs, available at ncbi.nlm.nih.gov on the World Wide Web. BLAST searches can be performed to determine percent sequence identity between a first nucleic acid and any other sequence or portion thereof aligned using the Altschul et al. algorithm. BLASTN is the program used to align and compare the identity between nucleic acid sequences, while BLASTP is the program used to align and compare the identity between amino acid sequences. When utilizing BLAST programs to calculate the percent identity between a sequence disclosed herein (e.g., SEQ ID NOs:1-10) and another sequence, the default parameters of the respective programs are used.

TABLE 1

Conservative Amino Acid Substitutions

Amino Acid	Representative Conservative Amino Acids
Ala	Ser, Gly, Cys
Arg	Lys, Gln, His
Asn	Gln, His, Glu, Asp
Asp	Glu, Asn, Gln
Cys	Ser, Met, Thr
Gln	Asn, Lys, Glu, Asp, Arg
Glu	Asp, Asn, Gln
Gly	Pro, Ala, Ser
His	Asn, Gln, Lys
Ile	Leu, Val, Met, Ala
Leu	Ile, Val, Met, Ala
Lys	Arg, Gln, His
Met	Leu, Ile, Val, Ala, Phe

TABLE 1

Phe	Met, Leu, Tyr, Trp, His
Ser	Thr, Cys, Ala
Thr	Ser, Val, Ala
Trp	Tyr, Phe
Tyr	Trp, Phe, His
Val	Ile, Leu, Met, Ala, Thr

[0048] Modifications, including substitutions, insertions or deletions are made by known methods. By way of example, modifications are made by site-specific mutagenesis of nucleotides in the DNA encoding the protein, thereby producing DNA encoding the modification, and thereafter expressing the DNA in recombinant cell culture. Techniques for making substitution mutations at predetermined sites in DNA having a known sequence are well known, for example M13 primer mutagenesis and PCR mutagenesis.

[0049] The presently-disclosed subject matter further includes, in some embodiments, a conductive channel-containing membrane, comprising (a) a membrane layer; and (b) a SPP1 connector polypeptide variant that is incorporated into the membrane layer to form an aperture through which conductance can occur when an electrical potential is applied across the membrane, wherein the SPP1 connector polypeptide variant is selected from among those disclosed herein.

[0050] Embodiments described herein find use in a variety of molecular analytical contexts, including, for example, sensitive detection and characterization of chemical and biochemical analytes for biomedical, clinical, industrial, chemical, pharmaceutical, environmental, forensic, national security, toxicological and other purposes, including any situation where rapid, specific and exquisitely sensitive detection and/or characterization of an analyte (*e.g.*, preferably a soluble analyte that is provided in solution) may be desired. Expressly contemplated are embodiments in which the presently disclosed compositions and methods are used for DNA sequencing, including dsDNA sequencing, high-throughput DNA sequencing, genomics, SNP detection, molecular diagnostics and other DNA sequencing applications, and polypeptide detection and identification. Additional utilities include those described in International Patent Application Publication No. WO2010/062697, which is incorporated herein in its entirety by this reference.

[0051] Exemplary analytes thus include nucleic acids such as DNA and RNA (including dsDNA and dsRNA), including for the detection and identification of single nucleotide polymorphisms (SNPs) and/or mutations in such nucleic acids, and/or nucleic acid sequence determination. Other exemplary analytes that may be detected and/or characterized using the

herein described compositions and methods include polypeptides and other biopolymers (*e.g.*, proteins, glycoproteins, peptides, glycopeptides, oligosaccharides, polysaccharides, lipids, glycolipids, phospholipids, etc.) and other biomolecules (*e.g.*, soluble mediators, cofactors, vitamins, bioactive lipids, metabolites, and the like), drugs and other pharmaceutical and pharmacological agents, including natural and synthetic compounds, food and cosmetics agents such as flavorants, odorants, preservatives, antioxidants, antimicrobial agents, stabilizers, carriers, excipients, modifying agents and the like, natural and synthetic toxins, dyes, and other compounds.

[0052] Accordingly and in certain embodiments, any analyte for which detection and/or characterization is desired may be used, where it will be recognized from the disclosure herein that the analyte is preferably soluble in a solvent that does not compromise the integrity of the particular membrane layer in which the SPP1 connector polypeptide variant is incorporated to form an aperture through which conductance can occur when an electrical potential is applied across the membrane. Analyte selection may thus vary as a function of the composition of the particular membrane layer being used, which may therefore influence solvent selection. Those skilled in the art will be familiar with criteria to be employed for selecting a solvent that is compatible with a membrane layer of any particular composition. In preferred embodiments, the membrane layer comprises a phospholipid bilayer and the solvent in which the analyte is provided comprises an aqueous solvent, *e.g.*, a solvent that comprises water.

[0053] In some embodiments of the conductive channel-containing membrane, the membrane layer comprises a lipid layer. In a further embodiment the lipid layer comprises amphipathic lipids, which in certain still further embodiments comprise phospholipids and the lipid layer comprises a lipid bilayer. In certain other embodiments the lipid layer is selected from a planar membrane layer and a liposome. In certain embodiments the liposome is selected from a multilamellar liposome and a unilamellar liposome. In certain other embodiments the incorporated SPP1 connector polypeptide variant is mobile in the membrane layer. In certain other embodiments the conductive channel-containing membrane is capable of translocating double-stranded DNA through the aperture when the electrical potential is applied. In certain other embodiments the conductive channel-containing membrane is capable of translocating polypeptides through the aperture when the electrical potential is applied. In certain embodiments conductance occurs without voltage gating when the electrical potential is applied.

[0054] In some embodiments, the SPP1 connector polypeptide variant comprises a detectable label. In some embodiments, the detectable label is selected from the group consisting of a colorimetric indicator, a GCMS tag compound, a fluorescent indicator, a luminescent indicator, a phosphorescent indicator, a radiometric indicator, a dye, an enzyme, a substrate of an enzyme, an energy transfer molecule, a quantum dot, a metal particle and an affinity label.

[0055] While the terms used herein are believed to be well understood by those of ordinary skill in the art, certain definitions are set forth to facilitate explanation of the presently-disclosed subject matter.

[0056] Unless defined otherwise, all technical and scientific terms used herein have the same meaning as is commonly understood by one of skill in the art to which the invention(s) belong.

[0057] All patents, patent applications, published applications and publications, GenBank sequences, databases, websites and other published materials referred to throughout the entire disclosure herein, unless noted otherwise, are incorporated by reference in their entirety.

[0058] Where reference is made to a URL or other such identifier or address, it is understood that such identifiers can change and particular information on the internet can come and go, but equivalent information can be found by searching the internet. Reference thereto evidences the availability and public dissemination of such information.

[0059] As used herein, the abbreviations for any protective groups, amino acids and other compounds, are, unless indicated otherwise, in accord with their common usage, recognized abbreviations, or the IUPAC-IUB Commission on Biochemical Nomenclature (see, Biochem. (1972) 11(9):1726-1732).

[0060] Although any methods, devices, and materials similar or equivalent to those described herein can be used in the practice or testing of the presently-disclosed subject matter, representative methods, devices, and materials are described herein.

[0061] In certain instances, nucleotides and polypeptides disclosed herein are included in publicly-available databases, such as GENBANK[®] and SWISSPROT. Information including sequences and other information related to such nucleotides and polypeptides included in such publicly-available databases are expressly incorporated by reference. Unless otherwise indicated or apparent the references to such publicly-available databases are references to the most recent version of the database as of the filing date of this Application.

[0062] The present application can “comprise” (open ended) or “consist essentially of” the components of the present invention as well as other ingredients or elements described

herein. As used herein, “comprising” is open ended and means the elements recited, or their equivalent in structure or function, plus any other element or elements which are not recited. The terms “having” and “including” are also to be construed as open ended unless the context suggests otherwise.

[0063] Following long-standing patent law convention, the terms “a”, “an”, and “the” refer to “one or more” when used in this application, including the claims. Thus, for example, reference to “a cell” includes a plurality of such cells, and so forth.

[0064] Unless otherwise indicated, all numbers expressing quantities of ingredients, properties such as reaction conditions, and so forth used in the specification and claims are to be understood as being modified in all instances by the term “about”. Accordingly, unless indicated to the contrary, the numerical parameters set forth in this specification and claims are approximations that can vary depending upon the desired properties sought to be obtained by the presently-disclosed subject matter.

[0065] As used herein, the term “about,” when referring to a value or to an amount of mass, weight, time, volume, concentration or percentage is meant to encompass variations of in some embodiments $\pm 20\%$, in some embodiments $\pm 10\%$, in some embodiments $\pm 5\%$, in some embodiments $\pm 1\%$, in some embodiments $\pm 0.5\%$, and in some embodiments $\pm 0.1\%$ from the specified amount, as such variations are appropriate to perform the disclosed method.

[0066] As used herein, ranges can be expressed as from “about” one particular value, and/or to “about” another particular value. It is also understood that there are a number of values disclosed herein, and that each value is also herein disclosed as “about” that particular value in addition to the value itself. For example, if the value “10” is disclosed, then “about 10” is also disclosed. It is also understood that each unit between two particular units are also disclosed. For example, if 10 and 15 are disclosed, then 11, 12, 13, and 14 are also disclosed.

[0067] As used herein, “optional” or “optionally” means that the subsequently described event or circumstance does or does not occur and that the description includes instances where said event or circumstance occurs and instances where it does not. For example, an optionally variant portion means that the portion is variant or non-variant.

[0068] Although any methods, devices, and materials similar or equivalent to those described herein can be used in the practice or testing of the presently-disclosed subject matter, representative methods, devices, and materials are now described.

[0069] The presently-disclosed subject matter is further illustrated by the following specific but non-limiting examples. The following examples may include compilations of data that are representative of data gathered at various times during the course of development and experimentation related to the present invention.

EXAMPLES

[0070] **Example 1:** Nanopore of SPP1 DNA Packaging Motor for Dynamic Analysis of Protein Conformation and Folding in Real Time

[0071] Information about conformation and folding of proteins can convey biological functions. A simple method to replace conventional sophisticated and labor-intensive methodology to detect protein conformation and kinetic folding in real time is desirable. While nanopore technology has become a powerful tool in single molecule sensing, protein nanopore appears more advantageous concerning channel amenability, structure homogeneity, and production reproducibility. However, most of the well-studied protein nanopores are ion channels with a pore size of only around 1.2-nm, which is too small to allow passage of protein or peptides that are typically in multiple nanometers scale. To acquire protein channels with larger pore sizes, we reengineered a robust portal channel from bacteriophage SPP1 that allowed the translocation of peptides with higher ordered structure to produce clear and reproducible electron signatures. By combining optical single molecule microscopy with resistive pulse technologies, we observed translocation of peptides quantitatively and analyzed their conformation and dynamic folding in real-time at single molecule level. The secondary structure and conformation, such as the oxidative and the reduced states, were directly and clearly differentiated. A similar ΔG^0 was obtained when different concentrations of substrates were applied, suggesting that the use of SPP1 nanopore for real-time quantification of peptide folding is feasible. With the intrinsic nature of size and conjugation amenability, this SPP1 nanopore has the potential for development into a tool for the quantification of protein structure and folding in real time.

[0072] Living systems contain wide varieties of nanomachines with diverse structures and functions. The ingenious design of viral DNA packaging motors and their intriguing mechanism of action has provoked a wide range of interests among scientists in many different areas¹⁻⁵. Tailed bacteriophages are viruses that infect and reproduce within a bacterium, and widely exist in the biosphere. The portal protein is one of the essential

components of the viral packaging motors with a turbine-like shape^{6,7}. SPP1 is a dsDNA phage that infects *Bacillus subtilis*. The DNA packaging motor of SPP1 consists of a terminase composed of small (gp1) and large (gp2) subunits, portal protein gp6, and a two head completion proteins gp15 and gp16^{8,9} that powers the encapsulation of 45.9 kbp genomic DNA¹⁰. The central core of the SPP1 motor is a portal channel, termed connector, which is composed of 12 or 13 copies of protein gp6^{9,11}. Explicit engineering of the SPP1 portal protein is possible due to its available crystal structure^{12,13}. The connector has an overall diameter of 16 nm and a height of 10.5 nm. The narrowest constriction of the internal channel is ~3 nm (**Fig. 1**)^{12,13}.

[0073] Nanopore technology has recently emerged as a real-time and high-throughput single molecule detection method, holding great potential for sensing a wide range of analytes, molecular diagnostics and DNA sequencing applications¹⁴⁻²³. Solid state nanopores generated by microfabrication generally have less reproducible pore sizes and lack chemical and location selectivity. Protein nanopores harvested in bacteria are homogenous in size and can be functionalized with probes, but commonly used nanopores derived from ion channels have an internal channel diameter of only ~1.2 nm²⁴⁻²⁷. In search for alternative larger sized protein channels, we have previously developed membrane-embedded phi29 motor channels²⁸⁻³⁰ for single molecule sensing of nucleic acids^{28,29,31-34}, chemicals³⁵, or binding assays of antibody based on channel conformational change^{31,36}. Herein, we reengineered the SPP1 connector and inserted it into a lipid bilayer to serve as a robust nanopore for detecting the translocation and conformational dynamics of peptides and proteins at single molecule level.

[0074] Elucidating the structural conformation and folding of proteins and peptides is critical for understanding their biological functions. A wide range of biophysical methods, such as X-ray crystallography, NMR, Circular Dichroism, Dual Polarization Interferometry, and Mass Spectrometry have been used to investigate folding and dynamic structural changes of proteins. However, these methods require expensive instrumentation and specialized labor. Nanopores offer an attractive alternative as they are intrinsically single molecule in nature requiring ultra-low sample volumes, are label free, amplification free, and function using a simple detection process requiring no specialized expertise. While the translocation of DNA and RNA have been studied extensively in biological nanopores^{14,16,24,37}, studies on translocation of protein or peptides have just been emerged. This is mainly due to the size limitation of the commonly used protein pores, which are not large enough to allow the passage of proteins. The proteome can be an accurate and direct indicator of current health

status of patients³⁸. For example, early diagnosis and monitoring the changes of amyloid- β peptide and α -synuclein are critical for the management of Alzheimer's and Parkinson's disease³⁹.

[0075] A handful of studies demonstrating peptide translocation using nanopores have been published, but quantitative analysis and translocation validation remain challenging⁴⁰⁻⁴⁷, since there is no technique available for amplifying protein substrates as in DNA or RNA qualification using PCR⁴⁸. The availability of only trace amounts of peptide or protein for analysis after translocation is far beyond the sensitivity threshold of classical protein detection methods. In this report, we combined single molecule fluorescence microscopy and resistive pulse technique to quantitatively study peptide translocation through SPP1 connectors and elucidate the structural conformations of peptides at the single molecule level. This new nanopore peptide sensing technology with explicit engineering capabilities has the potential to transform the proteomics field, including future biomarker analysis, early disease diagnosis and even protein sequencing.

[0076] RESULTS

[0077] *Characterization of SPP1 reengineered nanopore embedded in a lipid bilayer*

[0078] Structural analysis revealed that the central region of the SPP1 connector shows slight hydrophobicity compared with the flanking regions at the N- and C-terminal ends which are more hydrophilic (**Fig. 1a-b**). To facilitate SPP1 connector purification, a 6 $\times$ His-tag was inserted at the C-terminal end along with a 6 $\times$ glycine linker for end-flexibility. The presence of His-tag enhanced the hydrophilicity of the C-terminal, thus making the hydrophilic–hydrophobic–hydrophilic layers more distinct, which is necessary to mimic the lipid bilayer architecture. After His-tag column purification, the purified protein was further purified by 15-35% glycerol gradient ultracentrifugation to verify whether SPP1 subunit could assemble into the channel complex (**Fig. 7**). Bovine serum albumin (BSA) with molecular weight of 60 KDa, which is almost equal to a single subunit (58 KDa) of the SPP1 connector, centered at fraction 27, whereas the SPP1 portal complex was at fraction 17, indicating the formation of higher molecular weight complex.

[0079] To incorporate SPP1 portal channel into planar lipid membranes, a two-step procedure was employed²⁸. The connector was first reconstituted into liposomes and then the liposome-connector complex was fused with a planar lipid membrane. Direct incubation of the connector or liposome alone with a planar lipid bilayer did not result in connector insertion into the lipid bilayer. Single channel conductance assay was performed to measure the electrophysiological properties of membrane-embedded SPP1 connectors. The connector

insertion steps were observed as distinct stepwise increase in conductance as revealed in a continuous current trace (**Fig. 2a**). The insertion of single portal channel results in ~200 pA in current jump under an applied potential of 50 mV in conducting buffer (1 M KCl, 5 mM HEPES, pH 8). Occasionally ~400 pA current jumps were observed, attributed to simultaneous insertion of two connectors. The average conductance of reengineered SPP1 connectors is 4.27 ± 0.27 nS (**Fig. 2b**). The conductance is uniform without displaying any voltage gating phenomena under the reported conditions of ± 50 mV (**Fig. 2c**).

[0080] *Characterization of peptide translocation through reengineered SPP1 connector*

[0081] A positively charged 12 amino acid TAT peptide was used in the translocation studies with sequence Cys-Tyr-Gly-Arg-Lys-Lys-Arg-Arg-Gln-Arg-Arg-Arg. Due to the presence of cysteine at the N-terminus, the peptide forms a dimer by disulfide bond under physiological conditions, which is confirmed by mass spectroscopy. In the absence of peptide, the current trace was quiescent. In contrast, when the peptide was premixed with the conducting buffer in both *cis*- and *trans*-chambers, a burst of transient blockage events were observed immediately after the insertion of connector in the lipid membrane (**Fig. 3a**). As the peptide concentration was increased from 1.25, 2.5, 3.75, to 5 $\mu\text{g/mL}$, the density of current blockage events increased with a corresponding linear increase in peptide translocation rate from 2.33 ± 1.54 , 7.5 ± 1.66 , 10.17 ± 3.58 , to 16.96 ± 6.01 events per second (**Fig. 3b**). One parameter used to characterize the translocation was the current blockage percentage, calculated as the ratio of current blockade resulting from peptide translocation to the open current of one portal channel, expressed as $[(1 - \text{Current}_{\text{peptide}}) / \text{Current}_{\text{open_channel}}]$. The distribution of current blockage was relatively broad with a major peak at $\sim 55.1 \pm 3\%$ (**Fig. 3c**). Another parameter was the dwell time (τ), the time taken for the peptide to traverse from one end of the connector to the other end. The dwell time distribution followed an exponential decay with a rate constant of $0.84 \pm 0.09 \text{ s}^{-1}$ (**Fig. 3d**).

[0082] *Quantitative validation of peptide translocation by single molecule fluorescence imaging*

[0083] To validate the translocation of peptide through SPP1 connector, we conducted single molecule fluorescence imaging of samples collected from patch clamp experiments. HPLC purified Cy3 conjugated TAT peptide at a final concentration of 2.5 ng/ μL was added to the *cis*-chamber after stable insertion of SPP1 connectors in the lipid bilayer. Under an applied negative trans-membrane voltage, the positively charged peptide translocated through the portal channel to the *trans*-chamber. 50 μL samples were collected from the *trans*-

chamber at 0, 20, 40, and 60 minutes after addition of Cy3-TAT peptide and loaded onto glass coverslips. The positively charged peptide bound to the negatively charged glass surface through charge-charge interactions and appeared as individual fluorescent spots (**Fig. 5a**). Fluorescence imaging revealed that the number of Cy3 spots in the field of view increased over the time course of 60 mins (**Fig. 5b**). In contrast, in a control experiment under the same conditions but in the absence of portal channel, very few Cy3 spots were observed, compared to the sample containing SPP1 connector (**Fig. 5b**). Since TAT is a membrane penetrating peptide, it is conceivable that a small fraction could potentially pass through the lipid bilayer, contributing to the small increase in the fluorescence background signal.

[0084] *Kinetic analysis of TAT conformational status in real time*

[0085] The focus of this study was to investigate whether the large and robust SPP1 motor nanopore can be used for kinetic analysis of protein folding in real time. TAT peptide with and without a cysteine was used as a model system to study conformational changes. Structural analysis of TAT sequence using the computer program PEP-FOLD⁴⁹ revealed that without forming disulfide bond, the TAT only adopts one α -helical conformation (**Fig. 4**). However, in the presence of a cysteine at the N-terminus, the peptide can be oxidized into a dimer by forming a disulfide bond between two N-terminal cysteines and exhibiting two parallel α -helices. The presence of two states, the oxidized and the reduced conformations, were confirmed by translocation studies with the SPP1 portal channel. Under an oxidized state, the current blockage distribution by TAT displayed a major peak, centered at $55.1 \pm 3\%$ (**Fig. 4a**), with a minor peak centered at $28.5 \pm 1.9\%$. However, after adding the reducing agent TCEP [(tris(2-carboxyethyl)phosphine)] to break the disulfide bond, the blockage distribution significantly changed. The peak of $55.1 \pm 3\%$ decreased significantly with time, while the majority of the blockade distribution shifted to $28.5 \pm 1.9\%$. This indicated that most of the peptide passing through the channel is in its single α -helical conformation (**Fig. 4b**). Current blockage distribution of the reduced TAT peptide was similar to Cy3-TAT signature with TCEP treatment, since the conjugation of Cy3 prevented the disulfide bond formation and resulted in similar blockage, representing the single TAT helix (**Fig. 4c**).

[0086] To quantify the kinetics parameter, we further examined the conformational changes of TAT peptide upon addition of TCEP and recorded the change of the current blockage profile in real time (**Fig. 6**). The current blockade signature progressively shifted from predominantly $55.1 \pm 3\%$ to $28.5 \pm 1.9\%$, representing the real-time kinetic process of disulfide bond reduction (**Fig. 6a**). To derive the kinetic parameter of this reaction process,

two parameter γ (defined as the ratio of the number of peak A versus the sum of A and B) and reaction quotient Q_r were determined (**Fig. 6b, d**). The transition of γ and Q_r as shown in **Fig. 6b** and **5d** represents the timely evolution of the reduction reaction. The γ and Q_r is theoretically derived by applying second-order reaction dynamics and diffusion model, as follows:

$$\begin{array}{ccccccc}
 TAT & -TAT & +TCEP & \xrightarrow{H_2O} & 2TAT & +O-TCEP \\
 C_0 & & C(t) & & 0 & & 0 \\
 C_t & C(t) - (C_0 - C_t) & 2(C_0 - C_t) & & C_0 - C_t & & \\
 \frac{dC_t}{dt} = -kC_t[C(t) - (C_0 - C_t)], & & & & & & \\
 \text{since } C(t) \gg C_0, & & & & & & \\
 \text{then, } C_t = C_0 * \exp(-kC(t)t) & & & & & & (1)
 \end{array}$$

$C(t)$ is the TCEP concentration at the pore entrance. According to diffusion equation, $C(t)$ has

the following form: $C(t) = \frac{N}{(4\pi Dt)^{1.5}} \exp(-\frac{r^2}{4Dt})$.

[0087] In order to study the TAT reduction process quantitatively, we calculated the ratio

$$\gamma \left(\gamma \equiv \frac{[TAT - TAT]}{[TAT - TAT] + [TAT]} \right) \text{ and the reaction quotient } Q_r \left(Q_r \equiv \frac{[TAT]^2 * [O - TCEP]}{[TAT - TAT] * [TCEP]} \right), \text{ unit}$$

is normalized). Because there is a ε ($\varepsilon \equiv \frac{[TAT]}{[TAT - TAT]}$, prior to adding TCEP) portion of

TAT monomer at the beginning, the equation was solved to obtain the following equation:

$$\gamma = \frac{1}{2 \exp\left(\frac{Nk}{(4\pi D)^{1.5} t^{0.5}} \exp(-\frac{r^2}{4Dt})\right) - 1 + \varepsilon} \quad (2)$$

$$Q_r = \frac{2(1 - \gamma)^3 C_0^2}{\gamma C_r (1 + \gamma)^2}. \quad (3)$$

[0088] $C(t)$ is the equilibrium TCEP concentration. To evaluate whether the parameters derive from this real time quantification analysis represent the real parameters of the reaction, the ΔG° value were determined from different assays using constant TAT but varied concentration of TCEP from 100, 250, 500 μ M, and compared with the ΔG° derived by conventional methods.

[0089] The red line in **Fig. 6b** is the fitting curve with equation (2) and Q_r in **Fig. 6d** is derived from γ with equation (3). When the reaction approaches equilibrium, Q_r is equal to standard equilibrium constant K° . From **Fig. 6d**, K° is estimated to be 1.4×10^4 and corresponding to -23.3 kJ/mol Gibbs free energy (ΔG°).

[0090] As shown in **Fig. 6b**, γ decreased over the course of reaction time, indicating the decrease in oxidized dimer conformation and concurrent increase in reduced monomeric state. Fitting the data to a single exponential function revealed a rate constant, $k = 0.989 \pm 0.096 \text{ min}^{-1}$ and $t_0 = 5.2 \pm 0.04 \text{ min}$. The trend is also demonstrated in current blockage percentage versus dwell time distribution of events over the course of the reaction time (**Fig. 6c**). All the data from different concentrations of TCEP, show the same trends to approach equilibrium status. The related K° and ΔG° are depicted in table 1.

[0091] The assay revealed that, when different concentration of TCEP were used, the ΔG° were very similar (Table 1). When the concentration of TCEP was increased from 100, 250, to 500 μM , the ΔG° were -24.5, -23.1, and -23.3 kJ/mol respectively. The similarity of the ΔG° derived from different assays with variable TCEP concentration suggests that using SPP1 nanopore for real time quantification of peptide folding is feasible. In addition, ΔG° derived from these experiments is very close to the ΔG° of -27 kJ/mol for disulfide bond in protein obtained by conventional methods⁵⁰.

Table 1: Parameters for the quantification of dynamic folding of TAT peptide in real time

TAT concentration (nM)	TCEP concentration (μM)	Q_x (10^3)	K° (10^3)	ΔG° (kJ/mol)
300	100	26.6	26.6	-24.5
300	250	11.8	11.8	-23.1
300	500	13.9	13.9	-23.3

[0092] DISCUSSION

[0093] In this study, we showed that the SPP1 portal channel can be successfully incorporated into planar lipid bilayer to serve as a nanopore with robust electrophysiological properties. With a conductance of 4.27 nS, it is one of the larger channels and only the second viral protein channel (apart from phi29 connector)²⁸ to be incorporated into a lipid membrane. In the presence of TAT peptide, a burst of current blockage events with characteristic current amplitude and dwell time were observed indicating the translocation of the peptides through the SPP1 connector. This was further validated by single-molecule fluorescence imaging. The oxidized and reduced conformations of TAT peptide can be clearly distinguished in real time. The results demonstrate the potentials of nanopore-based proteomics.

[0094] Direct evidence of DNA translocation through α -hemolysin was demonstrated twenty years ago by quantitative PCR⁵¹. Although a handful of studies utilizing peptide and protein translocation through nanopore, have been published over the last several years including protein folding, unfolding, stability and disease detection^{41,44,45,52-61}. Here, we developed a very simple method to provide evidence of peptide translocation using single molecule fluorescence imaging of samples obtained from the nanopore setup. This method can be easily adapted and employed by other laboratories to prove protein translocation and quantitatively study peptide translocation through nanopores.

[0095] In this study, we further demonstrated that this new protein nanopore is capable of detecting kinetics of TCEP mediated reduction of disulfide bond and demonstrating peptide conformational changes in real time. From **Fig. 6**, the frequency of dimer forms (~55%) decreased gradually to monomer forms (30%), upon addition of TCEP, as the reaction time increased. Analyzing peptides and proteins is a complex task, since they fold into versatile and complex three dimensional structures composed of 20 building blocks compared to DNA or RNA with only 4 building blocks. Further improvements are necessary to develop the nanopore into a robust platform for proteomic profiling. The current blockage signature based on the translocation profile can be used to investigate the length, charge, hydrophobicity, secondary structures and ultimately the amino acid sequences of the peptides. The kinetics of protein folding and unfolding as well as entropic and energetic contributions can be further dissected at the single molecule level.

[0096] CONCLUSION

[0097] A robust portal channel of bacteriophage SPP1 has been reengineered and inserted into lipid bilayer to allow the translocation of peptides with higher ordered structure to produce clear and reproducible electron signatures. Optical single molecule fluorescence microscopy were combined with resistive pulse technologies to detect the translocation of peptides quantitatively and to analyzed their conformation and dynamic folding in real-time at single molecule level. The oxidative and the reduced conformation were clearly differentiated and their timely transition in the presence of reducing agent were directly observed. When different concentrations of substrates were applied, a similar ΔG^0 was obtained, suggesting that the use of SPP1 nanopore for real-time dynamic quantification of peptide folding is feasible.

[0098] MATERIALS AND METHODS

[0099] Materials: The phospholipid 1, 2-diphytanoyl-*sn*-glycerol-3-phosphocholine (DPhPC) was obtained from Avanti Polar Lipids, Inc. Organic solvents (*n*-decane and chloroform) were purchased from Fisher Scientific, Inc. and TEDIA, Inc., respectively. TAT peptide was custom-ordered from *GenScript*, Inc. All other reagents were purchased from Sigma or Fisher, if not specified.

[00100] Cloning and Purification of the SPP1 connector: Gene gp6 encoding SPP1 portal was synthesized and cloned into PET3 vector between *NheI* and *BamHI* by *GenScript*, Inc. His-tag was inserted into the C-terminal of the connector for purification. Then the plasmid was transformed into BL21 (DE3) for expression and purification. The purification procedure has been described previously^{28,30}. Briefly, the transformed bacteria were cultured in 10 mL LB medium overnight at 37°C. The bacteria were transferred to 1 L of fresh LB medium. When the OD₆₀₀ reached ~0.5-0.6, 5 mM IPTG was added to the medium to induce protein expression. The bacteria were collected by centrifugation after 3 hours continuous culture. Bacteria was lysed by passing through French press. Lysed cell was separated by centrifugation and supernatant containing expressed protein was collected. The supernatant was purified with Nickel affinity chromatography (Novagen)⁶². His Binding Buffer (15% glycerol, 0.5 M NaCl, 5 mM Imidazole, 10 mM ATP, 50 mM Tris-HCl, pH 8.0), and the cleared lysate was loaded onto a His•Bind® Resin Column and washed with His Washing Buffer (15% glycerol, 0.5 M NaCl, 50 mM Imidazole, 50 mM Tris-HCl, pH 8.0). The His-tagged connector was eluted by His Elution Buffer (15% glycerol, 0.5 M NaCl, 0.5 M Imidazole, 50 mM Tris-HCl, pH 8.0). The purified connector from chromatography was further purified by a 15-35% glycerol gradient ultracentrifugation at 35K for 2 hrs. Before incorporating into liposomes, the purified connector was dialyzed against buffer (0.5 M NaCl, 50 mM Tris-HCl, pH 8.0) to remove excess glycerol.

[00101] Insertion of the connector into planar lipid bilayer: The protocol for the incorporation of connectors into lipid bilayer has been reported^{28-31,63}. Briefly, planar bilayer lipid membranes (BLMs) were generated in a BCH-1A horizontal BLM cell (Eastern Scientific). A Teflon partition with a 200 µm aperture was placed in the apparatus to separate the BLM cell into cis- (top) and trans- (bottom) compartments. A planar lipid bilayer was formed by painting the aperture with 0.5 µL of 3% (w/v) DPhPC in *n*-decane. A conducting buffer (1 M KCl, 5 mM HEPES, pH 7.8) was added to both the top and bottom compartments of the BLM cell, and Ag/AgCl electrodes were placed in the buffer of each compartment. The electrode in the trans-compartment was connected to the headstage of an Axopatch 200B amplifier (Axon Instruments, Inc.), and the electrode in the top compartment was grounded.

Connector insertion was observed after addition of 1 μ L of the diluted Liposome/connector complex to the cis- compartment directly.

[00102] Electrophysiological measurements: The headstage and Axopatch 200B patch clamp amplifier were connected to a DigiData 1440 analog-digital converter (Axon Instruments, Inc.) to monitor and record electrochemical currents through BLMs²⁸⁻³⁰. The current recordings were low-pass filtered at a frequency of 1 or 5 kHz. The sampling frequency was 20-200 kHz in all experiments, unless otherwise specified. The data were recorded with pClamp 9.1 software (Axon Instruments, Inc.), and analyzed with the Clampfit module of pClamp 9.1 and OriginPro 8.1 (OriginLab Corporation).

[00103] Peptide translocation experiments: TAT peptide (Cys-Tyr-Gly-Arg-Lys-Lys-Arg-Arg-Gln-Arg-Arg-Arg) with a final concentration of 23 μ g/mL was premixed with the conducting buffer before the insertion of connector. For quantitative validation, Cy3 fluorophore was conjugated to TAT peptide by sulfhydryl-maleimide chemistry and purified by HPLC. Multiple channels were inserted into bilayer lipid membrane and then Cy3-TAT peptide was added into cis-chamber. Samples were collected from the trans-chamber after 0, 20, 40, and 60 min.

[00104] Single molecule fluorescence imaging: Samples collected from the patch clamp were incubated with the ozone pre-treated glass substrate for 10 min before imaging to ensure sufficient adsorption. A 532 nm green laser was used for the excitation of Cy3. A 60 $\times$ objective (N.A. = 1.4, oil immersion) was used for fluorescence imaging. The signals were recorded using Andor iXon 887 V electron multiplied CCD camera. Images were taken with an exposure time of 500 ms. The number of spots in the images were counted using iSMS software⁶⁴.

[00105] Example 2: Translocation of double strand DNA through C-his SPP1 phage nanomotor embedded in lipid bilayer.

[00106] During the process of assembly of tailed bacteriophages, their genome is packaged within a preformed procapsid through a tunnel formed by the portal protein assembly. Such entropically unfavorable process is accomplished by an ATP-driven molecular motor.⁶⁵⁻⁶⁸

[00107] For spp1 bacteriophage, the molecular motor consists of three proteins:gp1, gp2 (ATPase) and gp6 (portal protein). Such cyclical homooligomers are located at a single vertex of the icosahedral viral capsid that binds to the tail. During the process of assembly, portal protein translocate double-stranded DNA. After termination of packaging, portal protein is binded by head completion proteins (gp15 and gp16), which form a head-to-tail

connector stereoscopic structure(Andrey A Lebedev, The EMBO Journal 2007).In common with the other herpesvirus portal protein ⁶⁹, the portal protein of bacteriophage SPP1 can exist as a circular assembly. The novelty and ingenious design of viral DNA packaging motors and their mechanism of action have provoked a broad range of interest among scientists in nanotechnology, biophysics. The genome of linear double-stranded DNA(dsDNA) virus is packaged into a preformed delicate procapsid.

[00108] There exists debate about the molecular motor work mechanism of how each different component coordinate with each other. Hendrix proposed that DNA translocation is accompanied by rotation of portal protein based on the fact the low-energy barriers to rotation of symmetry mismatching protein rings relative to each other. Another models which involve the rotation of the portal protein DNA translocation is accompanied by sequential conformational changes propagating along the belt of tunnel loops with the porta protein ⁷⁰⁻⁷¹. Also, the structure of the most constricted part of the internal loops which is expected to be in close contact with the DNA during translocation has been characterized and defined. The portal protein has varying number of subunits: it exerts as a 13-subunit assembly in its isolated form but as a 12-subunit assembly active form when integrated into the functional viral capsid⁷². Based on the above mentioned models, a possible scenario of events during DNA translocation was discussed⁷³. There is a close interaction between the portal protein and DNA and such interactions are independent of whether the viral ATPase itself performs the power stroke pushing the DNA into the capsid or whether portal protein apply the force to the DNA in response to conformational changes. Also, a 4-steps model for DNA gating(a state from open state to closed state) was proposed ⁷⁴. During this model, the nascent prohead portal binds to the packaging proteins and then a conformational change causes termination of packaging and conformational switch. After the neck and tail assembly, DNA is delivered into the host.

[00109] The information encoded in DNA is the footing stone for the development of modern medicine and life sciences. The development of individualized medicine and related scientific research has prompted the development of new kind of sequencing techniques beyond the original Sanger sequencing. As one of potential technologies, nanopore sequencing is one the rapidly sequence approaches which is expected to sequence a human genome for under \$1000. So far, biological nanopores have been used to study the transport of DNA and other molecules like DNA, RNA, chemical agents, peptides, proteins and polymers⁷⁵⁻⁷⁹. But due to the limitation of pore dimension, most pores only allow small molecules and single-strand DNA or RNA, such as hemolysin and MSPA. α -haemolysin (α -

HL) pore has been used to study the transportation of single DNA or RNA by using the transient blockade of ionic current. Bacteriophage phi 29 motor, having a larger pore size of 3.6nm, have highly widen nanopore biomedical applications for potential DNA sequencing. There are a variety of engineered phi 29 channel have been developed to detect dsDNA, ssDNA, chemical agents and even carcinogenic antigens⁸⁰⁻⁸³. For bacteriophage spp1 molecular motor, it has similar pore size of 3.5nm which make it very suitable for dsDNA sequencing because dsDNA normally just have a diameter of about 2 nm. However, the electrophysiological measurements of spp1 connector channels for translocation of double strand DNA have not been reported so far. In this study, the wild type spp1 connector was cloned and purified. After that, the connector was inserted into liposomes and lipid bilayer. dsDNA translocation was confirmed by single channel conductance measurements and Q-PCR assay. Our study expands the future application of spp1 nanopore in nanomedicine and nanobiotechnology.

[00110] Results

[00111] *Cloning and purification of C-his spp1 connector*

[00112] To facilitate connector purification, a C-terminus was inserted with a poly histidine tag for improved affinity tag flexibility. The linker was included to provide end flexibility. After purified by his-tag affinity column, it was found that the C-his spp1 connector self-assembled a unique 13-fold symmetric structure, which is compared with 12-fold *in vivo* active form, as observed by transmission electron microscopy. The existence of the motor configuration was demonstrated through its ability to package the dsDNA into the procapsid and to assemble the resulting DNA-filled capsid into the infectious phi29 virion.

[00113] *Reconstituting the connector into liposomes*

[00114] A protocol for reconstituting the connector into liposomes was developed by co-incubation of the connector with the DOPC or DPhPC lipid and hydration-rehydration of the lipid film with proteoliposome containing the reengineered connector in the presence of sucrose. Such rehydration and co-incubation make the hydrophobic layer of the connector easier to interact with the hydrophobic domain of the lipid molecules. By such hydration-dehydration process, small unilamellar vesicles containing spp1 connectors(50 μ m) was developed. Fluorescence microscopy and sedimentation analysis was used to confirm the insertion of the connector protein into the lipid membrane. The presence of the connector in the membrane was showed a clear florescent ring around the lipsome. While as a negative control, no fluorescent ring was observed when fluorescently tagged connector was mixed

non-specifically with the non-connector inserted liposome. The free connectors were removed by 5-20% sucrose gradient ultracentrifugation.

[00115] *Incorporating the connector into planar lipid membranes*

[00116] The purified connector was then inserted into a planar lipid bilayer membrane (BLM). The insertion was demonstrated by the discrete stepwise increase of the current. The number of the channels inserted can be counted by counting of the number of the stepwise.

[00117] The step size of each current augment was almost identical, representing the even and uniform structure and stoichiometry. This was similar with the phi29 gp10 connector channel, which also can form quite uniform and even channels in the BLM^{75, 79}. The channel conductance of the connector was compared with that of phi29 gp10 using solutions of different ionic strength. Similarly, after a statistical analysis of conductance of two kinds of channel: wild type phi29 GP10 connector and C-his spp1 connector, two sharp Gaussian distributions were shown. Comparing to the conductance of phi29 gp10 connector, spp1 connector has a similar conductance of 4.4 nS in the presence of 1M KCl, 5mM HEPES, representing that both spp1 and phi29 gp10 connector have similar size. This can also be demonstrated by the crystal structure of two connector proteins^{70, 79}. In addition, the channel demonstrates a perfectly linear relationship with respect to the applied voltage of 50mV. The spp1 connector did not display voltage gating properties under the conditions described above but it showed gating at a higher voltage of 75mV.

[00118] *Translocation of double-strand DNA*

[00119] In this report, both linear and circular plasmid were used to examine the translocation to examine the translocation of dsDNA through the connector channel. In the absence of linear DNA, the current trace was quiescent and only a quite small number of unspecific blockades were observed with a long time trace. But when dsDNA (142bp) were added in to the cis chamber, numerous current blockades were observed, which led to the current jump of single connector insertion to be transiently reduced by 28-34%. When the voltage changed to negative potential, the blockades cannot be observed. This implies that the connector exerts one-way traffic for dsDNA translocation, which is similar with one-way traffic mechanism of phi29 gp10 connector. When circular plasmid dsDNA was used, no translocation of the circular plasmid was observed. Interestingly, the same amount of circular plasmid was digested by DNase and then added into the chamber, a burst of transient blockades occurred. All the above results pile up to prove that only the linear dsDNA can pass through the connector channels.

[00120] The blockade rates were affected by two factors: DNA concentration and transmembrane voltage. In the presence of 30 μ M DNA under one channel insertion, the blockade rate was approximately 18 blockades per second. When the concentration of DNA placed in the chamber was doubled and tripled, the blockades frequency per second almost doubled and tripled. This implies that the blockades frequency is direct proportion to the DNA concentration.

[00121] To calculate the dwell time(τ) for DNA translocation events, only the blockade episodes greater than 30% were grouped as this percentage of blockade seemed consistent with the ratio of the cross-sectional area between dsDNA and the pore. It should also be noted that 8(under 75mV) individual outlying events which scattered from 100ms to 10000ms were excluded in the graph for clarity. The dwell time distribution under 40mV seemed broader than that under 75mV. The average dwell time for DNA blockades under 75mV was 14.6 ms.

[00122] To verify the passage of dsDNA through the connector channels, quantitative polymerase chain(PCR) (Q-PCR) was used to quantify the translocation of 142-bp DNA under a voltage of 75mV. DNA was added to the trans side and samples were taken from the cis side for quantification at 30 minutes intervals. For comparison, control experiments were performed in the absence of connectors.

[00123] The SPP1 connector protein used in this example comprises the sequence of SEQ ID NO: 4.

[00124] **Example 3:** Translocation of peptides through membrane-embedded C-his SPP1 motor protein nanopores.

[00125] Living systems contain a wide range of nanomachines with diverse structures and functions. Tailed bacteriophages are a virus that infect and reproduce within a bacterium, and widely exist in the biosphere. Among the variety of bacterial viruses, viral connector protein is one of the essential components in packaging motor with turbine-like shape. SPP1 is a dsDNA phage which infects *bacillus subtilis* and the connector of SPP1 is composed of portal protein gp6, and two head completion protein gp15 and gp16.^{84,85} The packaging motor of SPP1, which consist of gp1, gp2, and gp6, powers encapsulation of the 45.9 kb genome.⁸⁶ The ingenious design of viral DNA packaging motors and the novel mechanism of action has provoked a wide range of interest among scientists in many different areas. Although various portal proteins share little homology in primary sequence, and varies in size

⁸⁷⁻⁸⁹, they exhibit significant morphological similarity. Portal protein of SPP1 and phi29 share similar secondary structure and 3-D structure. Our recent research also found that they even share similar role in packaging genome ⁹¹⁻⁹³. Explicit engineering of the SPP1 connector is possible due to its accessible crystal structure ^{94,95}. SPP1 portal protein have 13-subunit form and the pseudoatomic structure with 12-subunits. The overall diameter of 13-subunit form is ~1.65 nm, and the height is ~1.10nm. The narrow part is 2.77 nm in 13 subunit form and 1.81 in 12 subunit form. ^{94,95}

[00126] Nanopore technology has recently emerged as a new real-time single molecule sensing method, holding a great potential to reduce the cost of biosensing and DNA sequencing. Over the last decade, the α -hemolysin (α -HL) and MspA channel has been expansively investigated for sensing and DNA translocation. ⁹⁶⁻⁹⁹ Previously, we have successfully inserted phi29 connector channel from bacteriophages phi29 into lipid bilayer with robust properties. ^{100,101} A one-way traffic property for dsDNA translocation with a valve mechanism for DNA packaging has also been observed. ¹⁰² The reengineered connector channel is capable to discriminate single-strand DNA or RNA from double strand. By selectively engineering a probe on the connector, single chemicals and antibody can be identified based on their distinct fingerprints. ¹⁰³ The channel also exhibits gating (a transition from an open state to a closed state) at higher trans-membrane potentials, a process that induces a conformational change in the channel subunits. ¹⁰⁴

[00127] The translocation of DNA and RNA have been studied extensively in other biological nanopores, such as the α -hemolysin, MsPA channel. ¹⁰⁵ However, translocation of protein or peptide studies still have not attraction enough attention as it should be. Compared to the genome, the proteome is probably a much more accurate and direct indicator of the current health status of humans ¹⁰⁶. For examples, early diagnosis and monitoring the change of amyloid- β peptide is critical to the treatment of Alzheimer's and Parkinson's disease ¹⁰⁷. However, proteomic data practically when only trace amount of peptide or protein is present, is more difficult to acquire, because there is no amplified technique available for protein as DNA or RNA which can be readily amplified by the polymerase chain reaction ¹⁰⁸. In this report, we first time show that this reengineered SPP1 reengineered motor channel has been successfully incorporated into planar lipid bilayers, when reconstituted into liposomes, allows the translocation of various charged peptides with different level blockade. The peptides can be differentiated by the blockage of the channel. This engineered SPP1 connector channel has potential for peptide or protein related diagnosis and sequencing in the future.

[00128] RESULTS

[00129] Characterization of C-his SPP1 connector channel embedded into lipid bilayer
Portal protein of SPP1 and phi29 share high homology. Similarly to phi29 connector, the central surface region of SPP1 portal protein also shows slight hydrophobicity compared with the two flanking layers at both ends, which are hydrophilic. To facilitate SPP1 connector channel purification, a C-terminus his tag was inserted just downstream of a six-glycine linker for improved affinity tag flexibility. The linker was introduced to provide end flexibility. After purification to homogeneity, the molecular weight of single subunit of SPP1 is 56 kDa as revealed in SDS PAGE gel, in agreement with theoretical molecular weight. To verify whether C-his SPP1 subunit could assemble to complex correctly, 15-35% sucrose gradient ultracentrifugation was performed. The fraction of bovine serum albumin (BSA) with molecular weight 60 kDa, is located at fraction 27, which represented single subunit of C-his SPP1 connector, whereas the fraction of C-his SPP1 portal protein shift to fraction 17, demonstrating high molecular weight complex has been formed.

[00130] To incorporate C-his SPP1 connector channel into planar lipid membranes, two step procedures which was also used in phi29 connector channel, was employed that the connector channel was first reconstituted into liposome first and then the liposome connector complex was fused into lipid membrane. Direct incubation of the connector channel or liposome with a planar lipid bilayer did not lead to channel insertion into the lipid bilayer. Single channel conductance assay was performed to measure the property of SPP1 connector channel. The connector channel insertion was observed through a distinct stepwise increase in conductance. The insertion of single connector channel into lipid bilayer results in an approximately 200 pA in the current jump under a potential of 50 mV in 1 M KCl 5 mM HEPES pH 8. Occasionally 400 pA current jump was observed, attributed to two connector channels insertion. Similar results were also observed under different buffer condition (data not shown). As comparison, the conductance of C-his SPP1 connector channel was also measured under a scanning voltage. Uniform distribution of conductance under and linear I-V relationship without display any voltage gating phenomenon under the reported conditions, 50 and 65 mV. The average conductance for a single channel was 2.06 ± 0.08 nS 50 mV in 1 M NaCl 5 mM HEPES pH 8.

[00131] Translocation of peptides through C-his SPP1 connector channels and kinetic study

[00132] Several different peptides with various isoelectric point (PI) and length have been used to examine the translocation through SPP1 connector channel, including poly

histidine, poly lysine and CysTyrGlyArgLysLysArgArgGlnArgArgArg(TAT). In absence of peptide, the current trace was quiescent. Occasionally, unspecific blockades were detected with a minimum detectable time. Compared with peptide translocation events, these unspecific blockades happened rarely. Buffer pH was adjusted according to corresponding peptide pI to have peptide possessed positive charge in order to translocate through channel under voltage. A continuous current trace recording was shown before and after addition of peptide in real time. Before TAT was added, the current trace was quiescent with occasionally non-specific blockade events during long time recording time. Once TAT peptide was added to cis-chamber after X channel inserted into lipid bilayer, the blockage events were not immediately observed. After X minutes from the adding point, the blockade events gradually increased sparsely until reaching to a burst stage. This lagging time between the adding point and the events showing up, indicates the diffusion time from top of the solution to the nearby entrance of channel. In addition, the blockade events can only be observed under negative voltage, because the peptide was only added to cis-chamber and positive charged peptide can only translocate from cis- to trans-chamber under negative voltage. When switch the polarity to positive voltage, there is no translocation events observed. In the same experiment, however, when small amount of buffer was transferred from cis-chamber to trans-chamber, the translocation events have been observed under positive voltage. However, the translocation rate is much lower than that under negative voltage due to the concentration of TAT in trans-chamber is much lower than cis-chamber.

[00133] In comparison, when TAT peptide was premixed with buffer in both chamber before channel insertion, a burst of transient translocation events have been observed immediately after the first insertion occurred. One parameter used to characterize the translocation was the current blockage percentage, which represents the difference between the open connector channel and the reduced current caused by peptide translocation. Current blockage percentage was calculated as follows: size of current blockage resulting from peptide translocation to one connector channel divided by step size of the current for one connector insertion.

[00134] The reduced current caused by TAT translocation is about 115 or 170pA, and the blockade has two major distributions, $55\pm3\%$ and $81\pm4\%$ respectively. Translocation rate was influenced by peptide concentration. The current trace of single channel with different concentration of premixed TAT peptide under same time length. As the concentration increased from 1.25, 2.5 and 5 $\mu\text{g/ml}$, the density of blockade events has also increased correspondingly. The translocation rate versus the concentration. As concentration increases

from 1.25, 2.5, 3.75, 5 ug/ml, the translocation rate also increase linearly from 2.33 ± 1.54 , 7.5 ± 1.66 , 10.17 ± 3.58 , to 16.96 ± 6.01 events per second.

[00135] Theoretical PI of poly lysine is about 9.7, therefore here we used 1 M KCL 5 mM Hepes, pH 8. When poly lysine peptide was added to the chamber, similar results have been observed but with different level of blockade. The current trace of single channel under 1M KCl 5 mM Hepes, pH X, 65 mV and as the concentration increase from 2 ug/ml to 20 ug/ml, the translocation events increase dramatically. The reduced current is about and the blockade distribution is majored in 25%.

[00136] Similar to other peptides, the blockade events have been observed after adding poly histidines. Because poly histidine theoretical PI is only 7.6, we used 1 M KCL 5 mM Hepes, pH 6 in the experiments to ensure poly histidines possessed positive charge. Under 50 mV, translocation events have been observed immediately when poly histidines was premixed with buffer in both chamber. Compared with TAT and poly lysine, the blockade is not uniform and the major blockage distribution is majored in X. As concentration increase from X to 18 ug/ml, the translocation rate increase linearly from X.

[00137] DISCUSSION

[00138] In this study, we first time showed that C-his SPP1 connector channel successfully has been incorporated into planar lipid bilayers when reconstituted into liposomes. Similar to phi29 connector channel, SPP1 connector channel with a central tunnel plays crucial role during the DNA packaging or ejection¹⁰⁹. Although the connector proteins of different phage have no similarity in the primary sequence, available electron microscopy data has shown that they all share a common turbine-like structure (Carrascosa, 2012). SPP1 portal protein has 13-subunit in its isolated form, but as a 12 subunit from when assembled into the functional viral capsid (Orlova et al, 2003). The diameter of SPP1 connector channel as 13 subunit form in the constrict part is 2.8 nm, which is smaller than phi29 connector channel which is 3.6 nm. Similar to phi29 connector channel, structure analysis of SPP1 connector revealed that the central surface region of SPP1 connector also shows slight hydrophobicity compared with the two flanking layers at both ends.

[00139] Nanopore detection is an emerging technique that has been extensively used to study the DNA, RNA translocation given the promise of ultra-rapid nucleic acids sequencing. In contrast, study of polypeptide or protein translocation through a protein nanopore has not been pursued as aggressively. One of the biggest challenges probably is due to the complexity of protein or polypeptide which could fold into different structure and even interact with nanopore. Another problem is that neutral peptides cannot be driven through

the nanopore under voltage. In addition, sequencing peptides or protein with nanopores is even more challenging since there are 20 common amino acids in nature protein compared to 4 nucleotides in DNA.

[00140] The insertions result in a uniform conductance distribution, we first time Nanopores have been extensively utilized to study DNA, RNA and analytes. However, peptide or protein recognition or sequencing using nanopore technique is still not studies extensively and has been attract enough attention. Compared to the genome, the proteome is probably a much more accurate and direct indicator of the current health status of humans. Various modifications have been studied extensively on both biological nanopores and solid-state nanopores for use in nucleic acid sensing.^{110, 111} The modification of biological nanopores can be easily achieved by the insertion, deletion, or substitution of various amino acids.^{112, 113} Biological nanopores can also be modified by covalently linking molecular recognition agents in the interior lumen of the pores.¹⁰³ Solid-state nanopores, conversely, have been studied mainly by chemical modifications. Different assembly pathways from modified phage portal protein monomers have been reported on SPP1⁹⁴, phi29⁸⁷, T7¹¹⁴ and P2.¹¹⁵ Both 12-mer and 13-mer assembly of SPP1 connectors were obtained at different conditions⁹⁴. The mutations and modifications of these portal protein channels can be used to investigate and refine the nanopores for technical applications, such as DNA sequencing.

[00141] Translocation of single-stranded DNA or RNA by biological or solid state nanopores has drawn great attention and has been well studied by several research groups, because of its importance in DNA sequencing and many other applications in recent years.¹¹⁶⁻¹¹⁹ Discrimination between single- and double-stranded nucleic acid molecules by translocating single-stranded homopolymers poly (A), poly (U), and poly (C) through solid state nanopore has been reported.¹¹⁶ The translocation of long random sequence and short length ssDNA through solid state nanopores has been characterized compared with the dsDNA and the hybrid DNA/RNA.^{117, 118} The relatively large diameter of the connector channel is an ideal tool for studying double-stranded DNA transport,¹¹⁷ but detection of single-stranded nucleic acid remained difficult using the system reported previously.^{100, 101} In this paper, we generated smaller connector channels by deleting the internal loop segments of gp10 subunits and demonstrated its capability to detect ssDNA and ssRNA. The translocation events produced a uniform current blockade of 16-21%, with a peak value of ~19%. This blockade percentage is approximately half of the dsDNA blockade through the native channel, even though the cross-sectional area of the mutant channel is only 60% of the wild type.

[00142] CONCLUSION

[00143] The deletion of an internal flexible segment from the gp10 subunit of the phi29 DNA packaging nanomotor resulted in a smaller conductance channel with a pore diameter ~60% of the wild-type connector channel. While the single-stranded nucleic acids could not be detected by wild type channel, they showed a blockade of 16-21% while translocated through the smaller size channel. This mutant channel with smaller size is ideal for the detection and sensing of single-stranded nucleic acids, including ssDNA and ssRNA. Connector channels created from gp10 subunits with the internal loop deleted allowed for bidirectional translocation of single-stranded nucleic acids. This may also lead to the development of a hybrid system in which the smaller conductance channel can be reconstituted into a lipid bilayer, in conjunction with the wild type connector channel, for the simultaneous detection and sensing of both dsDNA and ssDNA.

[00144] MATERIALS AND METHODS**[00145] Materials**

[00146] The phospholipids 1, 2-diphytanoyl-sn-glycerol-3-phosphocholine (DPhPC) and 1, 2-dioleoyl-sn-glycerol-3-phosphocholine (DOPC) were obtained from Avanti Polar Lipids, Inc. Organic solvents (n-decane and chloroform) were purchased from Fisher Scientific, Inc. and TEDIA, Inc., respectively. The DNA and RNA oligonucleotides used in this study were obtained from Integrated DNA Technologies, Inc. All other reagents were purchased from Sigma-Aldrich, Inc., if not specified.

[00147] Cloning and Purification of the C-his SPP1 Connector Protein

[00148] Gene gp6 encoding SPP1 connector on the C-terminus was synthesized by Genescript company and then cloned into PET3 vector between NheI and BamHI. A His-tag was inserted into the C-terminal for purification. Then the plasmid was transformed into BL 21 (DE3) for expression and purification. The transformed bacteria were cultured in 10 ml Luria-Bertani medium overnight at 37 °C. Then the cultured bacteria were transferred to 1 L of fresh LB medium. 0.5mM IPTG was added to the medium to induce protein expression when OD₆₀₀ reached 0.5~0.6. The bacteria were collected by centrifugation after 3 hours continue culture. Bacteria was lysed by passing through French press. Lysed cell was differentiated by centrifugation and supernatant containing expressed protein was collected. The supernatant was purified with Nickel affinity chromatography (Novagen)¹²⁰. Cells were resuspended with His Binding Buffer (15% glycerol, 0.5 M NaCl, 5mM Imidazole, 10mM ATP, 50mM Tris-Cl, pH 8.0), and the cleared lysate was loaded onto a His•Bind® Resin Column and washed with His Washing Buffer (15% glycerol, 0.5 M NaCl,

50mM Imidazole, 10mM ATP, 50mM Tris-Cl, pH 8.0). The His-tagged connector was eluted by His Elution Buffer (15% glycerol, 0.5 M NaCl, 0.5M Imidazole, 50mM ATP, 50mM Na₂HPO₄ –Tris-Cl, pH 8.0).

[00149] Insertion of the Connector Protein into Preformed Lipid Bilayers

[00150] Procedures have been recently developed^{100-102,104,121} for the incorporation of connector channels into lipid bilayers. Planar bilayer lipid membranes (BLMs) were created in a BCH-1A horizontal BLM cell (Eastern Scientific, LLC). A Teflon® partition with a 0.2 mm aperture was placed in the apparatus to separate the top and bottom compartments of the BLM cell. A planar lipid bilayer of DPhPC was formed by painting the aperture with 0.5 µl of 3% (w/v) DPhPC in *n*-decane. A conducting buffer (1 M NaCl, 10 mM Tris-HCl, pH 7.8) was added to both the top and bottom compartments of the BLM cell, and a Ag/AgCl electrode was placed in the buffer of each compartment. The electrode in the bottom compartment was connected to the headstage of an Axopatch 200B amplifier (Axon Instruments, Inc.), and the electrode in the top compartment was grounded.

[00151] When a planar BLM was successfully formed in the aperture of the partition, separating the two compartments filled with buffer, the BLM acted as an insulator. Intact planar BLM exhibited no detectable electrical current between the two electrodes. GUVs (0.5 – 2.0 µl) containing connector channels were added to the top compartment. Membrane fusion between the liposomes and the BLM happened immediately after applying GUVs containing the connector protein, and it was found that a current passing through the connector channel could be recorded.

[00152] Electrophysiological Measurements The headstage and Axopatch 200B patch clamp amplifier were connected to a DigiData 1440 analog-digital converter (Axon Instruments, Inc.) to monitor and record electrochemical currents through BLMs^{17,18,38}. The current recordings were low-pass filtered at a frequency of 1 kHz. The sampling frequency was 2 kHz in all experiments, unless otherwise specified. The data were recorded and stored on a PC with pClamp 9.1 software (Axon Instruments, Inc.), and analyzed with the Clampfit module of pClamp 9.1 and OriginPro 8.1 (OriginLab Corporation).

[00153] Electrical potentials were applied across BLMs in experiments with two different modes, holding potential mode and ramping potential mode with the pClamp 9.1 software. The ramping potential mode was used in the experiments to examine connector channel stability and DNA/RNA translocation orientation. In the holding potential mode, a constant voltage (either +75 mV or -75 mV) was applied across the BLM. In the ramping

potential mode, a voltage which constantly increased over time, from either -100 mV or +100 mV to -200 mV or +200 mV, was applied across the BLM.

[00154] Purification of the DNA/RNA Used in the Experiment

[00155] A synthetic 20-nt DNA fragment was purified by mixing a 100 μ M stock of the oligonucleotide with an equal volume of 2x TBE buffer (178 mM Tris base, 178 mM boric acid, 2 mM EDTA, pH 8.3). The oligonucleotide/TBE mixture was loaded onto an 8% (w/v) polyacrylamide gel containing 8 M urea-TBE. The DNA was electrophoresed through the gel at 120 V for ~1 h. After electrophoresis, the single-stranded DNA band was visualized as a shadow on a TLC plate with a hand-held UV light and then cut out of the gel. The gel containing the ssDNA was minced and the ssDNA was eluted from the gel fragments by incubating them in a small volume of elution buffer (0.5 M ammonium acetate) at 37 °C for 4 h. The gel fragments were pelleted by centrifugation and the supernatant was removed. The ssDNA was precipitated from the supernatant by adding 1/10 volume of 3 M sodium acetate and 2.5 volumes of ethanol and storing the solution overnight at -20° C. The ssDNA was again pelleted by centrifugation, and excess salt was removed by rinsing the ssDNA pellet with 70% ethanol. The purified ssDNA was dissolved and stored in TMS buffer (100 mM NaCl, 10 mM MgCl₂, 50 mM Tris-HCl, pH 8.0). A synthetic 20-nt ssRNA fragment was purified in a similar manner.

[00156] Translocation Experiments of DNA and RNA

[00157] Liposomes containing connector channels were added to the top compartment of the BLM cell. Soon after, highly conductive connector channel(s) were inserted into the planar BLM and a stepwise increase in current was observed. For translocation experiments, the purified ssDNA or ssRNA was mixed with conducting buffer and added to the top compartment in the BLM cell. The translocation of ssDNA or ssRNA was observed as transient blockades of the electrochemical current.

[00158] The SPP1 connector protein used in this example comprises the sequence of SEQ ID NO: 4.

[00159] **Example 4:** One-way Traffic of SPP1 Viral Motor Channel for peptide Translocation.

[00160] A variety of bacterial viruses exist in diverse form in natural environment, however, many of them share lots of similarities from shape, structure, and life cycle, to packaging and infection mechanism. Recently, it was discovered a common mechanism of dsDNA translocation motors using one-way revolution mechanism, exists many bacteria and

virus, including phi29, SPP1, T4, T7 and others. Among the complex system of many different phages, viral connector protein is one of the key components with turbine-like shape in phage genome packaging and ejection, shared by many bacteriophages and other viral system. Previously we have successfully inserted phi29 connector channel into lipid bilayer and found the one way traffic of DNA. Here we reported that inserted reengineered bacteriophage SPP1 connector channel also exercised a one-way traffic property for peptide translocation, as demonstrated by holding and ramping, voltage, electrode polarity switching, and quantification of peptide translocation frequency. These results indicate that most of the DNA packaging motors probably utilize the same one way traffic mechanism during DNA packing process. In addition, the observed single direction peptide transportation provide a novel system with a natural valve to control dsDNA loading and gene delivery in liposomes or peptide sequencing without using of gating control found in other biological channels.

[00161] Introduction.

[00162] A variety of bacterial viruses exist in diverse form in natural environment, however, many of them share lots of similarities from shape, structure, and life cycle, to packaging and infection mechanism. Recently, it was discovered a common mechanism of dsDNA translocation motors using one-way revolution mechanism, exists many bacteria and virus, including phi29, SPP1, T4, T7 and others¹³³⁻¹³⁵. An unusual feature of many linear dsDNA viruses is the packaging of their DNA into a preformed capsid, which is entropically unfavorable process and accomplished by an ATP-driven motor. The packaging motor of SPP1, composed gp1, gp2, and gp6, powers encapsulation of the 45.9 kb genome.¹²⁸ The ingenious and dedicate design of viral DNA packaging motors and the novel mechanism of action has provoked a wide range of interest among scientists in many different areas. Various portal proteins share little homology in primary sequence, and varies in size¹²⁹⁻¹³¹, although, they exhibit considerable morphological similarity¹³². Portal protein of SPP1 and phi29 both have similar secondary structure and 3-D structure. Explicit engineering of the SPP1 connector is possible because of its accessible crystal structure¹³⁵⁻¹³⁷. SPP1 portal protein exists in 13-subunit form and the pseudoatomic structure with 12-subunits. The overall diameter of 13-subunit form is ~1.65 nm, and the height is ~1.10nm and the narrow part is 2.77 nm in 13 subunit form and 1.81 in 12 subunit form.^{136,137}.

[00163] In vitro, the biomimetic motor can be inserted into synthetic nanodevices^{138,139}. In vivo, the artificial nanomotors can be also applied for loading drugs, delivering DNA/RNA, pumpng ions, transporting cargoes, or driving the motion of components in the heart, eye or other sensing organs in the body. We have previously

reengineered the connector protein and inserted the connector into lipid bilayers¹⁹²⁰. The translocations of ions and peptide through the channel have been characterized to demonstrate the power of the SPP1 connector for use in single DNA sensing and fingerprinting. In this study, we report that the reengineered SPP1 motor channel exercises a one-way peptide traffic mechanism, which may related with the packaging mechanism.

[00164] Recently, chemically modified synthetic pores¹⁴¹⁻¹⁴⁴ and some reengineered protein channels¹⁴⁵⁻¹⁴⁷ have been found to have ion rectifying capabilities by allowing ions to flow in one direction. For example, a mutant 7R- α -haemolysin has been discovered to only allow ion transport under positive voltages even under high ion concentrations¹⁴⁵. Chemically modified synthetic pores can be made to exhibit electronic diode-like properties by a different mechanism.^{148,149}

[00165] Previously we have successfully inserted phi29 connector channel into lipid bilayer and found the one way traffic of DNA. Here we reported that inserted reengineered bacteriophage SPP1 connector channel also exercised a one-way traffic property for peptide translocation, as demonstrated by holding and ramping, voltage, electrode polarity switching, and quantification of peptide translocation frequency. These results indicate that most of the DNA packaging motors probably utilizes the same one way traffic mechanism during DNA packing process. In addition, the observed single direction peptide transportation provide a novel system with a natural valve to control dsDNA loading and gene delivery in liposomes or peptide sequencing without using of gating control found in other biological channels.

[00166] Results.

[00167] Characterization of reengineered SPP1 connector channel

[00168] As revealed in crystal structure, SPP1 connector channel share similar structure with phi29 connector channel. To purify SPP1 connector channel, 6X his tag was inserted downstream of a six-glycine linker which was introduced to provide end flexibility. After purification to homogeneity, the molecular weight of single subunit of SPP1 is 56 kDa, which is much higher than 35 KDa of phi29 connector as revealed in SDS PAGE gel, in agreement with theoretical molecular weight. The purified reengineered SPP1 connector channel was inserted into a planar lipid bilayer membrane (BLM) via two steps. First, the connector channel was incorporated into liposome and then connector complex was fused into lipid membrane. The discrete stepwise increase of the current was observed after the connectors inserted into the membrane. Each step size was about 200 pA under 1M KCL 5 mM HEPES, pH 8 under 50mV, and the conductance is uniform and did not display voltage gating under the applied voltage.

[00169] One-way traffic of peptide was probed by applying a constant holding voltage and by switching the voltage polarity to a membrane. There was no peptide translocation under either positive or negative voltage when no peptide is present in both chambers in a single channel study or multiple channel study. Peptide was pre-mixed in both chambers under a constant voltage. Under single channel, switching of voltage polarity showed that the connector channel allowed peptide a unidirectional translocation. Since the connector channel is incorporated into the lipid bilayer via the fusion of the liposome/connector complex, the direction of the connector cannot be controlled. The conformation of connector channel could be either C-terminus facing the *cis*-chamber or N-terminus facing the *cis*-chamber.

[00170] The correspondence to voltage switching was dependent upon the orientation of the connector in the BLM. For example, when there was no peptide translocation under positive potential, switching the voltage to negative potential resulted in peptide translocation. Similarly, translocation events appeared under positive voltage but disappeared when switch to negative voltage indicating the channel only allow one way traffic of peptide. Peptide translocation under multiple channels was shown. Under positive voltage, the first inserted channel has not shown any translocation events, however, when the second channel inserted into membrane, a burst of translocation events appeared immediately indicating the direction of two channel is opposite. Under positive voltage, the first channel has shown uniform blockade events. As expected, after switching the voltage to negative, there is no blockade events shown up, however, when the second channel inserted the blockade events suddenly appeared in a uniform manner. To further verify the one-way traffic, peptide was added to one side of chamber after multiple channel insertion. If is found that the blockade events was only appeared in one side. When the peptide added to *cis*-chamber, the translocation events only appeared under negative voltage since the peptide is positive charged. Similar phenomenon was also observed when peptide was added to the *trans*-chamber. In addition, after added different ratio of peptide in the two chambers, the blockade events rate is also different corresponding to the ratio of peptide in both chambers.

[00171] One-way traffic of peptide was probed by quantification of peptide translocation frequency in the presence of multiple channels. If connector channels allow peptide passing through only in one direction, the translocation frequency should be affected by a different conformation of connector orientation when multiple channels were inserted into the bilayer, regardless of the peptide concentration. Experiments were performed under a constant holding potential of -50mV and in the presence of varying concentrations of peptide.

The current traces of multiple connector insertions revealed the change in the frequency of peptide translocation events. When one additional connector was incorporated with an opposite orientation that did not allow peptide to pass, the frequency of DNA translocation did not change. The insertion of first channel allows the translocation of peptides from cis-chamber to trans-chamber under negative voltage. The additional insertion did not change the frequency due to the direction of second one is opposite to the first one which did not allow peptide translocation through that side. When one additional connector was incorporated with the same orientation to the previous one, the frequency of DNA translocation double depending if the direction allow peptide translocation. The second channel does not increase the translocation frequency because of the orientation is opposite to the first one, which allow peptide translocation. When the third channel inserted, the translocation frequency doubled because the direction is the same as the first connector channel.

[00172] One-way traffic of peptide was probed by applying a ramping potential to a membrane. Given that the connector channel is inserted into the BLM via the fusion of the liposome/connector complex with the BLM, the orientation of the connector channel is random. The application of a ramping potential to the BLM containing a single connector channel revealed a unidirectional translocation of peptide. A ramping voltage from -50 mV to +50 mV was applied to obtain the I-V curves.

[00173] Discussion.

[00174] Recently, Guo and Lee proposed a pushing or injection model¹⁵⁰. In this model, the connector remains static; DNA translocation is induced by a DNA packaging enzyme or terminase, which pushes a certain length of DNA into the procapsid, and then shifts to bind to a far distal region of the DNA and inserts an additional section. This model does not exclude the *socket wrench* rotating motion by the enzyme. The studies reported here provide direct evidence to prove the model¹⁵⁰.

[00175] It is very interesting to observe the counter-chirality between the left-handed spp1 connector channel and the right-handed dsDNA. The spp1 connector channel is composed of 12 copies of the protein gp10 with each existing as three-helices to encircle the channel, and exhibiting a left-handed configuration. However, the regular viral dsDNA genome is the normal B-form DNA exhibiting a right-handed configuration. As far as the energy is concerned, it is expected that such an opposite configuration would favor the ejection of the dsDNA during infection and would not favor the packaging during assembly. However, our findings revealed a contrasting phenomenon: under an external electrical force, the channel favored DNA entry but blocked DNA exit, suggesting a very intriguing structure

and motion mechanism for the motor. In this situation, the channel wall and the dsDNA constitute two counter-parallel helices. It is possible that the sequential attraction between the negative charge of the phosphate backbone and the positive charge of the channel wall, which contains two 12-lysine rings (aa #200, 209) separated by 2 nm (and possibly two more rings at locations K234, K235)¹²⁹, will produce a step wise propelling force for DNA translocation. It has been reported that the conformation of the spp1 connector is substantially changed after DNA packaging^{151,152}. Such significant rearrangements of the connector after DNA packaging, a similar feature reported in other phage systems as well^{136,137,153}, might change the channel configuration to reversely favor the DNA exit during infection.

[00176] MATERIALS AND METHODS

[00177] Materials

[00178] The phospholipid, 1,2-diphytanoyl-sn glycerol-3-phosphocholine (DPhPC) (Avanti Polar Lipids, Alabaster, AL), Nickel-NTA nanogold (1.8 nm; Nanoprobes), n-Decane (Fisher), chloroform (TEDIA) were used as instructed by the vendor. All other reagents were from Sigma, if not specified. The construction and purification of reengineered spp1 C-terminal tagged connectors have been reported^{140,154}.

[00179] Electrophysiological measurements

[00180] A bilayer was formed on a thin Teflon film partition (aperture 200 µm in diameter) which separates a BLM chamber into a *cis*- and *trans*-chamber (compartment). The *cis*-chamber refers to the grounded compartment to which the connector reconstituted liposome was added. Connector channel insertion into the bilayer has been described previously using vesicle fusion¹⁹. Briefly, connector reconstituted liposomes were prepared using a dehydration-rehydration method and further extruded to form unilamellar liposomes. The reconstituted liposomes were further diluted by 10-fold using the conducting buffer before applying to BLM chambers. The final concentration of added protein was 5-50 µg/mL.

[00181] For electrophysiological measurements, both compartments in the BLM chamber were filled with a conducting buffer (1 M NaCl/5 mM Tris, pH 7.8, if not specified). A pair of Ag/AgCl electrodes connected directly to both the compartments was used to measure the current traces across the bilayer lipid membrane. The current trace was recorded using an Axopatch 200B patch clamp amplifier coupled with the Axon DigiData 1322A analog-digital converter (Axon Instruments) or the BLM workstation (Warner Instruments). All voltages reported were those of the *trans*-compartment. Data was low band-pass filtered

at a frequency of 5 kHz or 1 KHz and acquired at a sampling frequency of 10 KHz. The PClamp 9.1 software (Axon Instruments) was used to collect the data, and the software Origin Pro 8.0 was used for data analysis.

[00182] The SPP1 connector protein used in this example comprises the sequence of SEQ ID NO: 4.

[00183] Example 5: Three-steps of Channel Conformational Twisting Related to Paused and Quantized Packaging during Directional Genome Translocation of Bacterial Virus SPP1.

[00184] The channel of the DNA-packaging motor of double-stranded DNA virus allows viral genomic to enter the protein procapsid shell during viral maturation and to exit during host infection. We recently demonstrated that DNA packaging motor of bacterial virus phi29 exercises a one-way traffic property, and uses a revolution mechanism without rotation for dsDNA packaging using a left-handed connector channel to gear the right-handed DNA. This raises a question of how dsDNA is ejected during infection using this one way valve. Previously, we reported the finding of three step gating in the channel of phi29 DNA packaging motor, and proposed a conformational changes of motor channel during DNA packaging process to prepare for the reversal of dsDNA. To determine whether the finding and proposed mechanism in phi29DNA packaging motor is universal, we cloned the gene gp-6 coding for SPP1 connector and, studied the conformational change of SPP1 connector inserted into the lipid bilayer. We found reengineered SPP1 connector also exhibited conformational changes during DNA packaging process and three step gating. We investigated and linked the gating and conformational change phenomenal to the quantized packaging studies of DNA in phi29 and T3 reported previously. It was found that the three step gating coincide with the three major steps of quantized DNA packaging intermediates. These finding lead to the conclusion that three step gating, is due to three step of motor conformational change that was revealed in three major step of quanlized packaging of DNA intermediate. This supports the speculation that the one way inbound channel were transformed into an outbound channel during the DNA packaging process and this finding will lead to the slight twisting of the dsDNA by a motor with revolution mechanism without rotation.

[00185] 1. INTRODUCTION

[00186] Tailed bacterial viruses are widely distributed in nature and they shared many similarities, from structure, shape, to life cycle, packaging and infection mechanism. Among

the complex system of many different phages, viral connector protein is one of the key components with turbine-like shape, shared by many bacteriophages and other viral system. It was inserted into the prohead as major part of portal complex and responsible for genome packaging and ejection. Linear dsDNA viruses package their genome into a preformed protein shell referred as procapsid¹⁵⁵. This DNA encapsulation process is completed by a nanomotor using ATP as energy¹⁵⁶⁻¹⁶⁰. During this process, connector acts as a docking point for ATPase which powers the DNA genome translocation. SPP1 is a virulent dsDNA phage which infects *bacillus subtilis* bacteriophages and the connector consists of portal protein gp6, and two head completion protein gp15 and gp16.¹⁶¹⁻¹⁶² The molecular motor of SPP1, composed of gp1, gp2 and gp6, powers encapsulation of the 45.9 kbp genome.¹⁶³ The ingenious design of viral DNA packaging motors and the novel mechanism of action has irritated a wide range of interest among scientists in virology, molecular biology, structure biology, nanotechnology and other areas. Although various connectors share little homology in primary sequence and varies in size¹⁶⁴⁻¹⁶⁶, they exhibit significant morphological similarity¹⁶⁷. Portal protein of SPP1 and phi29 share similar secondary structure and 3-D structure. Our recent research also found that they even share similar role in packaging genome¹⁶⁸⁻¹⁷⁰. The structure of the SPP1 connector has been previously solved by X-ray crystallography,^{171,172} showing SPP1 portal protein have 13-subunit form and the pseudoatomic structure with 12-subunits, The overall diameter of 13-subunit form is ~1.65 nm, and the height is ~1.10nm. The narrow part is 2.77 nm in 13 subunit form and 1.81 in 12 subunit form. Gating is very common in various ion channels or protein pores¹⁷³ and plays a critical role in regulating ion transportation through a membrane and DNA translocation direction. By the nature of gating, ion channels could be classified¹⁷⁴ as ligand-gated¹⁷⁵, stretch-gated¹⁷⁶, voltage-gated¹⁷⁷, or other gating¹⁷⁴. Voltage-gated ion channels are sensitive to potential change near the channel, whereas the ligand-gated ion channels are activated by the binding of a chemical messenger (i.e., a ligand).

[00187] Previous studies, the connector channel from bacteriophages phi29 has been inserted into lipid bilayers to serve as a membrane-embedded nanopore with robust properties.^{178,179} A one-way traffic property for dsDNA translocation with a valve mechanism for DNA packaging has also been observed.¹⁸⁰ The reengineered connector channel is capable to discriminate single-strand DNA or RNA from double strand. By selectively engineering a probe on the connector, single chemicals and antibody can be identified based on their distinct fingerprints.¹⁸¹

[00188] Based on previous study, we further want to verify whether this phenomenon is universal. Here, we first time expressed reengineered SPP1 connector and found this connector also exercises 3-step gating which is similar to phi29 connector. We also further investigate the effect of voltage polarity, pH, ion concentration, and type on the gating. This finding may indicate the phenomenon of connector conformation change is common in most of the bacteriophages and share similar role in genome packaging and ejection.

[00189] 2. MATERIALS AND METHODS

[00190] 2.1 Materials

[00191] The phospholipid, 1,2-diphytanoyl-sn glycerol-3-phosphocholine (DPhPC) (Avanti Polar Lipids, Alabaster, AL), , n-Decane (Fisher), chloroform (TEDIA) were used as instructed by the vendor. All other reagents were from Sigma, if not specified.

[00192] 2.2 Expression and purification of reengineered SPP1 connector

[00193] Gene gp6 encoding SPP1 connector on the C-terminus was syntheses by Genescript company and then cloned into PET3 vector between NheI and BamHI. A His-tag was inserted into the C-terminal for purification. Then the plasmid was transformed into BL 21 (DE3) for expression and purification. The transformed bacteria were cultured in 10 ml Luria-Bertani medium overnight at 37 °C. Then the cultured bacteria were transferred to 1 L of fresh LB medium. 0.5mM IPTG was added to the medium to induce protein expression when OD₆₀₀ reached 0.5~0.6. The bacteria were collected by centrifugation after 3 hours continue culture. Bacteria was lysed by passing through French press. Cells were resuspended with His Binding Buffer (15% glycerol, 0.5 M NaCl, 5mM Imidazole, 10mM ATP, 50mM Tris-Cl, pH 8.0), and the cleared lysate was loaded onto a His•Bind® Resin Column and washed with His Washing Buffer (15% glycerol, 0.5 M NaCl, 50mM Imidazole, 10mM ATP, 50mM Tris-Cl, pH 8.0). The His-tagged connector was eluted by His Elution Buffer (15% glycerol, 0.5 M NaCl, 0.5M Imidazole, 50mM ATP, 50mM Tris-Cl, pH 8.0).

[00194] 2.3 Electrophysiological measurements

[00195] A bilayer was formed on a thin Teflon film partition with aperture 200 µm in diameter. This partition separates the bilayer lipid membrane (BLM) chamber into two compartments: a cis- and trans-chamber. The connector reconstituted liposome was added to the cis-chamber referred as grounded compartment. Phi29 connector channel insertion into the lipid bilayer has been reported previously using vesicle fusion¹⁷⁸. Here we used the same procedure for reengineered SPP1 connector. Briefly, connector reconstituted liposomes were prepared using a dehydration-rehydration method and further extruded to form unilamellar

liposomes. The reconstituted liposomes were further diluted by 10-fold using the conducting buffer before applying to BLM chambers. For electrophysiological measurements, both compartments of BLM chamber were filled with a conducting buffer. A pair of Ag/AgCl electrodes inserted to both compartments was used to measure the current traces across the bilayer lipid membrane. The current trace was recorded using an Axopatch 200B patch clamp amplifier coupled with the BLM workstation (Warner Instruments). or the Axon DigiData 1322A analog-digital converter (Axon Instruments) All voltages reported were those of the trans-compartment from 100mV, 75mV, 50mV. Data was low band-pass filtered at a frequency of 5 kHz or 1 KHz and acquired at a sampling frequency of 10 KHz. The PClamp 9.1 software (Axon Instruments) was used to collect the data, and the software Origin Pro 8.0 was used for data analysis.

[00196] 3. RESULTS

[00197] 3.1 The lipid membrane-embedded C-his SPP1 connector display uniform conductance.

[00198] To facilitate connector purification, a C-terminal His tag was inserted downstream of the gp-6 gene. The reengineered connectors were purified to homogeneity and run on SDS-PAGE gel with C-His phi29 connector as controls. The molecular weights of C-His tagged SPP1 and phi29 connector channel are 56 kDa and 35 kDa, respectively. After purification, the connector channels was first incorporated into liposome and then fused into lipid bilayer. The connector channel embedded into the lipid bilayer was stable and displayed a uniform conductance. The conductance of reengineered SPP1 connector channel is 2.06 nS under 1M NaCL 5mM HEPES pH 8.0.

[00199] 3.2 C-his SPP1 connector exercise discrete and uniform three step conformational change

[00200] At a low constant holding potential (± 50 mV), the channel conductance is stable and there is no gating observed under both voltage in a long time recording. However, when a higher potential voltage is applied, a 3-step channel size reduction is often observed immediately thereafter, which is very similar to phi29 connector channel and other membrane gating behaviors. In the given experiment conduction (1 M KCl with 5 mM HEPES, pH 8), to induce gating, the positive voltage applied across the membrane need to be over 100 mV. Under this voltage, single channel current jump is around 380 pA, and each reduction step size was 120 pA accounting for 32% of the entire channel. Whereas to induce gating under negative potential, the threshold value just need to be over 75 mV and each

reduction step size was about 90 pA still accounting for the 32% of the open channel. This 3-step channel gating can be induced by the high voltage regardless the buffer conditions. It is also interesting to note that after the 3-step reduction, there was still a residual current of 20 pA opening, which is about~5.3% of one channel. Similar results were observed at phi29 connector channel. However, this closed channel can open immediately if an opposite potential is applied or a lower voltage applied.

[00201] 3.3 Polarity dependence gating of C-his SPP1 gating

[00202] When multiple channels inserted into lipid bilayer and reach a steady stage, the polarity was switched from positive potential to negative potential and it is interestingly found that the gating showed polarity dependence. A trace current of multiple channels at ± 50 , ± 75 and ± 100 mV. Under 50 mV, 1M NaCl, 5mM Hepes pH 6, multiple channels have been continually incorporated into lipid bilayer until reaching a steady stage without any gating. When varies the voltage to +75 mV under the same buffer condition, there was still no gating observed for several minutes recording. However, when switching to -75 mV, gating occurred immediately and continuously until the channels closing completely. Each reduction size is in agreement of the data in single channel study. When applied 100 mV voltage across the membrane under the same buffer condition, gating occurred immediately when switch to -100 mV. The gating step is still one third of one channel current jump which is the same as observed under single channel study. The data clearly shows that gating of C-his SPP1 channel is polarity dependence and gating occurs much often under negative voltage than that under positive voltage.

[00203] It was also observed that the gating of the connector channel was reversible. When a transmembrane voltage of 0 pA applied for 10 to 30 seconds, the gated channel was observed to re-open. After the channel shutting down was triggered by higher voltage (-100 mV and -150 mV), a lower voltage was applied to allow the connector protein to recover. It was found that the conductance of the channels restored to their original state within 10 seconds.

[00204] 3.3 pH effect on gating property

[00205] In order to further characterize the factors that affect the gating property of C-his SPP1 channel, a series of pH has been tested. When pH increase to 10 and 12 under 1M NaCl 5mM Hepes, 75mV applied across the membrane, similar gating trend has been observed as it under pH 6 that gating occurred immediately when switching from positive

voltage to negative voltage. When applied scanning voltage from -100 to 100 mV, similar phenomenon has been observed. However, the noise level change.

[00206] When pH decrease from 6 to below 3, incorporation channels under 75 mV is very difficult. So to increase the incorporation chance, the voltage has been decreased to ± 25 mV. Under this voltage, it's interesting observing the voltage polarity dependence become opposite that the channel become very unstable under positive voltage but when switch to negative voltage, the channels became stable. This phenomenon has also been confirmed under scanning voltage.

[00207] 3.4 Salt concentration effect on gating property

[00208] Salt concentration effect on the gating has also been investigated. In this study, we have investigated the gating property under 1 M, 0.5M, 0.2 M and 50 mM NaCl. As the concentration decrease to 0.5 M, the channels are stable under positive voltage but gating occurs immediately under -75 mV. However, when the concentration decreases to 0.2M, the channels are stable under both positive and negative 75 mV voltage. Similar phenomenon has also been observed under 50 mM NaCl.

[00209] 4. DISCUSSION

[00210] Viral DNA packaging has been extensively studied in many viral systems, but the actual mechanism remains elusive. Many models have been proposed to interpret the mechanism of packaging process during the last several decades, including: 1) Ratchet mechanism¹⁸² 2) Gyrase-driven supercoiled and relaxation^{159,183,184}; 3) Brownian motion¹⁸⁵; 4) Force of osmotic pressure¹⁸⁶ 5) Five-fold/six-fold mismatch connector rotating thread¹⁸⁷; 6) Supercoiled DNA wrapping¹⁸⁸; 7) Electro-dipole within central channel¹⁶⁴; 8) Sequential action of motor components^{189,190} and, 9) Connector contraction hypothesis¹⁹¹. All these models are intriguing, but none of them have been supported by conclusive experimental data; or in other cases, validated in one viral system but disproved in another. The five-fold/six-fold mismatch connector rotating thread model¹⁸⁷ has been popular for several decades, because this model could introduce a new mechanical motor prototype. Numerous laboratories, including our own, have persevered to search, interpret, even design a five-fold ring to fit findings to this fascinating model^{190,192-194}. Unfortunately, recent studies in many viral DNA packaging motors reveal that the stoichiometry of motor components is an even number rather than an odd number¹⁹⁵⁻¹⁹⁸. In 1998, Guo and his colleagues first proposed a DNA packaging mechanism which is similar to the process of the translocation of dsDNA in hexameric AAA+ ATPase during DNA replication and repair. The finding of even-number

subunits agree with the mechanism of many other well-studied DNA tracking motors¹⁹⁹⁻²⁰² and the AAA+ ATPase family. More recent publications^{194,203-205} also support our findings^{197,198,206} that hexameric structure is built by pRNA dimer²⁰⁶. X-ray crystallography study also confirmed that pRNA forms a dimer in solution which forms a tetramer without the procapsid^{194,206}, supporting the theory that the dimer is the building unit of the hexamer and the sequential action in hexamer assembly is $2 \rightarrow 4 \rightarrow 6$. It would be desirable if the mechanism of the sequential action among the channel subunits is elucidated in the future. Recently, Guo and his colleagues discovered a third type of biomotor with revolution mechanism without rotation in bacteriophages phi29. Later, they found this mechanism is very common in DNA translocation motors of viruses and bacteria, including HK97, SPP1, P22, T4, T7, cellular DNA translocase FtsK. By employing revolution mechanism the motor can translocate dsDNA free of coiling and torque which has solved many puzzles and debates occurred throughout the history of viral DNA packaging motor studies.

[00211] Revolution involves the left-handed connector channel to facilitate the entry of the dsDNA during packaging^{170,180}. It would be reasonable to believe that, after DNA packaging is completed, a conformational changes will occur in order to convert the left-handed connector into the right-handed configuration to facilitate DNA exit to infect host cells. Indeed, three steps of conformational changes of phi29 connector have been reported (Jia). In addition, previous studies have found that isolation of particles derived from DNA intermediates has shown three distinct peak, indicating that three step conformational change of connector may play an important role in the DNA packing process [Mary-Ann Bjornsti, 1983; Philip Serwer, 2014]. It is expected that a small right-handed twisting of the dsDNA will be observed since the dsDNA is aligned with the wall of the connector channel and the left- to right-handed conformational change. Biophysical studies to distinguish the very small angle right-handed twisting from the 3600/10.5 base pairs left-handed rotation would be very interesting.

[00212] In addition, many studies have shown the expansion of procapsid during DNA packaging of bacteriophages.²⁰⁷⁻²¹⁴ Connector is a crucial component in the regulation of procapsid shape and size²¹⁵. Therefore, it is reasonable to assume that procapsid expansion is related to the connector conformational change.

[00213] Our results support the proposal that these three steps of conformational change are responsible for packaging dsDNA and this is common in many viral DNA packaging motor. We demonstrated that the binding or contact of components to the C-terminal would result in the similar discrete steps of conformational change. Conductance

assay using mutant connector conjugated with a Strep-tag to its C-terminal and inculcated with anti-strep tag antibody or nanogold coated with streptavidin revealed a three discrete steps of channel change with blockade of about 32%. It is reasonable to believe that the translocation of DNA into the procapsid or the internal pressure of the fully packaged DNA within the procapsid might lead to the contact of DNA with the C-terminal flexible domain, inducing a conformation changes with two subsequent functions: 1) to prevent the dsDNA from exit; and 2) to prepare a new channel configuration to facilitate the injection of DNA during the host cell infection process. The proposed second step of action has also been evidence in the P22 system²¹³. To further support this, a connector mutant with the removal of a 25 amino acid segment at each C-terminus of the 12 subunits were constructed with a strep-tag conjugated to the C-terminus. With this mutant, the discrete steps of 32% conductance change were not observed in the presence of streptavidin. However, conductance assay of the mutant connector by electrical ramping revealed that three-discrete steps of conformational changes were also observed. The finding suggests that conformational change was a result of the transition of the entire connector structure, and the C-terminal only served as a trigger. This conclusion agrees with the finding in the connector structure of bacteriophage P22 12. A unique topology of the C-terminal domain was reported to be a ~200-Å-long α -helical barrel that inserts deeply into the virion and is highly conserved in the Podoviridae family. They proposed that the barrel domain would facilitates genome spooling into the interior of the procapsid during DNA packaging, and in analogy to a rifle barrel to increases the accuracy of DNA ejection during infection. During the course of our investigation, we also found that the batch of the polyclonal antibody made a difference, suggesting that not all binding to any location of the C-terminus would induce conformational changes and certain epitope at the C-terminus was important for triggering the conformation change.

[00214] The mechanism of ion channel gating has been extensively studied¹⁷⁴. The ligand-gated ion channels are regulated by a ligand and are usually very selective to one or more ions such as Na⁺, K⁺, Ca²⁺, or Cl⁻. Such receptors located at synapses convert the chemical signal of a presynaptically released neurotransmitter directly and very quickly into a postsynaptic electrical signal. Conformational changes of α helices in voltage-gated sodium channels and calcium channels have been proposed to explain the gating mechanism. From crystallographic structural studies of the SPP1 connector protein, it is possible to surmise that when a potential difference is introduced over the membrane, the associated electromagnetic field induces a conformational change in the protein channel. From the structural viewpoint

of a single chain in the connector protein, certain areas of the chain are flexible enough to induce a conformational change in response to an environmental stimuli. The interaction of dsDNA or SPP1 terminal protein gp3 induces a conformational change that distorts the shape of the channel proteins sufficiently such that the cavity, or channel, opens to admit ion influx or efflux across the membrane, down its electrochemical gradient. This conformational change leads to the opening or closing of the channel, which will help to control the packaging or release of the viral genome.

[00215] The controllable opening and closing of the connector protein has been achieved with different polarity, which also resembles the voltage-gated ion channels. However, they have different functions in the biological environment. The ion channels may have to open and close multiple times in their life cycle, whereas the SPP1 DNA connector protein might require less. The significantly different gating behavior of the internal flexible loop-cleaved connector suggests that the flexible loops may play a key role in the voltage gating. Results described above show that after the removal of these loops, both the occurrence and extent of gating reduced tremendously. These loops can induce conformational changes to adjust the channel size in response to an applied potential. Further studies are necessary to investigate if other domains of the connector protein also contribute to its gating and conformational changes.

[00216] 5. CONCLUSIONS

[00217] Three step gating and a conformational changes of motor channel in the channel of SPP1 connector has been found in this study, a phenomenal identical to the finding in phi29. We propose the gating and conformational change phenomenal may link to the quantized packaging studies of DNA in phi29 and T3 reported previously. It was found that the three step gating coincide with the three major steps of quantized DNA packaging intermediates. These finding lead to the conclusion that three step gating, is due to three step of motor conformational change resulting in three major step of quanlized packaging or DNA intermediate, and support the speculation that the one way inbound channel were transformed into an outbound channel during the DNA packaging process and this finding will lead to the slight twisting of the dsDNA by a motor with revolution mechanism without rotation.

[00218] The SPP1 connector protein used in this example comprises the sequence of SEQ ID NO: 4.

[00219] All publications, patents, and patent applications mentioned in this specification are herein incorporated by reference to the same extent as if each individual publication, patent, or patent application was specifically and individually indicated to be incorporated by reference, including the references set forth in the following list:

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[00220] It will be understood that various details of the presently disclosed subject matter can be changed without departing from the scope of the subject matter disclosed herein. Furthermore, the foregoing description is for the purpose of illustration only, and not for the purpose of limitation.

SEQUENCE LISTING**SEQ ID NO:1**

Wild type SPP1 connector channel:

DNA sequence:

ATGGCTGATATCTACCCACTAGGGAAAACACACACAGAGGAACTGAATGAAATCATCGTAGAGAGCGC
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SEQ ID NO:2

Protein sequence:

MADIYPLGKTHTEELNEIIVESAKEIAEPDTTMIQKLIDEHNPEPLLKGVRYMCENDIEKKRRTYYD
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DEGGDDLEEDDPNAGAAESGGAGQVS

SEQ ID NO:3

1. Mutant 1 with added Gly-Gly-His-His-His-His-His on C-terminal
(Gray labels indicate distinctions relative to wild type)

DNA sequence:

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SEQ ID NO:4

Protein sequence:

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VYYYEKIDGVYQMDYSYGENNPRPHMTKGGQAIGWGRVPIIPFKNNEEMVSDLKFYKDLIDNYDSITS
STMDSFSDFQQIVYVLKNYDGENPKEFTANLRYHSVIKVS DGGVDTLRAEIPVDSAAKELERIQDEL
YKSAQAVDNSPETIGGGATGPALENLYALLDLKANMAERKIRAGLRLFFWFFAEYLRNTGKGDFNPK
ELTMTFTRTRI QNDSEIVQSLVQGV TGGIMSKETAVARNPFVQDPEEELARIEEEMNQYAEMQGNLLD
DEGGDDLEEDDPNAGAAESGGAGQVS

SEQ ID NO:5

2. Mutant 2 loop deletion.

DNA sequence:

ATGGCTGATATCTACCCACTAGGGAAAACACACACAGAGGAACTGAATGAAATCATCGTAGAGAGCGC
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SEQ ID NO:6

Protein sequence:

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SEQ ID NO:7

3. Mutant 3 – grey labeled is position of mutation(s)

E461G/E462G/E463G

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SEQ ID NO:8

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4. Mutant 4 - grey labeled is position of mutation(s) **D314N/D318N**

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SEQ ID NO:10

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QGNLLDDEGGDDLEEDDPNAGAAESGGAGQVS

CLAIMS

What is claimed is :

1. A nucleic acid molecule encoding an SPP1 connector polypeptide variant, wherein the polypeptide comprises an amino acid sequence having at least 95% sequence identity to the sequence of SEQ ID NO:4, SEQ ID NO:6, SEQ ID NO:8, or SEQ ID NO:10.

2. The nucleic acid molecule of claim 1, wherein the polypeptide comprises an amino acid sequence having at least 99% sequence identity to the sequence of in SEQ ID NO:4, SEQ ID NO:6, SEQ ID NO:8, or SEQ ID NO:10.

3. The nucleic acid molecule of claim 1, wherein the polypeptide comprises the amino acid sequence of SEQ ID NO:4, SEQ ID NO:6, SEQ ID NO:8, or SEQ ID NO:10.

4. The nucleic acid molecule of claim 1, having the sequence of SEQ ID NO:3, SEQ ID NO:5, SEQ ID NO:7, or SEQ ID NO:9.

5. A SPP1 connector polypeptide variant, wherein the polypeptide comprises an amino acid sequence having at least 95% sequence identity to the sequence of SEQ ID NO:4, SEQ ID NO:6, SEQ ID NO:8, or SEQ ID NO:10.

6. The polypeptide of claim 5, wherein the polypeptide comprises an amino acid sequence having at least 99% sequence identity to the sequence of SEQ ID NO:4, SEQ ID NO:6, SEQ ID NO:8, or SEQ ID NO:10.

7. The polypeptide of claim 5, wherein the polypeptide comprises the amino acid sequence of SEQ ID NO:4, SEQ ID NO:6, SEQ ID NO:8, or SEQ ID NO:10.

8. A SPP1 connector polypeptide variant, wherein the polypeptide comprises an amino acid sequence encoded by a nucleic acid molecule having the sequence of SEQ ID NO:3, SEQ ID NO:5, SEQ ID NO:7, or SEQ ID NO:9.

9. A conductive channel-containing membrane, comprising:

(a) a membrane layer; and
(b) a SPP1 connector polypeptide variant that is incorporated into the membrane layer to form an aperture through which conductance can occur when an electrical potential is applied across the membrane, wherein the SPP1 connector polypeptide variant is selected from:

(i) a polypeptide comprising an amino acid sequence having at least 95% sequence identity to the sequence of SEQ ID NO:4, SEQ ID NO:6, SEQ ID NO:8, or SEQ ID NO:10;

(ii) a polypeptide comprising an amino acid sequence having at least 99% sequence identity to the sequence of SEQ ID NO:4, SEQ ID NO:6, SEQ ID NO:8, or SEQ ID NO:10;

(iii) a polypeptide comprising the amino acid sequence of SEQ ID NO:4, SEQ ID NO:6, SEQ ID NO:8, or SEQ ID NO:10; and

(iv) a polypeptide comprising an amino acid sequence encoded by a nucleic acid molecule having the sequence of SEQ ID NO:3, SEQ ID NO:5, SEQ ID NO:7, or SEQ ID NO:9.

10. The conductive channel-containing membrane of claim 9, wherein SPP1 connector polypeptide variant comprises a detectable label.

11. The conductive channel-containing membrane of claim 10, wherein the detectable label is selected from the group consisting of a colorimetric indicator, a GCMS tag compound, a fluorescent indicator, a luminescent indicator, a phosphorescent indicator, a radiometric indicator, a dye, an enzyme, a substrate of an enzyme, an energy transfer molecule, a quantum dot, a metal particle and an affinity label.

12. The conductive channel-containing membrane of claim 9, wherein the membrane layer comprises a lipid layer.

13. The conductive channel-containing membrane of claim 12, wherein the lipid layer comprises amphipathic lipids.

14. The conductive channel-containing membrane of claim 13, wherein the amphipathic lipids comprise phospholipids and the lipid layer comprises a lipid bilayer.

15. The conductive channel-containing membrane of claim 12, wherein the lipid layer is selected from the group consisting of a planar membrane layer and a liposome.

16. The conductive channel-containing membrane of claim 15, wherein the liposome is selected from the group consisting of a multilamellar liposome and a unilamellar liposome.

17. The conductive channel-containing membrane of claim 9, wherein the incorporated SPP1 connector polypeptide variant is mobile in the membrane layer.

18. The conductive channel-containing membrane of claim 9, which is capable of translocating a single stranded nucleic acid molecule, a double-stranded nucleic acid molecule, and/or a polypeptide through the aperture when the electrical potential is applied, wherein said conductive channel-containing membrane is capable of polypeptide detection, identification, sequencing, and discrimination.

19. A method of sensing a molecule using a conductive channel-containing membrane of any one of claims 9-18.

20. The method of claim 19, wherein the molecule is a polypeptide.

21. The method of claim 19, wherein the molecule is a nucleic acid molecule.

22. The method of claim 21, wherein the nucleic acid molecule is a double-stranded nucleic acid molecule.

23. A method of DNA sequencing using a conductive channel-containing membrane of any one of claims 9-18.

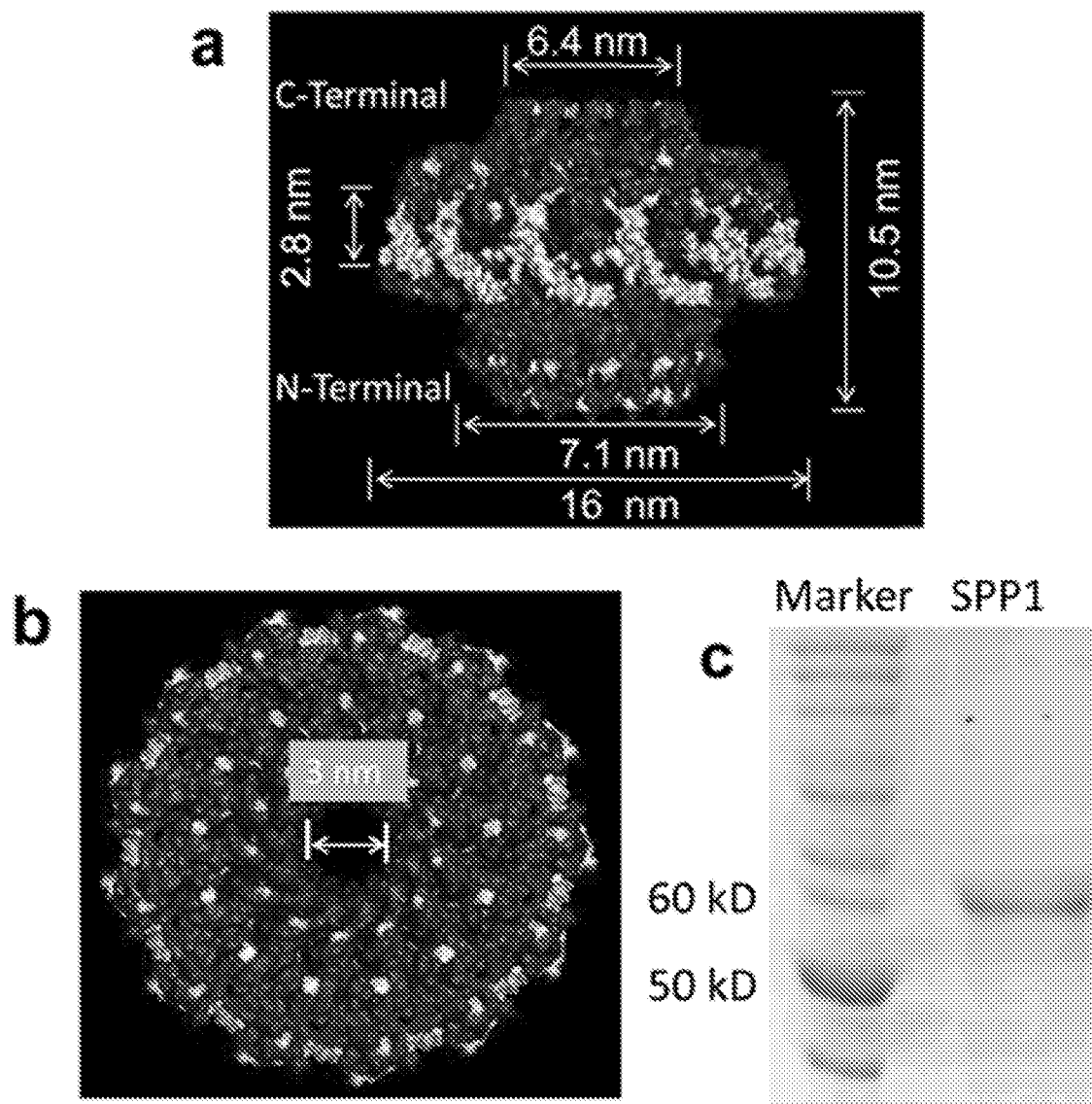


FIG. 1

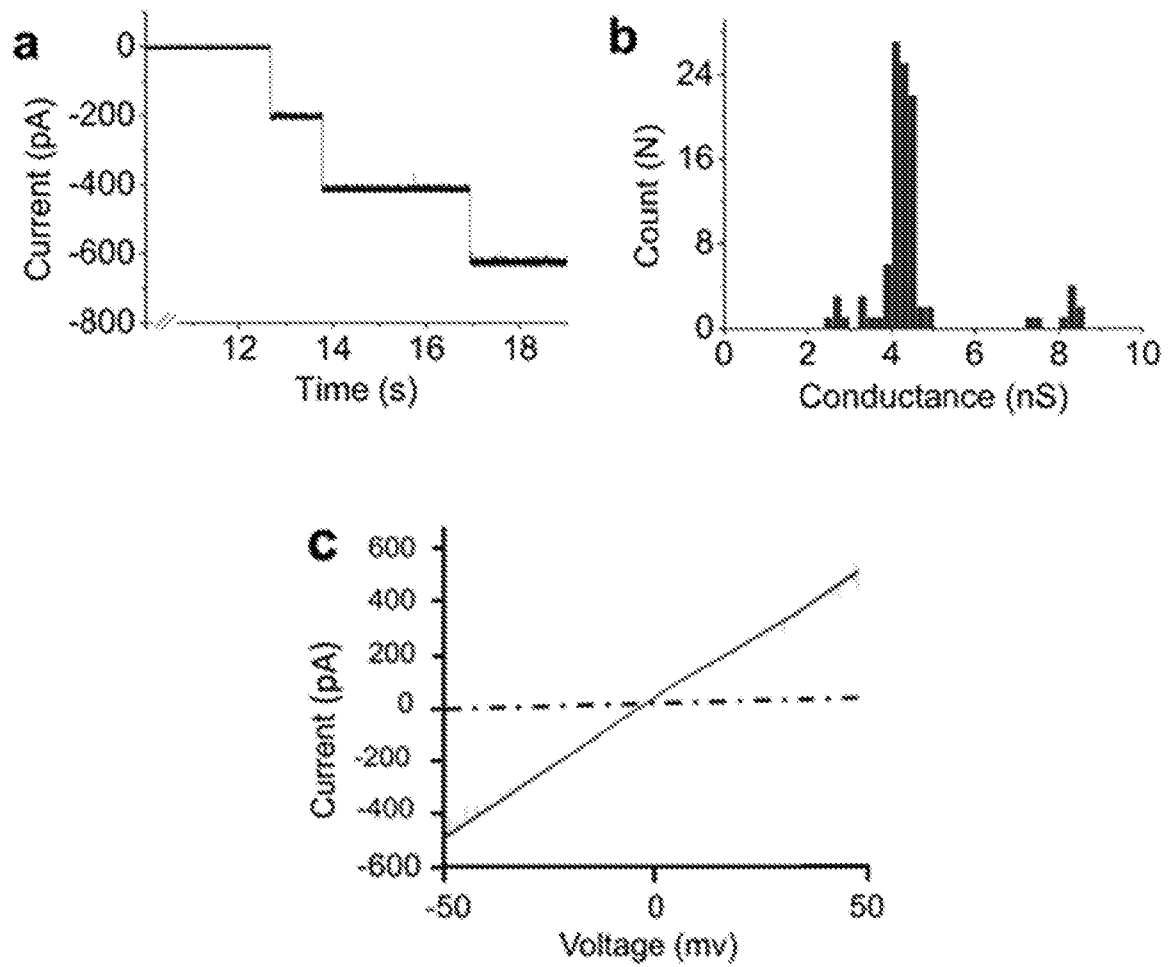


FIG. 2

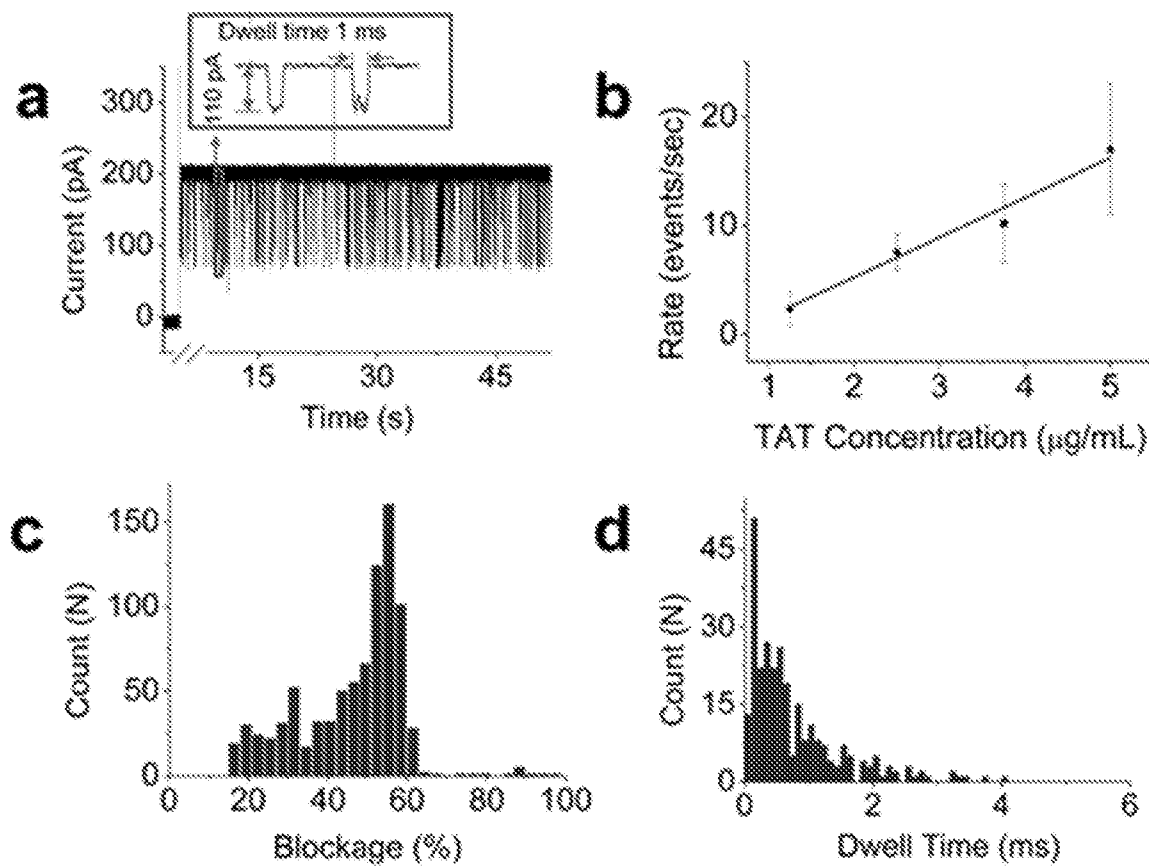


FIG. 3

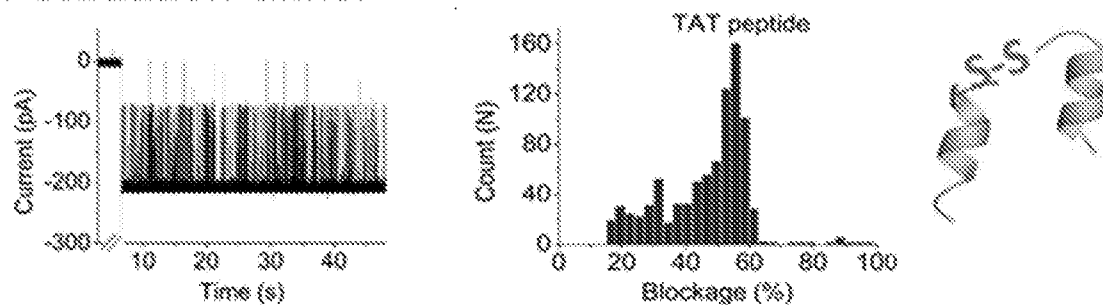
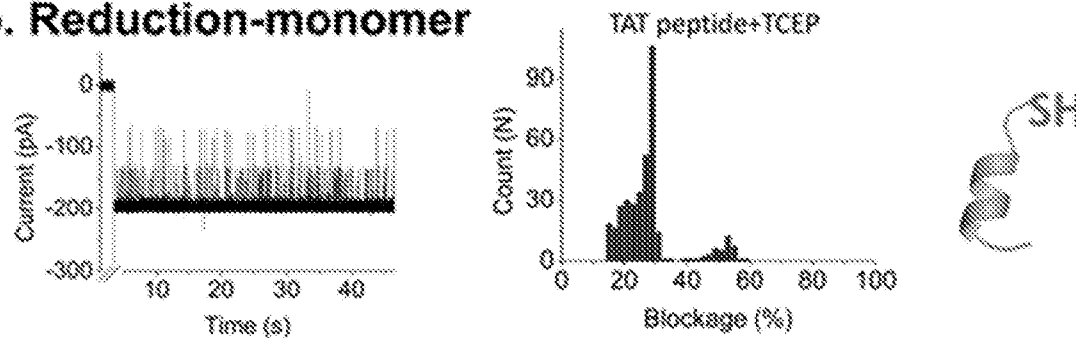
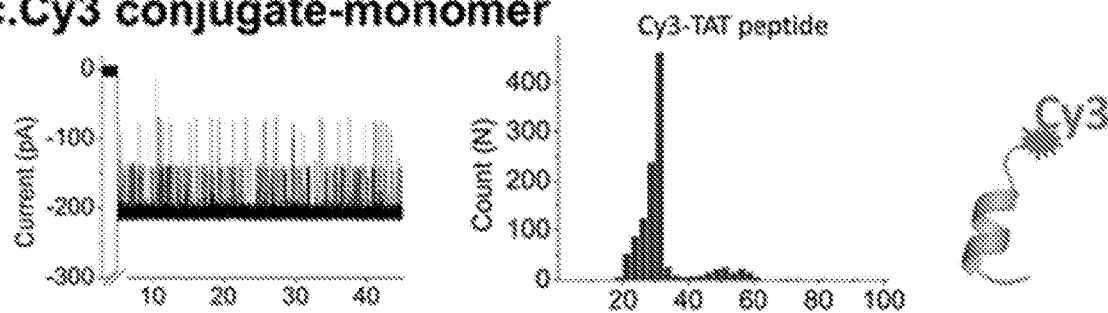
a. Oxidation-dimer**b. Reduction-monomer****c. Cy3 conjugate-monomer**

FIG. 4

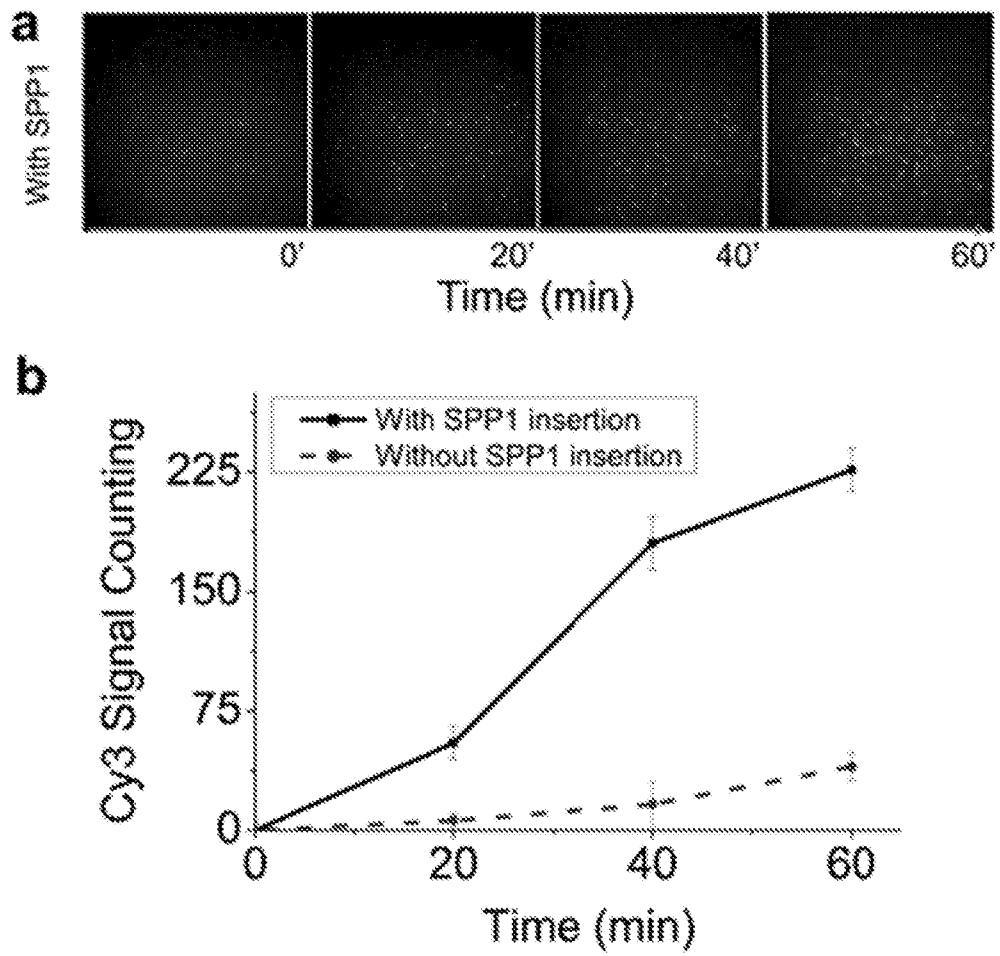


FIG. 5

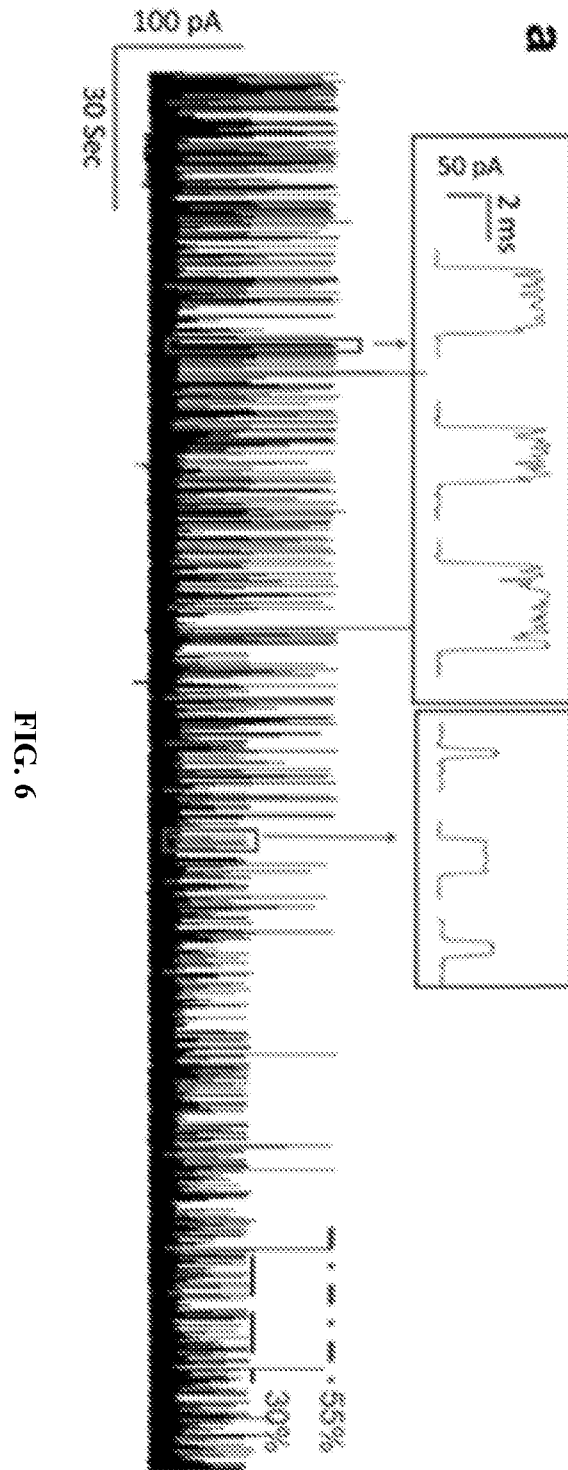


FIG. 6

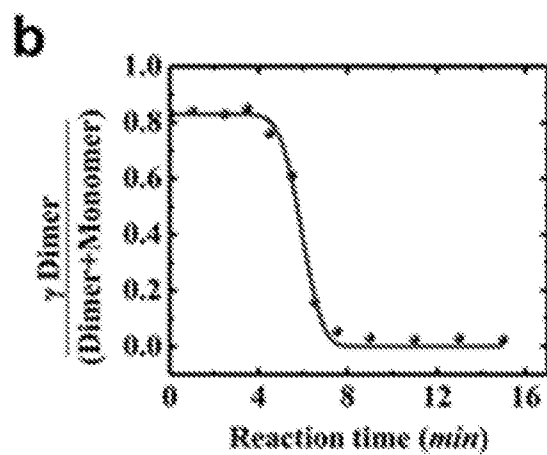
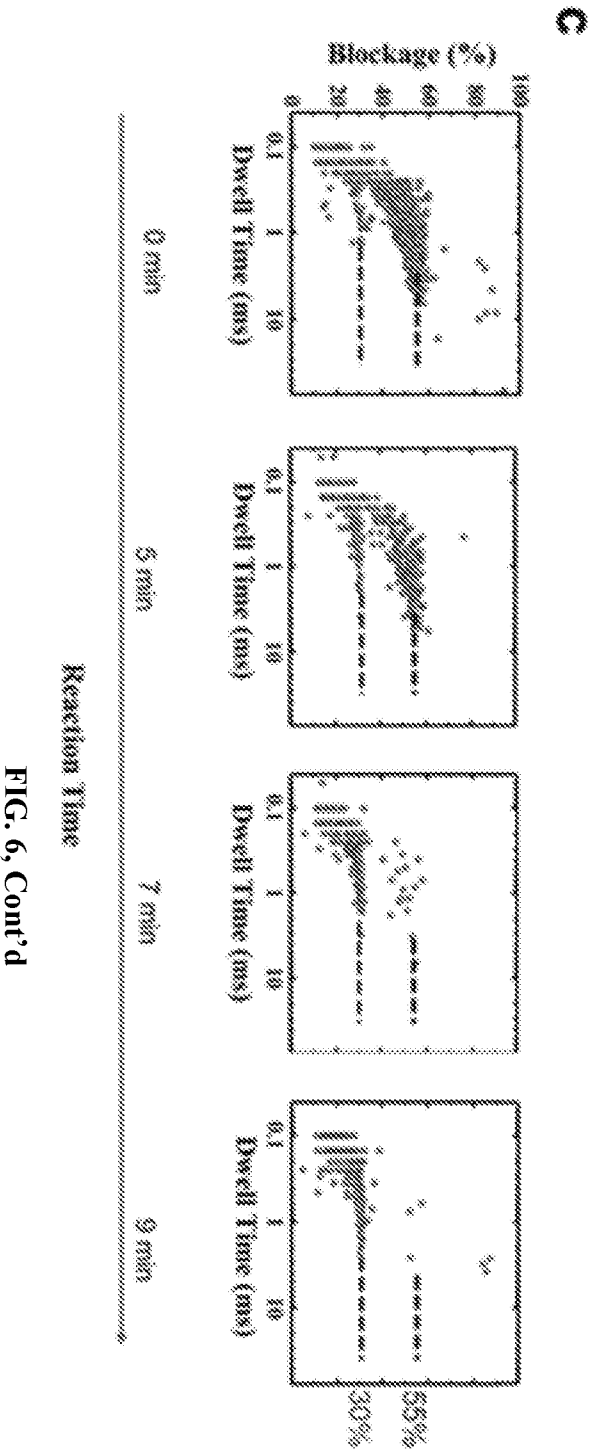


FIG. 6, Cont'd



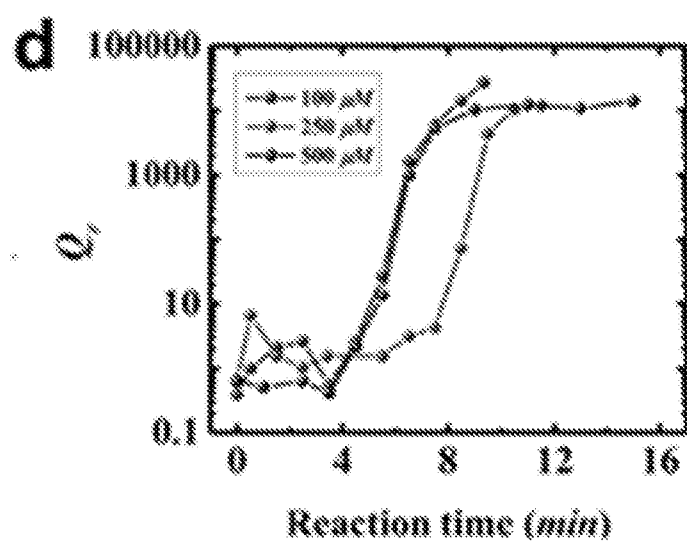


FIG. 6, Cont'd

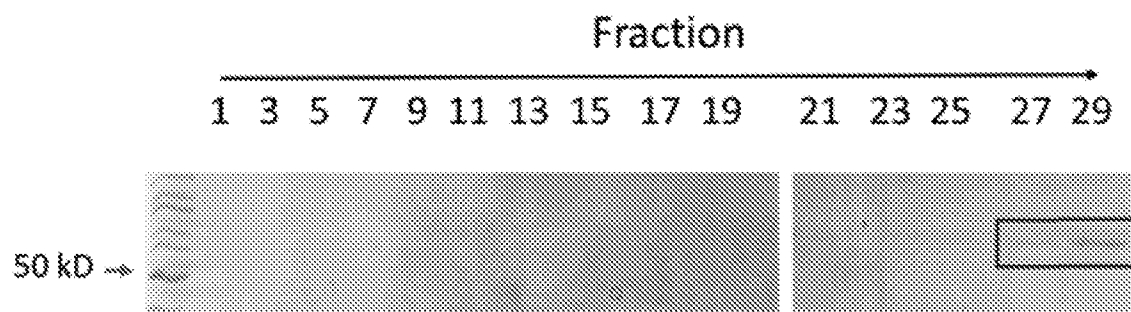
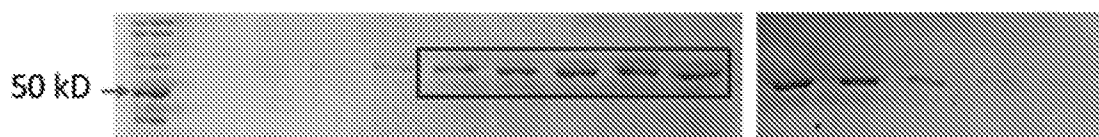
a. BSA (MW: 60 kD)**b. SPP1 connector (MW: 58 kD)**

FIG. 7

PATENT COOPERATION TREATY

PCT

DECLARATION OF NON-ESTABLISHMENT OF INTERNATIONAL SEARCH REPORT

(PCT Article 17(2)(a), Rules 13ter.1(c) and (d) and 39)

Applicant's or agent's file reference 13177-1981WO	IMPORTANT DECLARATION	Date of mailing <i>(day/month/year)</i> 01 APR 2016
International application No. PCT/US16/13702	International filing date <i>(day/month/year)</i> 15 January 2016	(Earliest) Priority Date <i>(day/month/year)</i> 16 January 2015
International Patent Classification (IPC) or both national classification and IPC IPC: C12Q 1/68; CPC: C12Q 2600/158		
Applicant UNIVERSITY OF KENTUCKY RESEARCH FOUNDATION		

This International Searching Authority hereby declares, according to Article 17(2)(a), that **no international search report will be established** on the international application for the reasons indicated below.

1. ☐ The subject matter of the international application relates to:
 - a. ☐ scientific theories
 - b. ☐ mathematical theories
 - c. ☐ plant varieties
 - d. ☐ animal varieties
 - e. ☐ essentially biological processes for the production of plants and animals, other than microbiological processes and the products of such processes
 - f. ☐ schemes, rules or methods of doing business
 - g. ☐ schemes, rules or methods of performing purely mental acts
 - h. ☐ schemes, rules or methods of playing games
 - i. ☐ methods for treatment of the human body by surgery or therapy
 - j. ☐ methods for treatment of the animal body by surgery or therapy
 - k. ☐ diagnostic methods practised on the human or animal body
 - l. ☐ mere presentations of information
 - m. ☐ computer programs for which this International Searching Authority is not equipped to search prior art
2. ☐ The failure of the following parts of the international application to comply with prescribed requirements prevents a meaningful search from being carried out:

☐ the description
☐ the claims
☐ the drawings
3. ☒ A meaningful search could not be carried out without the sequence listing; the applicant did not, within the prescribed time limit:

☒ furnish a sequence listing in the form of an Annex C/ST.25 text file, and such listing was not available to the International Searching Authority in a form and manner acceptable to it; or the sequence listing furnished did not comply with the standard provided for in Annex C of the Administrative Instructions.

☐ furnish a sequence listing on paper or in the form of an image file complying with the standard provided for in Annex C of the Administrative Instructions, and such listing was not available to the International Searching Authority in a form and manner acceptable to it; or the sequence listing furnished did not comply with the standard provided for in Annex C of the Administrative Instructions.

☐ pay the required late furnishing fee for the furnishing of a sequence listing in response to an invitation under Rule 13ter.1(a) or (b).
4. Further comments:

Applicant failed to submit a valid electronic seq. listing in response to the ISA/225.

Name and mailing address of the ISA/ Mail Stop PCT, Attn: ISA/US, Commissioner for Patents P.O. Box 1450, Alexandria, Virginia 22313-1450 Facsimile No. 571-273-8300	Authorized officer Blaine Copenheaver PCT Helpdesk: 571-272-4300 PCT OSP: 571-272-7774
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