

ATTACHING NUCLEOTIDES TO A POLYNUCLEOTIDE WITH A POLYMERASE

CROSS-REFERENCE TO RELATED APPLICATIONS

[0001] This application claims the benefit of U.S. Provisional Patent Application No. 62/890,065, filed on August 21, 2019 and entitled “Attaching Nucleotides to a Polynucleotide with a Polymerase,” the entire contents of which are incorporated by reference herein.

SEQUENCE LISTING

[0002] The instant application contains a Sequence Listing which has been submitted electronically in ASCII format and is hereby incorporated by reference in its entirety. Said ASCII copy, created on August 7, 2020, is named IP-1820-PCT_SL.txt and is 3,379 bytes in size.

BACKGROUND

[0003] Many current sequencing platforms use “sequencing by synthesis” (SBS) technology and fluorescence based methods for detection. Alternative sequencing methods that allow for more cost effective, rapid, and convenient sequencing and nucleic acid detection are desirable as complements to SBS. Charge based sequencing is an attractive approach.

[0004] Current sequencing by synthesis (SBS) technology uses nucleotides that are modified at two positions: 1) the 3' (or 3-prime) hydroxyl (3'-OH) of deoxyribose, and 2) the 5-position of pyrimidines or 7-position of purines of nitrogenous bases (A, T, C, G). The 3'-OH group is blocked with an azidomethyl group to create reversible nucleotide terminators. This may prevent further elongation after the addition of a single nucleotide. Each of the nitrogenous bases is separately modified with a fluorophore to provide a fluorescence readout which identifies the single base incorporation. Subsequently, the 3'-OH blocking group and the fluorophore are removed and the cycle repeats.

[0005] The current cost of the modified nucleotides may be high due to the synthetic challenges of modifying both the 3'-OH of deoxyribose and the nitrogenous base. There are several possible methods to reduce the cost of the modified nucleotides. One method is to move the readout label to the 5'-terminal (or 5-prime-terminal) phosphate instead of the nitrogenous base. In one example, this removes the need for a separate cleavage step, and allows for real time detection of the incoming nucleotide. During incorporation, the pyrophosphate together with the tag is released as a by-product of the elongation process, thus a cleavable linkage is not involved.

[0006] A method for assessing polymerase activity in a context relevant to or used in such a real-time SBS system without requiring detection of a tagged nucleotide per se may be beneficial. For example it may be advantageous to be able to assess a polymerase's ability to add a tagged nucleotide under different testing conditions independently of or separate from context and conditions of sequencing on a device. Because, by design, a tag on a nucleotide may be released from a nucleotide upon its incorporation into a nascent strand, measuring timing and kinetics of polymerase-mediated incorporation of such tagged nucleotides is challenging. An ability to determine how well or in any case at what pace a polymerase is able to incorporate a nucleotide bearing such a releasable tag under various conditions, in a high-throughput manner, may be helpful in deploying on-device real-time SBS systems using such tagged nucleotides and polymerases.

SUMMARY

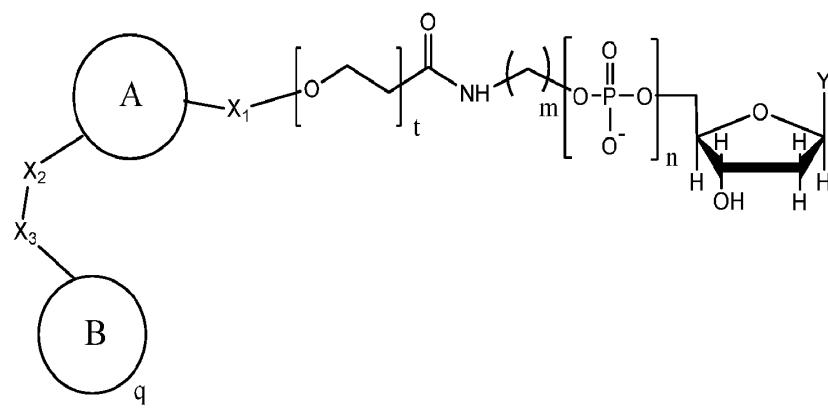
[0007] In an aspect, provided is a method, including hybridizing a polynucleotide to a template, and contacting a first template nucleotide 5-prime adjacent to a 5-prime-most nucleotide of the plurality of nucleotides that are complementary to the polynucleotide with a polymerase and a charge-tagged nucleotide, wherein the charge-tagged nucleotide is complementary to the first template nucleotide and includes a charge tag attached to a 5-prime polyphosphate of the charge-tagged nucleotide. Another example further includes attaching the charge-tagged nucleotide to the polynucleotide with the polymerase, wherein the attaching includes detachment of the charge tag from the charge-tagged nucleotide.

[0008] In an example, the template is bound to a substrate. In yet another example, the polymerase is bound to a substrate. In still another example, the template and the polymerase are bound to a substrate. In another example, the polymerase is selected from a Klenow fragment and a Phi29 polymerase.

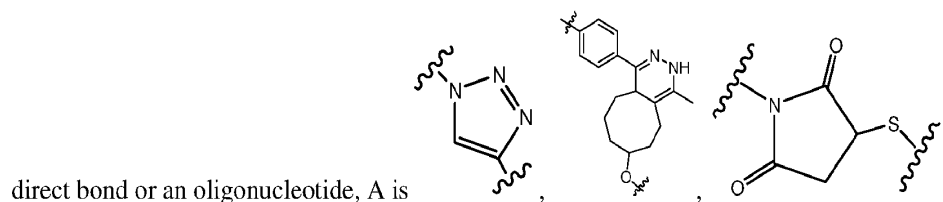
[0009] In yet another example, the template is attached to a substrate and includes linking nucleotides, and wherein the linking nucleotides are between the second template nucleotide and the substrate. In an example, there may be more than 20 linking nucleotides, or from 1 to 20 linking nucleotide, or from 10 to 20 linking nucleotides, or from 1 to 10 linking nucleotides, or from 1 to 5 linking nucleotides, or from 5 to 10 linking nucleotides, or from 10 to 15 linking nucleotides, or from 15 to 20 linking nucleotides. In still another example, the substrate is a solid support conductive channel and the conductive channel is to detect the charge-tagged nucleotide during the contacting the first template nucleotide with the charge-

tagged nucleotide and attaching the charge-tagged nucleotide to the polynucleotide. In still a further example, the method includes attaching the charge-tagged nucleotide in a solution wherein the charge tag includes a Debye length in the solution, and the Debye length is between about 0.5 nm and about 10 nm.

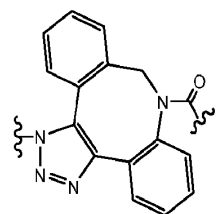
[0010] In an example, the charge-tagged nucleotide is a compound of Formula I



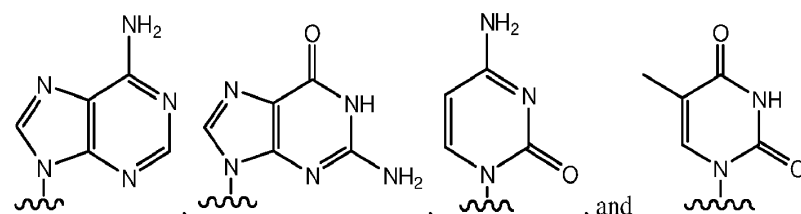
wherein n is an integer from 3 to 10, m is an integer from 1 to 10, t is an integer from 0 to 50, X₁ is a direct bond, a C₁-C₁₀ alkyl, a C₁-C₁₀ oxaalkyl, a C₁-C₁₀ thiaalkyl, or a C₁-C₁₀ azaalkyl, X₂ is C₁-C₂₀ alkyl wherein optionally one or more CH₂ residues are individually replaced with a peptide bond or (-O-CH₂-CH₂-)_a, wherein a is an integer from 1 to 24, X₃ is a



direct bond or an oligonucleotide, A is

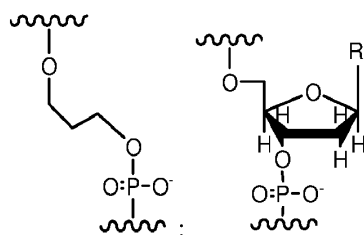


, or an amide bond, and Y is selected from the group consisting of



, q is an integer

from 1 to 100, and B is selected from the group consisting of: an amino acid; a nucleotide;



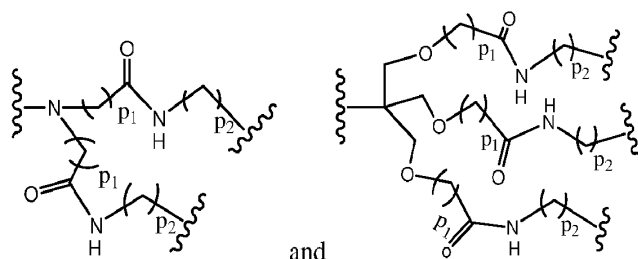
wherein each R is independently selected from Y and

hydrogen; and a dendron; and wherein q is equal to 1 when B is a dendron, and the q number of B has a charge and a charge density.

[0011] In another example the charge is between about -100e and about +100e. In yet another example, the charge density is between about -100e per cubic nanometer and about +100e per cubic nanometer. In still another example, the charge is between about -200e and about +200e. In still another example, the charge density is between about -200e per cubic nanometer and about +200e per cubic nanometer.

[0012] In a further example, B includes a q number of nucleotides and q is more than 1. In still a further example, the polynucleotide is selected from a branched polynucleotide and one or more hairpin loops. In yet a further example, the polynucleotide includes between two and five hairpin loops.

[0013] In another example, the q number of B includes a polypeptide. In still another example, the polypeptide is selected from a branched polypeptide, coiled polypeptide, and coiled-coil polypeptide. In yet another example, B includes an amino acid, and one or more of the q number of B include methyllysine, dimethyllysine, or trimethyllysine. In a further example, B is a dendron of z generations including one or more constitutional repeating unit and a plurality of end units, wherein z is an integer from 1 to 6, the constitutional end units are selected from



and

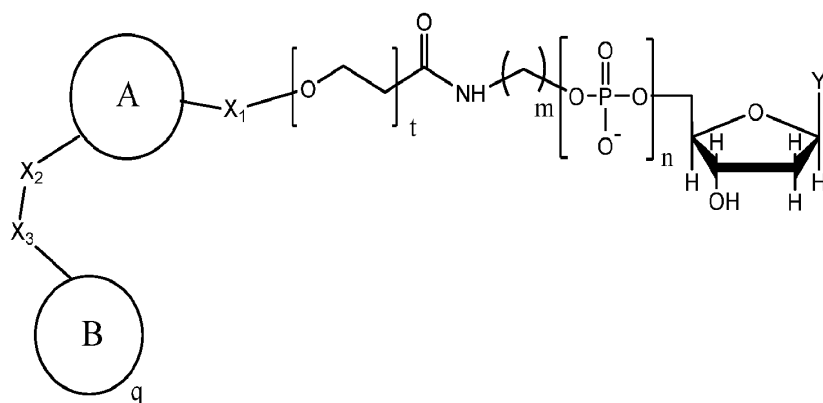
, wherein p1 is an integer from

1 to 3, wherein any one or more of the p1 -CH₂- groups is optionally replaced with from 1 to 3 -O-CH₂-CH₂- groups, p2 is an integer from 1 to 3, wherein any one or more of the p2 -CH₂-

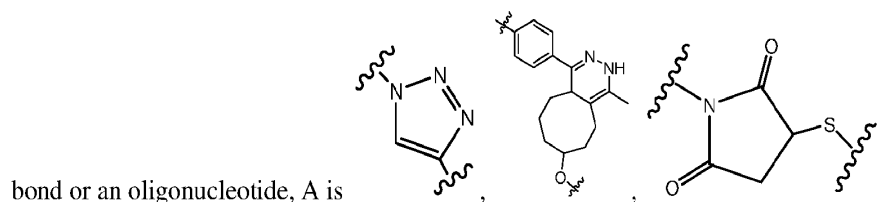
groups is optionally replaced with from 1 to 3 -O-CH₂-CH₂- groups, and the end groups are selected from carboxylic acid, sulfonic acid, phosphonic acid, sperminyl group, amino group, and quaternary ammonium group.

[0014] In still a further example, A was formed by a reaction including a linking reaction and the linking reaction is selecting from an azide-alkyne copper-assisted click reaction, a tetrazine-trans-cyclooctene ligation, an azide-dibenzocyclooctyne group copper-free click reaction, and a thiol-maleimide conjugation. In yet a further example, X₂ is (-O-CH₂-CH₂-)_a wherein a is an integer from 1 to 24, or wherein a is 24, or wherein a is 16, or wherein a is 12 or wherein a is 8, or wherein a is 4.

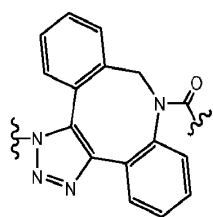
[0015] In an example, the template is attached to a substrate and includes linking nucleotides wherein the linking nucleotides are between the second template nucleotide and the substrate, and X₃ includes an oligonucleotide, wherein the oligonucleotide hybridizes to a plurality of the linking nucleotides when the charge tag is in proximity to the substrate. In another example, the charge-tagged nucleotide is a compound of Formula I



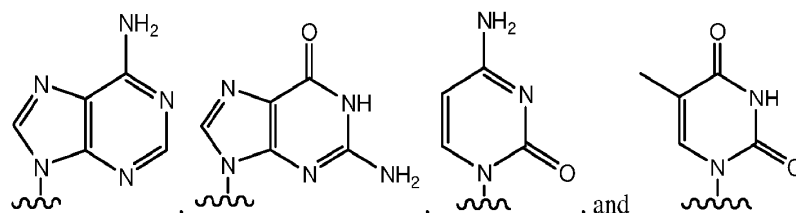
wherein n is an integer from 3 to 10, m is an integer from 1 to 10, t is an integer from 0 to 50, X₁ is a direct bond, a C₁-C₁₀ alkyl, a C₁-C₁₀ oxaalkyl, a C₁-C₁₀ thiaalkyl, or a C₁-C₁₀ azaalkyl, X₂ is C₁-C₂₀ alkyl wherein optionally one or more CH₂ residues are individually replaced with a peptide bond or (-O-CH₂-CH₂-)_a, wherein a is an integer from 1 to 24, X₃ is a direct



bond or an oligonucleotide, A is



, or an amide bond, and Y is selected from the group consisting of



, q is an integer

from 1 to 100, B includes an amino acid, and the q number of B has a charge and a charge density.

[0016] In yet another example, the charge is between about -100e and about +100e. In a further example, the charge density is between about -100e per cubic nanometer and about +100e per cubic nanometer. In still a further example, the charge is between about -200e and about +200e. In yet a further example, the charge density is between about -200e per cubic nanometer and about +200e per cubic nanometer. In still another example, the q number of B includes a polypeptide. In yet another example, the polypeptide is selected from a branched polypeptide, coiled polypeptide, and coiled-coil polypeptide. In a further example, one or more of the q number of B includes methyllysine, dimethyllysine, or trimethyllysine.

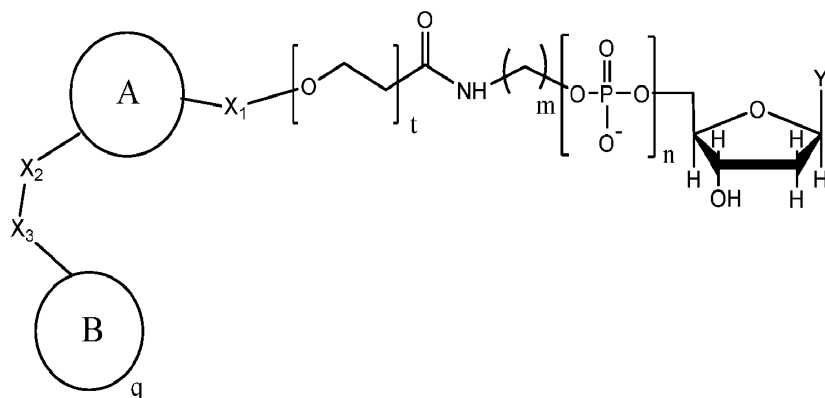
[0017] In still a further example, the polypeptide is a branched polypeptide including one or more forks each of the one or more forks including a plurality of branches, and one or more of the plurality of branches each independently includes a number of amino acids. In yet a further example, the polypeptide includes three or more forks and the number of amino acids of one or more of the plurality of branches is at least four. In another example, the polypeptide includes seven or more forks. In still another example, the number of amino acids of one or more of the plurality of branches is at least six.

[0018] In yet another example, A was formed by a reaction including a linking reaction and the linking reaction is selecting from an azide-alkyne copper-assisted click reaction, a tetrazine-trans-cyclooctene ligation, an azide-dibenzocyclooctyne group copper-free click reaction, and a thiol-maleimide conjugation. In a further example, X_2 is $(-O-CH_2-CH_2-)_a$ wherein a is an integer from 1 to 24. In still a further example, a is 24, or a is 16, or a is 12, or

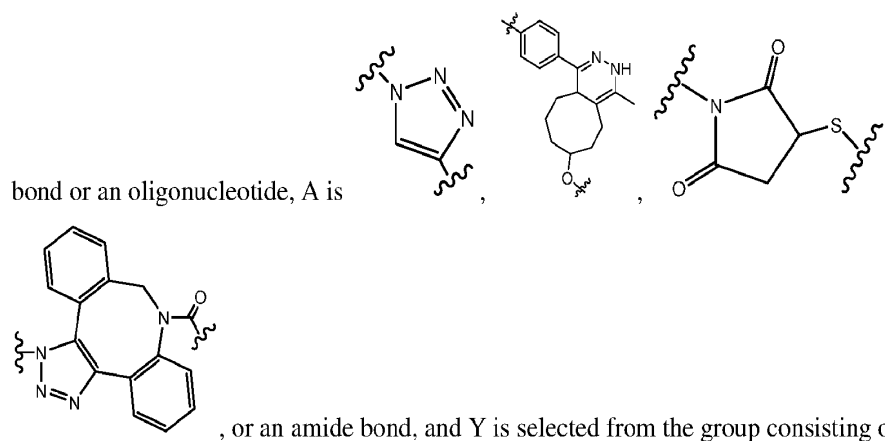
a is 8, or a is 4.

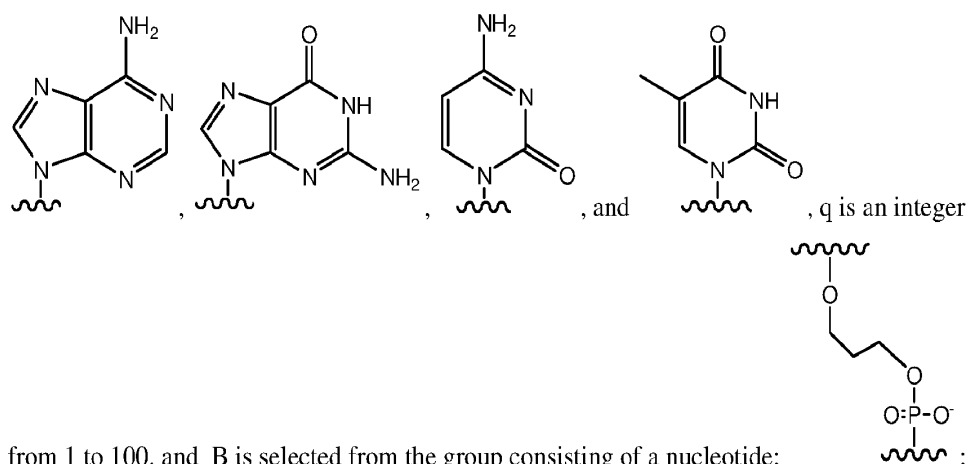
[0019] In yet a further example, the template is attached to a substrate and includes linking nucleotides wherein the linking nucleotides are between the second template nucleotide and the substrate, and X_3 includes an oligonucleotide, wherein the oligonucleotide hybridizes to a plurality of the linking nucleotides when the charge tag is in proximity to the substrate.

[0020] In another example, the charge-tagged nucleotide is a compound of Formula I

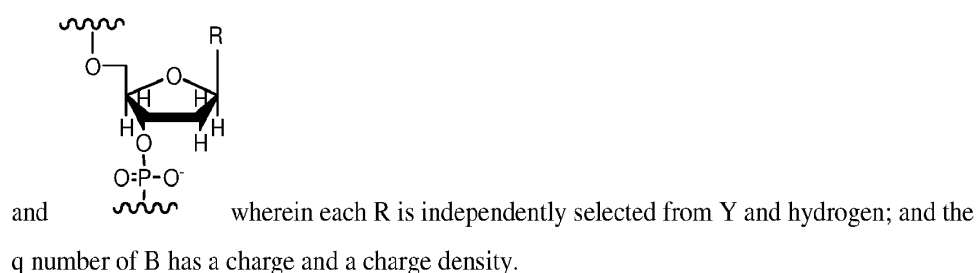


wherein n is an integer from 3 to 10, m is an integer from 1 to 10, t is an integer from 0 to 50, X_1 is a direct bond, a C_1 - C_{10} alkyl, a C_1 - C_{10} oxaalkyl, a C_1 - C_{10} thiaalkyl, or a C_1 - C_{10} azaalkyl, X_2 is C_1 - C_{10} alkyl wherein optionally one or more CH_2 residues are individually replaced with a peptide bond or $(-O-CH_2-CH_2-)a$ wherein a is an integer from 1 to 24, X_3 is a direct





from 1 to 100, and B is selected from the group consisting of a nucleotide;



[0021] In still another example, the charge is between about $-100e$ and about $+100e$. In yet another example, the charge density is between about $-100e$ per cubic nanometer and about $+100e$ per cubic nanometer. In a further example, the charge is between about $-200e$ and about $+200e$. In still a further example, the charge density is between about $-200e$ per cubic nanometer and about $+200e$ per cubic nanometer.

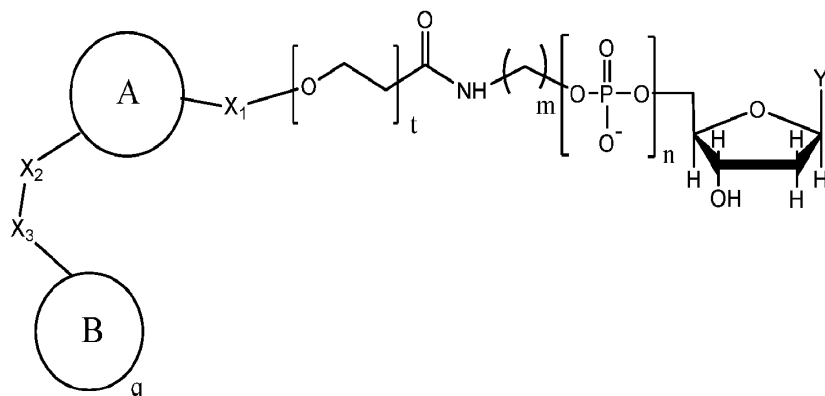
[0022] In another example, B includes a q number of nucleotides and q is more than 1. In still another example, the polynucleotide is selected from a branched polynucleotide and one or more hairpin loops. In yet another example, the polynucleotide includes between two and five hairpin loops.

[0023] In a further example, A was formed by a reaction including a linking reaction and the linking reaction is selecting from an azide-alkyne copper-assisted click reaction, a tetrazine-trans-cyclooctene ligation, an azide-dibenzocyclooctyne group copper-free click reaction, and a thiol-maleimide conjugation. In still a further example, X_2 is $(-O-CH_2-CH_2-)_a$ wherein a is an integer from 1 to 24, or a is 24, or a is 16, or a is 12 or a is 8, or a is 4.

[0024] In yet a further example, the template is attached to a substrate and includes linking nucleotides wherein the linking nucleotides are between the second template nucleotide and

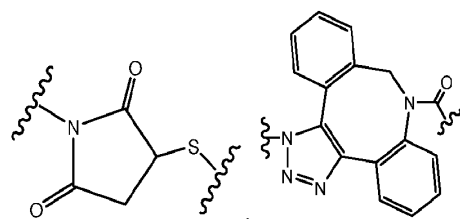
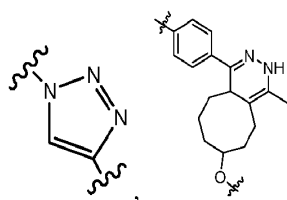
the substrate, and X₃ includes an oligonucleotide, wherein the oligonucleotide hybridizes to a plurality of the linking nucleotides when the charge tag is in proximity to the substrate.

[0025] In another example, the charge-tagged nucleotide is a compound of Formula I

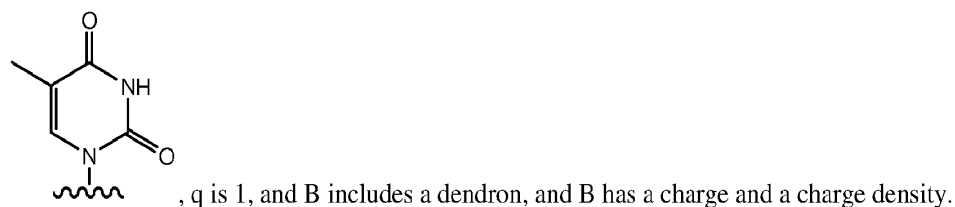
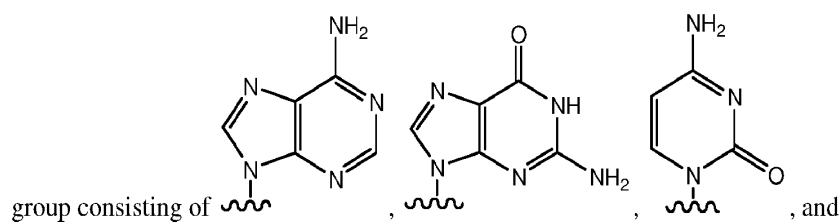


wherein n is an integer from 3 to 10, m is an integer from 1 to 10, t is an integer from 0 to 50, X₁ is a direct bond, a C₁-C₁₀ alkyl, a C₁-C₁₀ oxaalkyl, a C₁-C₁₀ thiaalkyl, or a C₁-C₁₀ azaalkyl, X₂ is C₁-C₂₀ alkyl wherein optionally one or more CH₂ residues are individually replaced with a peptide bond or (-O-CH₂-CH₂-)_a, wherein a is an integer from 1 to 24,

X₃ is a direct bond or an oligonucleotide, A is

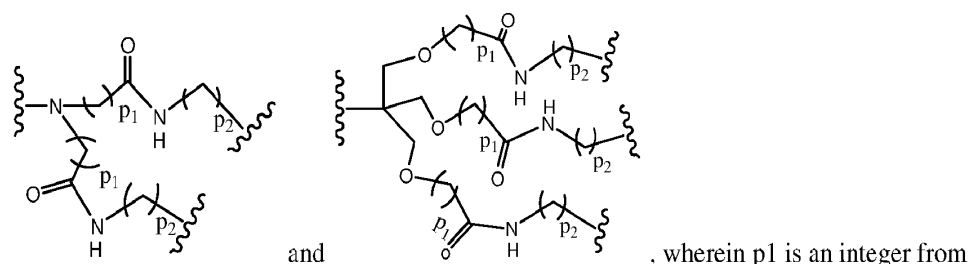


, or an amide bond, and Y is selected from the



[0026] In another example, the charge is between about -100e and about +100e. In still another example, the charge density is between about -100e per cubic nanometer and about +100e per cubic nanometer. In yet another example, the charge is between about -200e and about +200e. In a further example, the charge density is between about -200e per cubic nanometer and about +200e per cubic nanometer.

[0027] In still a further example, B is a dendron of z generations including one or more constitutional repeating unit and a plurality of end units, wherein z is an integer from 1 to 6, the constitutional end units are selected from:



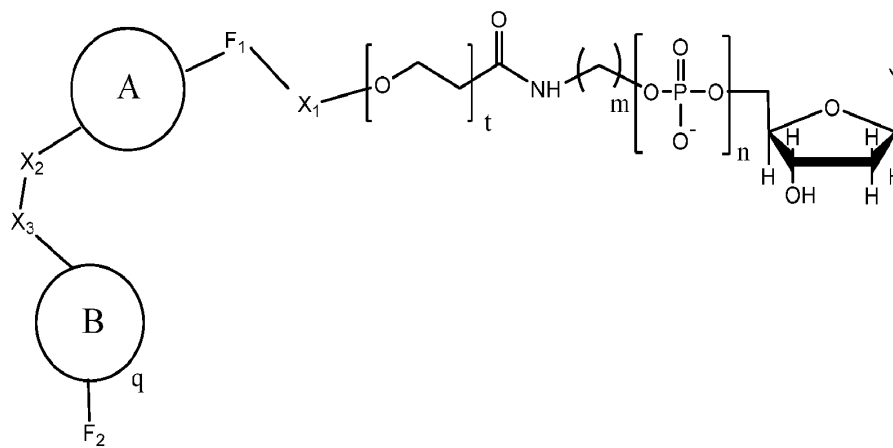
1 to 3, wherein any one or more of the p1 -CH₂- groups is optionally replaced with from 1 to 3 -O-CH₂-CH₂- groups, p2 is an integer from 1 to 3, wherein any one or more of the p2 -CH₂- groups is optionally replaced with from 1 to 3 -O-CH₂-CH₂- groups, and the end groups are selected from carboxylic acid, sulfonic acid, phosphonic acid, sperminyl group, amino group, and quaternary ammonium group.

[0028] In yet a further example, A was formed by a reaction including a linking reaction and the linking reaction is selecting from an azide-alkyne copper-assisted click reaction, a tetrazine-trans-cyclooctene ligation, an azide-dibenzocyclooctyne group copper-free click reaction, and a thiol-maleimide conjugation. In another example, X₂ is (-O-CH₂-CH₂-)_a

wherein a is an integer from 1 to 24 or a is 24, or a is 16, or a is 12, or a is 8, or a is 4. In still another example, the template is attached to a substrate and includes linking nucleotides wherein the linking nucleotides are between the second template nucleotide and the substrate, and X₃ includes an oligonucleotide, wherein the oligonucleotide hybridizes to a plurality of the linking nucleotides when the charge tag is in proximity to the substrate.

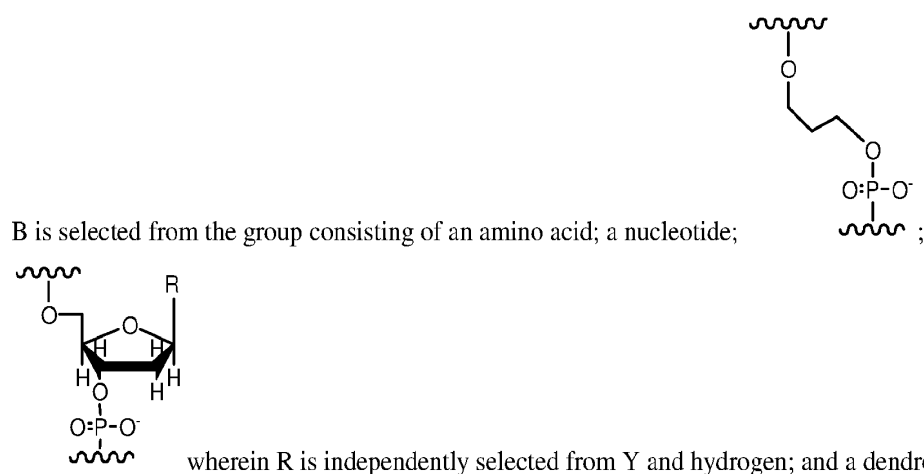
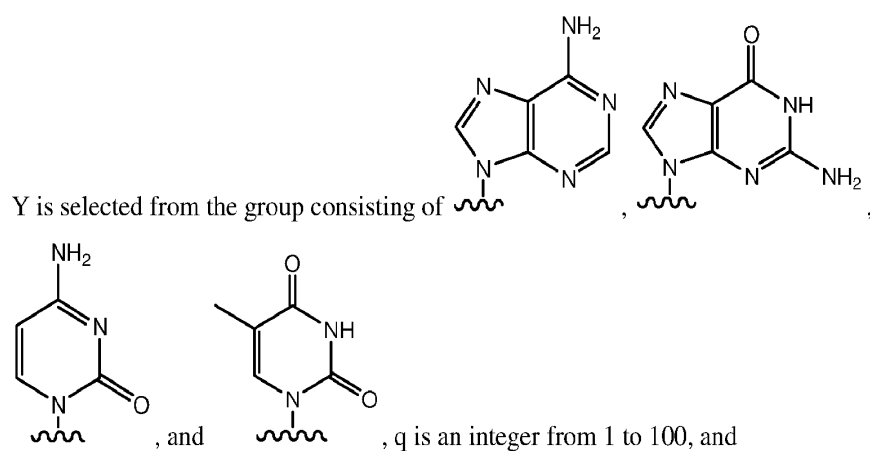
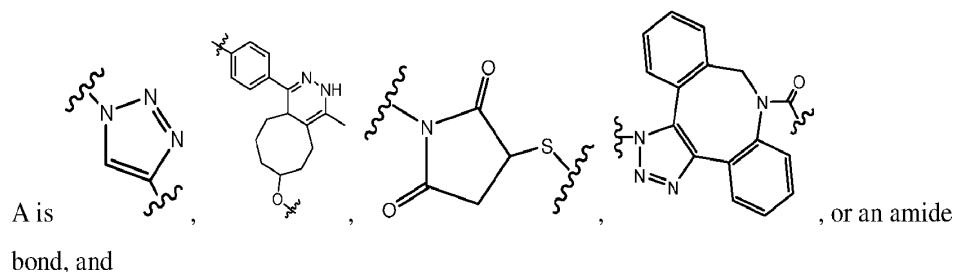
[0029] In another example, the template further includes a second template nucleotide 5-prime adjacent to the first template nucleotide, and the method further includes contacting the second template nucleotide with a fluorescently-tagged nucleotide wherein the fluorescently tagged nucleotide is complementary to the second template nucleotide but not to the first template nucleotide, and attaching the fluorescently-tagged nucleotide to the polynucleotide with a polymerase. In yet another example, the method further includes measuring attachment of the fluorescently tagged nucleotide to the polynucleotide by measuring fluorescence emitted from the polynucleotide. In still another example, the method further includes eluting the polynucleotide from the template before measuring.

[0030] In another aspect, provided is a method including detecting an incorporation of a labeled nucleotide into a nascent polynucleotide strand complementary to a template polynucleotide strand by a polymerase, wherein the polymerase is tethered to a solid support conductive channel by a tether, the labeled nucleotide is a compound of Formula I



wherein n is an integer from 3 to 10, m is an integer from 1 to 10, t is an integer from 0 to 50, X₁ is a direct bond, a C₁-C₁₀ alkyl, a C₁-C₁₀ oxaalkyl, a C₁-C₁₀ thiaalkyl, or a C₁-C₁₀ azaalkyl, X₂ is C₁-C₂₀ alkyl wherein optionally one or more CH₂ residues are replaced with a peptide bond or (-O-CH₂-CH₂-)_a, wherein a is an integer from 1 to 24, X₃ is a direct bond or an

oligonucleotide wherein the oligonucleotide hybridizes to an acceptor region of the tether when the label is in proximity to the conductive channel, F_1 is selected from a fluorophore and a direct bond and F_2 is absent or a fluorophore,



and wherein q is equal to 1 when B is a dendron, and the q number of B has a charge and a charge density, and

the conductive channel is to detect the labeled nucleotide during the incorporation.

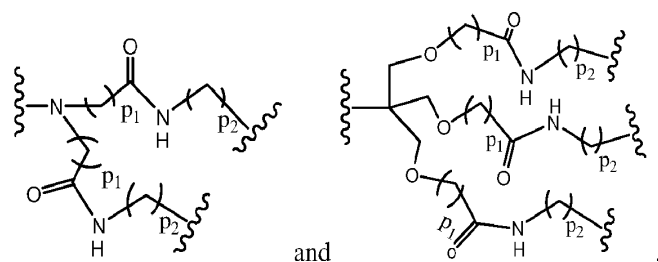
[0031] In an example, the charge is between about -100e and about +100e. In another

example, the charge density is between about -100e per cubic nanometer and about +100e per cubic nanometer. In yet another example, the charge is between about -200e and about +200e. In still a further example, the charge density is between about -200e per cubic nanometer and about +200e per cubic nanometer.

[0032] In a further example, B includes a q number of nucleotides and q is more than 1. In yet a further example, the polynucleotide is selected from a branched polynucleotide and one or more hairpin loops. In still another example, the polynucleotide includes between two and five hairpin loops.

[0033] In another example, the q number of B includes a polypeptide. In yet another example, the polypeptide is selected from the group consisting of branched polypeptide, coiled polypeptide, and coiled-coil polypeptide. In still another example, B includes an amino acid and one or more of the q number of B includes methyllysine, dimethyllysine, or trimethyllysine.

[0034] In another example, B is a dendron of z generations including one or more constitutional repeating unit and a plurality of end units, wherein z is an integer from 1 to 6, the constitutional end units are selected from:



wherein

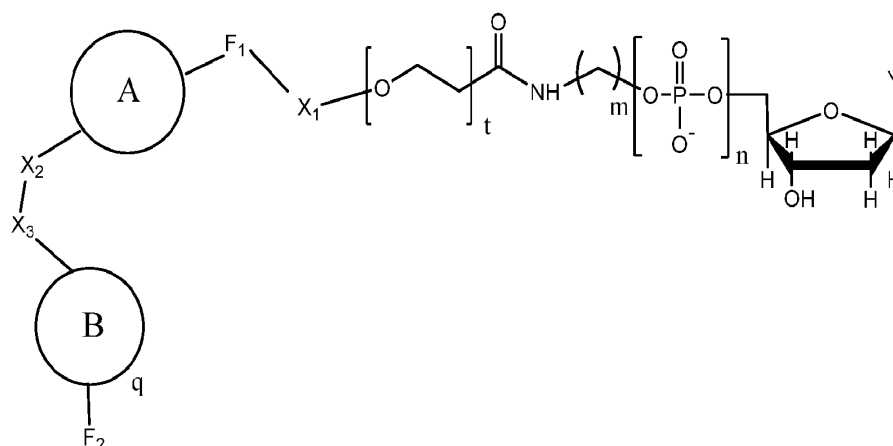
p_1 is an integer from 1 to 3, wherein any one or more of the p_1 -CH₂- groups is optionally replaced with from 1 to 3 -O-CH₂-CH₂- groups, p_2 is an integer from 1 to 3, wherein any one or more of the p_2 -CH₂- groups is optionally replaced with from 1 to 3 -O-CH₂-CH₂- groups, and the end groups are selected from carboxylic acid, sulfonic acid, phosphonic acid, sperminyl group, amino group, and quaternary ammonium group.

[0035] In yet another example, A was formed by a reaction including a linking reaction and the linking reaction is selecting from an azide-alkyne copper-assisted click reaction, a tetrazine-trans-cyclooctene ligation, an azide-dibenzocyclooctyne group copper-free click reaction, and a thiol-maleimide conjugation.

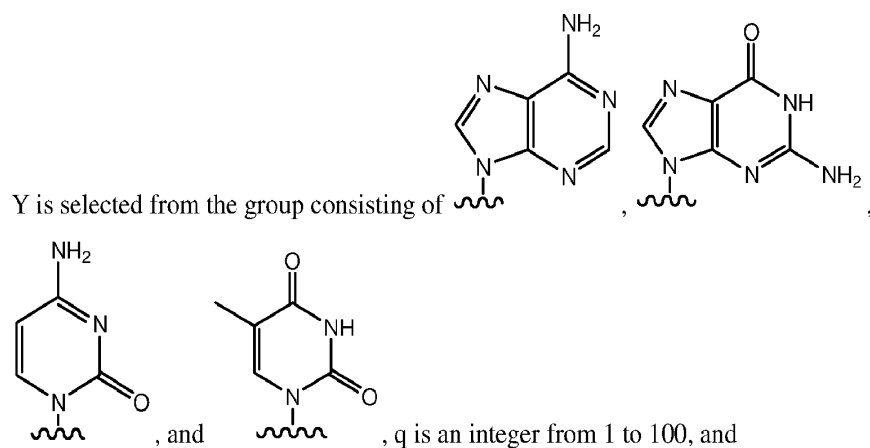
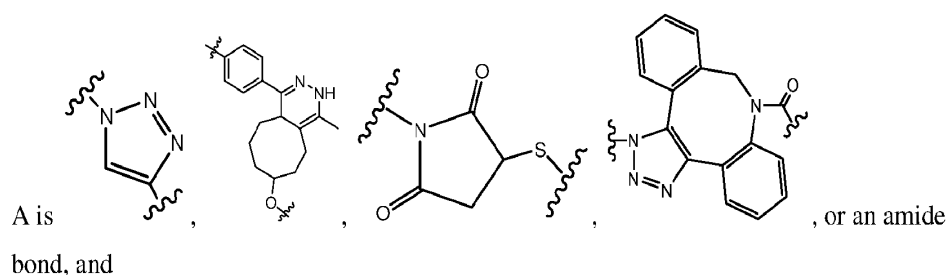
[0036] In still another example, the method further includes successively incorporating a plurality of labeled nucleotides wherein the charge of each of the plurality of labeled nucleotides differs from the charge of any other of the plurality of labeled nucleotides when the Y of the each and the Y of the any other differ from each other. In a further example, the method further includes identifying the Y of one or more labeled polynucleotide incorporated into the nascent polynucleotide strand based on the charge detected by the conductive channel.

[0037] In yet a further example, X_2 is $(-O-CH_2-CH_2-)_a$ wherein a is an integer from 1 to 24. In an example, a is 24. In another example, a is 12. In another example, a is 8. In still another example, a is 4.

[0038] In another aspect, provided is a method including detecting an incorporation of a labeled nucleotide into a nascent polynucleotide strand complementary to a template polynucleotide strand by a polymerase, wherein the polymerase is tethered to a solid support conductive channel by a tether, the labeled nucleotide is a compound of Formula I



wherein n is an integer from 3 to 10, m is an integer from 1 to 10, t is an integer from 0 to 50, X_1 is a direct bond, a C_1 - C_{10} alkyl, a C_1 - C_{10} oxaalkyl, a C_1 - C_{10} thiaalkyl, or a C_1 - C_{10} azaalkyl, X_2 is C_1 - C_{20} alkyl wherein optionally one or more CH_2 residues are individually replaced with a peptide bond or $(-O-CH_2-CH_2-)_a$, wherein a is an integer from 1 to 24, X_3 is a direct bond or an oligonucleotide wherein the oligonucleotide hybridizes to an acceptor region of the tether when the label is in proximity to the conductive channel, F_1 is selected from a fluorophore and a direct bond and F_2 is absent or a fluorophore,



B includes an amino acid, and the q number of B has a charge and a charge density, and the conductive channel is to detect the labeled nucleotide during the incorporation.

[0039] In an example, the charge is between about -100e and about +100e. In another example, the charge density is between about -100e per cubic nanometer and about +100e per cubic nanometer. In yet another example, the charge is between about -200e and about +200e. In still a further example, the charge density is between about -200e per cubic nanometer and about +200e per cubic nanometer.

[0040] In another example, the q number of B includes a polypeptide. In yet another example, the polypeptide is selected from the group consisting of branched polypeptide, coiled polypeptide, and coiled-coil polypeptide. In still another example, B includes an amino acid and one or more of the q number of B includes methyllysine, dimethyllysine, or trimethyllysine.

[0041] In still a further example, the polypeptide is a branched polypeptide including one or more forks and a plurality of branches, and one or more of the plurality of branches each independently includes a number of amino acids. In yet a further example, the polypeptide includes three or more forks and the number of amino acids of one or more of the plurality of

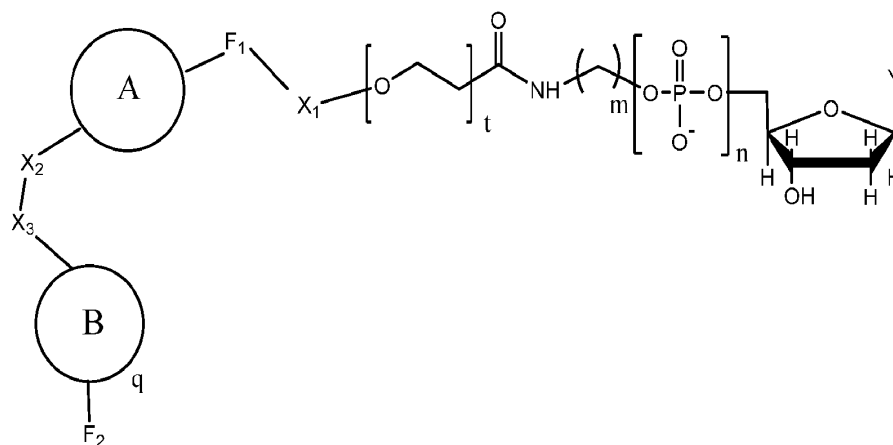
branches is at least four. In another example, the polypeptide includes seven or more forks. In still another example, the number of amino acids of one or more of the plurality of branches is at least six.

[0042] In yet another example, A was formed by a reaction including a linking reaction and the linking reaction is selecting from an azide-alkyne copper-assisted click reaction, a tetrazine-trans-cyclooctene ligation, an azide-dibenzocyclooctyne group copper-free click reaction, and a thiol-maleimide conjugation.

[0043] In still another example, the method further includes successively incorporating a plurality of labeled nucleotides wherein the charge of each of the plurality of labeled nucleotides differs from the charge of any other of the plurality of labeled nucleotides when the Y of the each and the Y of the any other differ from each other. In a further example, the method further includes identifying the Y of one or more labeled polynucleotide incorporated into the nascent polynucleotide strand based on the charge detected by the conductive channel.

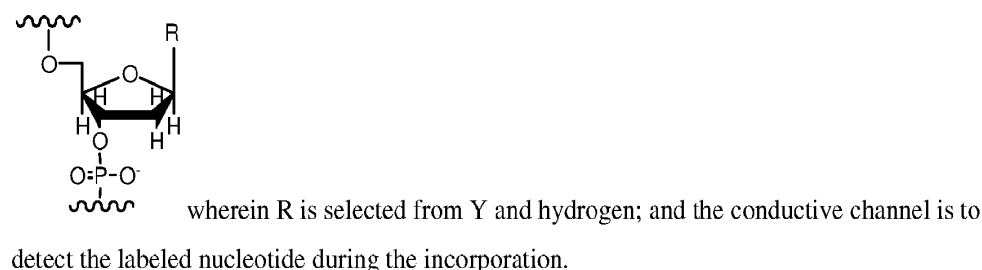
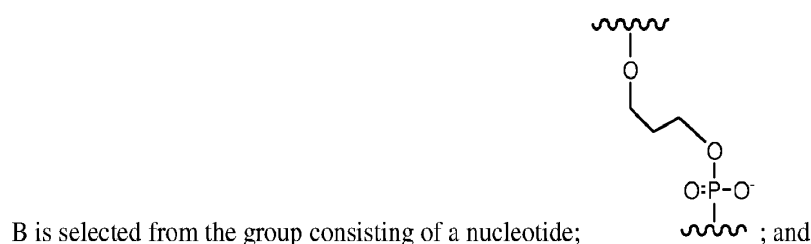
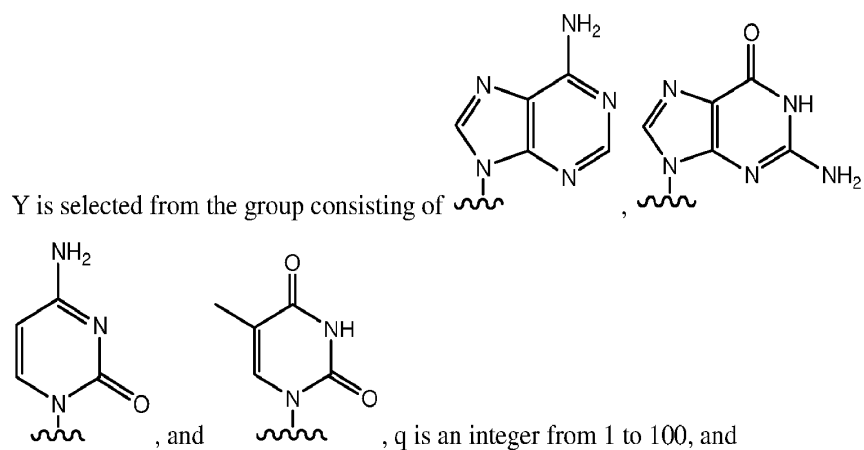
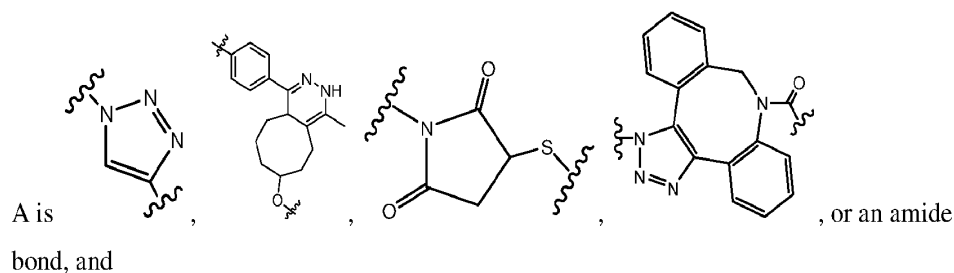
[0044] In yet a further example, X_2 is $(-O-CH_2-CH_2-)_a$ wherein a is an integer from 1 to 24. In an example, a is 24. In another example, a is 12. In another example, a is 8. In still another example, a is 4.

[0045] In still another aspect, provided is a method including detecting an incorporation of a labeled nucleotide into a nascent polynucleotide strand complementary to a template polynucleotide strand by a polymerase, wherein the polymerase is tethered to a solid support conductive channel by a tether, the labeled nucleotides is a compound of Formula I



wherein n is an integer from 3 to 10, m is an integer from 1 to 10, t is an integer from 0 to 50,

X_1 is a direct bond, a C_1 - C_{10} alkyl, a C_1 - C_{10} oxaalkyl, a C_1 - C_{10} thiaalkyl, or a C_1 - C_{10} azaalkyl, X_2 is C_1 - C_{20} alkyl wherein optionally one or more CH_2 residues are individually replaced with a peptide bond or $(-O-CH_2-CH_2-)_a$, wherein a is an integer from 1 to 24, X_3 is a direct bond or an oligonucleotide wherein the oligonucleotide hybridizes to an acceptor region of the tether when the label is in proximity to the conductive channel, F_1 is selected from a fluorophore and a direct bond and F_2 is absent or a fluorophore,



[0046] In an example, the charge is between about -100e and about +100e. In another example, the charge density is between about -100e per cubic nanometer and about +100e per cubic nanometer. In yet another example, the charge is between about -200e and about +200e. In still a further example, the charge density is between about -200e per cubic nanometer and about +200e per cubic nanometer.

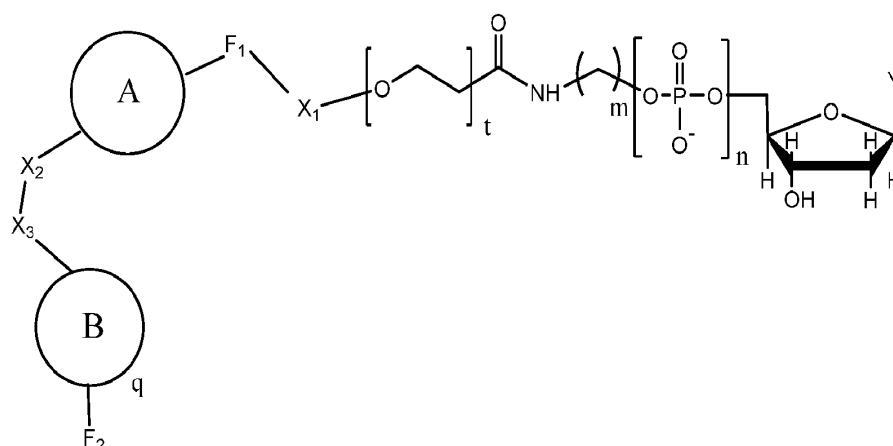
[0047] In a further example, B includes a q number of nucleotides and q is more than 1. In yet a further example, the polynucleotide is selected from a branched polynucleotide and one or more hairpin loops. In still another example, the polynucleotide includes between two and five hairpin loops.

[0048] In yet another example, A was formed by a reaction including a linking reaction and the linking reaction is selecting from an azide-alkyne copper-assisted click reaction, a tetrazine-trans-cyclooctene ligation, an azide-dibenzocyclooctyne group copper-free click reaction, and a thiol-maleimide conjugation.

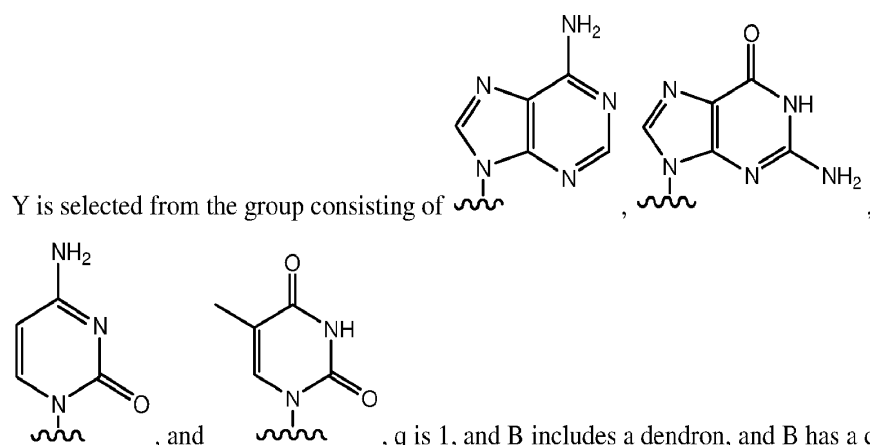
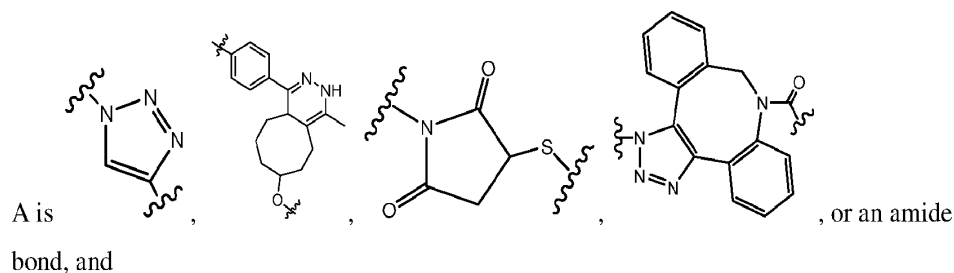
[0049] In still another example, the method further includes successively incorporating a plurality of labeled nucleotides wherein the charge of each of the plurality of labeled nucleotides differs from the charge of any other of the plurality of labeled nucleotides when the Y of the each and the Y of the any other differ from each other. In a further example, the method further includes identifying the Y of one or more labeled polynucleotide incorporated into the nascent polynucleotide strand based on the charge detected by the conductive channel.

[0050] In yet a further example, X_2 is $(-O-CH_2-CH_2-)_a$ wherein a is an integer from 1 to 24. In an example, a is 24. In another example, a is 12. In another example, a is 8. In still another example, a is 4.

[0051] In a further aspect, provided is a method including detecting an incorporation of a labeled nucleotide into a nascent polynucleotide strand complementary to a template polynucleotide strand by a polymerase, wherein the polymerase is tethered to a solid support conductive channel by a tether, the labeled nucleotide is a compound of Formula I



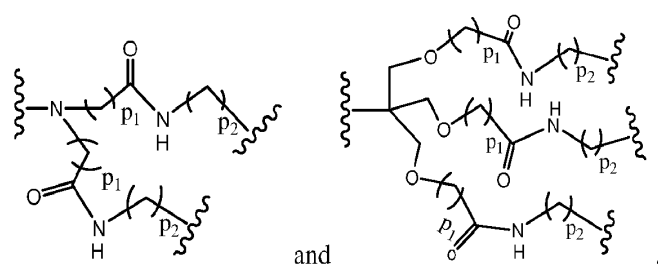
wherein n is an integer from 3 to 10, m is an integer from 1 to 10, t is an integer from 0 to 50, X_1 is a direct bond, a C_1 - C_{10} alkyl, a C_1 - C_{10} oxaalkyl, a C_1 - C_{10} thiaalkyl, or a C_1 - C_{10} azaalkyl, X_2 is C_1 - C_{20} alkyl wherein optionally one or more CH_2 residues are individually replaced with a peptide bond or $(-O-CH_2-CH_2-)_a$, wherein a is an integer from 1 to 24, X_3 is a direct bond or an oligonucleotide wherein the oligonucleotide hybridizes to an acceptor region of the tether when the label is in proximity to the conductive channel, F_1 is selected from a fluorophore and a direct bond and F_2 is absent or a fluorophore,



a charge density, and the conductive channel is to detect the labeled nucleotide during the incorporation.

[0052] In an example, the charge is between about -100e and about +100e. In another example, the charge density is between about -100e per cubic nanometer and about +100e per cubic nanometer. In yet another example, the charge is between about -200e and about +200e. In still a further example, the charge density is between about -200e per cubic nanometer and about +200e per cubic nanometer.

[0053] In another example, B is a dendron of z generations including one or more constitutional repeating unit and a plurality of end units, wherein z is an integer from 1 to 6, the constitutional end units are selected from:



wherein

p_1 is an integer from 1 to 3, wherein any one or more of the p_1 -CH₂- groups is optionally replaced with from 1 to 3 -O-CH₂-CH₂- groups, p_2 is an integer from 1 to 3, wherein any one or more of the p_2 -CH₂- groups is optionally replaced with from 1 to 3 -O-CH₂-CH₂- groups, and the end groups are selected from carboxylic acid, sulfonic acid, phosphonic acid, sperminyl group, amino group, and quaternary ammonium group.

[0054] In yet another example, A was formed by a reaction including a linking reaction and the linking reaction is selecting from an azide-alkyne copper-assisted click reaction, a tetrazine-trans-cyclooctene ligation, an azide-dibenzocyclooctyne group copper-free click reaction, and a thiol-maleimide conjugation.

[0055] In still another example, the method further includes successively incorporating a plurality of labeled nucleotides wherein the charge of each of the plurality of labeled nucleotides differs from the charge of any other of the plurality of labeled nucleotides when the Y of the each and the Y of the any other differ from each other. In a further example, the method further includes identifying the Y of one or more labeled polynucleotide incorporated into the nascent polynucleotide strand based on the charge detected by the conductive

channel.

[0056] In yet a further example, X_2 is $(-O-CH_2-CH_2-)_a$ wherein a is an integer from 1 to 24. In an example, a is 24. In another example, a is 12. In another example, a is 8. In still another example, a is 4.

[0057] It should be appreciated that all combinations of the foregoing concepts and additional concepts discussed in greater detail below (provided such concepts are not mutually inconsistent) are contemplated as being part of the inventive subject matter disclosed herein and contribute to the advantages and benefits as described herein.

BRIEF DESCRIPTION OF THE DRAWINGS

[0058] These and other features, aspects, and advantages of the present disclosure will become better understood when the following detailed description is read with reference to the accompanying drawings, wherein:

[0059] FIG. 1 shows a flow chart for performing a method in accordance with aspects of the present disclosure.

[0060] FIG. 2 shows an example of nucleotide incorporation into a polynucleotide in accordance with aspects of the present disclosure.

[0061] FIG. 3 shows, in an example, a polymerase attached to a conductive channel via a tether.

[0062] FIG. 4 shows, in one example, polymerases attached to conductive channels via nucleic acid tethers and bound to nucleotides that can be distinguished based on charge or proximity to the charge detector.

[0063] FIG. 5 shows, in one example, polymerases attached to conductive channels via nucleic acid tethers and bound to nucleotides that can be distinguished based on charge.

[0064] FIG. 6 shows, in one example, a polymerase tethered to a conductive channel, wherein the conductive channel is also attached to an acceptor region, including in this example a plurality of oligonucleotides capable of binding (e.g., hybridizing) to a specificity region within linkers on nucleotides.

[0065] FIG. 7 shows an illustration of a non-limiting example of a nucleotide analog bearing a charge tag in accordance with the present disclosure. A nucleotide analog may include a nucleotide polyphosphate (such as dT hexaphosphate as shown), a linker region optionally comprising a specificity region, and a charge tag. In this non-limiting example, a linker includes a covalent attachment formed by azide-alkyne click chemistry. As further described

below, a specificity region may be included in the linker and may assist in promoting charge tag proximity with a conductive channel during nucleotide incorporation by a polymerase.

[0066] FIG. 8 shows, in one example, a nucleotide label having negatively charged oxygens in the phosphodiester backbone of an oligonucleotide moiety of the label.

[0067] FIG. 9A, FIG. 9B, and FIG. 9C show, as examples, example multiplier units to construct branched charge tags that can be detected using a conductive channel.

[0068] FIG. 10 shows, in one example, a conductive channel that is attached to a polymerase (Pol) via a tether having a nucleic acid sequence (generically represented as a sequence of 10 Ns). The N nucleotides are selected from universal bases and bases that are complementary to nucleotides in a linker (e.g., a specificity region) attached to a charge tag.

[0069] FIG. 11 shows, in one example, a conductive channel that is attached to a polymerase (Pol) via a tether having an acceptor region, in this example a nucleic acid sequence (generically represented as a sequence of 7 Ns with an ABC region; charge tag portion not shown). The polymerase is complexed to a target nucleic acid and a labeled CTP analog. The linker on the CTP analog includes a nucleic acid region having inosines (I) and a specificity region (A'B'C') that hybridizes to an acceptor region on the tether (ABC).

[0070] FIG. 12 shows, in one example, a tethered polymerase in four different positional states relative to the conductive channel due to the binding of each of four different nucleotide analogs through a specificity region in each linker with an acceptor region in the tether. For this illustrative example, the nucleotide analogs are identified as ATP, GTP, CTP and TTP, but any nucleotide analogs could be used (e.g., deoxyribonucleotide analogs may be used). Each of the nucleotide analogs has an oligonucleotide moiety of the same length as the other 3 nucleotide analogs, but each nucleotide analog has a specific binding sequence that binds to a different region of the acceptor region in the tether compared to the regions where the other nucleotide analog linkers bind. The charge tag, being an oligonucleotide in this example or other phosphodiester-containing charge tag in other examples, extends outside the region of hybridization at the end of the linker opposite the nucleotide.

[0071] FIG. 13 shows, in one example, single nucleotide incorporation of phosphodiester based charge tags by polymerase phi29.

[0072] FIG. 14A, FIG. 14B, FIG. 14C, and FIG. 14D show examples of peptide-based charge tags in accordance with aspects of the present disclosure.

[0073] FIG. 15A, FIG. 15B, and FIG. 15C show, in an example, several structures of a

modified nucleotide with a structured oligonucleotide as a charge tag. Shown are modified nucleotides with a charge tag extending therefrom, wherein the charge tags include a specificity region bonded to an acceptor region (indicated as "Glue"). FIG. 15A shows a stem-and-loop shaped charge tag and FIG. 15C shows a cloverleaf-shaped charge tag (SEQ ID NO: 1).

[0074] FIG. 16A and FIG. 16B show an example of a cruciform charge tag. FIG. 16A shows a cruciform charge tag comprising four oligonucleotides bonded together in a Holliday structure-like configuration and single-stranded oligonucleotide overhangs. FIG. 16B shows the structure from FIG. 16A with sequences of peptide nucleic acids bound to the oligonucleotide overhangs and coiled polypeptide structures extending from the ends of the peptide nucleic acid sequences. In this example, the polypeptide sequences have a positive charge.

[0075] FIG. 17 shows several examples of polypeptide charge tags including coiled polypeptides and assembly thereof.

[0076] FIG. 18A and FIG. 18B show two views of an example of a charge tag including polypeptides arranged in a coiled-coil configuration.

[0077] FIG. 19A and FIG. 19B show examples of phosphodiester-based charge tags having a branched structure. FIG. 19A discloses "TTTTT TTTTT" as SEQ ID NO: 11.

[0078] FIG. 20A, FIG. 20B, FIG. 20C, FIG. 20D, and FIG. 20E, show non-limiting examples of phosphodiester charge tags in accordance with aspects of the present disclosure. Figures disclose SEQ ID NOS 2-6, respectively, in order of appearance.

[0079] FIG. 21A and FIG. 21B show examples of branched peptide-based charge tags.

[0080] FIG. 22 depicts a synthesis method for synthesizing examples of charge tags in accordance with aspects of the present disclosure.

[0081] FIG. 23A and FIG. 23B depict examples of branched peptide charge tags in accordance with aspects of the present disclosure.

[0082] FIG. 24A, FIG. 24B, and FIG. 24C depict examples of linear peptide charge tags (FIG. 24A) and branched peptide charge tags (FIGS. 24B and 24C) in accordance with aspects of the present disclosure.

[0083] FIG. 25A and FIG. 25B show examples of spermine-based charge tags in accordance with aspects of the present disclosure.

[0084] FIG. 26 depicts an apparatus with a conductive channel for detecting a charge tag in

accordance with aspects of the present disclosure.

[0085] FIG. 27 is a graph depicting charge detection of various charge tags by a conductive channel in accordance with aspects of the present disclosure.

[0086] FIG. 28 is a graph depicting charge detection of various charge tags by a conductive channel in accordance with aspects of the present disclosure.

[0087] FIG. 29A and FIG. 29B depict charge detection of an example charge tag according to various Debye lengths in different buffers.

DETAILED DESCRIPTION

[0088] This disclosure relates to a method for determining aspects of charge tagged nucleotide incorporation by polymerases under various conditions. A real-time method for SBS sequencing or other methods for detection of nucleotide incorporation into a nascent nucleotide strand with a conductive channel, including use of nucleotides for incorporation with a charge tag attached to a nucleotide by its 5-prime phosphate group, may include metrics for assessing rate of nucleotide incorporation across different stages of incorporation. An effect of modification of numerous features on an ability, rate, kinetics, or other characteristics, of nucleotide incorporation of a nucleotide bearing a charge tag into a nascent polynucleotide strand according to a template may be measured. Advantageously, and as provided for in the present disclosure, a method may include assessing such effects independently of detecting the charge tag with a conductive channel during incorporation, such as when incorporation conditions are assessed or used under conditions not including a conductive channel or its use in detecting a charge tag.

[0089] When an electrically charged tag is attached to a nucleotide by the nucleotide's 5-prime phosphate group, a charge tag may dissociate from the charge tag upon the nucleotide's incorporation into a nascent oligonucleotide strand when the nucleotide is attached to the free 3-prime end of the nascent strand via the nucleotide's 5-prime phosphate. For example, as disclosed herein, a nucleotide may be linked to a charge tag by a polyphosphate attached to the 5-prime carbon of the sugar moiety of the nucleotide. A polymerase may cleave the charge tag from the nucleotide upon removing a portion of the polyphosphate from the nucleotide, much as a polymerase cleaves a 5-prime pyrophosphate group of an incorporating nucleotide when attaching a nucleotide to a 3-prime hydroxide of a nascent polynucleotide strand. As disclosed herein, a real-time SBS-like process may include incorporation of the charge-tagged nucleotide in proximity to a conductive channel such that

the conductive channel may detect the charge tag in association with the incorporating and thereby reflect a base being incorporated.

[0090] But in some examples, it may be desirable to perform aspects of such a method without detecting a charge tag with a conductive channel while retaining the ability to measure whether a nucleotide that included a charge tag prior to incorporation was added to the nascent oligonucleotide strand. Even in cases where incorporation of a charge-tagged nucleotide may be detected or detectable by a conductive channel in proximity thereto, or may be done in proximity to a conductive channel whether or not the conductive channel is used to detect such incorporation, it may be beneficial to detect such incorporation by another method independently of using a conductive channel to detect the charge tag.

[0091] Metrics concerning features of a method of real-time base calling using a charge tagged nucleotide and a conductive channel may be obtained with such a method. Structural features of an apparatus or surface and effects thereof on nucleotide incorporation, abilities and kinetics of different polymerase enzymes on nucleotide incorporation, incorporation characteristics of different nucleotides such as nucleotides including different charge tags, or with different links between a charge tag and the nucleotide, different surface chemistries of a substrate to which a polymerase is tethered in proximity to a conductive channel, different characteristics of different tethers by which a polymerase may be attached to a surface, including in proximity to a conductive channel, or other features as disclosed herein all may be interrogated with a method as disclosed herein independently of detection of charge tagged nucleotide by charge detection by a conductive channel as disclosed herein.

[0092] An example is depicted in FIG. 1. In this example, a template oligonucleotide (upward-pointing arrow) is hybridized to a polynucleotide (downwardly pointing arrow). In this example, following hybridization, a wash step may be performed whereby unhybridized, free polynucleotide is removed from the reaction. The hybridized template and polynucleotide may then be further incubated with a charge-tagged nucleotide (represented by dN6P in FIG. 1) and a polymerase enzyme. A sequence of the template and hybridized polynucleotide may be known and designed such that a vacant template nucleotide immediately 5-prime to the 5-prime-most template nucleotide that hybridizes to the polynucleotide may be known. In an example, such nucleotide may differ from the 5-prime-next nucleotide of the template. In such an example, because of base-pairing rules, it is possible to perform a single base incorporation step by including only one species of

nucleotide (C, G, A, or T) known to be complementary to the vacant template nucleotide, in the reaction, which species bears a charge tag.

[0093] A template, a polynucleotide hybridized thereto, or both, as disclosed herein, may be of any suitable length. In some examples, the template, polynucleotide hybridized thereto, or both, may be relatively short polynucleotides, on the order of 5, 10, 15, 20, 25, 30, 35, 40, 45, 50, 55, 60, 65, 70, 75, 80, 85, 90, 95, 100, 110, 120, 130, 140, 150, 160, 170, 180, 190, 200, or more nucleotides in length. In an example, one, the other, or both may be a primer, a relatively short polynucleotide that, when hybridized to a complementary polynucleotide, may serve as a primer for a polymerase, to be extended by the polymerase by addition of a nucleotide complementary to the next, non-hybridized nucleotide of its complement. In an example, a polynucleotide hybridized to a template may be a primer. A primer may be any suitable length, depending on a length of a complementary polynucleotide, such as a template, to which it is to hybridize and serve as a primer for a polymerase. In an example, a primer may be 5, 10, 15, 20, 25, 30, 35, 40, 45, 50, 55, 60, 65, 70, or 75 nucleotides in length, or any intervening length between these lengths. In an example, a primer may be longer than such lengths.

[0094] In this example, once a single base is incorporated into the hybridized polynucleotide, a further, second nucleotide may not be incorporated, because the species of nucleotide included in the reaction is not complementary to the second, 5-prime-next nucleotide of the template. Thus, a polymerase may not incorporate it. In other examples, the single species of nucleotide bearing a charge tag may also bear a chemical modification to reduce, minimize, or in some instances prevent addition of a second nucleotide after a single base is incorporated into the hybridized polynucleotide. For example, the charge-tagged nucleotide may bear a reversible blocking moiety that may reduce, minimize, or in some instances prevent addition of a subsequent nucleotide (e.g., a 3-prime azidomethyl group or other reversible blocking group). In such examples, further nucleotide addition may be reduced, minimized, or in some instances prevented until, and may be possible after, chemical modification of the free, 3-prime carbon end of the hybridized polynucleotide after incorporation of the charge-tagged nucleotide.

[0095] When a charge tag is attached to a nucleotide through a polyphosphate connected to the 5-prime carbon of the nucleotide, as in examples disclosed herein, incorporation of the nucleotide in the hybridized polynucleotide may result in dissociation of the charge tag from

the template and hybridized polynucleotide. In such cases it may be difficult to determine whether an incorporated nucleotide had been charge tagged, particularly in a high-throughput manner that may allow for determining further metrics associated with the incorporation as described above. As disclosed herein, incorporation of a second nucleotide may follow incorporation of the first, charge-tagged nucleotide, and the second nucleotide may possess features permitting its detection by conventional detection methods.

[0096] Continuing with the example depicted in FIG. 1, in addition to contacting a template and hybridized polynucleotide with a polymerase and charge-tagged nucleotide, they may further be contacted with a fully functional nucleotide (depicted as FFN in FIG. 1). The FFN may be complementary to the 5-prime-next nucleotide on the template, one nucleotide adjacent, in the template's 5-prime direction, to the template nucleotide complementary to the charge tagged nucleotide that was incorporated as described above.

[0097] In this example, such FFN may differ from the 5-prime-next nucleotide of the template. In such an example, because of base-pairing rules, it is possible to perform another single base incorporation step, after that used to incorporate a charge-tagged nucleotide as disclosed above, by including only one species of nucleotide (C, G, A, or T) known to be complementary to the next vacant template nucleotide, in the reaction, which species bears a detectable moiety such as a fluorescent tag.

[0098] In this example, once an FFN is incorporated into the hybridized polynucleotide, a further nucleotide may not be incorporated, because the species of FFN included in the reaction is not complementary to the 5-prime-next nucleotide of the template. Thus, a polymerase may not incorporate it. In other examples, the single species of FFN may also bear a chemical modification to reduce, minimize, or in some instances prevent addition of another nucleotide after an FFN is incorporated into the hybridized polynucleotide. For example, the FFN may bear a reversible blocking moiety that may reduce, minimize, or in some instances prevent addition of a subsequent nucleotide (e.g., a 3-prime azidomethyl group or other reversible blocking group). In such examples, further nucleotide addition may be reduced, minimized, or in some instances prevented until, and may be possible after, chemical modification of the free, 3-prime carbon end of the hybridized polynucleotide after incorporation of the FFN. In another example, the FFN may lack a hydroxyl group on its 3-prime carbon (e.g., may be a dideoxynucleotide including a detectable label such as a fluorescent tag), which may reduce, minimize, or in some instances prevent, the incorporation

of a subsequent nucleotide.

[0099] In another example, a method could include incorporation of more than one FFN, or one or more non-fluorescently tagged nucleotide in addition to the one or more incorporated FFN, in keeping with the method disclosed herein. Although an example is described herein in which an FFN is incorporated immediately 3-prime to the incorporated, erstwhile charge-tagged nucleotide, and presence of such FFN may subsequently be detected, in other examples other nucleotides may be incorporated after charge-tagged nucleotide incorporation and before the FFN is incorporated, and/or more than one FFN may be incorporate, adjacent thereto and/or to each other or spaced therefrom and/or each other, and detected for various reasons. All such examples are included in the present disclosure and explicitly considered as interchangeable with variations thereto as further disclosed below.

[0100] In an example, a template and hybridized polynucleotide may be incubated with a charge-tagged nucleotide and FFN sequentially, with a wash step performed in between incubations to remove unincorporated nucleotide. For example, in an example, a charge-tagged nucleotide and FFN for incorporation into a hybridized polynucleotide may both be of the same species of nucleotide as each other (e.g., both G, C, A, or T), whereas nucleotides of the template available for pairing with said nucleotides and incorporation into the hybridized polynucleotide may also both be the same as each other and complementary to the charge-tagged nucleotide and FFN. In such an example, a charge-tagged nucleotide may be modified so as to reduce, or in some instances minimize, or even prevent, incorporation of a following nucleotide (e.g., to reduce, minimize, or prevent incorporation of multiple charge-tagged nucleotides), such as disclosed above (e.g., by including aa 3-prime azidomethyl or other reversible block). Modification thereafter may be performed to permit subsequent incorporation of an FFN during a subsequent incubation step, with free charge-tagged nucleotide being washed out in between incubation steps.

[0101] In another example, a charge-tagged nucleotide and an FFN may both be incubated simultaneously. For example, a charge-tagged nucleotide may be of a species (e.g., C, G, T, or A) different from an FFN. Though both nucleotides may be simultaneously incubated with a template and hybridized polynucleotide and polymerase, they may be incorporated only serially because they are not complementary to the same template nucleotides. In an example, one or the other of the charge-tagged nucleotides may possess a 3-prime reversible block (such as an azidomethyl group or other reversible block) to reduce, minimize, or in some

instances prevent incorporation of another nucleotide thereafter absent chemical removal of the block. In another example, the FFN may lack a hydroxyl group on the 3-prime carbon (such as a dideoxynucleotide), to reduce, minimize, or in some instances prevent further addition of a nucleotide thereafter. In such examples, a charge-tagged nucleotide may be incorporated, followed by incorporation of an FFN, even if both may have been present simultaneously during one or more polymerization reactions.

[0102] In an example, a template may be attached to a substrate. For example, a 3-prime end of a template may be bound to a substrate such as a solid support, directly or indirectly. In another example, a 5-prime end of a template may be bound to a substrate such as a solid support, directly or indirectly. In an example, a template may be attached, directly or indirectly, to a substrate by a series of nucleotides not intended or selected for coding nucleotides for incorporation to the hybridized polynucleotide. For example, no charge-tagged nucleotide or FFN complementary to such nucleotides may be incubated with the template, hybridized polynucleotide, and polymerase, at all or at least not at a time when the hybridized polynucleotide may be extendable so as to incorporate nucleotides complementary to and hybridizable to such template nucleotides attaching a template to a surface. For example, a 3-prime end of a hybridized polynucleotide may be complementarily hybridized to a template nucleotide that is too far 5-prime from such attaching nucleotides of the template for a polymerase to use such attaching nucleotide as a coding substrate for attachment of further nucleotides to the hybridized polynucleotide, whether before or after a charge-tagged nucleotide or FFN have been incorporated into the hybridized polynucleotide. A template may be attached to a substrate by other chemical attachments as well, such as inert polymers such as polyethylene glycol or others.

[0103] In some examples, a polymerase reaction may take place in the presence of a charge-tagged nucleotide for incorporation by a polymerase according to a template, in the absence of another, fluorescently tagged nucleotide for subsequent incorporation. A washing step may be performed after polymerization in the presence of the first nucleotide so as to remove excess, unincorporated nucleotide. Fluorescently tagged nucleotide may then be added, in the presence of polymerase (whether the same or different from that which was present during polymerase-catalyzed incorporation of the charge-tagged nucleotide), for a second polymerization reaction.

[0104] In other examples, a polymerase may be bound to a substrate such as a solid support,

directly or indirectly. In another example, a template may be in solution and not bound to a substrate. In another example, a polymerase may be in solution not be bound to a substrate. In still another example, a hybridized polynucleotide may be bound to a substrate such as a solid support, directly or indirectly. In other examples it may be in solution and not bound to a solid support. A solid support may include a conductive channel as further described herein. A tether attaching a polymerase to a substrate may be of any given length suitable for a given purpose. Examples of tethers of various usable lengths and chemical compositions are explained in further detail below. In some examples, a polymerase may be attached to a substrate by covalent attachments, such as through an inert polymer such as polyethylene glycol or other inert polymer. In some examples, a first polymerase, a second polymerase, or both polymerases may be in solution during a polymerization reaction with a template tethered to a solid support.

[0105] In an example, there may be between 1 and 10 linking nucleotides attaching a template primer to a substrate, directly or indirectly, or there may be between 1 and 5 such linking nucleotides attaching a template primer to a substrate, directly or indirectly, or there may be between approximately 5 and 10 such linking nucleotides attaching a template primer to a substrate, directly or indirectly, or there may be between approximately 10 and 15 such linking nucleotides attaching a template primer to a substrate, directly or indirectly, or there may be between approximately 15 and 20 such linking nucleotides attaching a template primer to a substrate, directly or indirectly, or there may be between approximately 1 and 20 such linking nucleotides attaching a template primer to a substrate, directly or indirectly, or there may be between approximately 20 and 30 such linking nucleotides attaching a template primer to a substrate, directly or indirectly, or there may be between approximately 20 and 25 such linking nucleotides attaching a template primer to a substrate, directly or indirectly, or there may be between approximately 25 and 30 such linking nucleotides attaching a template primer to a substrate, directly or indirectly, or there may be between approximately 15 and 30 such linking nucleotides attaching a template primer to a substrate, directly or indirectly, or there may be approximately 5, 10, 15, 20, 25, 30, or more such linking nucleotides attaching a template primer to a substrate, directly or indirectly, where approximately means within 10% of the numbers indicated thereafter.

[0106] Non-exclusive examples of a suitable substrate may include epoxy siloxane, glass and modified or functionalized glass, polyhedral silsequioxanes and derivatives thereof, plastics

(including acrylics, polystyrene and copolymers of styrene and other materials, polypropylene, polyethylene, polybutylene, polyurethanes, polytetrafluoroethylene (such as TEFLON® FROM Chemours), cyclic olefins/cyclo-olefin polymers (such as ZEONOR® from Zeon), polyimides, etc.), nylon, ceramics/ceramic oxides, silica, fused silica, or silica-based materials, aluminum silicate, silicon and modified silicon (e.g., boron doped p+ silicon), silicon nitride, silicon oxide, tantalum pentoxide or other tantalum oxide(s), hafnium oxide, carbon, metals, inorganic glass, or the like. The substrate may also be glass or silicon or a silicon-based polymer such as polyhedral silsesquioxane material, optionally with a coating layer of tantalum oxide or another ceramic oxide at the surface. A substrate may also include a conductive channel as described in more detail below.

[0107] In an example, a solid support may include a conductive channel as further explained below. In an example, a conductive channel may detect a charge tag during incorporation of a charge-tagged nucleotide into a polynucleotide strand, according to methodology further explained below. In an example, a tether connecting a polymerase to a solid support, such as a conductive channel, may include an acceptor region and an attachment between a charge tag and a charge-tagged nucleotide may include a specificity region, such that the acceptor region and specificity region may bond to one another during incorporation of a charge-tagged nucleotide as further explained below. In an example, presence of an acceptor region and specificity region that bonds thereto may promote association of a charge tag with a conductive channel during incorporation of the charge tag to a hybridized polynucleotide and thereby enhance detection of the charge tag by a conductive channel.

[0108] As also further disclosed below, in a given solution a charge tag of a charge-tagged nucleotide may have a given Debye length. A Debye length is a measure of a charge carrier's net electrostatic effect in a solution and how far its electrostatic effect persists. Calculation of a Debye length of a charge tag of a charge-tagged nucleotide may be calculated according to an equation taking into consideration features of the charge tag and the solution, as more fully described below. A Debye length may determine how proximal a charge tag may be to a conductive channel in order for the conductive channel to detect the charge tag during incorporation of the charge tag into the hybridized polynucleotide. In an example, a Debye length may be between approximately 0.5 and approximately 10 nm, approximately 0.5 and approximately 1 nm, approximately 1 and approximately 1.5 nm, approximately 1.5 and approximately 2 nm, approximately 2 and approximately 2.5 nm, approximately 2.5 and

approximately 3 nm, approximately 3 and approximately 3.5 nm, approximately 3.5 and approximately 4 nm, approximately 4 and approximately 4.5 nm, approximately 4.5 and approximately 5 nm, approximately 5 and approximately 5.5 nm, approximately 5.5 and approximately 6 nm, approximately 6 and approximately 6.5 nm, approximately 6.5 and approximately 7 nm, approximately 7.5 and approximately 8 nm, approximately 8 and approximately 8.5 nm, approximately 8.5 and approximately 9 nm, approximately 9 and approximately 9.5 nm, approximately 9.5 and approximately 10 nm, approximately 1 and approximately 2 nm, approximately 2 and approximately 3 nm, approximately 3 and approximately 4 nm, approximately 4 and approximately 5 nm, approximately 5 and approximately 6 nm, approximately 6 and approximately 7 nm, approximately 7 and approximately 8 nm, approximately 8 and approximately 9 nm, approximately 9 and approximately 10 nm, approximately 0.5 and approximately 5 nm, approximately 5 and approximately 10 nm, less than approximately 0.5 nm, or more than approximately 10 nm, where approximately means 10% more or less than the following numbers.

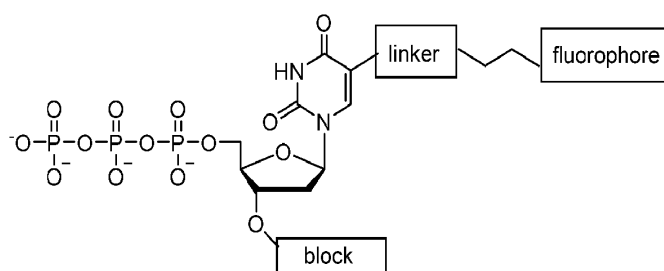
[0109] In an example, a first polymerase polymerizes incorporation of a charge-tagged nucleotide into a hybridized polynucleotide and a second polymerase polymerizes the incorporation of an FFN into the polymerase. In an example, the first polymerase and the second polymerase differ from each other. In another example, the first polymerase and the second polymerase may be the same as each other. In some examples, the first polymerase and the second polymerase may both be present simultaneously; whereas in other examples incorporation of a charge-tagged nucleotide by a first polymerase may occur in the presence of the first polymerase and absence of the second polymerase and the FFN may subsequently be incorporated in the presence of the second polymerase, whether or not the first polymerase is also present. In some examples, a first polymerase is washed out after incorporation of a charge-tagged nucleotide before incubation with the FFN, which occurs with incubation with the second polymerase. In an example, a first polymerase may incorporate a charge-tagged nucleotide with a given kinetics or under certain conditions such as pH, ionicity, or other parameter more preferable than the second polymerase may, and/or the second polymerase may incorporate the FFN with a given kinetics or under certain conditions such as pH, ionicity, or other parameter more preferable than the first polymerase may.

[0110] Any of a variety of suitable polymerases can be used in a method or composition set forth herein including, for example, protein-based enzymes isolated from biological systems

and functional variants thereof. Reference to a particular polymerase, such as those exemplified below, will be understood to include functional variants thereof unless indicated otherwise. A particularly useful function of a polymerase is to catalyze the polymerization of a nucleic acid strand using an existing nucleic acid as a template. Other functions that are useful are described elsewhere herein. Examples of useful polymerases include DNA polymerases and RNA polymerases, functional fragments thereof, and recombinant fusion peptides including them. Example DNA polymerases include those that have been classified by structural homology into families identified as A, B, C, D, X, Y, and RT. DNA Polymerases in Family A include, for example, T7 DNA polymerase, eukaryotic mitochondrial DNA Polymerase gamma., E. coli DNA Pol I (including Klenow fragment), *Thermus aquaticus* Pol I, and *Bacillus stearothermophilus* Pol I. DNA Polymerases in Family B include, for example, eukaryotic DNA polymerases α , β , and γ ; DNA polymerase C; T4 DNA polymerase, Phi29 DNA polymerase, *Thermococcus* sp. 9⁰N-7 archaeon polymerase (also known as 9⁰NTM) and variants thereof such as examples disclosed in U.S. Patent Application Publication No. 2016/0032377 A1, and RB69 bacteriophage DNA polymerase. Family C includes, for example, the E. coli DNA Polymerase III alpha subunit. Family D includes, for example, polymerases derived from the Euryarchaeota subdomain of Archaea. DNA Polymerases in Family X include, for example, eukaryotic polymerases Pol beta, Pol sigma, Pol lambda, and Pol mu, and *S. cerevisiae* Pol4. DNA Polymerases in Family Y include, for example, Pol eta, Pol iota, Pol kappa, E. coli Pol IV (DINB) and E. coli Pol V (UmuD'2C). The RT (reverse transcriptase) family of DNA polymerases includes, for example, retrovirus reverse transcriptases and eukaryotic telomerases. Example RNA polymerases include, but are not limited to, viral RNA polymerases such as T7 RNA polymerase; Eukaryotic RNA polymerases such as RNA polymerase I, RNA polymerase II, RNA polymerase III, RNA polymerase IV, and RNA polymerase V; and Archaea RNA polymerase. Any other suitable polymerase, including without limitation any disclosed in, for example, U.S Patent No. 8,460,910, are also included among polymerases as referred to herein, as are any other functional polymerases including those having sequences modified by comparison to any of the above mentioned polymerase enzymes, which are provided merely as a listing of non-limiting examples.

[0111] Returning to FIG. 1, following incorporation of the FFN, charge-tagged nucleotides, FFN, polymerase, etc., may be washed from the hybridized polynucleotide. In an example,

protein is degraded so as to denature or otherwise fully deactivate any residual polymerase not removed by a conventional wash step. Thereafter, FFN may be detected by known methodology. For example, FFN may be detected on surface using known optical fluorescence detection methodology, where fluorescence is elicited on and detected from surface where hybridized polynucleotide remains hybridized to template. Because FFN incorporation can only occur according to the disclosed method after charge-tagged nucleotide has been incorporated, measuring FFN incorporation may serve as a proxy for charge-tagged nucleotide incorporation. In another example, hybridized polynucleotide may be dehybridized from template and eluted in solution, with fluorescence detected in said elution solution rather than on surface. Any of various known methods for measuring fluorophores commonly attached to nucleotides for use in the disclosed method may be employed, on surface, in solution, or otherwise. In some examples, where a polymerase or polynucleotide are attached to a surface by a linker, the linker could be cleavable, such as a protease cleavage site (if the linker were a peptide linker) or other chemical moiety capable of being disrupted for release of the polymerase or polynucleotide from the surface if desired. [0112] Detection can be carried out by any suitable method, including fluorescence spectroscopy or by other optical means. The FFN label may be a fluorophore, which, after absorption of energy, emits radiation at a defined wavelength. Many suitable fluorescent labels are known. For example, Welch et al. (Chem. Eur. J. 5(3): 951-960, 1999) discloses dansyl-functionalised fluorescent moieties that can be used in the present invention. Zhu et al. (Cytometry 28:206-211, 1997) describes the use of the fluorescent labels Cy3 and Cy5, which can also be used according to aspects of the present disclosure. Labels suitable for use are also disclosed in Prober et al. (Science 238:336-341, 1987); Connell et al. (BioTechniques 5(4):342-384, 1987), Ansorge et al. (Nucl. Acids Res. 15(11):4593-4602, 1987) and Smith et al. (Nature 321:674, 1986). Other commercially available fluorescent labels include, but are not limited to, fluorescein, rhodamine (including TMR, Texas red and Rox), alexa, bodipy, acridine, coumarin, pyrene, benzantracene and the cyanins. Any suitable modification of any of the foregoing may be adopted for use in and employed in accordance with the method as disclosed herein. An FFN may include such a tag attached, for example. Commercially available fluorescently tagged nucleotides may be used in accordance with the present disclosure. A non-limiting, generalized example of an FFN may be depicted as follows:



[0113] In this example, a fluorophore is attached to a base of a nucleotide by a linker sequence. Such linker may include various chemical attachment moieties for attaching a fluorophore to a nucleotide. In other examples, a fluorophore may be attached to a different portion of a nucleotide, or a different base of a nucleotide. In the non-limiting example depicted, a reversible blocker is indicated on the 3-prime carbon, though such a blocker is not required or present in all examples included in the present disclosure.

[0114] In an example, incorporation of a charge-tagged nucleotide into a hybridized polynucleotide may be stopped by modification of the buffer or other polymerase reaction conditions (e.g., chelation of metal or other ions of solution components involved in nucleotide base pairing and/or polymerization by the first polymerase). In different samples, incorporation of the charge-tagged nucleotide may be stopped at various times after cessation of incorporation followed by incorporation of FFN under uniform conditions across all samples. By comparing the amount of fluorescence incorporated into the polynucleotide in different samples, incorporation of charge-tagged nucleotide in different samples can be comparatively determined. In other examples, rather than duration of polymerization by the first polymerase, different first polymerases may be compared to each other, different substrates, different charge tagged nucleotides, different surface chemistries of a surface to which a polymerase, template, and/or hybridized polynucleotide is attached, different tethers between a polymerase and a substrate, different attachments between a charge tag and nucleotide, different solutions with different buffers, pHs, and/or concentrations, and accordingly different Debye lengths for a given charge tag, or any other variable, may be modified across samples and effects on charge tag incorporation compared by comparing amounts of FFN (e.g., via fluorescence) incorporation between samples. In an example, charge tag detection by a conductive channel is not performed or is not required in order for such determinations to be made.

[0115] A nucleotide, template, or hybridized polynucleotide need not be a

deoxyribonucleotide. For example, a nucleotide, template, or hybridized polynucleotide may be a ribonucleotide. Similarly, a polymerase need not be a DNA polymerase and may instead be an RNA polymerase. A polymerase may also be a reverse transcriptase. In such examples, a charge-tagged nucleotide and FFN may be selected so as to appropriately complement a template nucleotide so as to be incorporated into the hybridized polymerase by the polymerase.

[0116] In an example, the hybridized polynucleotide may become dehybridized from and rehybridized to a template. Reference to a hybridized polynucleotide as such is in reference to hybridization during polymerized elongation by a polymerase as it incorporates a charge-tagged nucleotide or FFN. Dehybridization of the hybridized polynucleotide between such incorporations is included within aspects of the method as disclosed herein.

[0117] Examples of the present disclosure also provide compositions and methods for nucleotide incorporation events detected in nucleic acid sequencing procedures. There is a need for detection systems which provide a benefit of differential recognition of nucleotides on the basis of differences in charges, such as to permit long sequencing reads in high-throughput manner. Examples set forth herein may satisfy this need and provide other advantages as well. Charge-tagged nucleotides as described in more detail below are included as components and are equally suitable and intended for use in all of the foregoing examples as well.

[0118] As disclosed herein, an expensive and light-sensitive fluorescent label on a nucleotide with a different label for use with a different detection system. Detection of a conventional fluorescent label may involve expensive hardware such as lasers and detection optics which increases the size of a detection instrument. In addition, more powerful software is used to decode the multitude of information being generated. Importantly, as disclosed herein, expensive fluorophores are not needed. By replacing the fluorescent label with a charge label, the charge can be detected by a conductive channel which monitors the current in the system. This allows “real-time” sequencing to be performed and has the potential of achieving a faster turn-around time by reducing the cycle time of each nucleotide incorporation.

[0119] By enabling “real-time” sequencing, in one example the blocking group at the 3'-OH is not involved. This lowers the costs of the modified nucleotides as fewer synthetic steps are involved. An additional benefit is that polymerases are better suited to incorporating

nucleotides with 3' OH, that are closer to the native system, compared to a chemically modified bulky 3' protecting group.

[0120] A conductive channel for detecting a modified nucleotide including a charge may be responsive to a surrounding electric field. This field is modulated by positioning a modified nucleotide with a charge close proximity to a surface of the conductive channel. Close proximity of the charge tags to the surface may be important in some cases, such as if salt or other ions in the solution may screen a charge from detection by a conductive channel. A characteristic screening length is referred to as a Debye length, beyond which a conductive channel may be unable to detect charge.

[0121] A charge included in a modified nucleotide may be anywhere from between -200e to +200e, which may be in excess of 160 Angstroms when fully stretched linearly, whereas a Debye length around a conductive channel may be about 1 nm. Thus, structuring of a charge-carrying modification of a nucleotide to promote detection thereof by a conductive channel may be desirable.

[0122] Terms used herein will be understood to take on their ordinary meaning unless specified otherwise. Examples of several terms used herein and their definitions are set forth below.

[0123] As used herein, the term "array" refers to a population of conductive channels or molecules that are attached to one or more solid-phase substrates such that the conductive channels or molecules can be differentiated from each other according to their relative location. An array can include different molecules that are each located at a different addressable location (e.g. at different conductive channels) on a solid-phase substrate. Alternatively, an array can include separate solid-phase substrates each bearing a different molecule, wherein the different probe molecules can be identified according to the locations of the solid-phase substrates on a surface to which the solid-phase substrates are attached or according to the locations of the solid-phase substrates in a liquid such as a fluid stream. Molecules of the array can be nucleic acid primers, nucleic acid probes, nucleic acid templates or nucleic acid enzymes such as polymerases and exonucleases.

[0124] As used herein, the term "attached" refers to the state of two things being joined, fastened, adhered, connected or bound to each other. For example, a reaction component, such as a polymerase, can be attached to a solid phase component, such as a conductive channel, by a covalent or non-covalent bond. A covalent bond is characterized by the sharing

of pairs of electrons between atoms. A non-covalent bond is a chemical bond that does not involve the sharing of pairs of electrons and can include, for example, hydrogen bonds, ionic bonds, van der Waals forces, hydrophilic interactions and hydrophobic interactions.

[0125] As used herein, the term “electrically conductive channel” is intended to mean a portion of a detection device that translates perturbations at its surface or in its surrounding electrical field into an electrical signal. The conductive channel may be an electrically conductive channel. For example, as shown in FIG. 3, an electrically conductive channel 5 can translate the arrival or departure of a reaction component (e.g., the labeled nucleotide) into an electrical signal. In the examples disclosed herein, the electrically conductive channel 5 can also translate interactions between two reaction components (the template nucleic acid and a nucleotide of the labeled nucleotide) into a detectable signal through its interaction with the redox-active charge tag of the labeled nucleotide.

[0126] The electrically conductive channel 5 may be the channel of a conductive channel 2. The conductive channel 2 may include source and drain terminals S, D and the channel 5 connecting the terminals S, D. The channel may have any suitable geometries – e.g., tube, wire, plate, etc.

[0127] As used herein, the term “conductive channel” is intended to mean a detection device that translates perturbations at its surface or in its surrounding electrical field into an electrical signal. For example, a conductive channel can translate the arrival or departure of a reaction component into an electrical signal. A conductive channel can also translate interactions between two reaction components, or conformational changes in a single reaction component, into an electrical signal. An example conductive channel is a field effect transistor (FET) such as a carbon nanotube (CNT), single-walled carbon nanotube (SWNT) based FET, silicon nanowire (SiNW) FET, graphene nanoribbon FET (and related nanoribbon FETs fabricated from 2D materials such as MoS₂, silicene, etc.), tunnel FET (TFET), and steep subthreshold slope devices (see, for example, Swaminathan et al., *Proceedings of the 51st Annual Design Automation Conference on Design Automation Conference*, pg 1-6, ISBN: 978-1-4503-2730-5 (2014) and Ionescu et al., *Nature* 479, 329–337 (2011)). Examples of FET and SWNT conductive channels that can be used in the methods and apparatus of the present disclosure are set forth in US Pat. App. Pub. No. 2013/0078622 A1.

[0128] The terminals S, D may be any suitable electrically conductive material, including an

electrical conductor or a semiconductor. Examples of suitable source and drain materials include cobalt, cobalt silicide, nickel, nickel silicide, aluminum, tungsten, copper, titanium, molybdenum, indium tin oxide (ITO), indium zinc oxide, gold, platinum, carbon, etc.

[0129] The conductive channel 5 may include any conductive or semi-conductive material that can oxidize or reduce the redox-active charge tag. The material may comprise an organic material, an inorganic material, or both. Some examples of suitable channel materials include silicon, carbon (e.g., glassy carbon, graphene, etc.), polymers, such as conductive polymers (e.g., polypyrrole, polyaniline, polythiophene, poly(3,4-ethylenedioxythiophene) doped with poly(4-styrenesulfonate) (PEDOT-PSS), etc.), metals, biomolecules, etc.

[0130] In some examples, the conductive channel 5 may also be a nanostructure that has at least one dimension on the nanoscale (ranging from 1 nm to less than 1 μm). In one example, this dimension refers to the largest dimension. As examples, the electrically conductive channel 5 may be a semi-conducting nanostructure, a graphene nanostructure, a metallic nanostructure, and a conducting polymer nanostructure. The nanostructure may be a multi- or single-walled nanotube, a nanowire, a nanoribbon, etc.

[0131] As used herein, the term “different”, when used in reference to nucleic acids, means that the nucleic acids have nucleotide sequences that are not the same as each other. Two or more different nucleic acids can have nucleotide sequences that are different along their entire length. Alternatively, two or more different nucleic acids can have nucleotide sequences that are different along a substantial portion of their length. For example, two or more different nucleic acids can have target nucleotide sequence portions that are different for the two or more molecules while also having a universal sequence portion that is the same on the two or more molecules. The term “different” can be similarly applied to other molecules, such as polymerases and nucleic acid enzymes.

[0132] As used herein, the term “each,” when used in reference to a collection of items, is intended to identify an individual item in the collection but does not necessarily refer to every item in the collection. Exceptions can occur if explicit disclosure or context clearly dictates otherwise.

[0133] As used herein, the term “label,” when used in reference to a reaction component, is intended to mean a detectable reaction component or detectable moiety of a reaction component. A useful label is a charge label (also called a charge tag) that can be detected by a conductive channel. A label can be intrinsic to a reaction component that is to be detected

(e.g. a charged amino acid of a polymerase) or the label can be extrinsic to the reaction component (e.g. a non-naturally occurring modification of an amino acid). In some examples a label can include multiple moieties having separate functions. For example a label can include a linker component (such as a nucleic acid) and a charge tag component.

[0134] As used herein, the term “non-natural,” when used in reference to a moiety of a molecule, is intended to refer to a moiety that is not found attached to the molecule in its natural milieu or in a biological system unperturbed by human, technical intervention. In some examples, non-natural moieties are synthetic modifications of molecules that render the molecules structurally or chemically distinct from the unmodified molecule or from molecules having natural modifications. As used herein, the term “non-natural,” when used in reference to an analog used for a process, is intended to mean an analog that is not found in the natural milieu where the process occurs. In some examples, non-natural analogs are synthetic analogs that are structurally or chemically distinct from other types of molecules in the class to which the analog belongs.

[0135] As used herein, the term “nucleic acid” is intended to be consistent with its use in the art and includes naturally occurring nucleic acids or functional analogs thereof. Particularly useful functional analogs are capable of hybridizing to a nucleic acid in a sequence specific fashion or capable of being used as a template for replication of a particular nucleotide sequence. Naturally occurring nucleic acids generally have a backbone containing phosphodiester bonds. An analog structure can have an alternate backbone linkage including any of a variety of those known in the art such as peptide nucleic acid (PNA) or locked nucleic acid (LNA). Naturally occurring nucleic acids generally have a deoxyribose sugar (e.g. found in deoxyribonucleic acid (DNA)) or a ribose sugar (e.g. found in ribonucleic acid (RNA)).

[0136] A nucleic acid can contain any of a variety of analogs of these sugar moieties that are known in the art. A nucleic acid can include native or non-native bases. In this regard, a native deoxyribonucleic acid can have one or more bases selected from the group consisting of adenine, thymine, cytosine, and guanine; and a ribonucleic acid can have one or more bases selected from the group consisting of uracil, adenine, cytosine and guanine. Useful non-native bases that can be included in a nucleic acid are known in the art.

[0137] As used herein, the term “nucleotide” is intended to include natural nucleotides, analogs thereof, ribonucleotides, deoxyribonucleotides, dideoxyribonucleotides and other

molecules known as nucleotides. The term can be used to refer to a monomeric unit that is present in a polymer, for example to identify a subunit present in a DNA or RNA strand. The term can also be used to refer to a molecule that is not necessarily present in a polymer, for example, a molecule that is capable of being incorporated into a polynucleotide in a template dependent manner by a polymerase. The term can refer to a nucleoside unit having, for example, 0, 1, 2, 3 or more phosphates on the 5' carbon. For example, tetraphosphate nucleotides, pentaphosphate nucleotides, and hexaphosphate nucleotides can be particularly useful, as can nucleotides with more than 6 phosphates, such as 7, 8, 9, 10, or more phosphates, on the 5' carbon. Example natural nucleotides include, without limitation, ATP, UTP, CTP, and GTP (collectively NTP), and ADP, UDP, CDP, and GDP (collectively NDP), or AMP, UMP, CMP, or GMP (collectively NMP), or dATP, dTTP, dCTP, and dGTP (collectively dNTP), and dADP, dTDP, dCDP, and dGDP (collectively dNDP), and dAMP, dTMP, dCMP, and dGMP (dNMP). Example nucleotides may include, without exception, any NMP, dNMP, NDP, dNDP, NTP, dNTP, and other NXP and dNXP where X represents a number from 2 to 10 (collectively NPP).

[0138] Non-natural nucleotides also referred to herein as nucleotide analogs, include those that are not present in a natural biological system or not substantially incorporated into polynucleotides by a polymerase in its natural milieu, for example, in a non-recombinant cell that expresses the polymerase. Particularly useful non-natural nucleotides include those that are incorporated into a polynucleotide strand by a polymerase at a rate that is substantially faster or slower than the rate at which another nucleotide, such as a natural nucleotide that base-pairs with the same Watson-Crick complementary base, is incorporated into the strand by the polymerase. For example, a non-natural nucleotide may be incorporated at a rate that is at least 2 fold different – e.g., at least 5 fold different, 10 fold different, 25 fold different, 50 fold different, 100 fold different, 1000 fold different, 10000 fold different, or more, when compared to the incorporation rate of a natural nucleotide. A non-natural nucleotide can be capable of being further extended after being incorporated into a polynucleotide. Examples include, nucleotide analogs having a 3' hydroxyl or nucleotide analogs having a reversible terminator moiety at the 3' position that can be removed to allow further extension of a polynucleotide that has incorporated the nucleotide analog. Examples of reversible terminator moieties that can be used are described, for example, in U.S. Pat nos. 7,427,673; 7,414,116; and 7,057,026 and PCT publications WO 91/06678 and WO 07/123744. It will be understood

that in some examples a nucleotide analog having a 3' terminator moiety or lacking a 3' hydroxyl (such as a dideoxynucleotide analog) can be used under conditions where the polynucleotide that has incorporated the nucleotide analog is not further extended. In some examples, nucleotide(s) may not include a reversible terminator moiety, or the nucleotide(s) will not include a non-reversible terminator moiety or the nucleotide(s) will not include any terminator moiety at all. Nucleotide analogs with modifications at the 5' position are also useful.

[0139] As used herein, the term "protection moiety" is intended to mean a compound or portion thereof that is attached to a reaction component to reduce, minimize, or in some instances prevent the reaction component from undergoing a particular reaction. For example, a nucleic acid molecule can be bound to a nucleic acid enzyme such that the nucleic acid molecule reduces, minimizes, or in some instances prevents the nucleic acid enzyme from degradation or modification by a treatment that may otherwise cause degradation or modification of the enzyme. An antibody can also serve to bind a reaction component to protect the reaction component from degradation, inactivation or other reaction.

[0140] As used herein, the term "reaction component" is intended to mean a molecule that takes part in a reaction. Examples include, reactants that are consumed in a reaction, products that are created by a reaction, catalysts such as enzymes that facilitate a reaction, solvents, salts, buffers and other molecules.

[0141] As used herein, the term "repellant moiety" is intended to mean a molecule or portion thereof that will occupy a space to prevent or inhibit occupancy of another molecule at the space or to inhibit juxtaposition of another molecule near the space. A repellant moiety can act via steric exclusion, charge repulsion, hydrophobic-hydrophilic repulsion or other forces.

[0142] As used herein, the term "terminator moiety," when used in reference to a nucleotide, means a part of the nucleotide that inhibits or prevents the nucleotide from forming a covalent linkage to a second nucleotide. For example, in the case of nucleotides having a pentose moiety, a terminator moiety can reduce, minimize, or in some instances prevent formation of a phosphodiester bond between the 3' oxygen of the nucleotide and the 5' phosphate of the second nucleotide. The terminator moiety can be part of a nucleotide that is a monomer unit present in a nucleic acid polymer or the terminator moiety can be a part of a free nucleotide (e.g. a nucleotide triphosphate). The terminator moiety that is part of a nucleotide can be reversible, such that the terminator moiety can be modified to render the

nucleotide capable of forming a covalent linkage to a second nucleotide. In particular examples, a terminator moiety, such as a reversible terminator moiety, can be attached to the 3' position or 2' position of a pentose moiety of a nucleotide analog.

[0143] The examples set forth below and recited in the claims can be understood in view of the above definitions.

[0144] The present disclosure provides compositions useful for, among other things, nucleotide incorporation events detected in nucleic acid sequencing procedures, methods of making such compositions, and methods of using them in such procedures. The compositions and methods set forth herein are particularly useful, for example, in single molecule nucleic acid sequencing reactions, such as sequencing by synthesis. However, it will be appreciated that the compositions and methods set forth herein can be used for any other suitable detection schemes, including, but not limited to single molecule detection. Apparatuses and methods for nucleic acid sequencing in which compositions as disclosed herein may be used are disclosed in, for example, U.S. Patent Application Number 14/798,762.

[0145] For example, a method of nucleic acid sequencing can include the processes of (a) providing a polymerase tethered to a solid support conductive channel; (b) providing one or more labeled nucleotides, whereby the presence of the label can be detected by the conductive channel when the label is in proximity to the conductive channel; and (c) detecting incorporation of the labeled nucleotide into a nascent strand complementary to a template nucleic acid.

[0146] In some examples of a method of nucleic acid sequencing, the polymerase is held in proximity of less than 10, 9, 8, 7, 6, 5, 4, 3, 2, or 1 nm to the conductive channel.

[0147] In some examples, a label or a portion thereof (e.g., a charge tag) may be cleaved from a nucleotide after incorporation, for example, by a polymerase.

[0148] As provided herein, the one or more labeled nucleotides may include a plurality of charge tags. For example, one or more labeled nucleotides can comprise a unique charge tag for each type of nucleotide. For example, nucleotides bearing charge tags may be used in synthesizing a strand of DNA by a polymerase according to a template sequence, which template sequence may include a string of nucleotides, including the bases adenine, thymine, guanine, and cytosine, for example. Nucleotides bearing charge tags as disclosed herein may be incorporated into a string of nucleotides complementary to the template sequence by a polymerase enzyme. As disclosed herein, as a nucleotide bearing a charge tag is so

incorporated, a conductive channel may detect a charge of a given valence (meaning positivity or negativity) and magnitude specifically and differentially associated with each species of nucleotide, permitting recordation of an identity of successive nucleotides incorporated into a growing strand and thereby a sequence of nucleotides present in a template strand to which the growing strand is complementary. The charge tag can be a negative charge tag or a positive charge tag, and can have a charge anywhere from $-200e$ to $+200e$, such as from $-175e$ to $+175e$, or from $-150e$ to $+150e$, or from $-125e$ to $+125e$, or from $-100e$ to $+100e$, or from $-75e$ to $+75e$, or from $-50e$ to $+50e$.

[0149] A conductive channel used in a method of nucleic acid sequencing can include a nanowire FET. Optionally, a conductive channel may include a carbon nanotube. A conductive channel can be part of an array of conductive channels. A detecting process can include detecting a plurality of incorporation events in succession.

[0150] Compositions, apparatus, and methods set forth herein can provide long nucleic acid sequencing reads; fast reads; high throughput capability for sequencing; and a scalable platform for sequencing. In some examples, any compromises in single read accuracy can be mitigated by performing multiple overlapping reads due to the ability of the methods and apparatus set forth herein to provide throughput in the number of reads performed in parallel.

[0151] An example conductive channel is shown in FIG. 3. Here a polymerase 1 creates a reaction site where nucleotides can be incorporated into a primed DNA template 4. The polymerase 1 is attached to a nanowire FET 2 via a tether 3. The apparatus provides single molecule sensitivity. Changes in charge distribution at the reaction site (e.g. polymerase conformation changes, nucleotide incorporation, arrival or departure of charged tags, changes in proximity of the polymerase to the conductive channel etc.) transmit to the gate and can be detected.

[0152] In particular examples, an apparatus or method of the present disclosure may use deeply scaled FinFET transistors as single-molecule conductive channels. FinFET conductive channels benefit from technology already under development by leading edge semiconductor manufacturers. Furthermore, previously published components can be used, including but not limited to (1) those used for immobilization of lysozyme on CNT to observe enzyme processivity in real time as described in Choi et al, *Science*, 335, 319 (2012), (2) those used to immobilize the Pol 1 Klenow fragment on CNT and observe DNA processivity in real time as described in Olsen et al, *J. Amer. Chem. Soc.*, 135, 7885 (2013), (3) those used to elucidate a

transduction mechanism as moving charged residues due to protein allosteric motion as described in Choi et al, *NanoLett* 13, 625 (2013). The present methods can also employ the apparatus, components of the apparatus, and methods set forth in US Pat. App. Pub. No. 2013/0078622 A1.

[0153] Some examples of a labeled nucleotide may also include a specificity region. Thus, a labeled nucleotide may include a nucleotide, a linking molecule or linker attached to a phosphate group of the nucleotide, and a charge tag attached to the linker. A linking molecule or linker may comprise a specificity region that may hybridize to an acceptor region on a tether bound to a conductive channel. As examples, a specificity region may be any nucleotide sequence or peptide that is capable of temporarily attaching or bonding to an acceptor region on a tether. For example, a specificity region may include a sequence of nucleotides and an acceptor region may include a sequence of nucleotides such that pair bonding forms between nucleotides in a sequence of a specificity region and an acceptor region. Pair bonding in this instance refers to standard pair bonding between nucleotides, such as between a G and a C residue, or between an A and a T or U residue.

[0154] A specificity region may include a sequence of nucleotides and an acceptor region a correspondingly complementary sequence of nucleotides. In an example, when a polymerase accepts a nucleotide for incorporation into a growing polynucleotide strand, complementary to a template polynucleotide, a specificity region and an acceptor region may be brought into sufficient proximity to each other for pair bonding to form therebetween. Such pair bonding between a specificity region and an acceptor region may promote sufficient proximity between a charged tag and a conductive channel, promoting detection of the charge tag by the conductive channel during incorporation of the nucleotide.

[0155] In an example, a specificity region may include a nucleotide sequence including from about one nucleotide to about six nucleotides. In another example, a specificity region may further include inosine(s) flanking both sides of a nucleotide sequence. In some examples, a specificity region is included in part of a charge tag. For example, a specificity region may consist of segments or portions of a sequence of nucleotides or amino acids that are separated from each other along a linear sequence, such as by portions of a charge tag, wherein bonding to an acceptor region may induce the separate regions of the specificity region to come into proximity with each other while permitting adoption of a given three-dimensional structure by a charge tag.

[0156] In an example of a labeled nucleotide associated with a tether, specific binding affinity between a labeled nucleotide and a tether is combined with weak affinity produced by non-specific binding interactions. A labeled nucleotide may include a specificity region which is complementary to a portion of a tether. Specific binding between these regions can result from standard Watson-Crick base pairing or other non-covalent bonding. A specificity region, in this example, can also include inosines (I) flanking a nucleotide sequence. Inosines are universal bases, and thus can pair with all four native nucleotides of DNA. Additional binding interactions can result from interactions of the universal bases (e.g., inosine I) with native nucleotides on the tether. Thus, when a labeled nucleotide is bound to polymerase during incorporation, synergistic binding may occur between a specificity region of the labeled nucleotide and the acceptor region of the tether, which may greatly increase the stability of the interaction between the labeled nucleotide and the tether.

[0157] An interaction between a labeled nucleotide and polymerase, or polymerase and a tether, may cause the charge tag to come within a sensing zone of a conductive channel. Such interaction(s) may also aid in maintaining a charge tag within a sensing zone for a time sufficient for efficient and complete charge detection. Such time may be up to tens of milliseconds. Such relatively long interaction is unlike that for other labeled nucleotides present in the solution, which in theory may diffuse and briefly touch or approach the conductive channel. Such brief interaction may not be long enough for sufficient charge detection to take place, and thus in such instances, a charge tag is not detected by the conductive channel.

[0158] As disclosed herein, a charge tag may include polypeptides, oligonucleotides, oligomeric peptide nucleic acids, or any combination of two or more of the foregoing. In some examples, a charge tag may include a plurality of elements selected from amino acids, nucleotides, and linkers. Such molecules may adopt a three-dimensional structure to permit condensation of charges carried by aspects of the charge tag such that the total charge can be condensed into a smaller region. Such increased charge density may increase a charge detected by a conductive channel during incorporation of a nucleotide analog in a growing strand by a polymerase such that presence of a given species of nucleotide in such synthesis can be determined. A charge tag that adopts such a condensed conformation may minimize dispersal of its charge away from a conductive channel or over a large surface area of a conductive channel, or both. As a consequence, a conductive channel may be more likely to

detect a greater amount or proportion of charge of a charge tag.

[0159] Some examples disclosed herein exploit synergistic binding of a labeled nucleotide to a polymerase, alone or in combination with a tether, in order to bring and hold a charge tag in proximity of a sensing zone of a conductive channel. Stability of a complex formed with a tether can be relatively low such that a complex does not form for labeled nucleotides that are not also bound to a polymerase (i.e., labeled nucleotides that are free in solution may not substantially bind to a tether). In other words, the off rate of such a complex can be sufficiently high that a lifetime is short. However, when a stable association is formed between a labeled nucleotide and a polymerase, a local concentration of a linking molecule may increase around a tether, thus resulting in a high on rate. In this manner, an overall association time may be greatly increased in a polymerase-associated state compared to a non-associated state. Synergistic effect of the affinities of a labeled nucleotide for a polymerase, alone or in combination with a tether, may add up to allow substantial binding affinity overall. After cleaving by a polymerase, a synergistic effect is lost and a charge tag may also dissociate from the conductive channel.

[0160] Particular examples can exploit synergistic binding of a gamma-phosphate labeled nucleotide to a polymerase and to a tether. Stability of an oligonucleotide moiety:tether, or specificity region:acceptor region, complex can be relatively low such that the complex does not form for gamma-phosphate labeled nucleotide that are not also bound to polymerase, such that gamma-phosphate labeled nucleotides that are free in solution do not substantially bind to the tether. However, a synergistic effect of affinities of a nucleotide moiety for a polymerase and a specificity region, such as an oligonucleotide moiety, for an acceptor region of a tether may add up to allow substantial binding affinity overall. In some examples, a synergistic effect can exploit a combination of specific binding affinity between a nucleotide label and tether along with weak affinity produced by non-specific binding interactions. For example, as stated above, in some examples specific binding can result from standard Watson-Crick base pairing and non-specific binding interactions can result from interactions of promiscuous bases (e.g. inosine) with native nucleotides. Thus, when a gamma-phosphate labeled nucleotide is bound to polymerase during incorporation, synergistic binding may occur which may greatly increase stability of interaction between oligonucleotide moiety and tether. After the gamma phosphate is cleaved by the polymerase, the synergistic effect may be lost and the oligonucleotide moiety will dissociate from the

tether. Other types of nucleotide moiety:tether bonding, such as through non-covalent interactions between DNA, RNA, PNA, amino acids, or analogs or combinations thereof to contribute to such synergistic effect.

[0161] As shown in FIG. 4, a polymerase can be immobilized to a conductive channel such as a single walled carbon nanotube, silicon nanowire or FinFET. Immobilization can be via tethers that include DNA, RNA, PNA, amino acids, or analogs or combinations thereof. For convenience of demonstration FIG.4 shows four polymerases tethered to a conductive channel, each polymerase also being bound to a different gamma-phosphate labeled nucleotide type. As shown, nucleotides may have an oligonucleotide moiety attached to the gamma-phosphate. A beta- or gamma-phosphate-labeled nucleotide that is properly matched to a template strand of a target nucleic acid may be held in place by a polymerase that may also be bound to the template long enough to temporarily hybridize an oligonucleotide moiety or other specificity region to an acceptor region of a tether (e.g. via Watson-Crick base complementarity or other non-covalent bonding). The hybridization may cause a charge tag to perturb a field around a conductive channel which may produce a detectable signal due to a change in transistor current through the conductive channel. The diagram shows a charge tag entering a field that is within 1-2 nm of the conductive channel. The properly matched beta- or gamma-phosphate-labeled nucleotide may be incorporated into a nascent strand hybridized to the template nucleic acid. This may, in turn, break the bond between the beta phosphate and the newly incorporated nucleotide. As a result, the charge tag (whether attached at the beta- or gamma-position of the nucleotide) may be free to dissociate from the tether and diffuse away from the conductive channel, thereby returning the field around the conductive channel to its unperturbed state. The appearance and disappearance of signal as the field around the conductive channel is perturbed and returned to the unperturbed state, respectively, can be correlated with incorporation of a nucleotide into the nascent strand of the target nucleic acid.

[0162] The type of nucleotide that is incorporated into the nascent strand at each position of the template strand can be determined based on unique properties of labels incorporated into each type of nucleotide. For example, four types of dNTPs can be distinguished by the position where a specificity region hybridizes to an association region of a tether, the length of the specificity region and/or the presence of a charged moiety on the label, the valence of the charge, and the magnitude of the charge. For example, a given nucleotide may have a

charge of a given valence and magnitude which is not shared by other nucleotides, which have a charge with a different valence and/or magnitude. A conductive channel may be capable of detecting differences in valence and/or magnitude of a charge. During incorporation of a nucleotide with a charged tag into a nascent polynucleotide by a polymerase tethered to a conductive channel the conductive channel may detect the valence and/or magnitude of the tag of the nucleotide incorporated as the complement to a nucleotide of a template strand. When the polymerase moves on to incorporate the next species of nucleotide, in turn complementary to the next nucleotide of the template, the valence and/or magnitude of charge of such next species of nucleotide incorporated into the nascent strand may also be detected by the conductive channel. And so on as consecutive nucleotides with charge tags are incorporated into the nascent strand.

[0163] As successive charge tags are detected by the conductive channel, the differences in current flow through the conductive channel resulting from differences in charge tags may be recorded and stored such as in a computer-readable storage medium, which may be programmed so as to record a given, identified species of nucleotide for each incorporation polymerized by the polymerase as the growing nascent strand is synthesized on the basis of the valence and/or magnitude of charge detected by the conductive channel for each such incorporation.

[0164] FIG. 4 provides an example where four-state discrimination between bases G, A, C, and T is achieved using 2 charge tags and two tether hybridization positions. Specifically, dCTP is uniquely labeled with a negatively charged extrinsic moiety, dTTP is uniquely labeled with a positively charged extrinsic moiety, dATP and dGTP are distinguished from the other two nucleotide types based on absence of any extrinsic charge moiety, and dATP is distinguished from dGTP based on differential proximity of the oligonucleotide moieties to the conductive channel when they are hybridized to the tether.

[0165] It will be understood that different nucleotide types can be distinguished based on any of a variety of combinations of positive charge moieties, negative charge moieties and/or tether hybridization locations. Alternatively or additionally, charge moieties used to distinguish different types of nucleotides can differ in strengths of the charges, even if the charges have the same sign. An example configuration shown in FIG. 5 provides four-state discrimination between bases G, A, C, and T based on a single tether hybridization position and four different charge moieties. Specifically, in this non-limiting example, dGTP and

dCTP both contain negatively charged moieties that distinguish them from dATP and dTTP, and dGTP can be distinguished from dCTP due to charge that is distinguishably higher than the charge on dCTP. Similarly, dATP and dTTP can be distinguished from each other due to the higher positive charge on the dATP moiety compared to the dTTP moiety.

[0166] As noted previously herein, the precision of tag placement at specific hybridization positions along a tether can be enhanced through the use of a tether having ribonucleotides and a nucleotide label having 2'-O-Methyl (2'-O-Me) and 2'-Fluoro (2'F) modified RNA bases. Alternative configurations can use a tether that contains 2'-O-Me and 2'F modified ribonucleotides with label having ribonucleotides, or both the tether and label can include a mixture of native ribonucleotides and 2'-O-Me and 2'F modified ribonucleotides. Although it is possible to use a tether and/or oligonucleotide moiety that is primarily composed of RNA, it may be desirable to use a DNA-based or PNA-based or amino acid-based tether and/or oligonucleotide to avoid nuclease sensitivity that is associated with RNA. For example, a DNA-based or PNA-based tether or amino acid-based tether and/or oligonucleotide can include native ribonucleotides or non-native ribonucleotide analogs to achieve binding advantages set forth herein while reducing risk of unwanted nuclease digestion. In further examples, a tether can include one or more deoxyribonucleotides that are complementary to deoxyribonucleotides in a nucleotide label or alternatively the tether can include deoxyribonucleotides that are complementary to deoxyribonucleotides in a nucleotide label.

[0167] A tether that attaches a polymerase to a conductive channel can have different binding positions (e.g., acceptor regions) for different nucleotide sequences as set forth in several examples disclosed herein. Binding positions for two or more nucleotide sequences can overlap or they can be discrete with no overlap. For purposes of illustration, a tether sequence is depicted in FIG. 10 as a series of generic "N" nucleotides. Any of a variety of sequences can be used in accordance with rules of complementarity and desired hybridization strengths and specificities. Depending on the length of a tether, length of an acceptor region, and length of a specificity region, some, all, or no binding sites on a tether may overlap. In some aspects, the complementary bases are standard DNA bases, but any nucleotide analogs could be used (e.g., deoxyribonucleotide analogs may be used).

[0168] A tether-binding oligonucleotide moiety of a specificity region of a nucleotide analog can have a sequence of nucleotides that hybridizes specifically to a complementary sequence on a tether's acceptor region. In some examples a tether-binding oligonucleotide moiety can

also include promiscuous nucleotide positions that bind non-specifically to a tether. Such positions can provide a weak interaction between the tether-binding oligonucleotide moiety and tether that facilitates the formation of a specific hybrid structure. For example, as shown in FIG. 11, an oligonucleotide moiety can include several inosines (I) that are known to bind promiscuously, albeit weakly, with all four native nucleotides of DNA. A tether-binding oligonucleotide moiety (e.g., a specificity region) and tether (e.g., acceptor region) can form a weak complex via interactions between inosines in the tether-binding oligonucleotide moiety and native nucleotides in the tether. This can allow the specific portions of the sequence (e.g. indicated as ABC and its complement A'B'C' in the figure) to associate more rapidly than if required to diffuse absent formation of a weak complex. Furthermore, once a specific complex has formed inosines can provide further stability.

[0169] The non-limiting, example tether-binding oligonucleotide moieties in FIG. 11 include promiscuous nucleotide positions flanking both sides of a specific sequence. However, it will be understood that one or more promiscuous nucleotide positions can be located on only the 5' or 3' side of a specific sequence. Other examples of promiscuous nucleotide positions include those formed by degenerate oligonucleotide synthesis or those formed with other nucleotide analogs known in the art to hybridize promiscuously with 2 or more types of nucleotides.

[0170] Several examples set forth herein have exemplified the use of a plurality of different nucleotide analogs having oligonucleotide specificity regions of differing lengths. In such examples, different nucleotide analog types may be distinguishable based on different lengths of their specificity regions. Alternatively, different nucleotide analogs can have tether-binding oligonucleotide moieties of the same or similar lengths that may not permit of distinguishing one from another. However, each nucleotide analog can have a specificity sequence that binds to a different acceptor region of a tether compared to an acceptor region or regions where specificity regions of other nucleotide analogs bind. An example configuration is shown in FIG. 12 where binding of a polymerase to different nucleotide analogs places the polymerase in one of four distinguishable states. In the non-limiting example shown in FIG. 12, a tether-binding oligonucleotide moiety of an ATP analog binds to a location on the tether that is nearest to the attachment point of the tether to the polymerase, a tether-binding oligonucleotide moiety of a TTP analog binds to a location on the tether that is furthest from the attachment point of the tether to the polymerase, and a

tether-binding oligonucleotide moiety of GTP and CTP analogs bind to respectively distinct locations on the tether that are at intermediate distances from the binding sites for the tether-binding oligonucleotide moieties other two nucleotide analogs. Binding of different nucleotide analogs to the polymerase may position a polymerase at different distances from a conductive channel (e.g. causing different size loops to form in the tether as shown in the figure). In examples where one or more of the nucleotide analogs includes a charge tag or other detectable moiety (e.g. extending from an end of a tether-binding oligonucleotide moiety distal to the end that extends from the nucleotide to be incorporated into a nucleotide sequence by the polymerase), the binding between the tether-binding oligonucleotide moiety and tether may position the charge tag moiety at different distances from the conductive channel. In such cases, different types of nucleotide analogs can be distinguished at least in part based on differences in signals produced for the different distances of the detectable charge tag moieties from the conductive channel. For this illustrative example, the nucleotide analogs are identified as ATP, GTP, CTP and TTP, but any nucleotide analogs could be used (e.g., deoxyribonucleotide analogs may be used).

[0171] In other examples, such as illustrated in FIGs. 15A and 15B, a specificity region of a tagged nucleotide as disclosed herein may include polynucleotide sequences that each hybridize to a different section of an acceptor region of a tether. Between such sequences of the specificity region may be a span of nucleotides that do not hybridize to a portion of the acceptor region. The two sequences may therefore hybridize to the correspondingly complementary portions of the acceptor region of the tether and the intervening portion of the specificity region, with the intervening sequence free to hybridize elsewhere (such as two complementary portions of such intervening sequence of a specificity region hybridizing to each other to form a hairpin structure as shown in FIG. 15A) or free to hybridize or itself to remain unbound specifically (such as shown in FIG. 15B). In FIGs. 15A and 15B “Acceptor region” indicates an acceptor portion of a tether that hybridizes or otherwise transiently bonds to a specificity region of a tagged nucleotide. In some examples, such bonding may increase detection of a charged tag by a conductive channel (represented in FIGs. 15A and 15B by the wire to which the tether/acceptor region is attached).

[0172] As demonstrated by the example diagrammed in FIG. 6, a tether that attaches a polymerase to a conductive channel need not be capable of hybridizing to a charge tag or specificity sequence that may be present on an analog nucleotide. Rather, a conductive

channel can be functionalized by attachment of an acceptor region separate from a polymerase's tether, to which a specificity region of a nucleotide analog may bind. Discrimination of different nucleotides can be achieved based on valence of charge of a charge tag, strength of the charge, length of a specificity region:acceptor region binding complex, or proximity or location of an acceptor region:specificity region complex formation to or in relation to a conductive channel, or a combination thereof, whether the acceptor region is part of a polymerase tether or otherwise attached to the conductive channel.

[0173] An illustrative example of a nucleotide analog bearing a charge tag in accordance with the present disclosure is shown in FIG. 7. This is but one of many examples of a nucleotide analog as described and disclosed herein and is not limiting of the scope of the present disclosure. In this non-limiting example, a dT hexaphosphate is connected to a charge tag via a linker region comprising a specificity region. The linker in this non-limiting example includes covalent bonds formed by an azide-alkyne click reaction, though other chemistries may be employed instead, as further disclosed herein. For ease of reference, when describing portions of a nucleotide analog herein, the region towards the right of the molecule as illustrated in FIG. 7, will be referred to as the 3' end, according to a convention of referring to a free 3' hydroxyl group on the deoxyribose of the nucleotide. Correspondingly, the region towards the left of the molecule as illustrated in FIG. 7, where the charge tag is located in this example, will be referred to as the 5' end, as an extension of a phosphate group bound to the 5' carbon of the ribose of the nucleotide.

[0174] An examples of charge tags that can be useful in the apparatus and methods set forth herein is a phosphate moiety, for example, located at the 5' end of a nucleic acid moiety. This moiety, containing a phosphodiester group, can be readily added during available oligonucleotide synthesis protocols and may result in two negatively charged oxygens at the end of the oligonucleotide moiety as shown in FIG. 8. A polynucleotide chain or oligomer, by nature of its phosphate backbone, may also possess a negative charge, roughly proportional to the number of nucleotides in the oligonucleotide chain, and may be included in a charge tag. Chemical phosphorylation during oligonucleotide synthesis can be achieved by converting a DMT protecting group into a 5' phosphate group using 2-[2-(4,4'-Dimethoxytrityloxy)ethylsulfonyl]ethyl-(2-cyanoethyl)-(N,N-diisopropyl)-phosphoramidite (available from Glen Research, Sterling VA, catalog No. CPR 10-1900). A series of charge tags having different numbers of negative charges can be made using Tris-2,2,2-[3-(4,4'-

dimethoxytrityloxy)propyloxymethyl]ethyl-[(2-cyanoethyl)-(N,N-diisopropyl)]-phosphoramidite (available from Glen Research, Sterling VA, catalog No. 10-1922-xx), 1,3-bis-[5-(4,4'-dimethoxytrityloxy)pentylamido]propyl-2-[(2-cyanoethyl)-(N,N-diisopropyl)]-phosphoramidite (available from Glen Research, Sterling VA, catalog No. 10-1920-xx), 1-[5-(4,4'-dimethoxytrityloxy)pentylamido]-3-[5-fluorenomethoxycarbonyloxypentylamido]-propyl-2-[(2-cyanoethyl)-(N,N-diisopropyl)]-phosphoramidite (available from Glen Research, Sterling VA, catalog No. 10-1921-xx), or oligonucleotide dendrimers which contain various numbers of DMT (4,4'-dimethoxytrityl) or Fmoc (Fluorenylmethyloxycarbonyl) moieties, such as those available from Glen Research or such as those shown in FIGs. 9A and 9B (for a doubling branch) or FIG. 9C (for a trebling branch). A useful positively charged tag is 2-[2-(4-Monomethoxytrityl)aminoethoxy]ethyl-(2-cyanoethyl)-N,N-diisopropyl)-phosphoramidite (available from Glen Research, Sterling VA, catalog No. 10-1905-xx). Another useful positively charged moiety is a 5' primary amine which may have a single positive charge at the appropriate pH.

[0175] Table 1 provides a non-limiting listing of some useful modifications and charges that may be used as labels in an apparatus or method set forth herein.

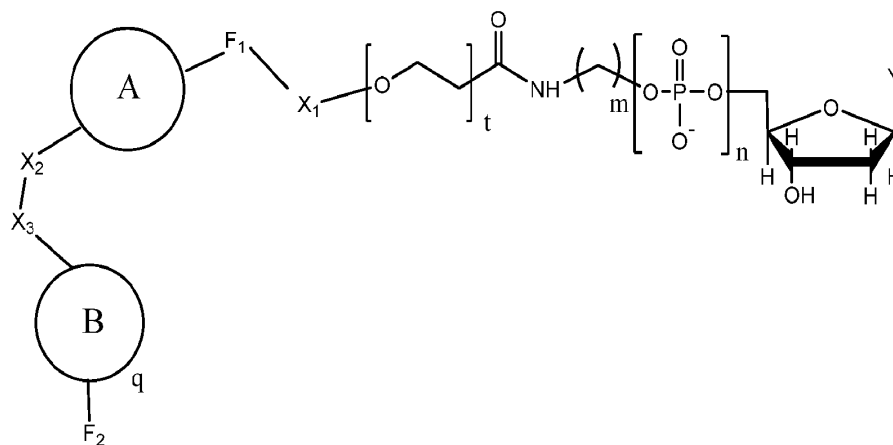
Table 1

5' Terminus	Reagents	Final Charge State
5' OH	N/A	Neutral
5' Phosphate	CPR 10-1900 (Glen Res.)	-2
5' Phosphate (x2)	CPR 10-1900 and symmetric doubler (Glen Res.)	-4
5' Phosphate (x3)	CPR 10-1900 and symmetric trebler (Glen Res.)	-6
5' primary amine	5' amino-modifier 5	+1

[0176] In an aspect, the present disclosure relates to a modified nucleotide including: a nucleotide; a linking molecule attached to a phosphate group of the nucleotide; and a charge

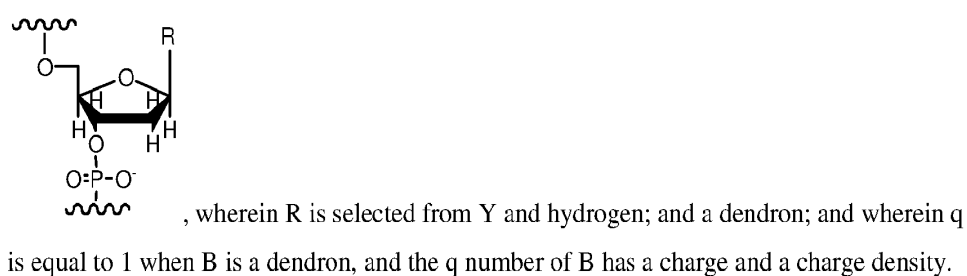
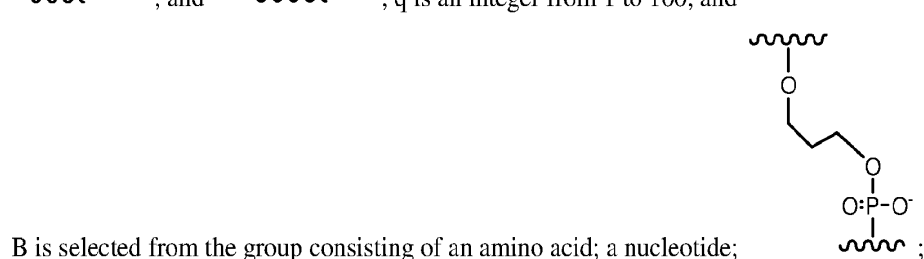
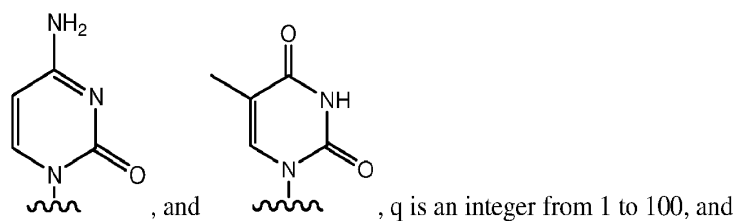
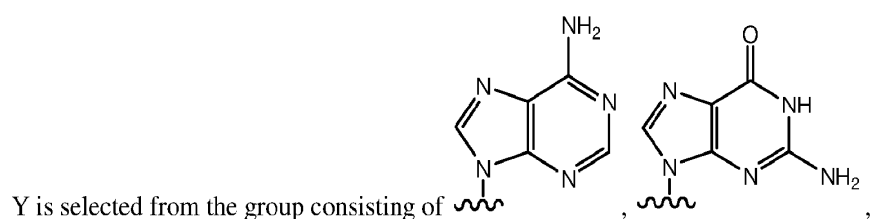
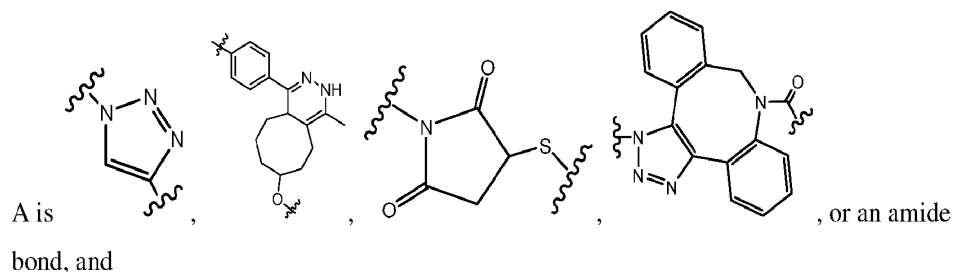
tag attached to the linking molecule, wherein the charge tag includes a plurality of elements selected from the group consisting of nucleotides and amino acids, and optional linkers between elements, and wherein the charge tag comprises an internal folded or secondary structure. In an example, wherein the charge tag comprises one or more phosphodiester groups, and optional linkers between elements. In some aspects, the nucleotide may be a natural nucleotide or a modified nucleotide. Modified nucleotide structures are known to one of ordinary skill in the art and may include structural modifications to the base or the sugar moiety (e.g., alkylation, amino groups, or protecting groups). In some examples, the linking molecule comprises a specificity region. In some examples, the specificity region comprises a nucleotide sequence including from one to six nucleotides. In some examples, the charge tag includes from about 1 charge to about 100 or about 200 charges. In some examples, the linking molecule comprises a structure as shown below in Formula I from $-X_2$ through the $(CH_2)_m$ group. In one example, the charge tag does not bind to a polymerase (e.g., Phi29) used in the methods herein. In some examples, the charge tag comprises a plurality of nucleotides comprising two noncontiguous regions that bind to an acceptor region in a polymerase tether, thereby forming a hairpin structure in the charge tag.

[0177] An example of a nucleotide analog, or a labeled nucleotide, is represented by a compound of the following Formula I:



wherein n is an integer from 3 to 10, m is an integer from 1 to 10, t is an integer from 0 to 50, X_1 is a direct bond, a C_1 - C_{10} alkyl, a C_1 - C_{10} oxaalkyl, a C_1 - C_{10} thiaalkyl, or a C_1 - C_{10} azaalkyl, X_2 is C_1 - C_{20} alkyl wherein optionally one or more individual CH_2 residue is replaced with one or more of a peptide bond and $(-O-CH_2-CH_2-)_a$ wherein a is an integer from 1 to 24, X_3 is

a direct bond or an oligonucleotide wherein the oligonucleotide hybridizes to an acceptor region of the tether when the label is in proximity to the conductive channel, F_1 is selected from a fluorophore and a direct bond and F_2 is absent or a fluorophore,



In an example, provided is a method including detecting an incorporation of a labeled nucleotide into a nascent polynucleotide strand complementary to a template polynucleotide strand by a polymerase, wherein the polymerase is tethered to a solid support conductive

channel by a tether, the labeled nucleotide is a compound of Formula I, and the conductive channel is to detect the labeled nucleotide during the incorporation.

[0178] In an example, B includes a charge tag, and the charge tag includes nucleotides, oligonucleotides, amino acids, peptide nucleic acids, or combinations thereof, wherein the charge tag has an internal folded or secondary structure.

[0179] As explained further herein, making a compound of Formula I may include forming A by a reaction including a linking reaction and the linking reaction is selecting from the group consisting of an azide-alkyne copper-assisted click reaction, a tetrazine-trans-cyclooctene ligation, an azide-dibenzocyclooctyne group copper-free click reaction, and a thiol-maleimide conjugation.

[0180] Also provided is a method of detecting, with a charge detector, a charge tag of a compound of Formula I, such as during incorporation of a nucleotide portion of a compound of Formula I into a nascent strand of a polynucleotide. In a non-limiting example, detecting may occur during sequencing a nucleic acid, including (a) providing a polymerase tethered to a solid support conductive channel; (b) providing one or more compounds of Formula I, whereby the presence of the compound can be detected by the conductive channel when the label is in proximity to the conductive channel; and (c) detecting incorporation of the compound into a nascent strand complementary to a template nucleic acid using the conductive channel.

[0181] Also provided is a compound of Formula I, wherein B includes one or more oligonucleotides with one or more stem-and-loop shapes, one or more cloverleaf shapes, one or more tubular shapes, one or more annular shapes, one or more cuboidal shapes, one or more cruciform shapes, one or more spherical shapes, one or more rectangular shapes, one or more pyramidal shapes, one or more diamond shapes, one or more laminar shapes, one or more columnar shapes, one or more corrugated shapes, or any combination of two or more of the foregoing. In another example of a compound of Formula I, B includes one or more polypeptides with one or more coiled shapes. Also provided is a compound of Formula I, wherein B includes one or more oligonucleotides forming a cruciform shape, one or more peptide nucleic acid molecules bonded to one or more of the oligonucleotides, and one or more polypeptides bonded to the one or more peptide nucleic acid molecules.

[0182] Also provided is a compound of Formula I wherein B has a charge of between $-100e$ and $+100e$. Also provided is a compound of Formula I wherein B has a charge of between -

100e and +100e and a charge density of between -100e per cubic nanometer and +100e per cubic nanometer. Also provided is a compound of Formula I wherein B has a charge of between -200e and +200e. Also provided is a compound of Formula I wherein B has a charge of between -200e and +200e and a charge density of between -200e per cubic nanometer and +200e per cubic nanometer.

[0183] In some examples, a compound of Formula I may optionally include a fluorophore, such as represented by F₁, F₂ or both. Some non-limiting examples of fluorophores include cyanine dyes (e.g., Cy2, Cy3, or Cy5), fluorescein isothiocyanate, rhodamine fluorophores (e.g., tetramethyl rhodamine), or others. Optional presence of a fluorophore in a compound of Formula I may provide additional uses such as for detection of a tagged nucleotide including a fluorophore. For example, presence of a fluorophore-containing charge tag may be detected not only through detection of a presence, valence, and magnitude of a charge carried by the tag but by methods for detecting fluorescence emission, such as fluorescence resonance energy transfer.

[0184] Also provided is a tagged nucleotide wherein the charge tag includes one or more peptide nucleic acids. In some examples, the charge tag includes one or more peptide nucleic acids, and one or more of the peptide nucleic acids is attached to one or more charged amino acids.

[0185] Also provided is a method of forming a compound of Formula I, wherein the charge tag includes oligonucleotides and is formed by DNA origami. DNA origami may involve folding DNA in creation of non-arbitrary shapes at the nanoscale. Compacted, origami DNA structures may permit high charge density, permitting variations in charge density in different compounds of Formula I. Higher charge, higher charge density, and greater flexibility in varying charge and charge density of a charge tag may increase probability of detection of a charge tag by a conductive channel and also permit discriminating between different charge tags detected by a conductive channel. A greater range of charges that may be carried by a charge tag allows for greater differentiation between charges carried by different examples of compounds of Formula I. In some examples, different nucleotides may be differentiated from each other by the charge carried by a tag to which they are linked as a compound of Formula I. A conductive channel may thereby be able to differentially detect different nucleotides constituting a portion of a compound of Formula I based on differences in the magnitude of the charge carried by different such nucleotides.

[0186] B may include positively charged amino acids such as arginine, histidine, and lysine, yielding a charge tag with a positive charge. B may instead include aspartic acid and glutamic acid, yielding a charge tag with a negative charge. In some examples, B is a branched polypeptide, or a linear polypeptide, or a cyclic polypeptide. In some examples, B may be a single amino acid or a polypeptide with anywhere from 2 to 10, or 11 to 20 amino acids. In some examples, some of the amino acids of B may be uncharged and in other examples B may contain some amino acids that are oppositely charged from other amino acids of B, yet B may retain an overall positive or negative charge.

[0187] In this non-limiting example of Formula I, a nucleotide directly bonded to the n phosphate groups may be a nucleotide recognizable by a polymerase, and incorporated into a nucleotide sequence synthesized thereby complementarily to a template sequence. While the nucleotide analog is held in place by the polymerase during addition to a growing synthesized polynucleotide sequence, the remainder of the analog may extend therefrom and, as disclosed herein, for a polymerase in proximity to a conductive channel, a charge tag (such as represented by B in Formula I and in a charged peptide to which it is directly bound) may move or be brought into proximity with the conductive channel such that the conductive channel may sense the valence and magnitude of the charge. Different nucleotide analogs may contain different nucleotides at the 3' end of the analog, and correspondingly different peptide charge tags at the 5' end of the analog, such that a conductive channel tethered to a polymerase may detect differences in charge valence and magnitude when the polymerase associates with different nucleotide analogs to incorporate a nucleotide in a nucleotide sequence being synthesized. In these examples, a polymerase may cleave all but one phosphate group bound directly to the 5' nucleotide of the nucleotide analog such that the dNMP portion of the nucleotide analog remains in a synthesized nucleotide sequence with a 5' phosphate group free to bind to the next nucleotide to be incorporated and the cleaved remainder of the nucleotide analog free to dissociate from the complex.

[0188] A phosphate or series of phosphate groups bound directly to the 3' nucleotide may include 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 phosphate groups. Other examples may include more than 10 phosphate groups. This portion of a nucleotide analog may then be connected by an alkyl linkage including 1-10 $-\text{CH}_2-$ groups. Other examples may include from 11-20 such groups at this position. In other examples, one or more of these 1-10, or 11-20, $-\text{CH}_2-$ groups may be substituted by a C_1 to C_{20} hydrocarbon.

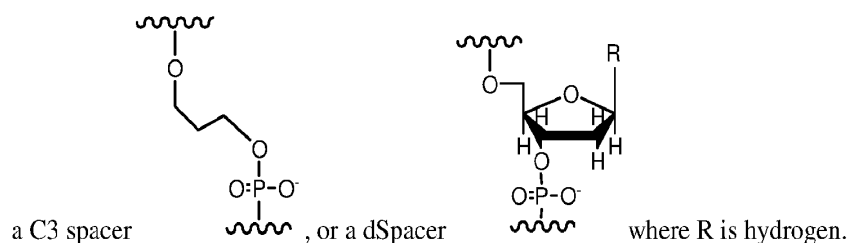
[0189] This portion of the nucleotide analog may be further connected by 0 to 50 oxaalkyl groups, such as -O-CH₂-CH₂- groups. In other examples, one or more of these 0-50 -O-CH₂-CH₂- groups may be substituted by a C₁ to C₁₅₀ hydrocarbon. This portion of the nucleotide analog may be further connected to by an alkyl linkage including 0-10 -CH₂- groups as represented by X₁. Other examples of X₁ may include from 11-20 such groups at this position. In other examples of X₁, one or more of these 1-10, or 11-20, -CH₂ groups may be substituted by a C₁ to C₂₀ hydrocarbon. X₁ may also be a direct bond, a C₁-C₁₀ alkyl, a C₁-C₁₀ oxaalkyl, a C₁-C₁₀ thiaalkyl, or a C₁-C₁₀ azaalkyl,

[0190] As described more fully below, A represents a linking group by which a 5' end of a nucleotide analog may be connected to a charge tag towards a 3' end of the nucleotide analog. For example, a nucleotide polyphosphate may have functional groups appended to the 5' phosphate group most distal to the deoxyribose (or ribose), at the end of which functional groups may be a reactive group. A reactive group is a chemical group capable of reacting with another chemical group--together being two reactive groups--to form a covalent bond or bonds therebetween, under controlled conditions such as in the presence of a specific reagent or reagents, or at a predetermined pH or temperature, etc. For example, compositions resembling or example of portions of Formula I from the 3' nucleotide up to or some number of bonds short of A may be commercially available or synthesized according to known methods. A reactive group may then be appended to the end of such compound such that a charge tag with another reactive group, with which the first can react to form a covalent bond, may be reacted together thereby covalently linking a charge tag to a 3' nucleotide to form a compound of Formula I.

[0191] Attached to A may be X₂. X₂ may be C₁-C₂₀ alkyl wherein individual CH₂ residues may be independently replaced with one or more of a peptide bond and (-O-CH₂-CH₂-)_a wherein a is an integer from 1 to 24. In other examples of X₂, a may be an integer from 6 to 20. In still other examples, one or more of the 1-20 alkyl groups of X₂ may be substituted by a C₁ to C₂₀ hydrocarbon.

[0192] In an example, B may represent a charge tag connected to X₂ by a phosphate linkage. B may include from 1 to 100 moieties containing phosphodiester groups. In an example, B may include from 1 to 200 moieties containing phosphodiester groups. Negative charges carried by oxygen atoms in such phosphodiester groups may confer a negative charge on a B charge tag, with magnitude proportional to the number of moieties. Each of the q moieties of

B may be a different moiety from any of the other moieties of B, or they may all be the same as each other. Any one or more moieties of B may be a dNMP with an adenine, thymine, cytosine, or guanosine base, for example. Any one or more moieties of B may be:

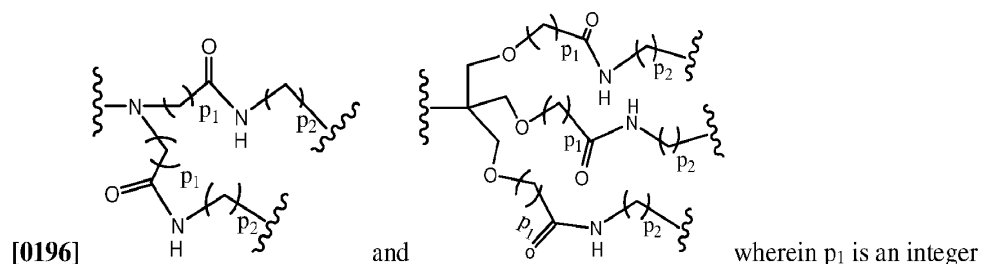


[0193] In some examples, any moiety of B may include any NPP (nucleotide polyphosphate). Charge tags whose charge valence, magnitude, or both differ from those of other charge tags to which different 3' nucleotides are bound permits differentiated identification of nucleotide analogs by a conductive channel as they are held by a polymerase tethered thereto during polynucleotide synthesis. In some examples, some nucleotide analogs are a compound of Formula I or similar compound as disclosed herein. In some examples, all nucleotides used in a sequencing-by-synthesis reaction contain a charged tag that includes one or more phosphodiester groups as disclosed herein, such as examples of compounds of Formula I or related compounds. In other examples, some nucleotides used in a sequencing-by-synthesis reaction contain a charged tag that includes one or more phosphodiester groups as disclosed herein, such as examples of compounds of Formula I or related compounds, whereas other nucleotides used in a sequencing-by-synthesis reaction contain a charged tag that do not include such compounds.

[0194] In other examples, each B may independently be selected from arginine, histidine, and lysine, yielding a charge tag with a positive charge. In another example, each B may independently be selected from aspartic acid and glutamic acid, yielding a charge tag with a negative charge. In some examples, the q number of B is a branched polypeptide, or a linear polypeptide, or a cyclic polypeptide. In some examples, the q number of B may be a single amino acid or a polypeptide with anywhere from 2 to 10, or 11 to 20 amino acids. In some examples, some of the amino acids of B may be uncharged and in other examples B may contain some amino acids that are oppositely charged from other amino acids of B, yet B may retain an overall positive or negative charge. In some examples, B may include non-natural amino acids.

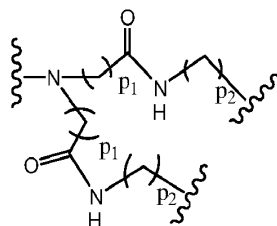
[0195] In still other examples, B may be a dendron of z generations comprising one or more

constitutional repeating unit and a plurality of end units, wherein z is an integer from 1 to 6, the constitutional end units are selected from the group consisting of:

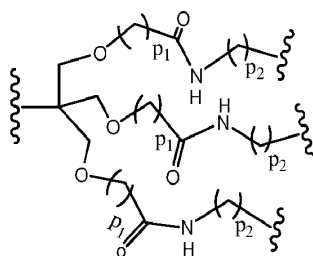


from 1 to 3 wherein any one or more of the p_1 -CH₂- groups is optionally replaced with from 1 to 3 -O-CH₂-CH₂- groups, p_2 is an integer from 1 to 3 wherein any one or more of the p_2 -CH₂- groups is optionally replaced with from 1 to 3 -O-CH₂-CH₂- groups, and the end groups are selected from the group consisting of carboxylic acid, sulfonic acid, phosphonic acid, amino group, or quaternary ammonium group.

[0197] B may represent a dendron charge tag connected to X₂ by its free valence end. In some examples, a dendron disclosed herein may be unattached to a nucleotide analog, such as before it has been chemically bonded thereto. B may include a constitutional repeating unit with 2 degrees of branching, such as represented by the following:



[0198] Or, B may include a constitutional repeating unit with 3 degrees of branching, such as represented by the following:



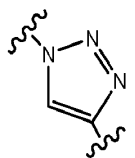
[0199] As further disclosed herein, dendron charge tags may be anywhere from 1 to 6 generations in size. End groups on terminal constitutional repeating units may be charged,

either positively or negatively. Dendrons with 2 degrees of branching may therefore yield a charge tag with a charge of 2^z and dendrons with 3 degrees of branching may yield a charge tag with a charge of 3^z (where the magnitude of charge per end group is 1 and z represents the number of generations).

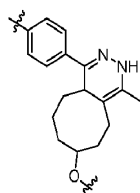
[0200] In an example where B represents a dendron, end groups may be any of a number of charged functional groups, such as, for example, carboxylic acid, sulfonic acid, phosphonic acid, amino group, or quaternary ammonium group, or any other charged functional group. In some examples, constitutional repeating units of a dendron may include a charge on an atom other than on an end group of the terminal constitutional repeating units. For example, as one non-limiting example, a constitutional repeating unit may contain a quaternary ammonium group at a branch point, which could carry a positive charge. Unlike a charged end group, which may only be present on a terminal constitutional repeating unit, such internal charge may be present on every instance of a constitutional repeating unit in the dendron.

[0201] A peptide bond may be present, such as represented optionally at X_2 in Formula I. In other examples, in place of the peptide bond shown in Formula I, a C_1 to C_{20} hydrocarbon may be present, or a direct bond.

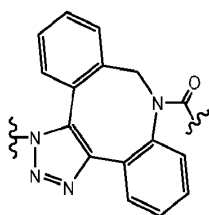
[0202] For A, a linker linking a nucleotide to a charge tag may be formed by a linking reaction between reactive groups. For example, A may be formed by an azide-alkyne copper-assisted click reaction between a nucleotide with an azide (or alkyne) group and a charge tag with an alkyne (or azide) group, yielding a chemical structure such as the following or an equivalent thereof:



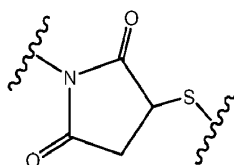
[0203] Or, A may be formed by a tetrazine (TET)-trans-cyclooctene (TCO) ligation between a nucleotide with a tetrazine (or trans-cyclooctene) group and a charge tag with a trans-cyclooctene (or tetrazine) group, yielding a chemical structure such as the following or an equivalent thereof:



[0204] Or, A may be formed by an azide-dibenzocyclooctyne (DBCO) group copper-free click reaction between a nucleotide with an azide (or dibenzocyclooctyne) group and a charge tag with a dibenzocyclooctyl (or azide) group, yielding a chemical structure such as the following or an equivalent thereof:



[0205] Or, A may be formed by a thiol-maleimide conjugation between a nucleotide with a thiol (or maleimide) group and a charged tag with a maleimide (or thiol) group, yielding a chemical structure such as the following or an equivalent thereof:



[0206] Or, A may be formed by an N-hydroxysuccinimide ester-amine linkage reaction between a nucleotide with an amine (or N-hydroxysuccinimide ester) group and a charged tag with an N-hydroxysuccinimide ester (or amine) group, yielding an amide bond.

[0207] Other suitable linking groups, formed by other ligation chemistries between suitable reactive groups, may be incorporated into the present disclosure to form other structures for A by which a 3' nucleotide may be linked to a charge tag.

[0208] B of Formula I represents a charge tag. As disclosed herein, a charge tag may include polypeptides, oligonucleotides, oligomeric peptide nucleic acids, or a dendron, or combinations of at least two of the foregoing. Charges of a charge tag may be carried by charged functional groups of such moieties, such as phosphodiester bonds, amide groups, carboxylic acid groups, or other charged functional groups that may be added to such compounds such as one or more sulfonic acid, phosphonic acid, or quaternary ammonium

groups. As disclosed herein, a charge tag may adopt a particular three-dimensional orientation such that the charges carried by elements thereof are held together and inhibited or in some instances prevented from splaying out and away from a conductive channel. Such condensation of charge by increasing charge density of a charge tag may increase charge detected by a conductive channel.

[0209] A charge tag may be synthesized so as to have a reactive group suitable for forming a click chemistry or ligation reaction according to the foregoing. For example, a charge tag may have an azide or alkyne group (such as for covalent attachment to and inclusion in a nucleotide analog as a charge tag by an azide-alkyne copper-assisted click reaction), or a tetrazine (TET) or trans-cyclooctene group (such as for covalent attachment to and inclusion in a nucleotide analog as a charge tag by a tetrazine (TET)-trans-cyclooctene (TCO) ligation), or an azide group or DBCO group (such as for covalent attachment to and inclusion in a nucleotide analog as a charge tag by an azide- DBCO group copper-free click reaction), or a thiol (e.g., a cysteine residue) or maleimide group (such as for covalent attachment to and inclusion in a nucleotide analog as a charge tag by a thiol-maleimide conjugation). Other known ligation, click chemistry, or other covalent attachment chemistries may also be employed, with corresponding reactive groups attached to the charge tag permitting its covalent attachment to a nucleotide analog.

[0210] A peptide bond may be present, such as is shown in between A and the 5' nucleotide in Formula I. In other examples, in place of the peptide bond shown in Formula I, a C₁ to C₂₀ hydrocarbon may be present, or a direct bond.

[0211] Some suitable examples include modifications to or variations of a compound of Formula I that incorporate features discussed above related to how an acceptor region of a tether (by which a polymerase is tethered to a conductive channel) may hybridize or otherwise form non-covalent bonds with a specificity region of nucleotide analogs. For example, some portion of an analog nucleotide between the 3' nucleotide and the 5' charge tag may incorporate nucleotides, PNA residues, or amino acids capable of forming non-covalent bonds with a tether by which a polymerase is connected to a conductive channel, or to a portion functionalized with an acceptor region extending from and attached to a conductive channel that itself may not be a portion of such tether, and may also include nucleotides, PNA residues, or amino acids, or combinations thereof. The foregoing may be substituted for or added to regions of the compound of Formula I as disclosed herein between

the 5' charge tag and 3' nucleotide. Such substitution or addition may contribute to a synergistic binding of an analog nucleotide to a polymerase and to a tether (or a functionalized portion of a conductive channel apart from a tether for purpose of binding to such substitution or addition) to promote association of a charge tag with a detection region of a charge detector of suitably long duration to permit detection of a charge tag to register and signify incorporation of a nucleotide analog bearing such charge, as disclosed herein.

[0212] In some examples, a charge tag's adoption of a three-dimensional structure may lead to formation of a specificity region by bringing together otherwise spatially disparate elements of a specificity region allowing for bonding of the so-assembled specificity region to an acceptor region. Such specificity region formation and acceptor region binding might not occur or might be unlikely to occur or to occur only very transiently in the absence of adoption of a particular three-dimensional structure of a charge tag. In other example, the bringing together of otherwise disparate elements of a specificity region upon binding to an acceptor region may induce or promote a charge tag's adoption of a given three-dimensional conformation. In some examples, adoption of a charge tag's three dimensional conformation and the coming together of otherwise spatially distal elements of a specificity region may be synergistic such that each promotes the other. In some cases, the three-dimensional conformation so adopted by the charge tag leads to a higher charge density than would otherwise be likely to occur and may increase detection of a charge tag by a conductive channel.

[0213] Various designs of peptide charge tags can be used. Using solid phase peptide synthesis, any of the 21 amino acids can be included in a charge tag. In addition, modified amino acids are also available commercially and can be added to a peptide charge tag to further modulate its properties. Besides using amino acids with electronically charged side chains such as arginine, histidine, and lysine (positive), and aspartic acid and glutamic acid (negative), other amino acids can be incorporated in the peptide charge tag to tweak its hydrophilicity, length and size.

[0214] As disclosed above, in one example, a peptide charge tag may be presented in the form of a linear (see FIG. 14A), branched (see FIGs. 14B and 14C) or cyclic chains (see FIG. 14D).

[0215] By using different combination of amino acids, such as KKKKK (SEQ ID NO: 8) or EEEEE (SEQ ID NO: 9) (or other combinations of charged amino acids, with or without

additional uncharged amino acids), of various lengths, 4 different nucleotide analogs may be distinguished for sequencing or various nucleotide analogs may be distinguished based on characteristic current signature from each peptide charge tag. Other more complex three-dimensional conformations are also possible. For example, a peptide charge tag may adopt a coiled conformation, such as an α -helix. Such a structure may include positive and negative amino acids, but an overall positive or negative charge. For example, placement of oppositely charged amino acids may induce bonding therebetween and adoption of an α -helical or other structure, wherein excess positive or excess negative charge is held together in proximity, increasing charge density. In other examples, similar bonding may promote adoption of a coiled coil structure including density of net positive or negative charge.

[0216] A charge tag may also include an oligonucleotide. An oligonucleotide charge tag may be attached to a nucleotide analog using click chemistry and ligation chemistry reactions described above for attaching a peptide charge tag to a nucleotide analog.

[0217] An oligonucleotide charge tag may adopt various three-dimensional orientations that promote compressing its charge at an elevated charge density. For example, phosphodiester bonds between nucleotides of an oligonucleotide may have a negative charge. By adopting a condensed three dimensional structure, negative charges of an oligonucleotide may be held in proximity to one another, increasing detection of such charge tag be a conductive channel. For example, an oligonucleotide may adopt well-known structures such as a step-and-loop structure, a cloverleaf structure, or a cruciform structure (such as a Holliday junction). Polynucleotide origami techniques may also be used to design polynucleotide charge tags that adopt other conformations that increase charge density. A polynucleotide charge tag may adopt a tubular shapes, an annular shapes, a cuboidal shapes, or a spherical shape. Such shapes may result in an oligonucleotide charge tag with a higher charge density than an oligonucleotide with the same nucleotide composition but not adopting the three-dimensional conformation, such as if it were stretched out into a linear conformation, would have.

[0218] For convenience and clarity, certain terms employed in the specification, examples, and claims are described herein.

[0219] Unless otherwise specified, alkyl is intended to include linear or branched saturated hydrocarbon structures and combinations thereof. Alkyl refers to alkyl groups of from 1 to 20 carbon atoms – e.g., 1 to 10 carbon atoms, such as 1 to 6 carbon atoms, etc. Examples of alkyl groups include methyl, ethyl, propyl, isopropyl, n-butyl, s-butyl, t-butyl and the like.

[0220] Cycloalkyl is a subset of hydrocarbon and includes cyclic hydrocarbon groups of from 3 to 8 carbon atoms. Examples of cycloalkyl groups include c-propyl, c-butyl, c-pentyl, norbornyl and the like.

[0221] C₁ to C₂₀ hydrocarbon includes alkyl, cycloalkyl, polycycloalkyl, alkenyl, alkynyl, aryl and combinations thereof. Examples include benzyl, phenethyl, propargyl, allyl, cyclohexylmethyl, adamantyl, camphoryl and naphthylethyl. Hydrocarbon refers to any substituent comprised of hydrogen and carbon as the only elemental constituents.

[0222] Unless otherwise specified, the term “carbocycle” is intended to include ring systems in which the ring atoms are all carbon but of any oxidation state. Thus (C₃-C₁₂) carbocycle refers to both non-aromatic and aromatic systems, including such systems as cyclopropane, benzene and cyclohexene. Carbocycle, if not otherwise limited, refers to monocycles, bicycles and polycycles. (C₈-C₁₂) Carbopolycycle refers to such systems as norbornane, decalin, indane and naphthalene.

[0223] Alkoxy or alkoxyl refers to groups of from 1 to 20 carbon atoms – e.g., 1 to 10 carbon atoms, such as 1 to 6 carbon atoms, etc. of a straight or branched configuration attached to the parent structure through an oxygen. Examples include methoxy, ethoxy, propoxy, isopropoxy and the like.

[0224] Oxaalkyl refers to alkyl residues in which one or more carbons (and their associated hydrogens) have been replaced by oxygen. Examples include methoxypropoxy, 3,6,9-trioxadecyl and the like. The term oxaalkyl is intended as it is understood in the art [see Naming and Indexing of Chemical Substances for Chemical Abstracts, published by the American Chemical Society, 2002 edition, ¶196, but without the restriction of 127(a) – the reference is incorporated by reference in its entirety] – it refers to compounds in which the oxygen is bonded via a single bond to its adjacent atoms (forming ether bonds); it does not refer to doubly bonded oxygen, as would be found in carbonyl groups. Similarly, thiaalkyl and azaalkyl refer to alkyl residues in which one or more carbons has been replaced by sulfur or nitrogen, respectively. Examples of azaalkyl include ethylaminoethyl and aminohexyl.

[0225] Heterocycle means a cycloalkyl or aryl carbocyclic residue in which from one to four carbons is replaced by a heteroatom selected from the group consisting of N, O and S. Heteroaryl is a subset of heterocycle in which the heterocycle is aromatic. Examples of heteroaromatic rings include: furan, benzofuran, isobenzofuran, pyrrole, indole, isoindole, thiophene, benzothiophene, imidazole, benzimidazole, purine, pyrazole, indazole, oxazole,

benzoxazole, isoxazole, benzisoxazole, thiazole, benzothiazole, triazole, tetrazole, pyridine, quinoline, isoquinoline, pyrazine, quinoxaline, acridine, pyrimidine, quinazoline, pyridazine, cinnoline, phthalazine, and triazine.

[0226] As used herein, the term “optionally substituted” may be used interchangeably with “unsubstituted or substituted”. The term “substituted” refers to the replacement of one or more hydrogen atoms in a specified group with a specified radical. For example, substituted alkyl, aryl, cycloalkyl, heterocyclyl etc. refer to alkyl, aryl, cycloalkyl, or heterocyclyl wherein one or more H atoms in each residue are replaced with halogen, haloalkyl, alkyl, acyl, alkoxyalkyl, hydroxyloweralkyl, carbonyl, phenyl, heteroaryl, benzenesulfonyl, hydroxy, loweralkoxy, haloalkoxy, oxaalkyl, carboxy, alkoxycarbonyl [-C(=O)O-alkyl], alkoxycarbonylamino [HNC(=O)O-alkyl], carboxamido [-C(=O)NH₂], alkylaminocarbonyl [-C(=O)NH-alkyl], cyano, acetoxy, nitro, amino, alkylamino, dialkylamino, (alkyl)(aryl)aminoalkyl, alkylaminoalkyl (including cycloalkylaminoalkyl), dialkylaminoalkyl, dialkylaminoalkoxy, heterocyclylalkoxy, mercapto, alkylthio, sulfoxide, sulfone, sulfonylamino, alkylsulfinyl, alkylsulfonyl, alkylsulfonylamino, arylsulfonyl, arylsulfonylamino, acylaminoalkyl, acylaminoalkoxy, acylamino, amidino, aryl, benzyl, heterocyclyl, heterocyclylalkyl, phenoxy, benzyloxy, heteroaryloxy, hydroxyimino, alkoxyimino, oxaalkyl, aminosulfonyl, trityl, amidino, guanidino, ureido, benzyloxyphenyl, and benzyloxy. “Oxo” is also included among the substituents referred to in “optionally substituted”; it will be appreciated by persons of skill in the art that, because oxo is a divalent radical, there are circumstances in which it will not be appropriate as a substituent (e.g. on phenyl). In one example, 1, 2, or 3 hydrogen atoms may be replaced with a specified radical. In the case of alkyl and cycloalkyl, more than three hydrogen atoms can be replaced by fluorine; indeed, all available hydrogen atoms could be replaced by fluorine. Such compounds (e.g., perfluoroalkyl) fall within the class of “fluorohydrocarbons”. To be clear, a generic term may encompass more than one substituent, that is, for example, “haloalkyl” or “halophenyl” refers to an alkyl or phenyl in which at least one, but perhaps more than one, hydrogen is replaced by halogen. In some examples, substituents are halogen, haloalkyl, alkyl, acyl, hydroxyalkyl, hydroxy, alkoxy, haloalkoxy, oxaalkyl, carboxy, cyano, acetoxy, nitro, amino, alkylamino, dialkylamino, alkylthio, alkylsulfinyl, alkylsulfonyl, alkylsulfonylamino arylsulfonyl, arylsulfonylamino and benzyloxy.

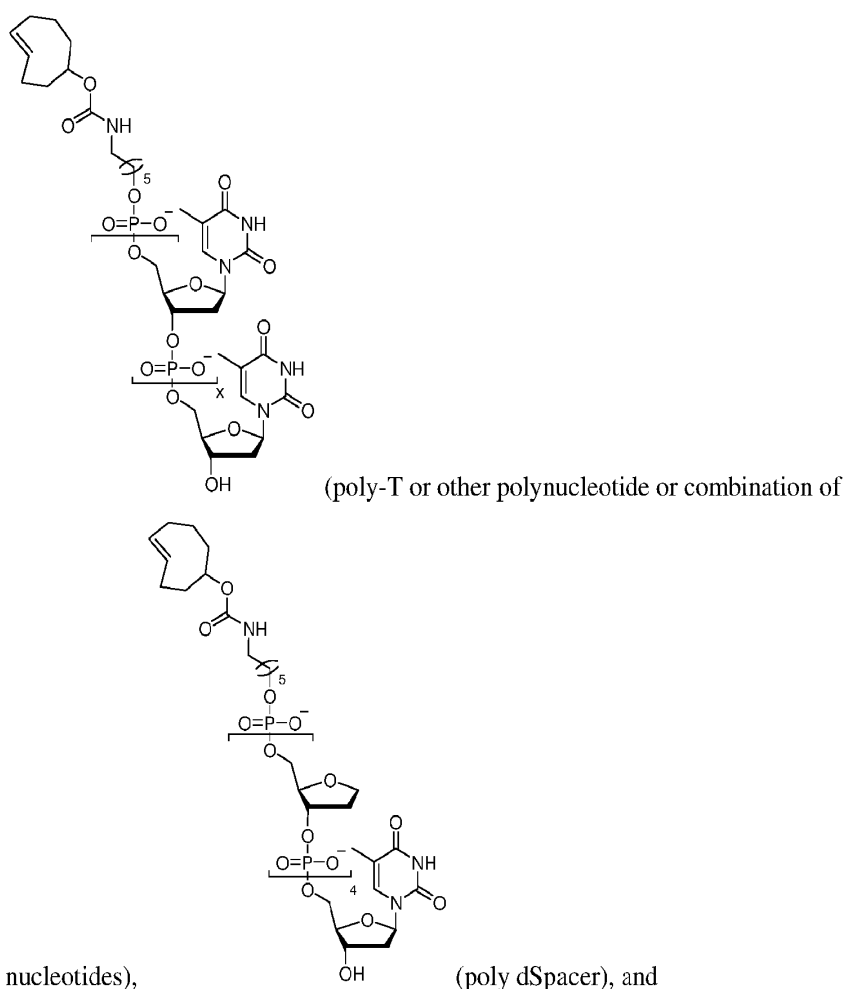
[0227] In describing compounds herein, the terminology “substituted with at least one

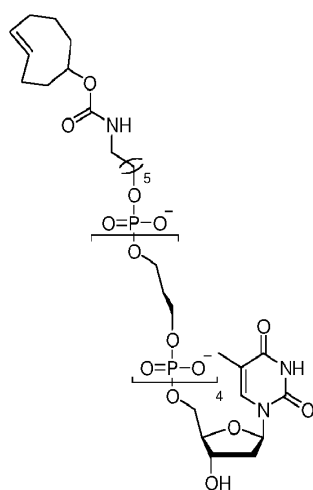
oxygenated substituent” is used. An oxygenated substituent is a substituent that contains oxygen in addition to carbon and hydrogen; an oxygenated substituent may also include additional heteroatoms, such as nitrogen (for example, a carboxamide or methanesulfonyl). Typical examples of oxygenated substituents include alkoxy, hydroxy, fluoroalkoxy, formyl, acetyl and other C₁ to C₆ acyl chains.

NON-LIMITING WORKING EXAMPLES

[0228] The following examples are intended to illustrate particular examples of the present disclosure, but are by no means intended to limit the scope thereof.

[0229] Some examples of charge tags for incorporation into a nucleotide that were made in accordance with the present disclosure include the following:





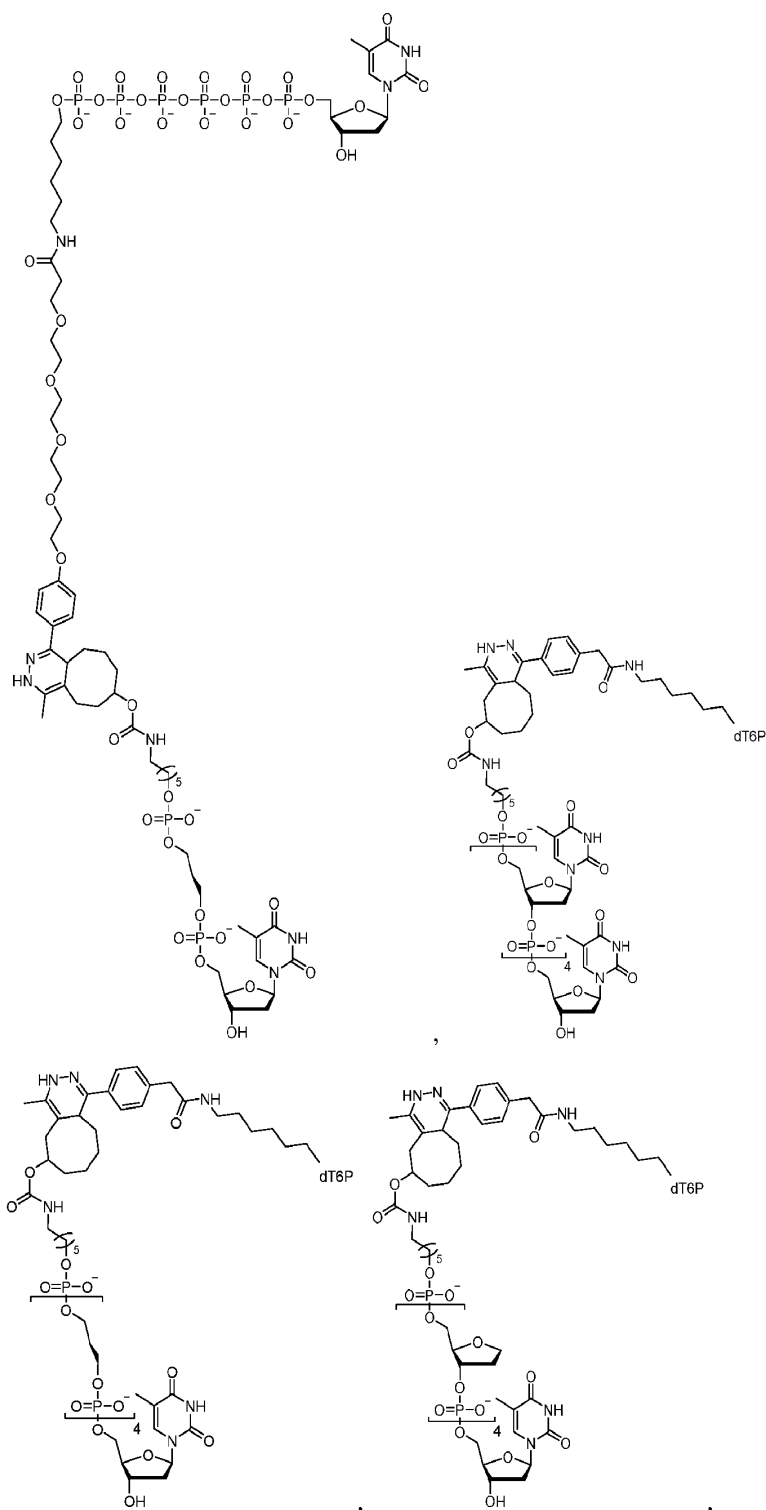
(poly C spacer), or combinations of any of the foregoing.

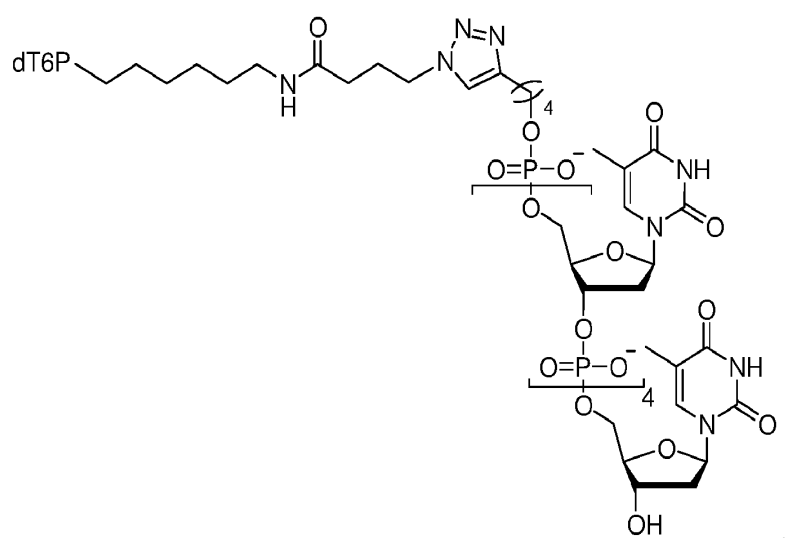
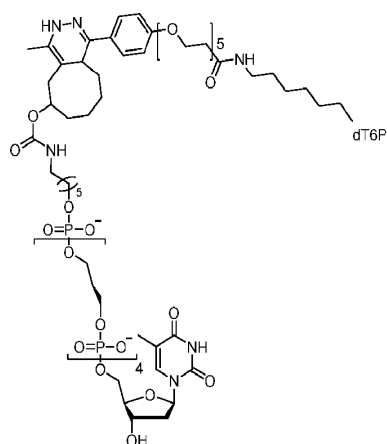
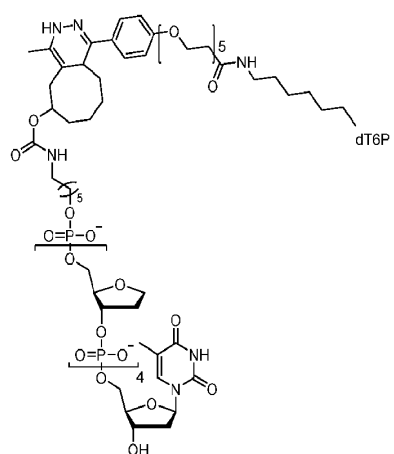
[0230] Charges for such charge tags may be varied by altering the number of phosphate group-containing moieties, e.g. 5, 10, 15, 20, 25, 30, 35, 40, or any number or range therebetween. As many as 40 may be included, or any number from 1 to 40. More than 40 may be included. Suitable reactive groups other than the trans-cyclooctene group shown in these examples may be used, in accordance with the present disclosure.

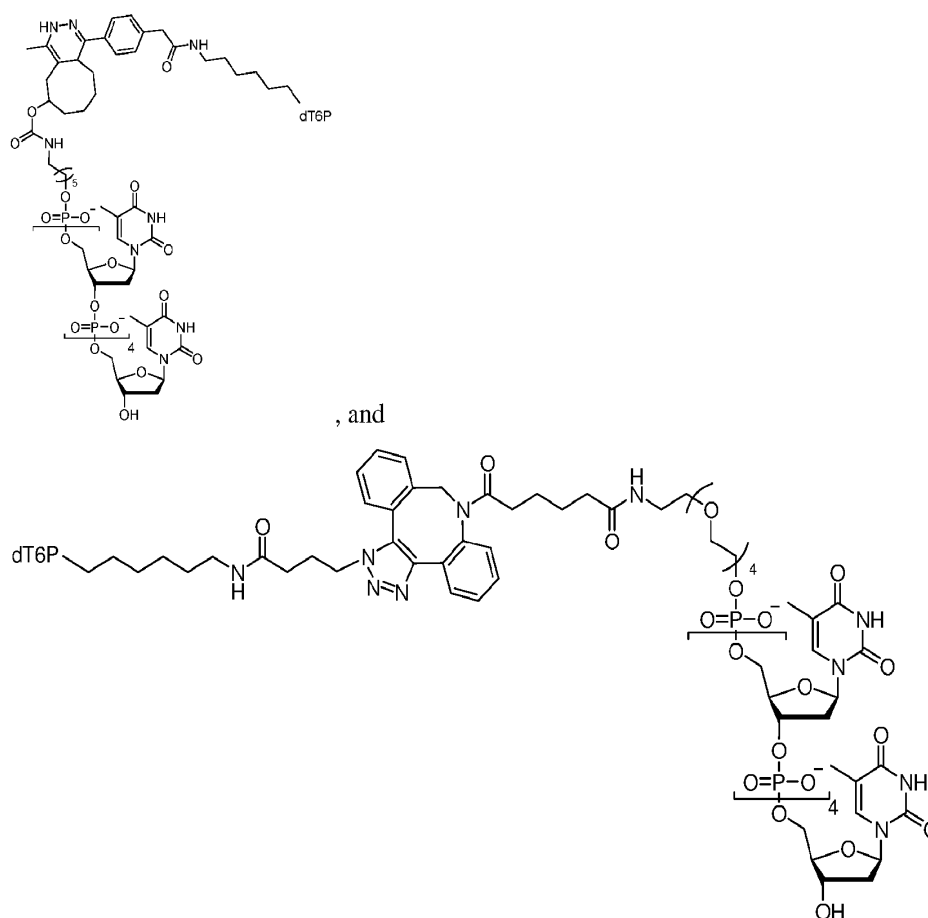
[0231] An oligonucleotide sequence can be used as a charge tag, with various lengths of charges conferred by phosphates in phosphodiester linkages. In addition, modified oligonucleotides such as dSpacer and C3 Spacer nucleotides can also be used to create charge tags with different hydrophilicity and size. An oligonucleotide sequence can be modified using different bases and hydrophobic modifications to modulate sequence specificity, minimize inhibition to polymerases and optimize interactions with the surface and linkers.

[0232] Phosphodiester based charge tags may be attached to a 5'-terminal phosphate of a nucleotide. Upon incorporation of each nucleotide by a polymerase into a growing strand during synthesis of a complement to a template strand, the charge label may be released as part of the pyrophosphate by-product. The charge on the label is detected by the detection system on the conductive channel. Based on a characteristic current signature from each tag (e.g., charge magnitude), bases incorporated into the synthesizing strand can be distinguished using differential magnitude of charge conferred by the tags.

[0233] Examples of analog nucleotides according to the present disclosure included the following, without limitation:

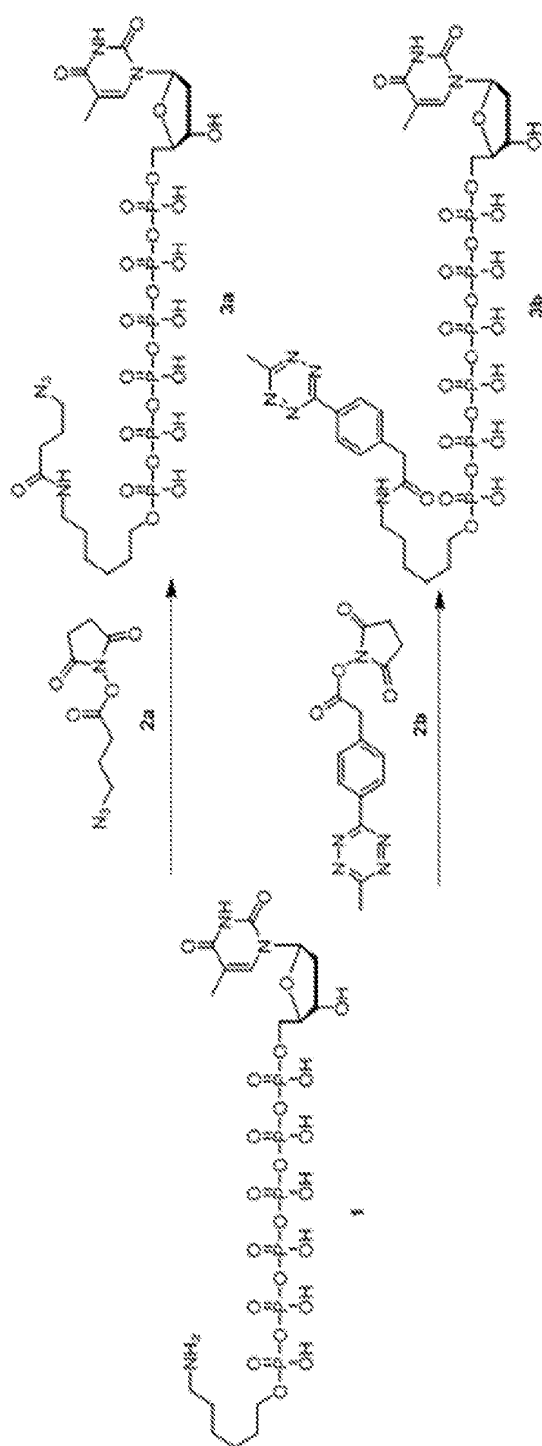






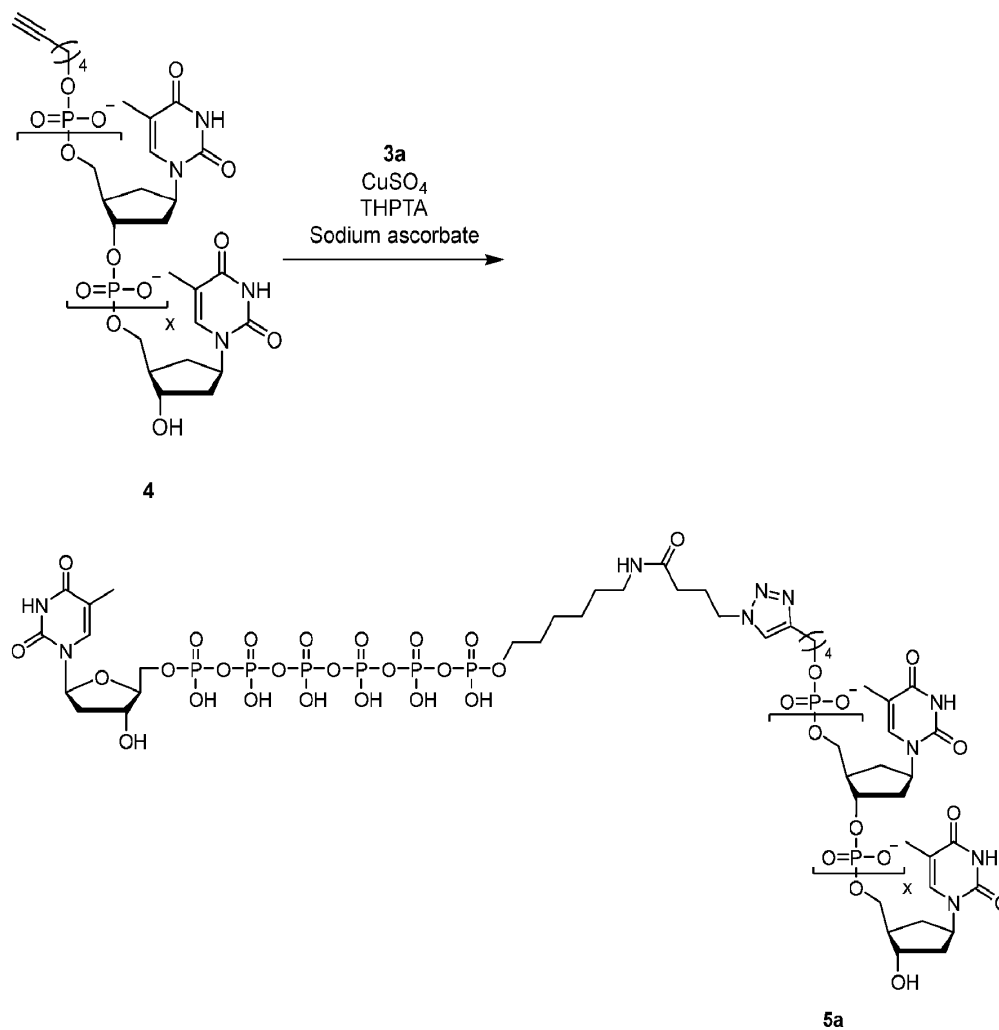
[0234] Phosphodiester based charge tags were synthesized using phosphoramidite chemistry and automated oligonucleotide synthesis. They were purified after synthesis, and then attached to a specific nucleotide via orthogonal chemistry methods. Orthogonal chemistry methods included, without limitation, copper catalyzed alkyne-azide, copper free click chemistry with DBCO and azide, TCO-tetrazine ligation, or thiol-maleimide ligation.

[0235] The non-limiting examples below show the modification of a 5' amino nucleotide hexaphosphate with various linkers to allow for orthogonal attachment chemistry to the phosphodiester charge tags. A 5'-amine deoxy-thymine hexaphosphate (dT6P) (or other NPP) (1) may be functionalized with azido-butyric N-hydroxysuccinimide (NHS) ester (2a) or methyltetrazine NHS ester (2b) to form azide dT6P (3a) or methyltetrazine dT6P (3b) respectively (Scheme 1).



Scheme 1. Functionalization of 5'-amine dT6P.

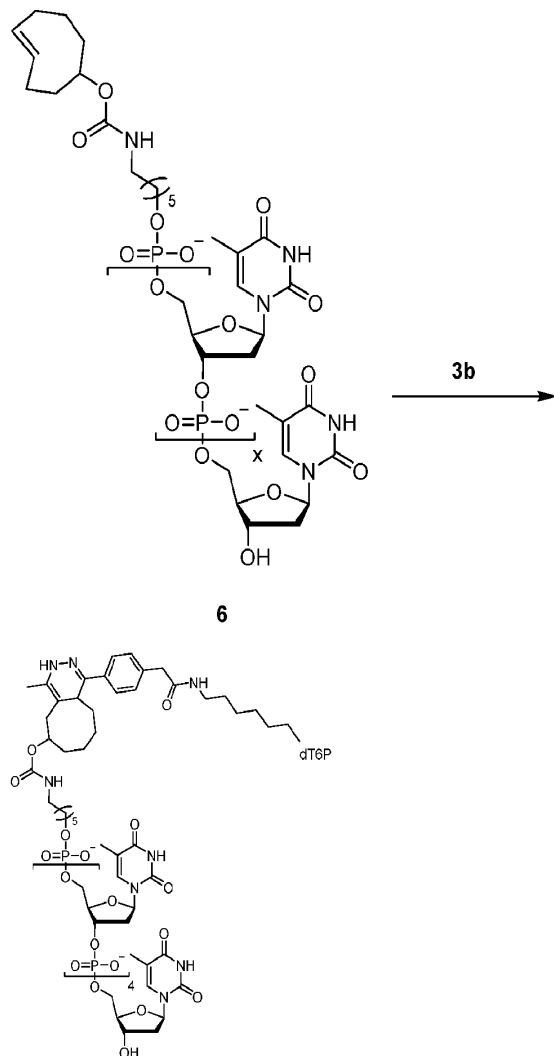
[0236] An azide dT6P (3a) may be conjugated to a linear strand of poly-T oligonucleotide (4) with a 5'-hexynyl group via copper(I)-assisted azide-alkyne cycloaddition (CuAAC) in the presence of CuSO_4 , tris-hydroxypropyltriazolylmethylamine (THPTA) ligand and sodium ascorbate to form an oligonucleotide conjugate (5a). Purification was performed on C18 reverse-phase High Performance Liquid Chromatography (HPLC) and eluted with 50 mM TEAA (pH 7.5) and acetonitrile. A representative example of the CuAAC reaction with poly-T oligonucleotide is shown in Scheme 2.



Scheme2. Representative CuCCA reaction.

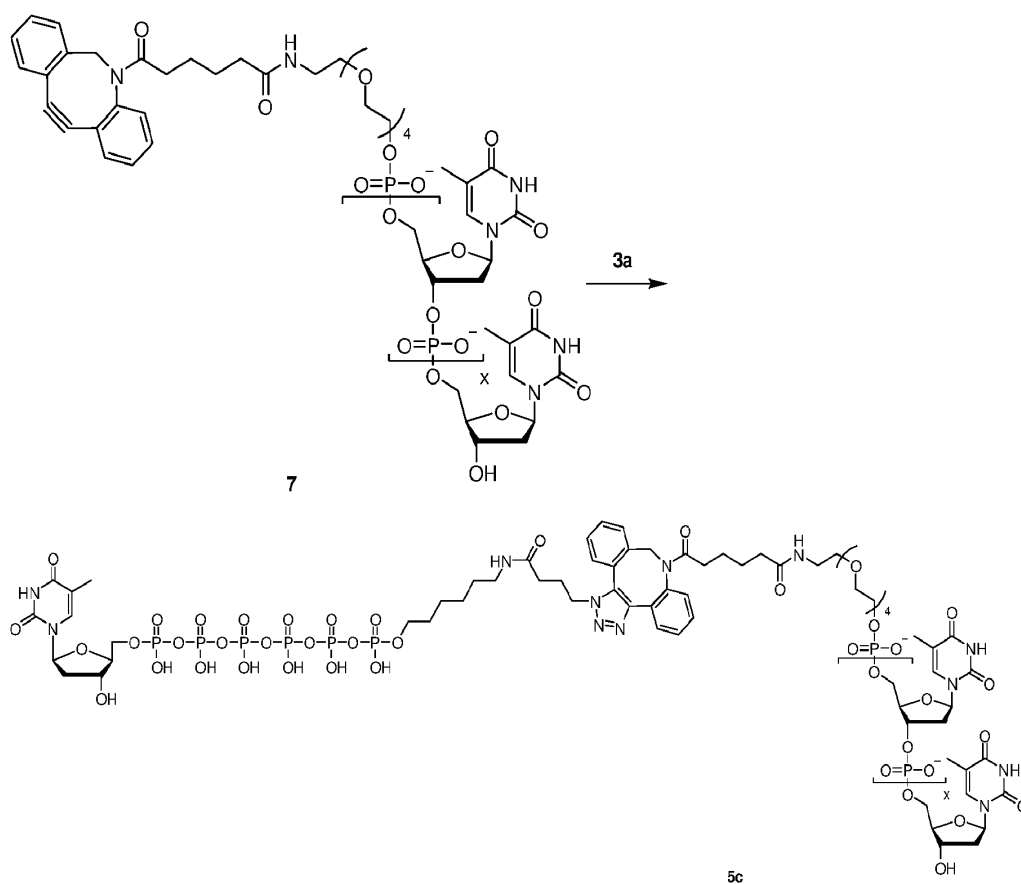
[0237] A methyltetrazine dT6P (3b) was conjugated to a linear strand of poly-T oligonucleotide (6) with a 5'-transcyclooctene (TCO) group in 50 mM phosphate buffer (pH

7.4) to form an oligonucleotide conjugate (5b). The purification was performed on C18 reverse-phase HPLC and eluted with 50 mM TEAA (pH 7.5) and acetonitrile. A representative example of the methyltetrazine-TCO ligation is shown in Scheme 3.



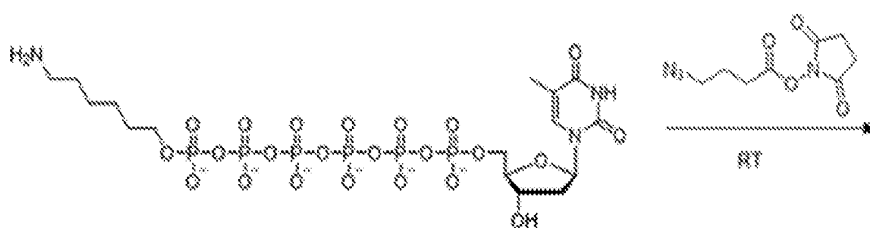
Scheme 3. Representative methyltetrazine-TCO ligation.

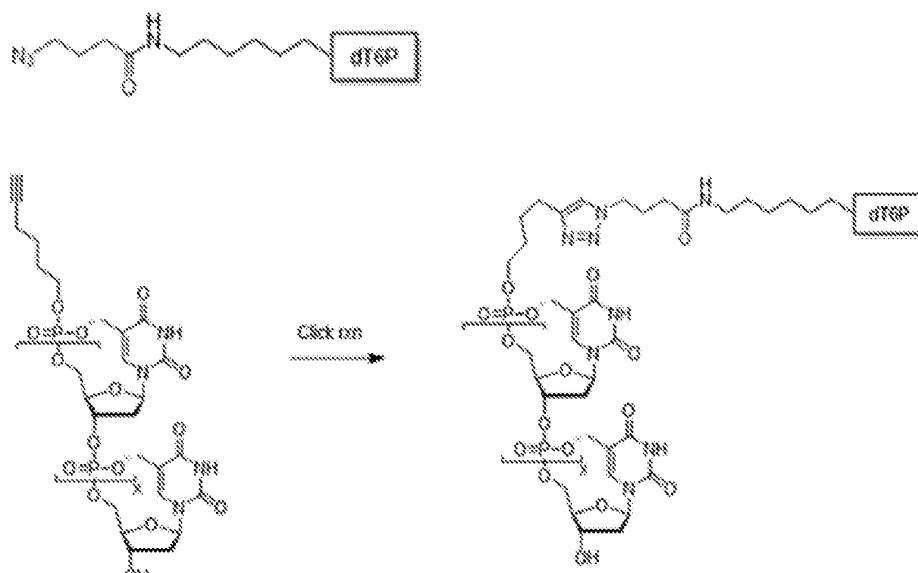
[0238] An azide dT6P (3a) was conjugated to a linear strand of poly-T oligonucleotide (7) with a 5'-dibenzocyclooctyl (DBCO) group via copper-free strain promoted azide-alkyne cycloaddition (SPAAC) in 50 mM phosphate buffer (pH 7.4) to form an oligonucleotide conjugate (5c). The purification was performed on C18 reverse-phase HPLC and eluted with 50 mM TEAA (pH 7.5) and acetonitrile. A representative example of the SPAAC reaction with poly-T oligonucleotide is shown in Scheme 4.



Scheme 4. Representative DBCO-azide conjugation.

[0239] In the following scheme, an azide-alkyne click reaction linked a nucleotide polyphosphate to a charge tag:

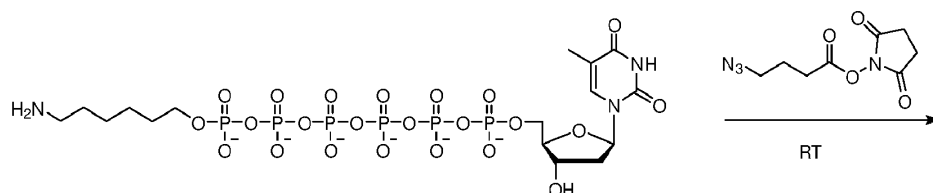


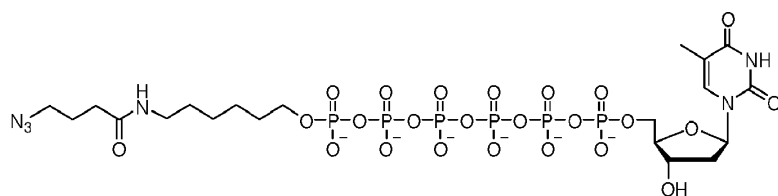


[0240] Scheme 5. Representative conjugation by click chemistry.

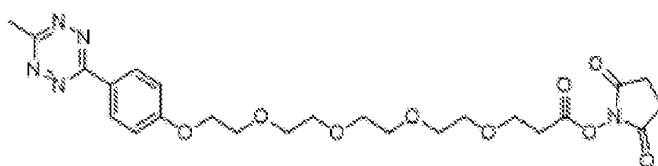
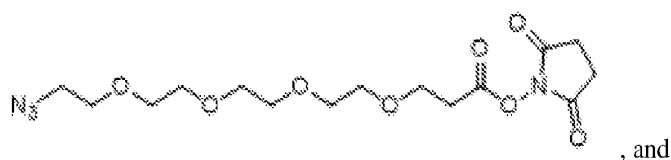
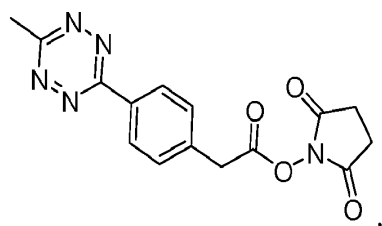
[0241] The foregoing examples may be modified, such as by reversing the placement of each reactive group of a ligation reaction or click chemistry reaction, yielding the foregoing linkages but oriented in the opposite direction with regard to the 5' and 3' ends of the analog nucleotides.

[0242] Reactive groups and linker chemistries may be appended to nucleotides and charge tags according to various applicable chemistries in accordance with the present disclosure. In some non-limiting examples, an azide or methyltetrazine tail may be added to an aminated NPP by reaction with an appropriate NHS residue, which may include linker portions of various lengths such as PEG4 linker, or PEG linker of varying lengths. Non-limiting examples of such synthesis schemes include the following and variations thereof:

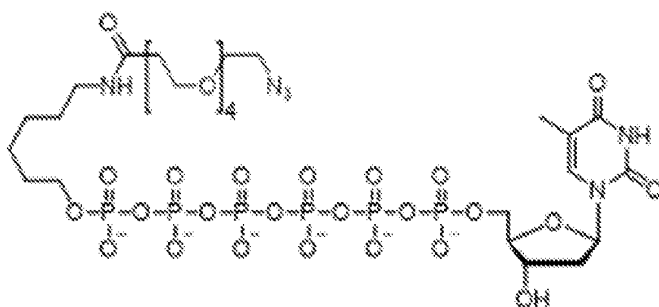




[0243] Different NHS-moieties were used to add an azide or methyltetrazine reactive group, and with various linker lengths. Non-limiting examples include:



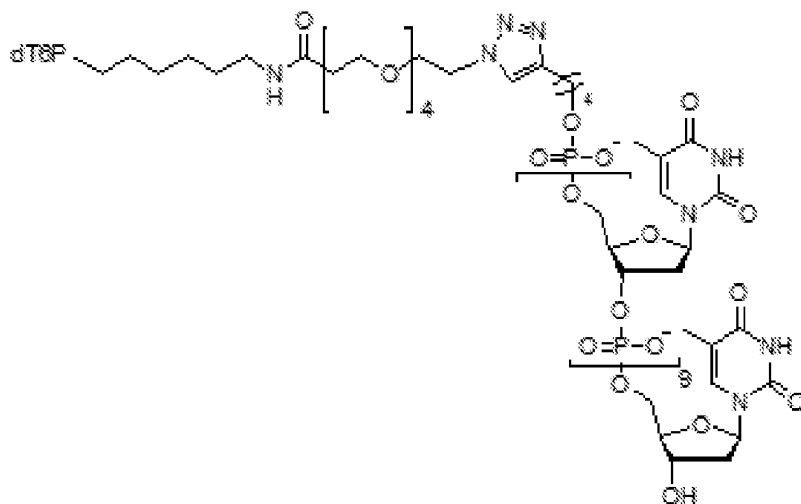
[0244] Various NPPs were formed with different reactive groups for click or ligation chemistry reactions to connect them covalently with charge tags. Some non-limiting examples included:



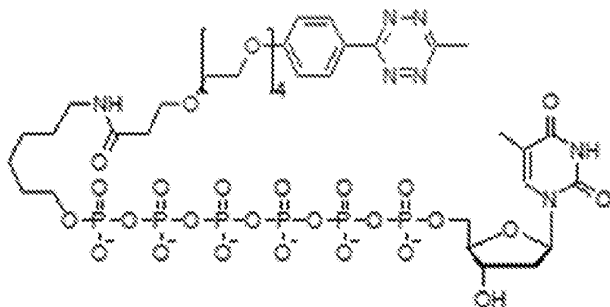
[0245]

reacted with an

alkyne-containing charge tag to create, for example, the following:

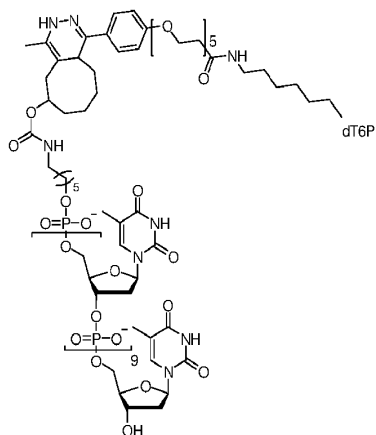


[0246] Alternatively, a methyltetrazine containing NPP such as

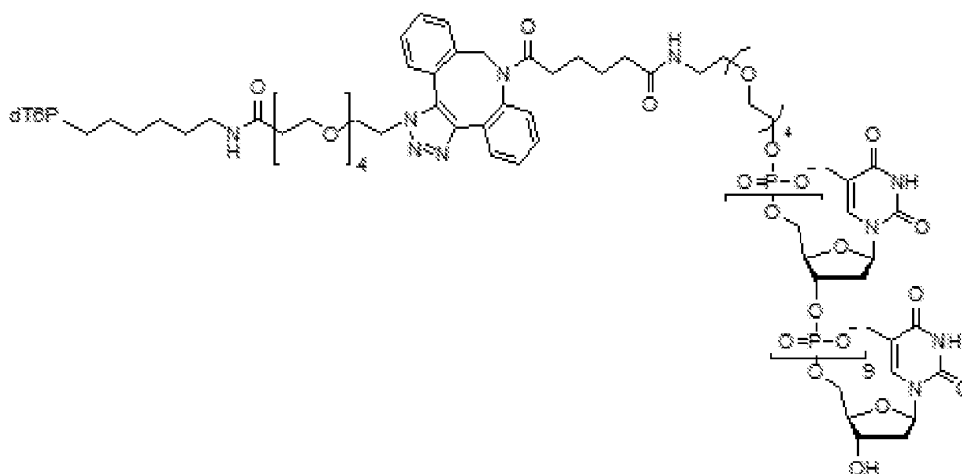


was reacted with a TCO-

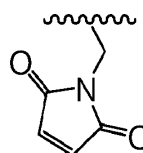
containing charge tag to form the following:



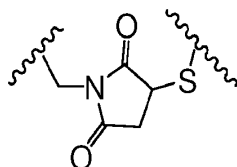
[0247] In other examples, DBCO-azide click chemistry between an NPP and a charge tag was used to form compounds such as the following:



[0248] In other examples, a maleimide group on a nucleotide or charge tag may be reacted with a thiol group on a charge tag or nucleotide, respectively, to link the two via a maleimide-thiol reaction:



[0249] An NPP or charge tag containing a maleimide group reacted with a charge tag or NPP containing a thiol-containing group, respectively, in the presence of a reducing agent such as (tris(2-carboxyethyl)phosphine) resulted in covalent bonding between



the two, for example

[0250] As shown in Table 2, various copper salts, ligands, additives, solvents, reaction durations, and reaction temperatures may be used for different copper-assisted click chemistries.

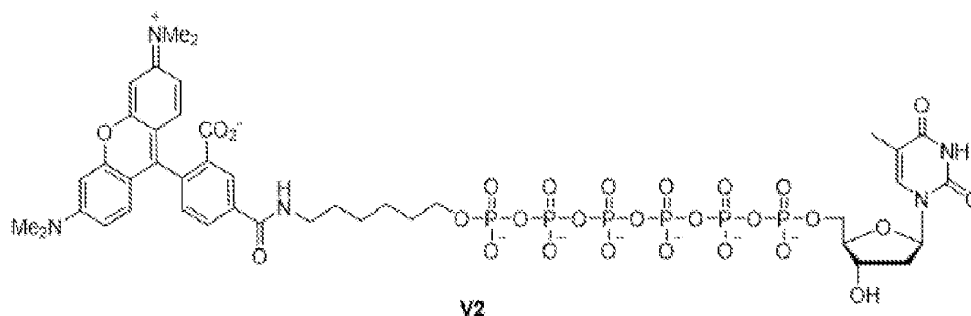
[0251] Table 2: Cu-assisted click chemistries

Cu salt	Ligand	Additive	Solvent	Duration	T Deg. C	Remarks
CuBr (10)	TBTA (20)	-	DMSO/t- BuOH	Overnight	40	No pdt, N ₃ -dT6P recovered
CuSO ₄ (25)	THPTA (50)	Na Asc (50)	H ₂ O	2h	RT	Incomplete rxn, pdt formed
CuSO ₄ (500)	PMDETA (3500)	Na Asc (10000)	H ₂ O	1h	RT	N ₃ -dT6P recovered
CuSO ₄ (500)	THPTA (3500)	Na Asc (10000)	H ₂ O	Overnight	RT	Pdt formed in low yield
CuSO ₄ (25)	THPTA (50)	Na Asc (eq)	H ₂ O	Overnight	RT	Pdt formed, incomplete rxn
					4	Pdt formed, incomplete rxn, highest yield in series
					-20	Pdt formed, incomplete rxn, lowest yield in series
CuSO ₄ (10)	TBTA (20)	Na Asc (200)	DMSO/t- BuOH	Overnight	RT	No pdt, both SMs present
CuSO ₄ (10)	THPTA (20)	Na Asc (200)	H ₂ O	Overnight	RT	Pdt formed, incomplete rxn
CuSO ₄ (100)	THPTA (300)	Na Asc (1000)	H ₂ O	24 h	RT	Pdt formed, complete rxn

[0252] As can be seen, in general terms, in high Cu loading, a reaction may run to completion but with low yield. Comparatively, with low Cu loading, a reaction may not run to full completion but yield may be higher. With intermediate Cu loading, a reaction may run

to completion and reaction product may be isolated in 86% yield by HPLC.

[0253] Incorporation of phosphodiester based charge tag modified nucleotides have been demonstrated. Incorporation may be carried out with different polymerases such as phi29 (and variants thereof) and Klenow fragment, or others used in sequencing-by-synthesis processes. Both polymerases can incorporate the charge tags successfully. Incorporation by phi29 for this example is shown in FIG. 13. In this example, single-stranded DNA template polynucleotide sequences were in solution were incubated with a buffer solution (50 mM Tris pH 7.5, 5 mM MnCl₂, 4 mM DTT) containing 100 nM 5'-Cy5-labeled DNA primers (16-mers) complementary to a portion of such template sequences, 1 μM phi29, and 10 μM of a given nucleotide for single-nucleotide incorporation into the primers based on the template. Following incubation for various durations at 30 degrees Celsius, to allow 5' incorporation of a charge-tagged thymidine (complementary to adenosine residue on the template strand immediately 5' to the portion complementary to the primers), polymerase reaction was quenched, primers were dehybridized and separated on a gel for detection of single-nucleotide incorporation. Linkages between deoxyribo-thymidine 5'-hexaphosphate (dT6P) included T5, T10 and T15, having the indicated repeats of thymidine nucleotides as charge tags; T5, T10 and T15 are attached via click chemistry, while T5-Tet is attached via tetrazine-TCO ligation. C3 Spacer (C3) and dSpacer (d) oligos were also used as charged tags, attached via TCO-tetrazine ligation. TMR is a tetramethylrhodamine-labeled dT6P with the following formula:



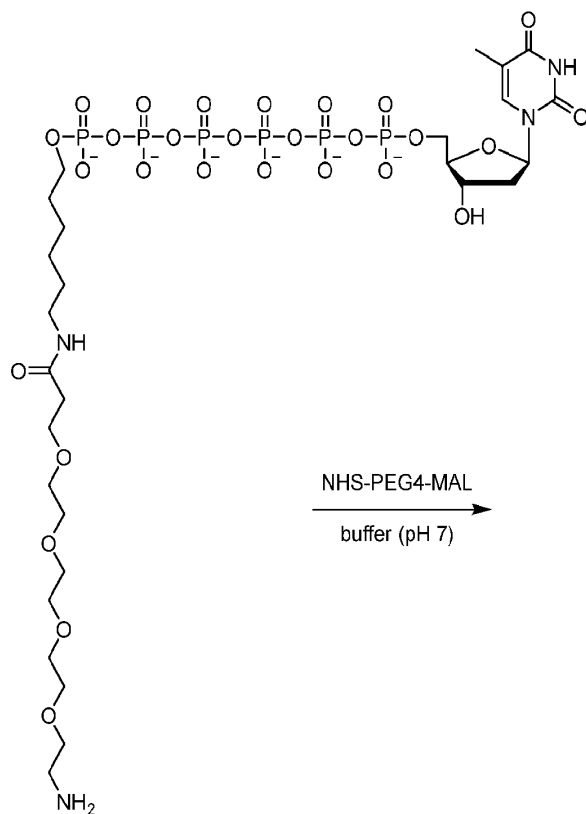
and dTTP is deoxy-thymidine triphosphate without a charge or label to serve as a control.

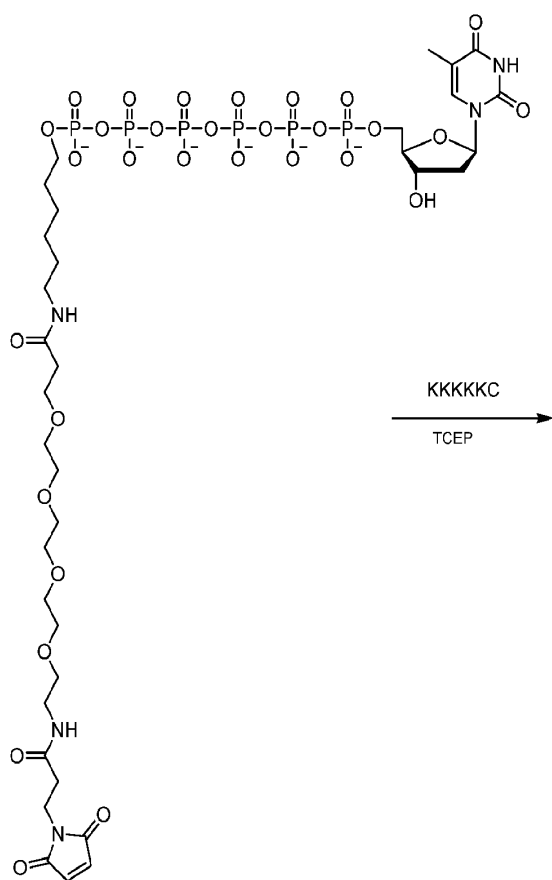
[0254] Referring to FIG. 13, 1310, 1320, and 1330 each individually represents % incorporation of dTTP, T10, or T5, respectively (whose plots overlap with each other and are therefore nearly indistinguishable from each other in FIG. 13), 1340 represents incorporation by T15, 1350 represents incorporation by T5-Tet, and 1360, 1370, and 1380 each

individually represents % incorporation of TMR, d, and C3, respectively (whose plots overlap with each other and are therefore nearly indistinguishable from each other in FIG.

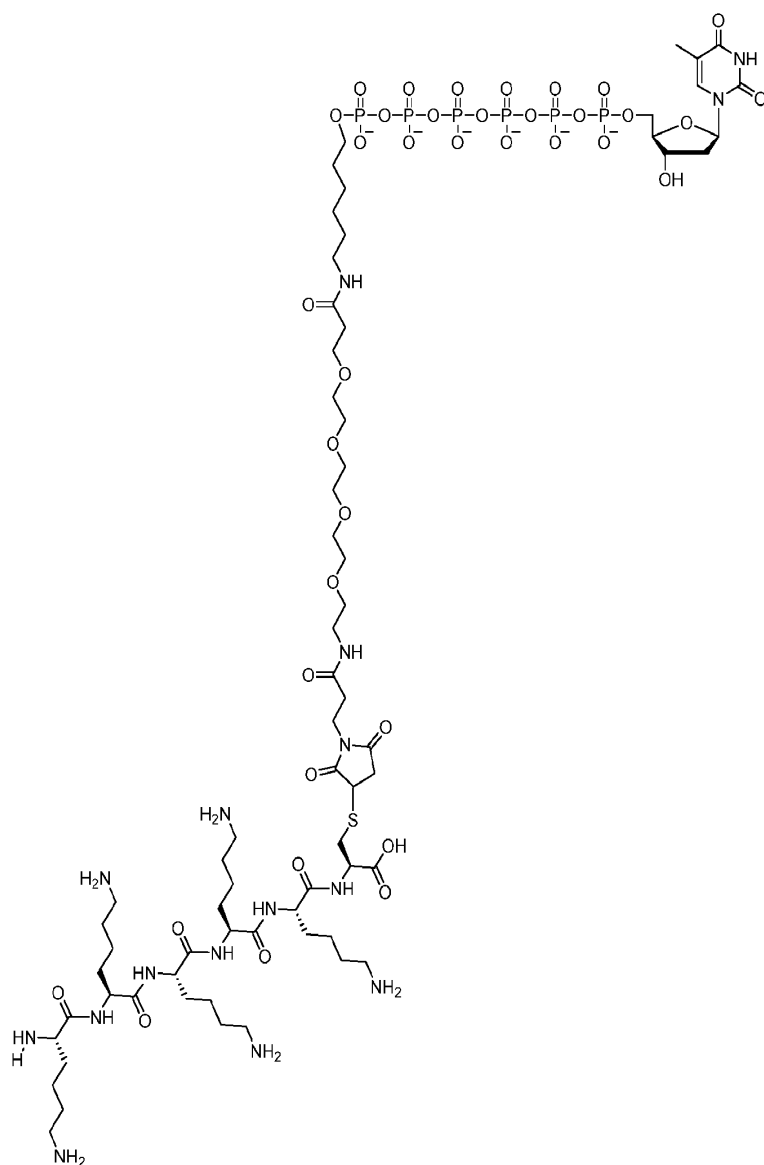
13). Incorporation for all examples exceeded 80% within 5 seconds.

[0255] A non-limiting example of a synthesis scheme used to synthesize a nucleotide analog with a peptide charge tag in accordance with the present disclosure is shown below

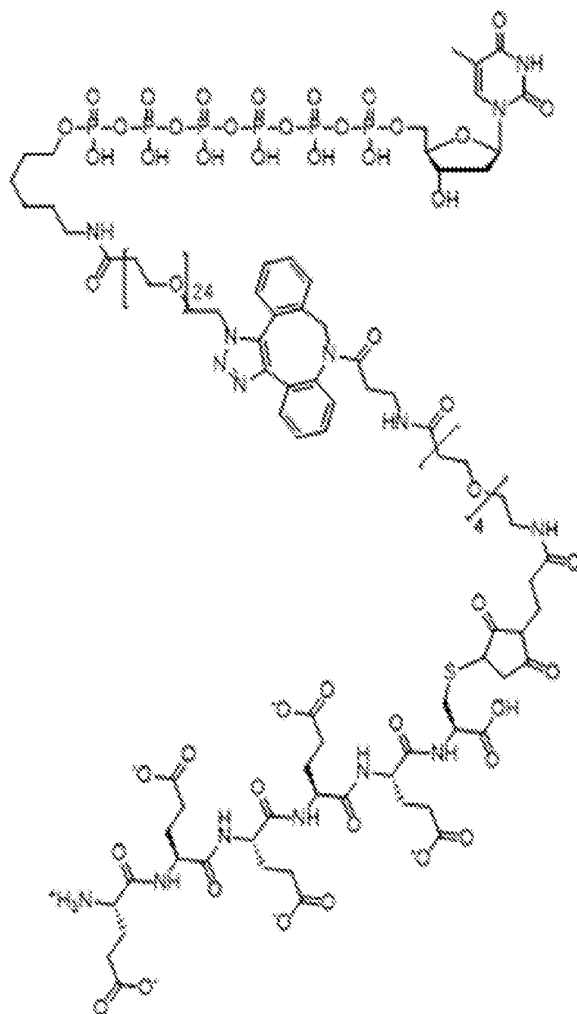




("KKKKKC" disclosed as SEQ ID NO: 10)

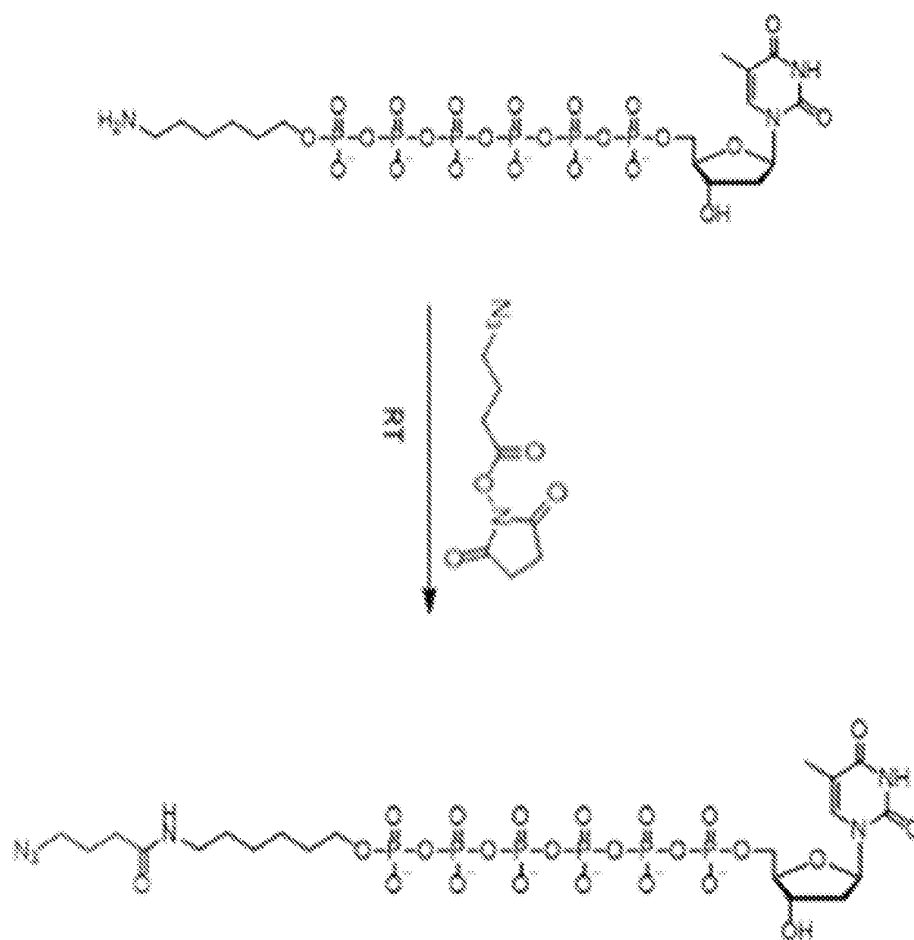


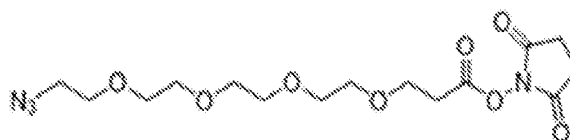
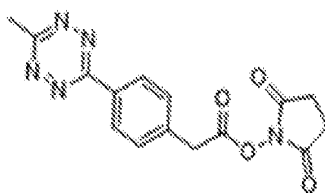
[0256] In another example, rather than positively charged lysine residues as illustrated above, a comparable scheme was used for synthesizing a charge tag with negatively charged amino acids such as:



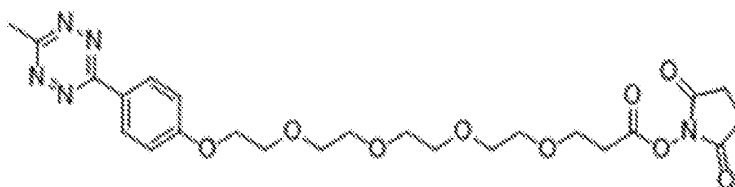
[0257] Many variations on this scheme are possible within keeping with the teachings of the present disclosure. For example, amino acids other than lysine, including any of the charged or uncharged amino acids described above may be employed, and they may number more or less than the peptide charge tag length shown in this example. Peptide charge tags with charges of different valences and magnitudes may therefore be employed.

[0258] Furthermore, different reactive groups may be added to the 5' end of a nucleotide analog, and with different types and lengths of linker portions, such as, for additional non-limiting examples:



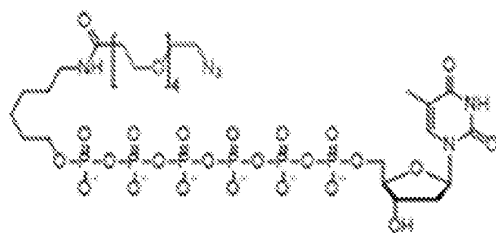
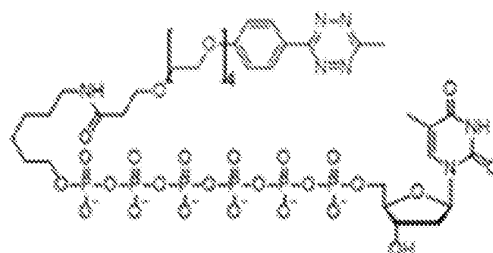


Azido-PEG4 NHS Ester



Methyltetrazine-PEG4 NHS Ester

[0259] These additions may yield nucleotide analogs with various reactive groups including azide or methyltetrazine reactive groups, appended by linkers of various types and lengths, including, as non-limiting examples:

dT6P-PEG4-N₃

dT6P-PEG4-methyltetrazine

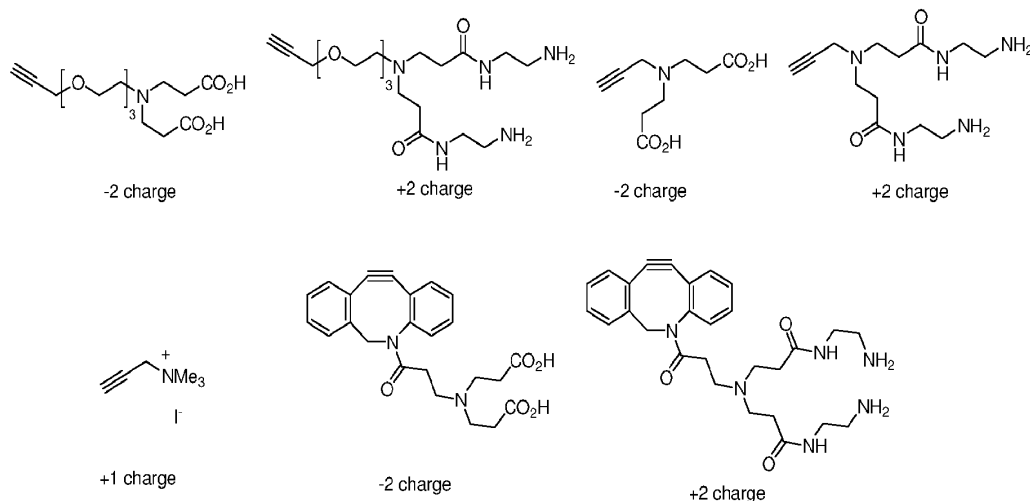
[0260] In still other examples, an alkyne, TCO, or DBCO group was similarly added, or a thiol group. A corresponding reactive group could then be added to a peptide charge tag such that the two could be joined by the above-disclosed click or ligation chemistries, or others known to skilled artisans. Peptide based charge tags can be synthesized using

fluorenylmethyloxycarbonyl (Fmoc) and tert-butyloxycarbonyl (Boc) protecting group chemistry for solid phase peptide synthesis. An orthogonal “handle” reactive group can be introduced in the peptide synthesis at the terminal end to allow conjugation to a nucleotide or nucleic acid. Orthogonal chemistry methods include azide-alkyne copper-assisted click reaction, copper free click chemistry with DBCO and azide, and TCO-tetrazine ligation. Reactive side chains of amino acids such as thiol of cysteine can also be used in thiol-maleimide chemistry.

[0261] The availability of amino acids containing side chains with different pK_as also allow peptide charge tags charged at different pHs. For instance, histidine has a pK_a of 6.04, while Lysine has a side chain with a pK_a of 10.54. Thus, at neutral pH, only lysine could be charged. This also allows further modulation of the number of charges and charge density by modifying the pH of the buffer environment.

[0262] In addition, peptide charge tags can be easily appended to peptide nucleic acid (PNA) oligomers, since both peptides and PNAs are synthesized with the same solid phase peptide chemistry. This may be used to further modify the properties of the peptide charge tag, or add association properties of the charge tag to linkers such as nucleic acid based linkers.

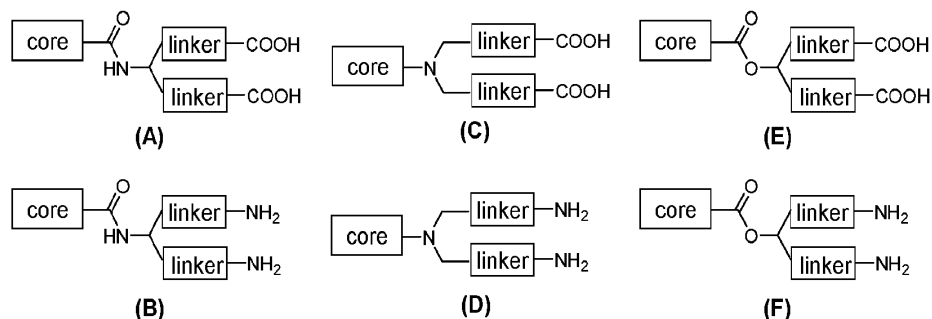
[0263] Examples of compounds used in the synthesis of a dendron charge tag, and corresponding charges per terminal constitutional repeating unit, include the following:



[0264] In these examples, different reactive groups are shown at the free valence end of the dendron, as well as different potential stem lengths between a branch point and a free terminal end of an individual constitutional repeating unit, but these are merely non-limiting

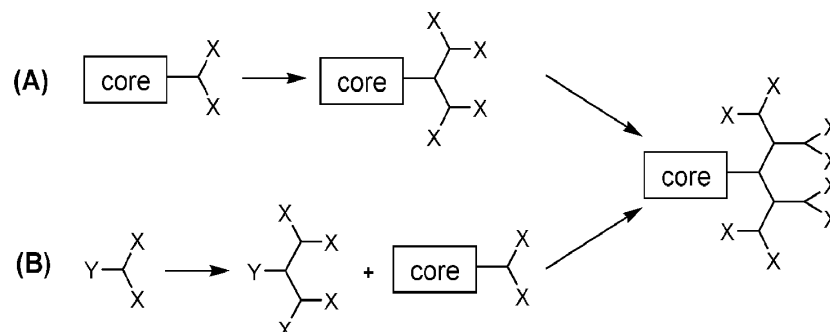
examples.

[0265] The following scheme provides illustrative examples of possible dendron charge tag structure:



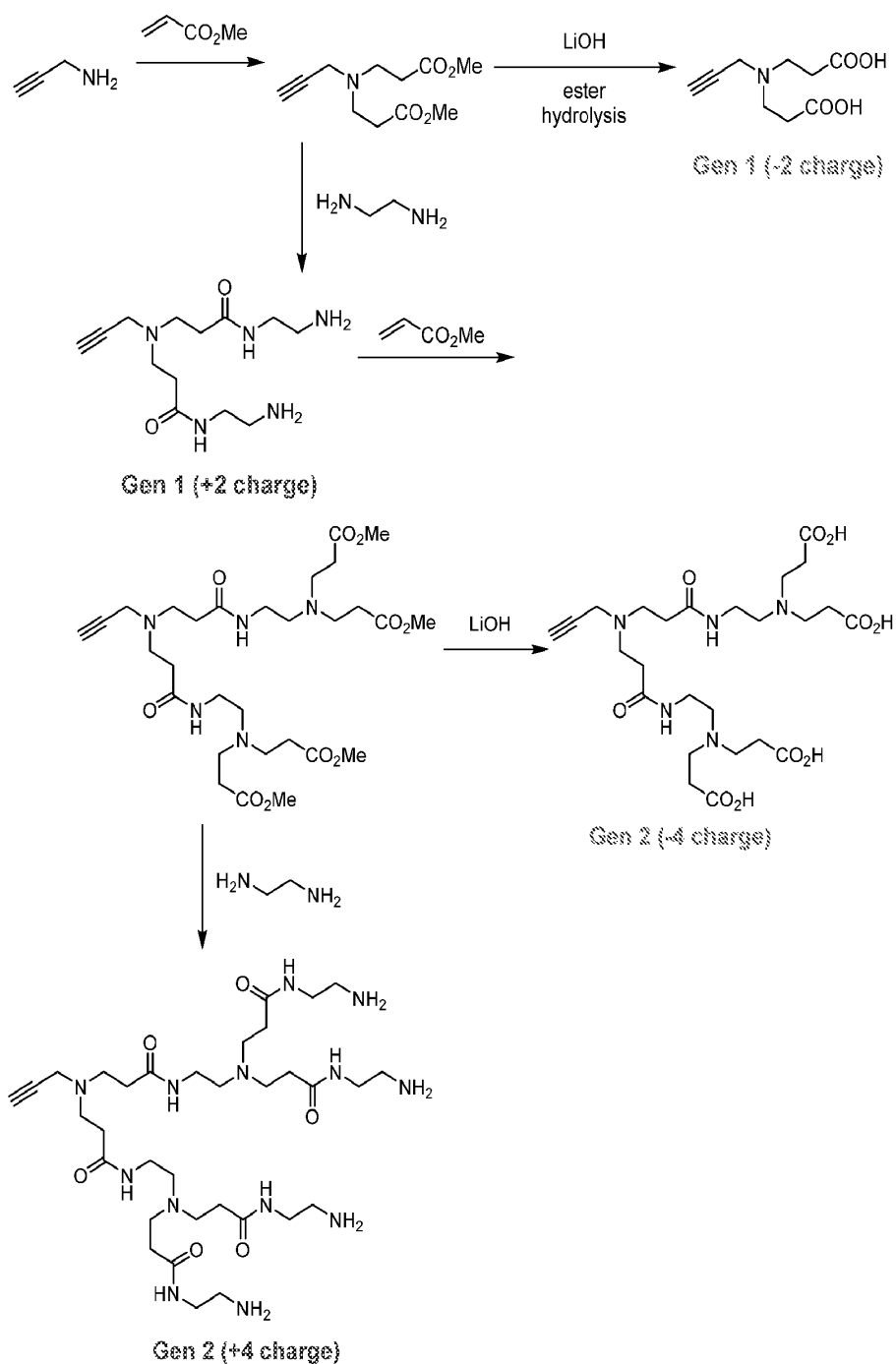
[0266] Shown are, for example, dendron with amide linkages and (A) terminal carboxylic acid or (B) amino groups; dendron with poly(propylene imine) (PPI) linkages and (C) terminal carboxylic acid or (D) amino groups; and dendrons with ester linkages and (E) terminal carboxylic acid or (F) amino groups.

[0267] Generally, dendron charge tags may be synthesized according to divergent or convergent synthesis methods, according to the following representative schemes:



[0268] In divergent synthesis (A), a dendron is assembled by a series of outwards extending reactions from the core, usually by repetitive Michael addition. In convergent synthesis (B), a dendron is constructed by a series of inwards building reactions from the peripheral and eventually attached to the core.

[0269] Some examples of such divergent synthesis schemes in accordance with the present disclosure were as follows:

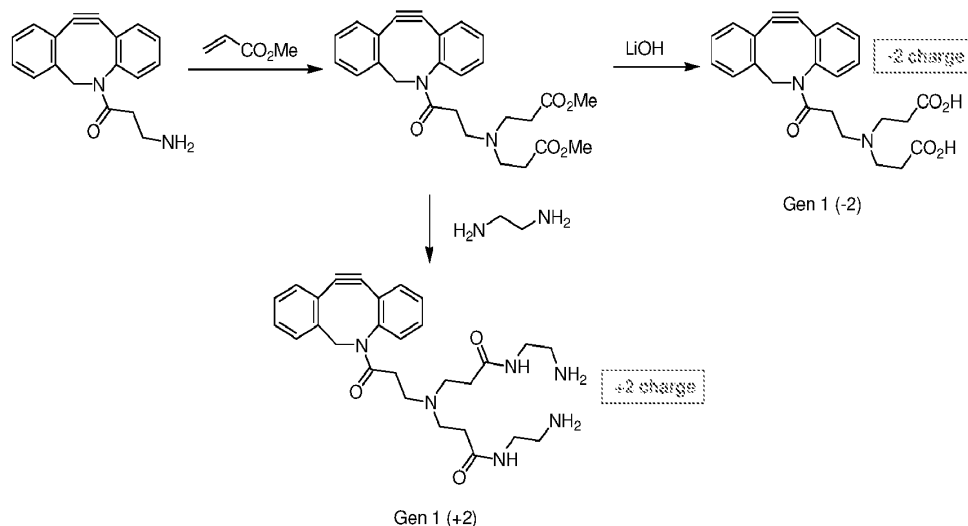


[0270] In these examples, a methacrylate group was added by Michael addition to an alkyne stem, followed by either deprotection of acetyl groups to form the carboxylic acid groups, or addition of ethylenediamine to form the amino groups. Repetitive cycles of Michael addition

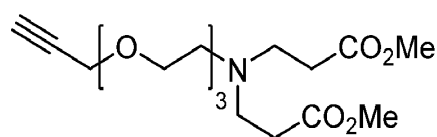
resulted in successive generation of dendrons with twice the number of terminal functional groups compared to the previous generation. Additional generations may be added, and a different reactive group could be used at the stem/free valence end. In some examples, an additional generation or more may be iteratively added according to the foregoing synthesis schemes to increase charge carried by a tag. Valence (positive or negative) of a charge may be varied by incorporating a positively or negatively charged amino acid at an end group. Examples are shown in FIGs. 21A and 21B. In both non-limiting examples, a charge tag terminating in a cysteine residue is shown, which could be linked to a linker section for charging a nucleotide as disclosed herein, though other chemistries such as disclosed herein are also intended as examples. In FIG. 21A, positively charged lysine residues for the end groups following either 2, 3, or 4 branchings, yielding different terminal charge magnitudes. Alternatively, as shown in FIG. 21B, a negatively charged amino acid such as glutamate could form end groups after various generations of branching, again yielding different magnitudes of terminal charge.

[0271] In another example, one or more lysine residues in a charge tag may be methylated (e.g., trimethylated). Unlike unmethylated lysine, the charge of trimethylated lysine is not pH-dependent.

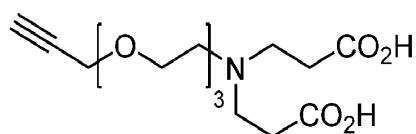
[0272] Another example, with a DBCO at the free valence end, is as follows:



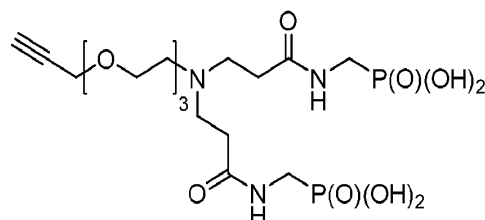
[0273] Some examples of amide-based and PPI dendron designs for dendron charge tags and their synthesis include the following:



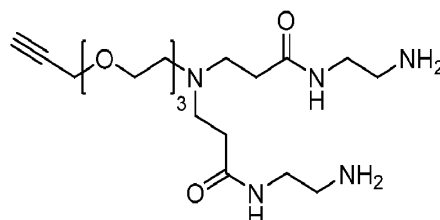
A-1



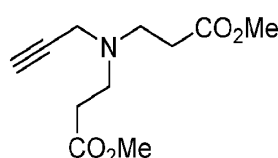
A-2



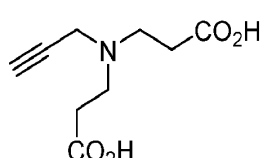
A-3



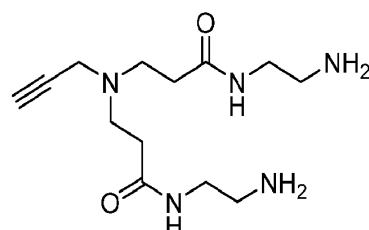
A-4



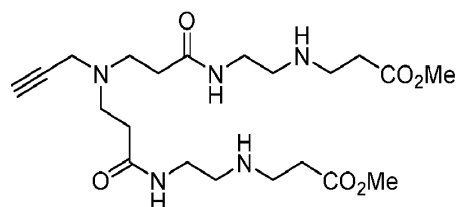
B-1



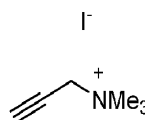
B-2



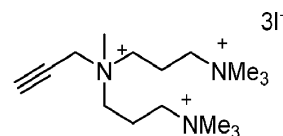
B-3



B-4



C-1

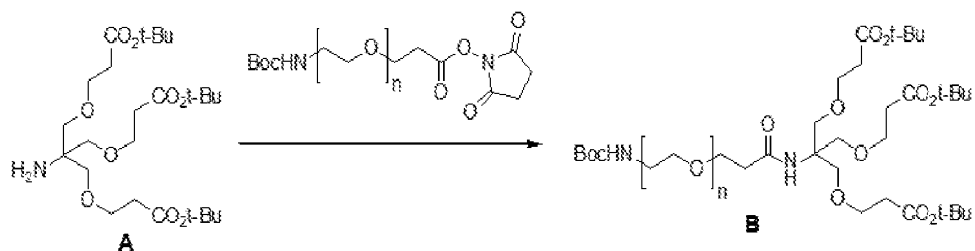


C-2

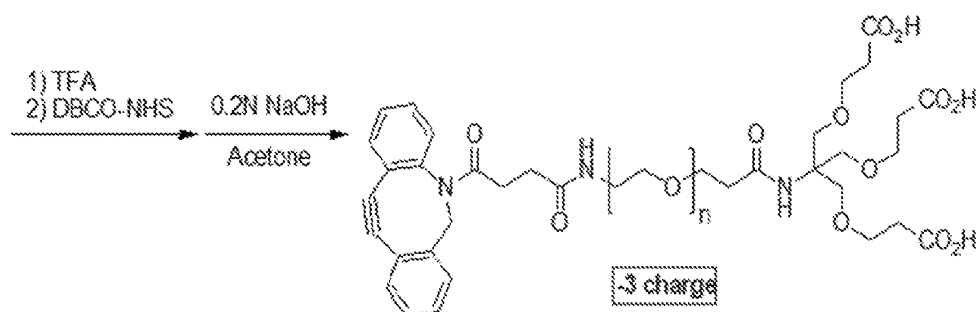
[0274] Some advantages of quaternary ammonium groups included in examples C-1 and C-2 are that they may not be affected by pH, may not coordinate metals, and may be less likely to attach to poly(vinyl phosphonic acid) (PVPA) during synthesis and handling.

[0275] In another example, a constitutional repeating unit with three degrees of branching may be used. In yet a further example, convergent synthesis may be used rather than divergent synthesis. A benefit of using a unit with three degrees of branching is that more charges may be added per generation, compared to a dendron with units having only two degrees of branching, resulting in fewer generations required to attain a given preferred charge. An

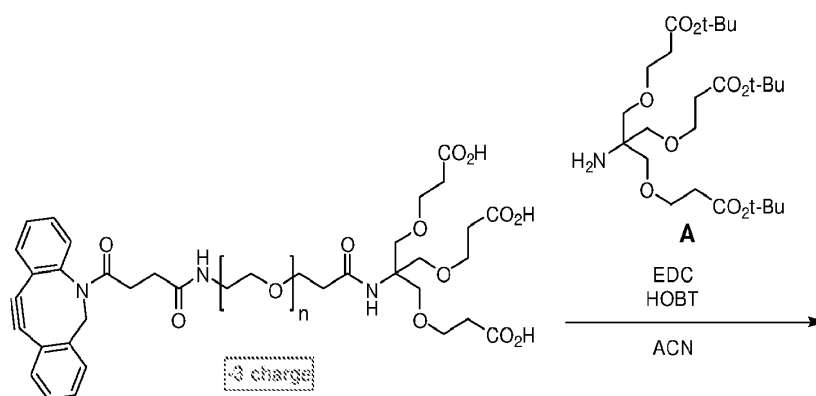
example was as follows:

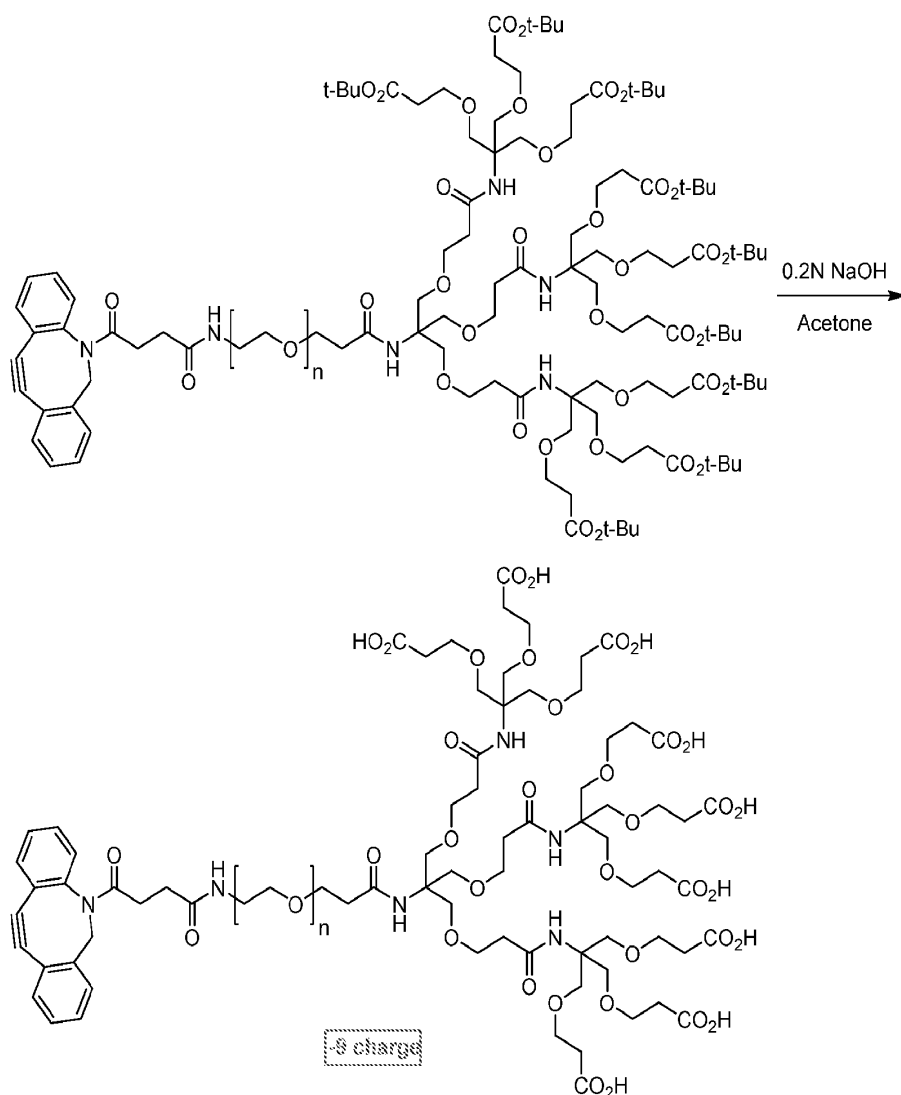


[0276] In this example, a constitutional repeating unit is functionalized with a tert-butyloxycarbonyl (Boc) group. Subsequently,

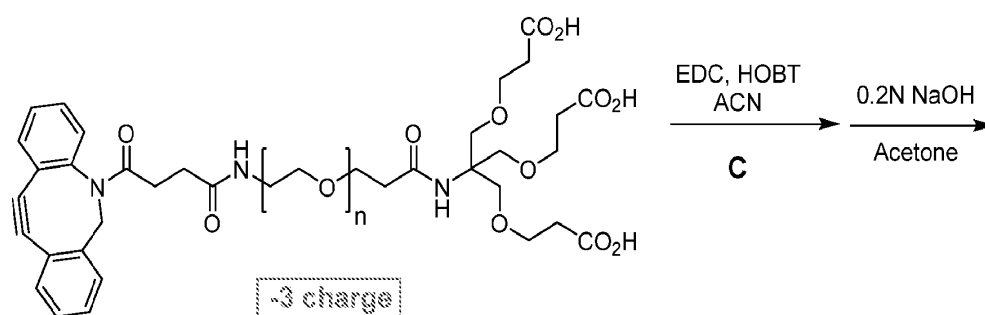
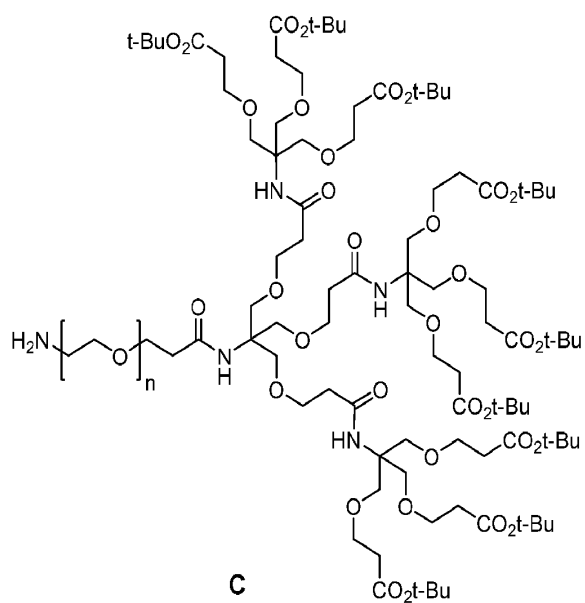
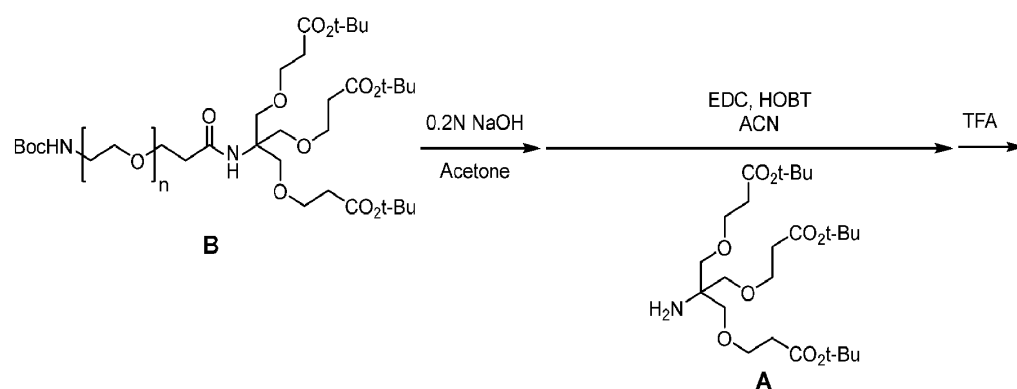


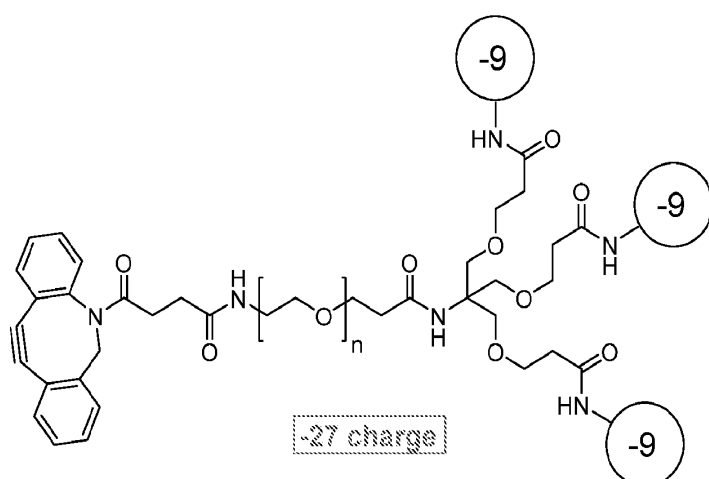
a DBCO group may be added and, upon deprotection of acetyl groups to form the carboxylic acid groups. The resulting compound has, in this case, a -3 charge. In a subsequent reaction, compound the compound A above was added in a second generation dendron, to give a charge of -9, as follows:



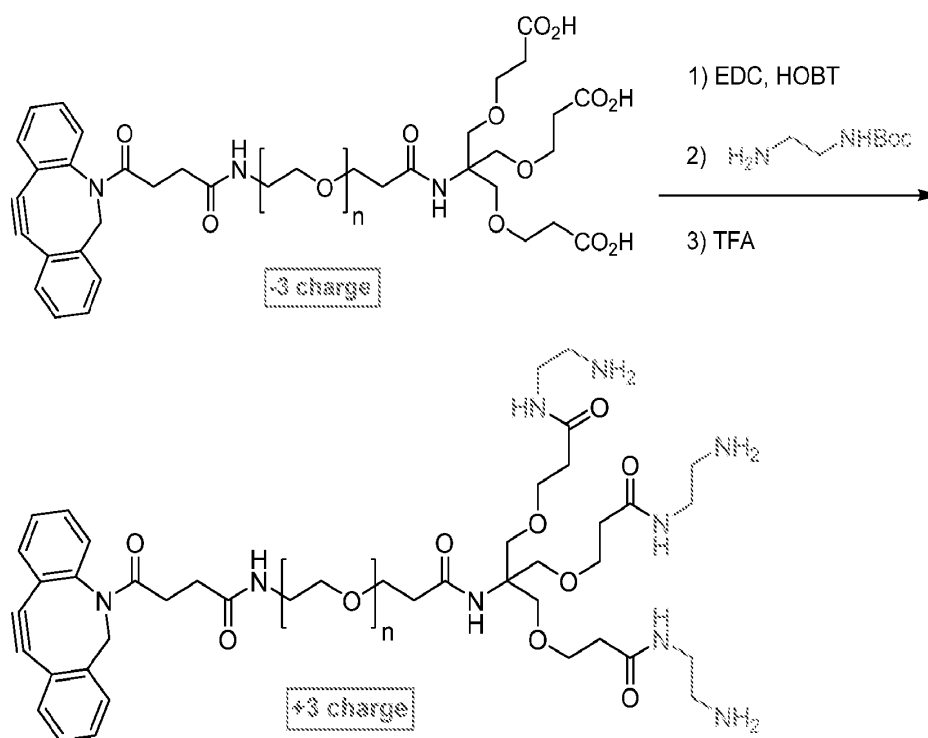


[0277] By iteratively combining the foregoing steps, three second degree dendrons can be combined, to create a third generation dendron with a charge of -27, via convergent synthesis, according to the following example:

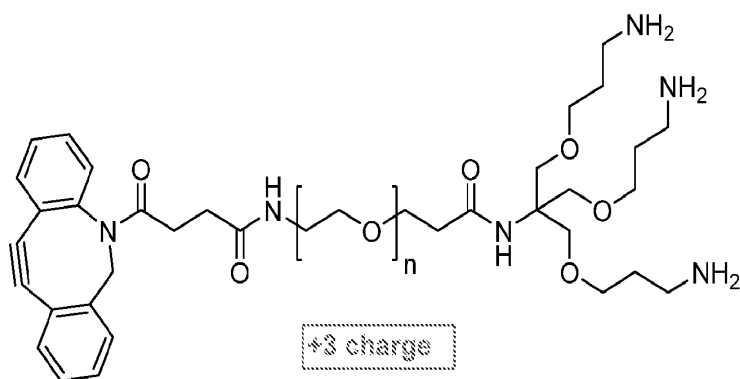
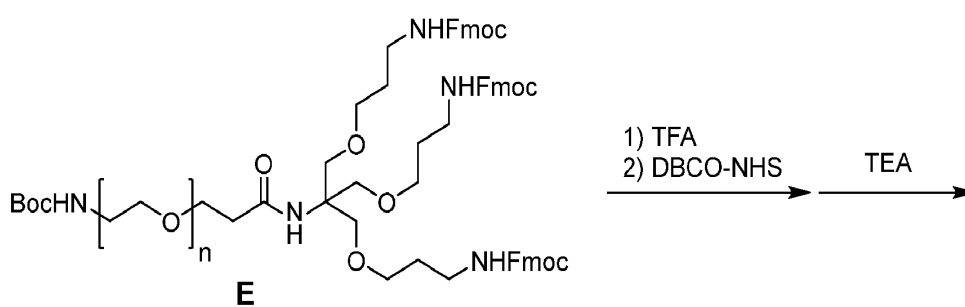
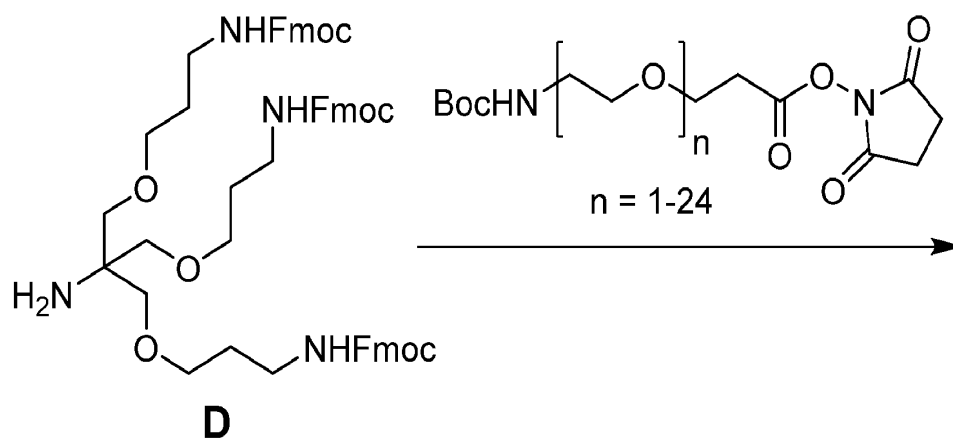




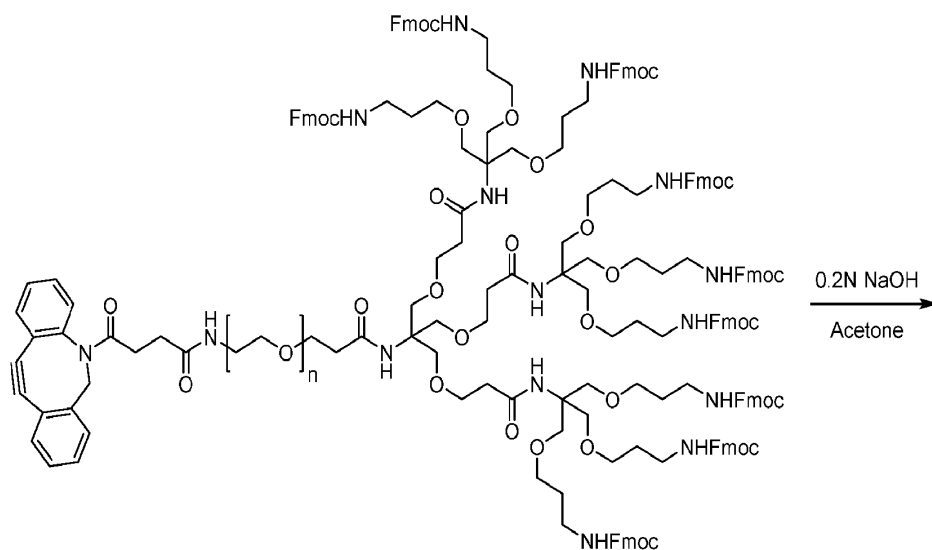
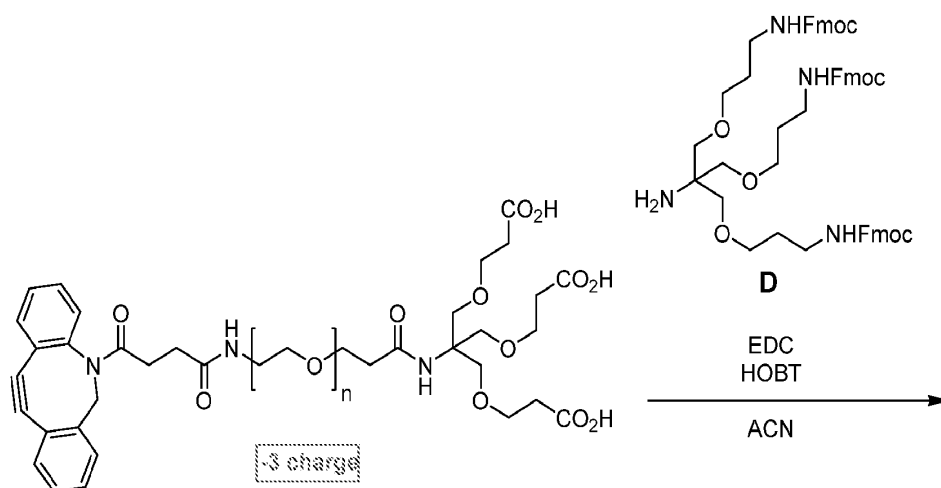
[0278] A dendron bearing negatively charged carboxylic acid groups was converted to a dendron bearing positively charged amine groups as follows:

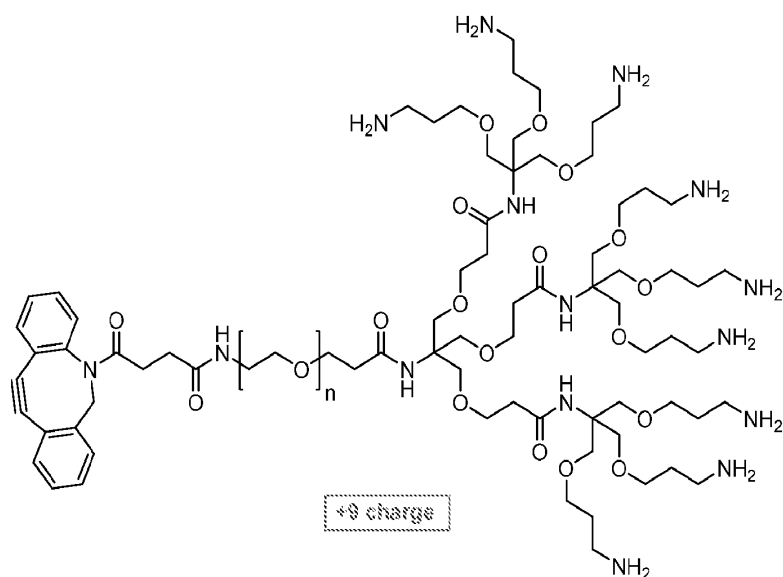


[0279] In another example, carboxylic acid groups was converted to amine groups according to the following scheme:

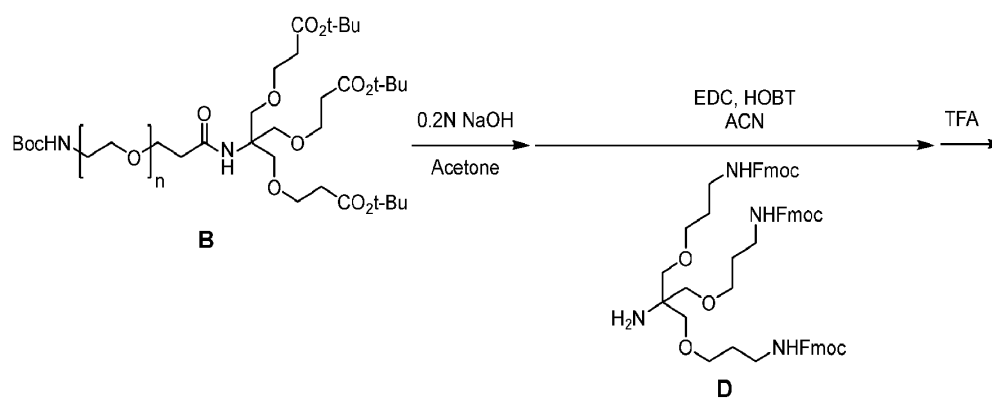


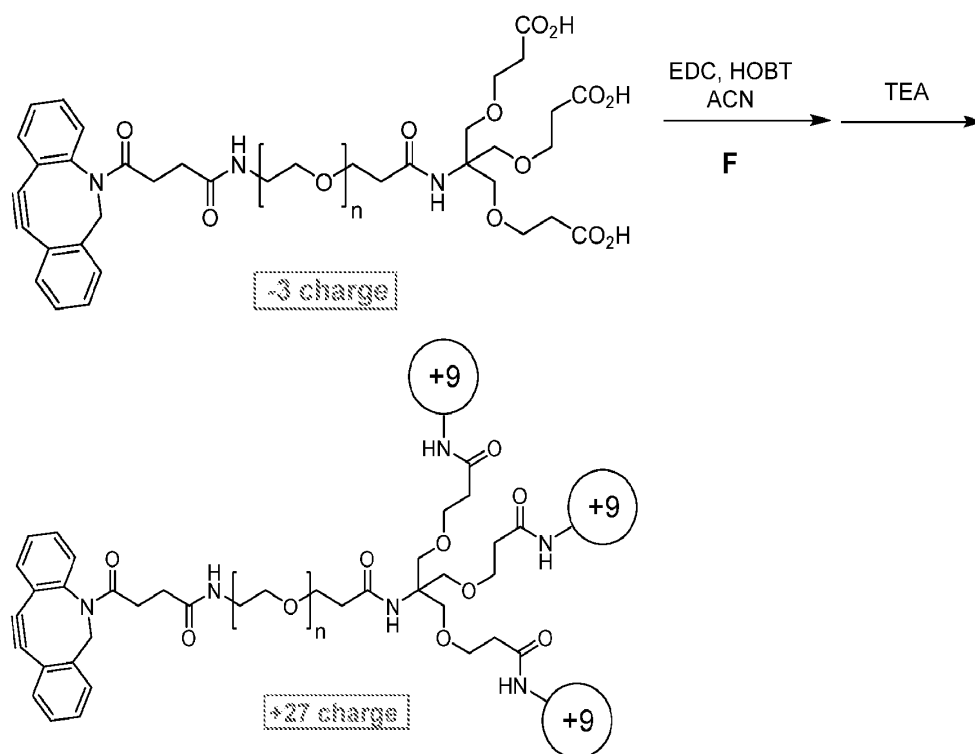
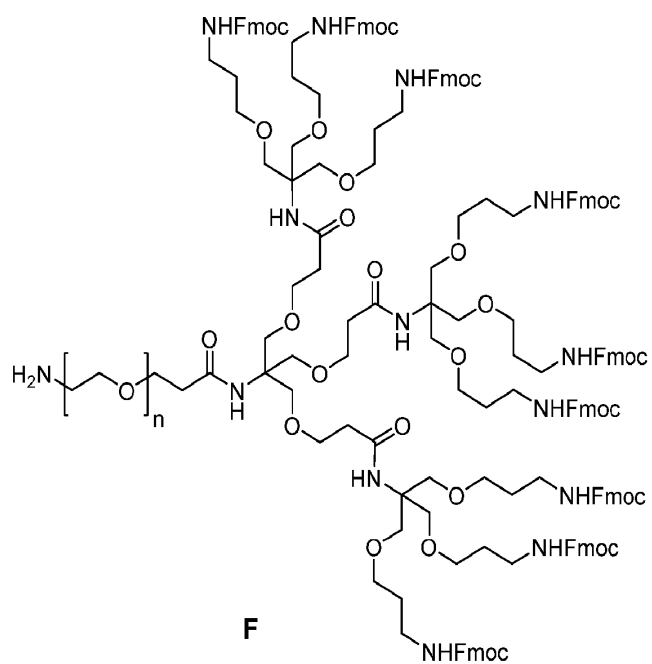
[0280] For a second generation, with a +9 charge, the following scheme may be used:





[0281] And, a third generation dendron may be synthesized, by a convergent synthesis scheme, to generate a dendron with +27 charge, as follows:





[0282] Depending on the reactive groups at the free valence end of a dendron synthesized in

accordance with the present disclosure, which may include without limitation any of the examples described above, a corresponding paired reactive group may be appended to a nucleotide analog to allow ligation of the charge tag dendron to the nucleotide analog. According to the foregoing, a wide range of charges may be included in a nucleotide analog, including -32, -27, -16, -9, -8, -4, -3, -2, +2, +3, +4, +8, +9, +16, +27, and +32. Charged functional groups other than those illustrated in the foregoing non-limiting, example synthesis schemes may also be used.

[0283] In some examples, such branching structure may be used to add multiples of phosphodiester-based charges to a charge tag. For example, rather than a single linear strand of polynucleotide or other phosphodiester-containing charge as disclosed herein, a branched structure such as according to a dendron structure as shown here may include as an end group a nucleotide or polynucleotide. By basing branching of such phosphodiester-containing tags in successive generations in accordance with a dendron structure as disclosed herein, multiple polynucleotides or other phosphodiester-based charges may be combined into a single charge tag. For example, dendron-based structures such as shown in FIG. 19A and B. FIG. 19A shows an example of a tag combining three poly-T sequences into a single tag, which can be incorporated into a compound of Formula I according to methods as disclosed herein. In this example, the tag carries a charge of -30. FIG. 19B illustrates several ways of combining phosphodiester-containing tags to yield a given charge (in this example, -30): a linear sequence of 30 phosphodiester charges, a triply-branched structure terminating in three phosphodiester sequences of 10, or a structure twice branched trebly and terminating in 6 phosphodiester sequences of five. An advantage of increased branching, such as in the last example as compared to the first, may be a higher density of charge, with a higher concentration of short charged sequences in proximity to each other as opposed to a single extended sequence which could extend away from a conductive channel.

[0284] In other examples, a branched fork structure based on an amino acid may be included in a charge tag. An example of a synthesis scheme for such fork and branch, amino acid based charge tag structure is depicted in FIG. 22. Solid phase peptide synthesis may be used to a sequence of amino acids together in, for example, a linear polypeptide chain according to the upper panel of FIG. 22. By iterative protection of the free amino group, followed by its removal and addition of an activated amino acid, a linear chain polypeptide may be synthesized. Linear strands of charged amino acids (positive or negative) may thereby be

generated as charge tags or part of charge tags for attachment to a nucleotide. A non-limiting example is depicted in FIG. 24A, which shows a strand of 16 negatively charged glutamate residues (SEQ ID NO: 12). Other examples could include other charged amino acids and in different numbers.

[0285] As further depicted in the bottom panel of FIG. 22, a convergent solid phase protein synthesis method may be used wherein lengths of individually formed polypeptide chains may be added as polymer sets during a synthesis step. In some examples, not depicted in FIG. 22, amino acids or polypeptides may be independently added as branches to a fork where the fork is a structure containing two amino groups, rather than concatenated linearly from a single amino group as depicted in FIG. 22. For example, a lysine amino acid, having two amino groups as shown in the left structure in FIG. 23A, may serve as a forked attachment points to which two branches of a linear polypeptide may be attached, one to each amino group according to the solid phase protein synthesis scheme depicted in FIG. 22.

[0286] By adding two strands of amino acids, each synthesized according to, for example, a solid phase protein synthesis scheme as per the upper panel of FIG. 22, to the two amino groups of a lysine residue, a charge tag including a fork with two branches may be synthesized where each branch is a polypeptide. The branches may be positively charged or negatively charged depending on the charge of the amino acids of which they are constituted. Two non-limiting examples are depicted in FIG. 23. On the left, two penta-lysine branches are attached to a lysine fork to form a positively charged charge tag, whereas on the right, two penta-glutamate branches are attached to a lysine fork to form a negatively charged charge tag. Other similar examples may have longer or shorter branches, with anywhere from 1 to 20 amino acids, or more. In an example, each branch may have a number of amino acids that differs from the number of amino acids in the other. The species of amino acids in each branch may also differ, between and/or within branches, as well.

[0287] In still other examples, multiple forks may be attached to a central fork to create trees with more than two branches. For example, two lysines may be attached to two amino groups of a single lysine fork, resulting in a central structure with four available amino groups for subsequent addition of charged amino acids. An example is depicted in the central molecule shown in FIG. 23. Here, a lysine has been added to each amino group of a central lysine by solid phase protein synthesis, yielding four reactive amino groups for subsequent addition of charged amino acids or peptides. Polypeptides including strands of charged amino acids

may then be added to each of the amino groups of this molecule. A non-limiting example is depicted in FIG. 24B. In this example, a tetra-glutamate strand has been added to each of the four available amino groups of a four-branch forked lysine structure such as depicted in the central panel of FIG. 23B. In other examples, other charged amino acids could be added as one or more of the branches. Branches could also each have a different number of amino acids from the example depicted in FIG. 24B, and/or could have different number of amino acids from each other. Examples include some or all branches each having, independently, anywhere from 1 to 20 or more amino acids. Species of amino acids in a branch may also differ, from other amino acids within a branch and/or from amino acids found in other branches.

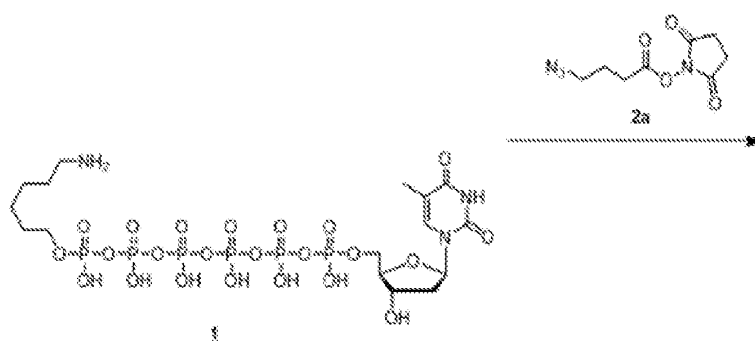
[0288] Continuing with this convergent synthesis scheme, two four-branch lysine forks could be attached to the amino groups of a central lysine fork to form an 8-branched structure as depicted in FIG. 23A, right-hand panel. As in the previous examples, strands of charged amino acids may be added to each of the 8 free amino groups to form a charge tag, again with branches that are the same or different lengths from each other (with from 1 to 20 amino acids or more each, independently), and different charged amino acids from branch to branch or within one or more branches, or the same species of charged amino acid in each branch. In a still further example, two 8-branch forks may be added to the two amino groups of a lysine group to form a 16-branch structure. A non-limiting example is depicted in FIG. 24C. In this example, a glutamate residue has been added to each of the 16 amino groups on the structure. In other example, polypeptides may be added to some or more of the amino acid groups, again having the same or different lengths from each other (with from 1 to 20 amino acids or more per branch, independently) or the same number of amino acids per branch, and the same or different species of charged amino acid, within a branch and/or between branches.

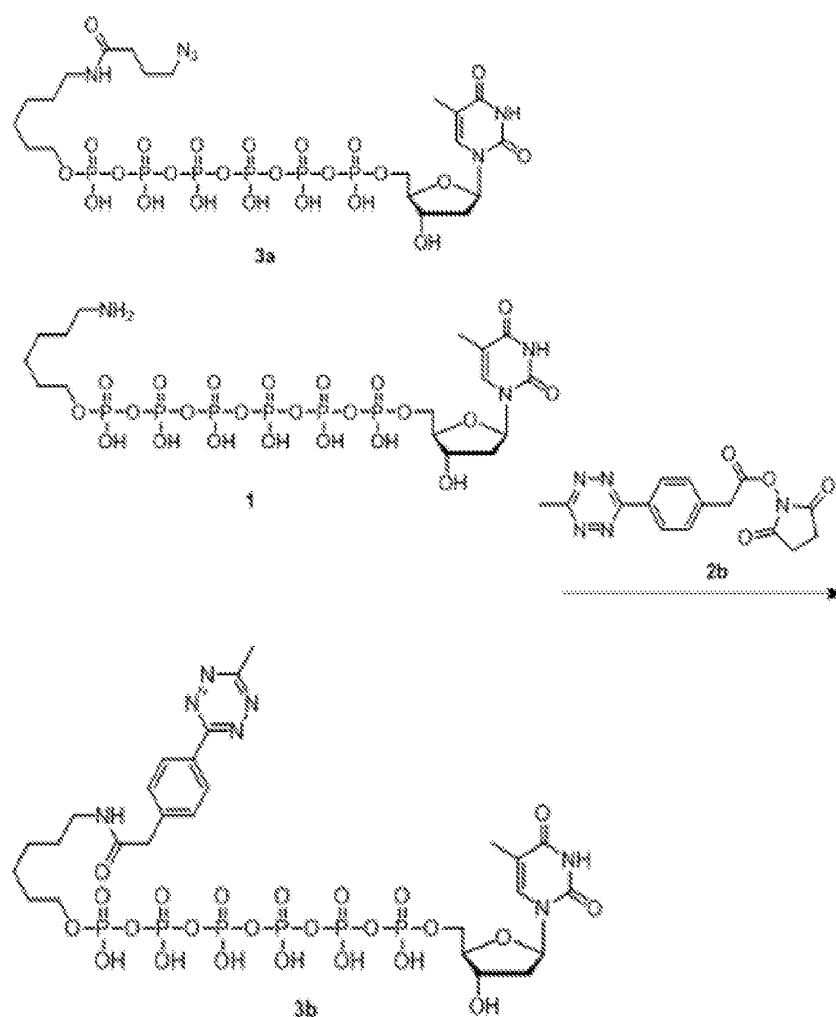
[0289] Note that the non-limiting examples of charge tags depicted in FIGs. 24A-24C each have the same net charge, with 16 negative amino acid moieties (in this example glutamate). However, the structure of the charge tags differs such that the density and distribution of the charge borne by the charge tag is carried differently by each of the three examples. All of the previously noted combinations may also be adopted in a forked charge tag with branches including charged amino acids as disclosed herein, providing many examples of charge tags that differ from each other not in the valence (positive or negative) of charge that they include, but also the value of the charge and the distribution or density of the charge within

the charge tag.

[0290] In another example, charge may be provided by a spermine-based component of a charge tag. For example, a spermine-based oligocationic charge may be added to a nucleotide and provide a positive charge as a charge tag in accordance with the present disclosure. An oligo-spermine conjugate has approximately 2.5 protonated amines at pH 7. An example of such a charge-tagged nucleotide is shown in FIGs. 25A and 25B. FIG. 25A shows an example of an oligo-spermine conjugate in accordance with an aspect of the present disclosure, and FIG. 25B shows a dendron-structured tag with spermine-derived end groups for magnifying the amount of charge that can be located at the end terminals of a charge tag. In both examples, chemistries disclosed herein for attaching charge tags to nucleotides could be adapted by skilled artisans for attaching such spermine-derived charge tags to nucleotides in accordance with aspects of the present disclosure.

[0291] The non-limiting examples below show the modification of a 5' amino nucleotide hexaphosphate with various linkers to allow for orthogonal attachment chemistry to dendron charge tags. A 5'-amine deoxy-thymine hexaphosphate (dT6P) (or other NPP) (1) may be functionalized with azido-butyrac N-hydroxysuccinimide (NHS) ester (2a) or methyltetrazine NHS ester (2b) to form azide dT6P (3a) or methyltetrazine dT6P (3b) respectively (Scheme 6).





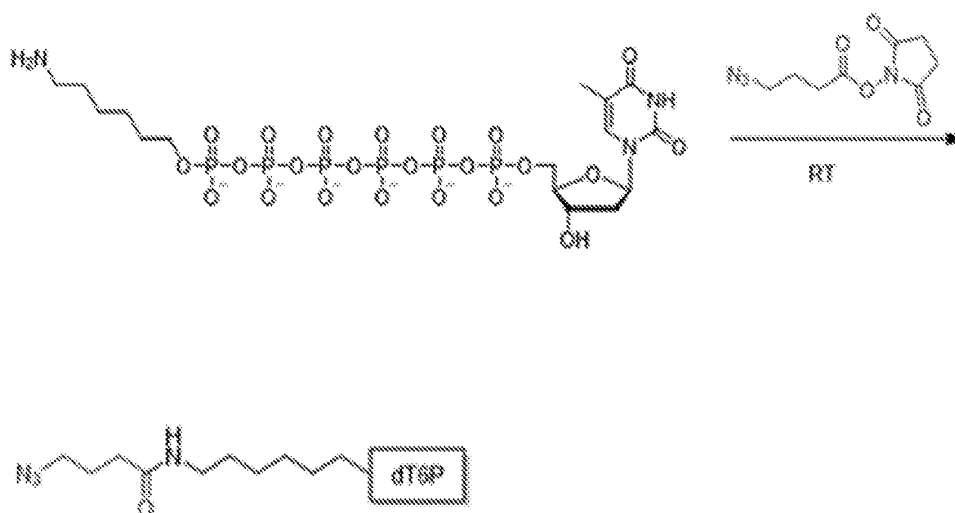
Scheme 6. Functionalization of 5'-amine dT6P.

[0292] An azide dT6P (3a) may be conjugated to a linear strand of poly-T oligonucleotide (4) with a 5'-hexynyl group via copper(I)-assisted azide-alkyne cycloaddition (CuAAC) in the presence of CuSO₄, tris-hydroxypropyltriazolylmethylamine (THPTA) ligand and sodium ascorbate to form an oligonucleotide conjugate (5a). Purification was performed on C18 reverse-phase HPLC and eluted with 50 mM TEAA (pH 7.5) and acetonitrile. A methyltetrazine dT6P (3b) may then be conjugated to a dendron with a transcyclooctene (TCO) group in 50 mM phosphate buffer (pH 7.4) to form a nucleotide analog with a dendron charge tag.

[0293] Alternatively, an azide dT6P (3a) may also be conjugated to a dendron charge tag

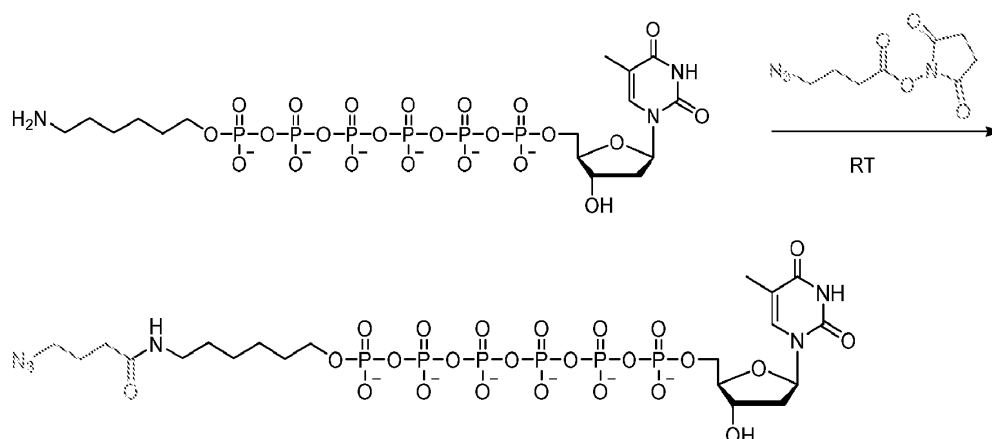
with a dibenzocyclooctyl (DBCO) group via copper-free strain promoted azide-alkyne cycloaddition (SPAAC) in 50 mM phosphate buffer (pH 7.4) to form a nucleotide analog with a dendron charge tag.

[0294] In the following scheme, an azide-alkyne click reaction may be made to link a nucleotide polyphosphate to a charge tag, such as a dendron charge tag with an alkyne group at its free valence end:

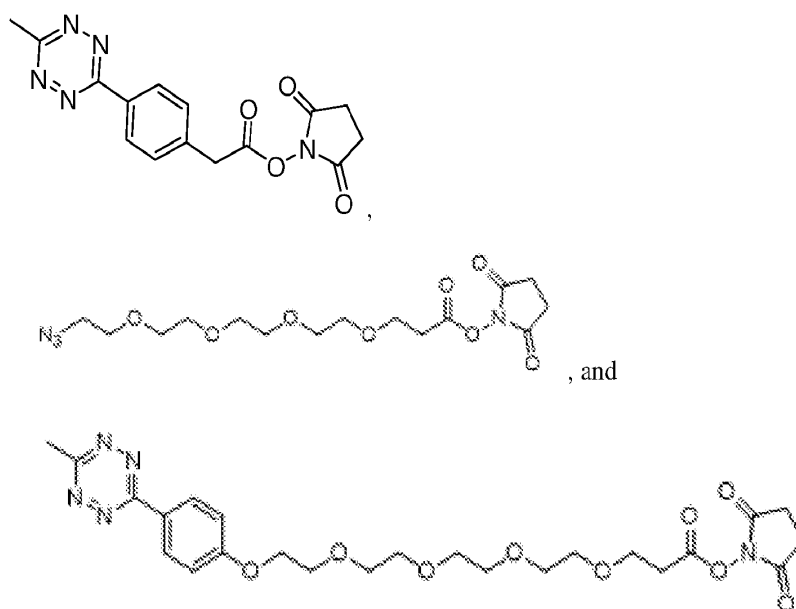


[0295] The foregoing examples may be modified, such as by reversing the placement of each reactive group of a ligation reaction or click chemistry reaction, yielding the foregoing linkages but oriented in the opposite direction with regard to the 5' and 3' ends of the analog nucleotides.

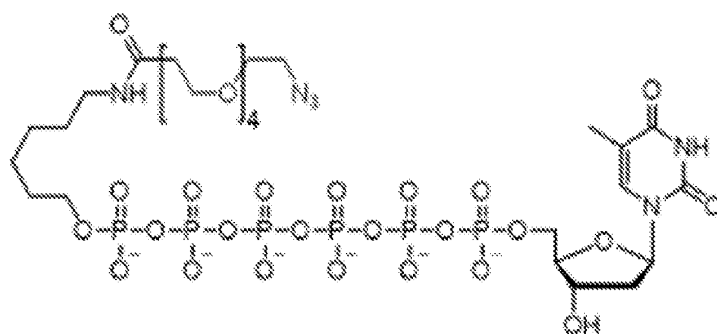
[0296] Reactive groups and linker chemistries may be appended to nucleotides and charge tags according to various applicable chemistries in accordance with the present disclosure. In some non-limiting examples, an azide or methyltetrazine tail may be added to an aminated NPP by reaction with an appropriate NHS residue, which may include linker portions of various lengths such as PEG4 linker, or PEG linker of varying lengths. Non-limiting examples of such synthesis schemes include the following and variations thereof:



[0297] Different NHS-moieties may be used, to add an azide or methyltetrazine reactive group, and with various linker lengths. Non-limiting examples include:

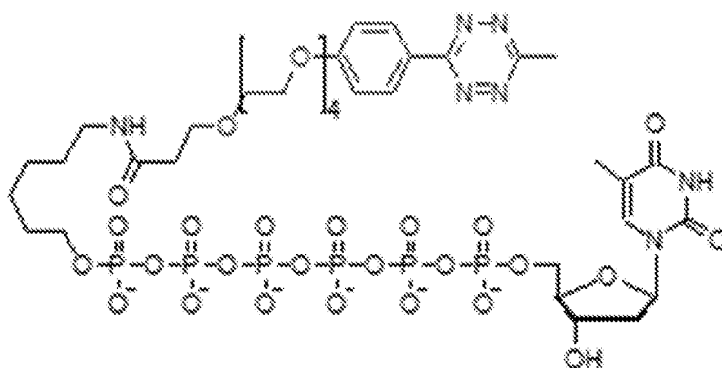


[0298] Various NPPs may be formed with different reactive groups for click or ligation chemistry reactions to connect them covalently with charge tags. Some non-limiting examples include:



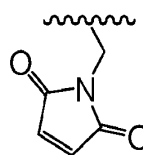
which could be reacted with an alkyne-containing charge tag, such as a dendron charge tag with an alkyne group at its free valence end.

[0299] Alternatively, a methyltetrazine containing NPP such as

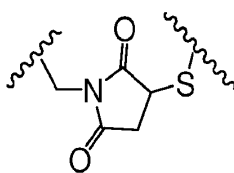


may be reacted with a TCO-containing charge tag, such as a dendron charge tag with a TCO group at its free valence end.

[0300] In other examples, DBCO-azide click chemistry between an NPP and a dendron charge tag may be used. In other examples, a maleimide group on a nucleotide or dendron charge tag may be reacted with a thiol group on a charge tag or nucleotide, respectively, to link the two via a maleimide-thiol reaction:



[0301] An NPP or charge tag containing a maleimide group reacted with a charge tag or NPP containing a thiol-containing group, respectively, in the presence of a reducing agent such as (tris(2-carboxyethyl)phosphine) may result in covalent bonding



between the two, for example

[0302] Some non-limiting, illustrative examples of charge tags with three-dimensional conformations that may cause high charge density are shown in FIGs. 15A-C, 16A, 16B, 17, and 18. FIGs. 15A-C show three examples of nucleotide analogs with oligonucleotide charge tags. For example, an oligonucleotide charge tag may contain 5, 10, 15, 20, 25, 30, 35, 40, or more oligonucleotides. Also shown are a conductive channel, in this case a nanowire, and a functionalized attachment to the conductive channel, specifically an accepting region. The accepting region is as indicated. The oligonucleotide charge tags are shown as dashed lines extending from the 5' end of the modified nucleotide. Shown are three different conformations the charge tags may take. FIG. 15A, for example, illustrates a recognizable stem-and-loop structure. In such a structure, nucleotides along the stem portion base pair with each other, leaving a loop portion therebetween, in this example illustrated as orienting away from the acceptor region. Negative charges from the phosphodiester bonds between nucleotides of the oligonucleotide charge tag may thereby be maintained in close proximity with each other, maintaining a higher charge density than may be obtained if they adopted a linear, stretched-out conformation.

[0303] FIG. 15B, for example, shows another example, with not a stem and loop structure but a bulge region of the charge tag. In this case, as in FIG. 15A, the charge tag includes a specificity region, shown bonding to the acceptor region. Here, the specificity region includes segments of the oligonucleotide that are disparate from each other spatially under circumstances when the oligonucleotide is stretched linearly. But, when induced by electrostatic attraction to associate with the acceptor region, the portions of the specificity region draw closer together. This conformation is consistent with adoption of a stem and loop conformation (FIG. 15A) or bulge conformation (FIG. 15B), in both case causing an increase of charge density of the charge tag.

[0304] FIGs. 15C and 20A-E show phosphodiester-bearing tags including polynucleotides adopting various three-dimensional configurations such as a stem-and-loop (e.g., 20A, 20B) or cloverleaf-like (e.g., 15C, 20C, 20D, or 20E) shape. Structured oligonucleotide charge tags were detectable by a conductive channel (see apparatus depicted in FIG. 26 as a structure for

conductive-channel detection of charge tags, as described below). Sequences for these charge tags are as follows:

Table 3: Example phosphodiester charge tags

FIG.	SEQ ID NO.	Sequence	# of bases
15C	1	CAGCGGAGCGGTATTTTTACCGCCAACGCTGTTTTTCAGCG TAGCACCGTTTTTCGGTGCGC	60
20A	2	CGAGACATCGTCGTGTCTCG	20
20B	3	CGAGACATCGTGCATATCGTACGATATGCACGATGTCTCG	40
20C	4	CGCCCGGGGATGAGTATCCCCGCGCTGAGTAGCGCGGGC G	40
20D	5	CGCCCGGGGATGAGTATCCCCGCGCATGAGTATGCGCT TGCTATGAGTATAGCAAGGGCG	60
20E	6	CGCCCTTGGGGATGAGTATCCCCAGCGCATGAGTATGCGC TTGCTATGAGTATAGCAAGTGCATGAGTATGCACAGGGCG	80

[0305] Similar to a stem and loop conformation, stems extending from a central hub may be formed by strands of nucleotides that are attracted to one another by Watson-Crick pair bonding rules, held together by a loop therebetween. Stems radiating from a central hub may be connecting strands of oligonucleotide oriented towards or around the central hub. As with other examples, the pair-bonding of bases of the nucleotides in the charge tag induce the tag to adopt a conformation that causes the negative charges of the phosphodiester bonds between nucleotides to condense together resulting in an increase in charge density compared to what the density may be if the oligonucleotide were stretched out linearly.

[0306] Other three-dimensional conformations of oligonucleotide charge tags are possible. Negative charges of phosphodiester bonds between nucleotides can be induced to come together at a high charge density because of Watson-Crick base-pairing. Various three-dimensional shapes can be adopted, using, for example, DNA origami methodology, creating oligonucleotide charge tags in tubular, circular, cuboid, helical, condensed helical, spherical or spheroid, or other conformations yielding high charge density.

[0307] FIG. 16 shows an example of two charge tags, one including oligonucleotide

sequences (16A, on the left) and the other including such sequences in addition to peptide nucleic acid sequences and polypeptides (16B, on the right). Not shown are connections between these charge tags and nucleotide analogs, but such attachment may be performed by chemical linking techniques such as those disclosed herein or otherwise known. The conformation of 16A on the left is a cruciform shape. Four oligonucleotide sequences are bound together in a conformation resembling a Holliday structure (as may occur during DNA recombination events). As shown in 16A, portions of the four polynucleotides bond to each other according to Watson-Crick base pairing. Each oligonucleotide also extends from the pair-bonded central portion into single-stranded overhangs. The pair bonding holds negative charges of the phosphodiester linkages within the oligonucleotides in proximity to each other, increasing charge density.

[0308] On the right, in 16B, peptide nucleic acid and polypeptide sequences are added to the charge tag shown in 14A, resulting in another non-limiting example of a charge tag. In this example, four sequences of peptide nucleic acids each connect, at their ends, polypeptide sequences. The polypeptide sequences form helical structures because of electrostatic attraction between some of the amino acids within the polypeptides. However, in these examples, the polypeptides have a net positive charge (notwithstanding the inclusion of some negatively charged amino acids therein which assist in adoption of a helical conformation). Portions of the peptide nucleic acid sequences connecting pairs of polypeptides are also hybridized to single-stranded portions of the polynucleotides that extend from the base-paired core. The strong bonds between the peptide nucleic acids and tightened coil conformation of the positively charged polypeptides allow for a net-positive charge of the charge tag and with a high charge density. Other examples of charge tags adopting similar architectures may have a net negative charge.

[0309] FIG. 17 shows some examples of polypeptide charge tags in which polypeptides adopt different three-dimensional architectures that result in high charge density. Coiled portions of polypeptide may be connected by linker sequences. When the linker sequences are fairly short, the coiled structures may be able to bind to one another in roughly overall linear arrays. Such conformation is possible because of electrostatic attraction between positively and negatively charged amino acids within the polypeptides. Overall, however, the polypeptide charge tags may have a net positive or net negative charge. With longer linkers between coiled portions of polypeptides of a charge tag, however, decreased steric hindrance

permits greater bending between adjacent coiled portions, permitting adoption of more complicated architectures such as shown in the lower portion of FIG. 17. These possibilities may result in even higher charge density. As with the examples shown in FIGs. 16A and 16B, these example charge tags could be attached to nucleotide analogs (not shown).

[0310] FIGs. 18A-18B show two views (a side view and a longitudinal view, respectively) of an example of a polypeptide charge tag adopting a coiled coil architecture, wherein electrostatic attraction between amino acids within a helix, and between amino acids of different helices, may induce the polypeptides to form a condensed structure. A result may be that a coiled coil may have a net negative or net positive charge, with the net charge held together at a high charge density (compared to what the charge density may be if the polypeptide sequences were stretched linearly).

[0311] For each example described herein, with a type of click chemistry or ligation chemistry described for attaching an example of a charge tag to an example of a nucleotide, each such type of click chemistry or ligation chemistry may also be used for attaching any described example of a charge tag to any described example of a nucleotide. In other words, a click chemistry or ligation chemistry is useful for irrespective of the type of charge tag or type of nucleotide to which it is attached.

[0312] Charge tags including oligonucleotides, polypeptides, or both, with or without peptide nucleic acids, may therefore adopt different three-dimensional architectures with elevated charge density compared to linear charge tags stretched linearly. A charge tag may have a net negative charge, such as if it contains an excess of phosphodiester or negative amino acids relative to positive charges, or a net positive charge such as if it has more positively charged amino acids than negatively charged groups. Coiled coils can be computationally designed to adopt specific compact structures, based on well characterized molecular interactions between amino acid components. An example of coiled coils that can be used include leucine zippers, which may be in, for example, dimeric or trimeric forms, of controlled length and diameter. Furthermore, because interactions that govern coiled coil compact structure are localized in the interior, the surface can be independently engineered to carry a wide range of charges.

[0313] A charge tag as disclosed herein may have a charge from anywhere between $-200e$ and $+200e$, or between $-100e$ and $+100e$, or between $-40e$ and $+40e$, or between $-20e$ and $+20e$, or any range therein. In some examples, net charge or partial net charge of a

charge tag may be packed into a density of from $-200e$ to $+200e$ per cubic nanometer, or from $-100e$ to $+100e$ per cubic nanometer, or from $-40e$ to $+40e$ per cubic nanometer, or from $-20e$ to $+20e$ per cubic nanometer, or any range therein.

[0314] In an example, a test apparatus was used for detection of a charge tag by a conductive channel. A schematic of such an apparatus is depicted in FIG. 26. Shown is a silicon nanowire (NW) field effect transistor capable of detecting an electric charge when in proximity thereto. Optionally, as depicted in FIG. 26, a surface modifier may be attached to the NW, such as may be adopted in a flow cell for SBS or other related methods. For example, a surface modifier may be poly(N-(5-azidoacetamidyl)pentyl)acrylamide-co-acrylamide), also known as PAZAM, or another related surface modifier polymer. In this example, an oligonucleotide is grafted to the surface modifier. To a solution surrounding the grafted oligo and NW, another oligonucleotide complementary to the grafted oligonucleotide is added. A sequence of the grafted oligonucleotide and the in-solution oligonucleotide are determined so as to bind to one another according to standard Watson-Crick base pairing rules. Also attached to the in-solution oligo is a charge tag. In the example depicted in FIG. 26, the charge tag is a phosphodiester-based tag. However, any of the examples of charge tags disclosed herein could be used in such a system instead. Binding of the in-solution nucleotide to the grafted nucleotide brings the charge tag in proximity with the NW such that the NW can detect the presence and magnitude of charge in the charge tag. In some control conditions (non-comp, as opposed to comp when a complementary in-solution oligonucleotide is used), an in-solution oligonucleotide not complementary to the grafted nucleotide, bearing a charge tag, was used. Examples of using such an apparatus and system for testing a conductive channel's ability to detect different charge tags are disclosed herein.

[0315] Tests of a conductive channel's ability to detect a charge tag were performed using an apparatus as depicted in FIG. 26 as follows. For baseline, in 1 mL of 10 mM phosphate buffer pH 5 was flowed over the NW, electrically measured for 5 minutes. This is followed by rinsing twice in blank PBS, and a further 5 min baseline measurement in 10 mM Phosphate Buffer, pH 5. Hybridization with a charge-tagged in solution oligonucleotide was then performed, by incubating in 1 mL of 500 nM comp or non-comp oligo in 2xPBS and incubating for 5 minutes, followed by rinsing with 1 mL 2xPBS, then rinsing in 1 mL 10 mM Phosphate Buffer pH 5. Another 5 min measurement was then taken in 10 mM Phosphate Buffer pH 5 for 5 min. In-solution oligonucleotides were then eluted with 1 mL of 100%

formamide and incubated for 15 minutes, then rinsed 1 mL 10 mM Phosphate Buffer pH 5, followed by another 5 min baseline measurement in 10 mM Phosphate Buffer pH 5. The process was then repeated with next samples containing different charge tags affixed to in-solution oligonucleotides.

[0316] Results of some examples are depicted in FIG. 27. Electrical current was measured as detected across multiple nanowires and normalized to voltage (mV) by each NW's conductance following incubation with in-solution oligonucleotides bearing the following charge tags: C20 (a phosphodiester charge tag with the following sequence (SEQ ID NO: 7): GAACAATTCCAGCCTTGATATCAACACTATTGATA), a linear sequence of 16 glutamate amino acids (SEQ ID NO: 12) ("LinearE16"), a branched charge tag with 16 glutamate amino acids with a structure as depicted in FIG. 24B ("BranchedE16"), a "dendritic" branched charge tag with 16 glutamate amino acids with a structure as depicted in FIG. 24C ("DendriticE16"), a linear tag of five trimethylated lysine residues (SEQ ID NO: 13), and again C20 ("K5-Me3") (in that order). As can be seen, a conductive channel was able to detect the charges with responses of between 10-14 mV. No current was detected in non-comp conditions.

[0317] In another example, similar measurements were taken but in this case with 50 mM phosphate buffer (pH 5). Results are shown in FIG. 28. In this buffer, differences between mV detected by the different charge tags are evident (even between charge tags that have the same overall amount of charge as each other), verifying the ability to distinguish between different charge tags.

[0318] A Debye length for a charge tag may be calculated as a function of a buffer solution in which it is sensed by a conductive channel. Debye length (κ^{-1}) in an ionic solution may be calculated according to the following equation:

$$\kappa^{-1} = \sqrt{\frac{\epsilon_r \epsilon_0 k_B T}{2 \times 10^3 N_A e^2 I}} \quad (\text{Eq. 1})$$

where $T = \sim 298\text{K}$; $k_B = 1.28\text{e-}23 \text{ J/K}$; $e = 1.6\text{e-}19\text{C}$; $\epsilon_0 = 8.85\text{e-}12 \text{ F/m}$; ϵ_r for water ~ 78 ; $N_A = 6.022\text{e}23$; $I = \text{ionic strength} = \sum c_i z_i^2$ where c is concentration of ions and z is charge number (depending on the solution). In an example, different Debye lengths for charge tag C20 were tested in Tris buffer (pH 7) with the following values for c and z and Debye lengths:

Table 4

Buffer	c1 (M)	z1	c2 (M)	z2	c3 (M)	z3	c4 (M)	z4	κ^{-1} nm
10 mM Tris7	1.00E-05	1	1.00E-05	1	1.00E-06	1	1.00E-06	1	2.71
5 mM Tris7	5.00E-06	1	5.00E-06	1	5.00E-07	1	5.00E-07	1	3.83
2 mM Tris 7	2.00E-06	1	2.00E-06	1	2.00E-07	1	2.00E-07	1	6.1
1 mM Tris 7	1.00E-06	1	1.00E-06	1	1.00E-07	1	1.00E-07	1	8.57

Results are shown in FIG. 29A.

[0319] In another example, different Debye lengths for charge tag C20 were tested in citrate buffer, with and without KCl, with the following values for c and z and Debye lengths:

Table 5

	c1 (M)	z1	c2 (M)	z2	c3 (M)	z3	c4 (M)	z4	κ^{-1} nm
10mM Citrate Buffer	1.00E-05	2	2.00E-05	1					2.01
1mM Citrate Buffer	1.00E-06	2	2.00E-06	1					6.35
5mM Citrate Buffer	5.00E-06	2	1.00E-05	1					2.84
50mM Citrate Buffer	5.00E-05	2	1.00E-04	1					0.90
10 mM Citrate Buffer +5 mM KCl	1.00E-05	2	2.00E-05	1	5.00E-06	1	5.00E-06	1	1.80
10 mM Citrate Buffer +10 mM KCl	1.00E-05	2	2.00E-05	1	1.00E-05	1	1.00E-05	1	1.64
10 mM Citrate Buffer + 15mM KCl	1.00E-05	2	2.00E-05	1	1.50E-05	1	1.50E-05	1	1.52
10 mM Citrate Buffer +50 mM KCl	1.00E-05	2	2.00E-05	1	5.00E-05	1	5.00E-05	1	1.07

Results are shown in FIG. 29B.

[0320] Thus, Debye length for a given charge tag may be modified by modifying aspects of the solution in which a charge tag is detected by a conductive channel such that proximity of a charge tag to a conductive channel is sufficient to permit detection of the charge tag under various conditions. In the examples depicted in FIGs. 29A and 29B, Debye lengths of 0.9 to 8.57 nm were tested for a given charge tag such that proximity of a charge tag may be determined. In an example, a charge tag may be in proximity to a conductive channel for purposes of and to permit charge detection of the charge tag by the conductive channel when the charge tag is within a Debye length from the conductive channel.

[0321] A non-limiting example of measuring incorporation of a nucleotide bearing a charge tag is shown in FIG. 2. A phosphodiester charge-tagged thymidine nucleotide (TetTCO-dT6P) for 5s, 1m, or 10m, alone or followed by a fluorescently tagged, azidomethyl 3-prime blocked adenosine nucleotide (ffA-AZM), were incorporated into a polynucleotide hybridized to an in-solution template oligonucleotide using a Phi29 polymerase. Following incorporation, polynucleotide was dehybridized from the template and separated by gel electrophoresis and incorporation of ffA-AZM measured. As shown in FIG. 2, incubation with charge-tagged nucleotide followed by ffA-AZM led to detection of ffA-AZM incorporated into the polynucleotide, as a proxy for measuring incorporation of charge-tagged nucleotide. No ffA-AZM incorporation was detected in the absence of either nucleotide (No nuc), or when only one or the other was present (TetTCO-dT6P alone, left, or ffA-AZM alone, right). In other examples, incorporation of fluorescently tagged nucleotide may be detected and measured by various other techniques, including on-surface fluorescence or in-solution fluorescence measurement following polynucleotide dehybridization from template.

[0322] In other examples, a charge-tagged nucleotide was incorporated into a polynucleotide complementary to a surface-attached template, followed by incorporation of a fluorescently tagged nucleotide (not shown). In one example, one polymerase was used to incorporate the charge-tagged nucleotide and a different polymerase was used to incorporate the fluorescently tagged nucleotide. For example, Klenow fragment incubated with a first, charge-tagged T nucleotide and polynucleotide hybridized to surface-attached template was used to incorporate the charge-tagged nucleotide. After washing to remove polymerase, excess nucleotide, etc., a second incubation in Phi29 and a fluorescently tagged A nucleotide was used to add a fluorescent nucleotide adjacent to the first-incorporated charge-tagged nucleotide. Template and polynucleotide sequences were designed such that the template requires addition of first a T then an A to the 3-prime end of the polynucleotide in the presence of polymerase. Extended polynucleotide was then dehybridized then run on an electrophoresis gel to detect incorporation of nucleotides. Phi29 catalyzed incorporation of A, indicating that Klenow fragment also had incorporated T (because Phi29 could not have incorporated an A unless a T had first been added to the 3-prime end of the polynucleotide). In other examples, Phi29 was used for incorporating both the T and the A, with a wash step in between, and did so. In another example, Phi29 was used in a single polymerase reaction, with both charge-tagged T and fluorescent A present, and again Phi 29 was able to

incorporate fluorescent A indicating that it had also incorporated charge-tagged T, indicating that Phi29 can incorporate both charge-tagged and fluorescent nucleotides.

[0323] In some examples of the technology disclosed herein, one or more computer readable storage devices or memory storing computer-readable instructions that when executed by a computer, cause the computer to perform at least any one of the methods disclosed herein. “Computer” herein may refer to any processor or processor-containing device. In some examples, a system is configured to perform at least a portion of any one of the methods disclosed herein. In some examples, a system is coupled to computer readable storage devices or memory storing computer-readable instructions that when executed, cause the system to perform at least any one of the methods disclosed herein.

[0324] All literature and similar material cited in this application, including, but not limited to, patents, patent applications, articles, books, treatises, and web pages, regardless of the format of such literature and similar materials, are expressly incorporated by reference in their entirety. In the event that one or more of the incorporated literature and similar materials differs from or contradicts this application, including but not limited to defined terms, term usage, described techniques, or the like, this application controls.

[0325] It should be appreciated that all combinations of the foregoing concepts and additional concepts discussed in greater detail below (provided such concepts are not mutually inconsistent) are contemplated as being part of the inventive subject matter disclosed herein. In particular, all combinations of claimed subject matter appearing at the end of this disclosure are contemplated as being part of the inventive subject matter disclosed herein. It should also be appreciated that terminology explicitly employed herein that also may appear in any disclosure incorporated by reference should be accorded a meaning most consistent with the particular concepts disclosed herein.

[0326] Reference throughout the specification to “one example”, “another example”, “an example”, and so forth, means that a particular element (*e.g.*, feature, structure, and/or characteristic) described in connection with the example is included in at least one example described herein, and may or may not be present in other examples. In addition, it is to be understood that the described elements for any example may be combined in any suitable manner in the various examples unless the context clearly dictates otherwise.

[0327] While several examples have been described in detail, it is to be understood that the disclosed examples may be modified. Therefore, the foregoing description is to be considered

non-limiting. Although some examples may have been depicted and described in detail herein, it will be apparent to those skilled in the relevant art that various modifications, additions, substitutions, and the like can be made without departing from the spirit of the present disclosure and these are therefore considered to be within the scope of the present disclosure as defined in the claims that follow.