NON-MASS DETERMINED BASE COMPOSITIONS FOR NUCLEIC ACID DETECTION

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The present invention provides systems, methods, and compositions for nucleic acid detection based on non-mass determined base compositions. For example, in certain embodiments, base count data for a template nucleic acid is generated using an approach that does not measure molecular mass of the template nucleic acid (e.g., by sequencing the template nucleic acid) and a database comprising base count entries is queried to identify the target nucleic acid. In particular embodiments, sequencing is employed which is conducted in substantially real-time.
FIGURE 1

Exemplary Process Embodiment

Obtain sample with template nucleic acids

sample preparation

Amplify template nucleic acids using one or more lysis primer pairs

* For partial base counts, not all nucleotide types in a given rxn mixture are labeled (i.e., only those that are to be detected)

* Determine amplicon (one or both strands) partial base counts, complete base counts and/or sequence

Zero-mode Waveguide array (ZMW)

Partial Base Count or Complete Base Count or Sequence

[A_{10}G_{15}]
[A_{12}G_{14}]
[A_{19}G_{15}C_{12}T_{17}]
[A_{12}G_{44}C_{15}T_{10}]
AGGTGCC... GGCTAAT...

Query database

Detect, identify, characterize, etc. the template nucleic acids
FIGURE 2

Exemplary Amplicon Analysis

- magbead
- Streptavidin
- biotin

Denature and wash away impurities including strand (b)

Wash, but don’t denature

(a) Purified amplicon with strand (a) and strand (b)

Separate

(a) Analyze as in Figure 1

(b) Analyze as in Figure 1

Note: When both amplicon strands are analyzed and only partial base counts determined, in certain cases complete base counts can be derived. For example, if one counts the number of As and Gs in both strands, then the number of Ts and Cs in the corresponding opposite strands are also known without the T or C nucleotides having been labeled or detected.
FIGURE 4

For each group of organisms

200 Identify candidate target sequences

210 Create/curate nucleotide alignment

220 Analyze nucleotide alignment

230 Primer Design

240 Select candidate primer pairs

300 in-silico analysis

310 Seq. database

320 Probability model for unknown organism identification

330 Product base composition database

340 in-silico specificity check

400 Lab PCR analysis

410 Organism collection

420 Validate primer sets
   - Sensitivity
   - Specificity
   - Reproducibility
**FIGURE 5**

- Primer pair candidates for testing
- Make T7/synthetic construct
  - Generate quantitate test material
  - Test Primers on subset of quantitated stocks and/or isolated genomes
  - Select primers meeting desired criteria
  - Test additional isolates
  - Design plasmid calibrant
  - Determine optimal calibrant level
- Redesign/optimize primers if needed
- Isolate nucleic acids
- Ibis strain repository
- Select final primer pairs
- Test in relevant background
- Validated Primer Set
FIGURE 6

[Diagram showing components labeled 1002, 1004, 1006, and 1008 connected to a computer with an arrow labeled 1000.]
The present invention provides systems, methods, and compositions for nucleic acid detection based on non-mass determined base compositions. For example, in certain embodiments, base count data for a template nucleic acid is generated using an approach that does not measure molecular mass of the template nucleic acid (e.g., by sequencing the template nucleic acid) and a database comprising base count entries is queried to identify the target nucleic acid. Essentially any sequencing approach is optionally utilized in the present invention, including, for example, pyrosequencing, zero-mode waveguide type sequencing, nanopore sequencing, and the like. In particular embodiments, sequencing is employed which is conducted in substantially real-time.

In certain embodiments, the base count or partial base count is determined by sequencing. In other embodiments, the base count or partial base count is determined by liquid chromatography, such as High Performance Liquid Chromatography (HPLC). In further embodiments, the base count or partial base count is determined by paper chromatography (e.g., 2-D paper chromatography).

In particular embodiments, at least a partial base count only includes the number of A’s present in a template nucleic acid (e.g., A1). In particular embodiments, at least a partial base count only includes the number of C’s present in a template nucleic acid (e.g., C1). In particular embodiments, at least a partial base count only includes the number of G’s present in a template nucleic acid (e.g., G1). In particular embodiments, at least a partial base count only includes the number of T’s present in a template nucleic acid (e.g., T1). In other embodiments, at least a partial base count only includes two of the canonical bases (e.g., A1, C1). In further embodiments, at least a partial base count only includes three of the canonical bases (e.g., A1, C1, G1). In additional exemplary embodiments, at least a partial base count includes all four canonical bases (e.g., A1, C1, G1, T1).

In certain embodiments, the template nucleic acid comprises DNA or RNA. In other embodiments, the determining is performed in real-time or substantially real-time (e.g., using a device comprising a zero-mode waveguide). In particular embodiments, the template nucleic acid comprises a mammalian nucleic acid, a bacterial nucleic acid, a viral nucleic acid, a fungal nucleic acid, or a protozoal nucleic acid. In further embodiments, the template nucleic acid is attached to a solid support. In additional embodiments, the method further comprises amplifying the template nucleic acid prior to or during (a). In certain embodiments, the method further comprises obtaining the template nucleic acid from one or
more sample sources selected from the group consisting of: an environmental sample and a sample derived from a subject.  

[0010] In particular embodiments, step (a) comprises contacting the template nucleic acid with at least one nucleotide incorporating biocatalyst, labeled nucleotides, and at least one primer nucleic acid that is at least partially complementary to at least a subsequence of the template nucleic acid, under conditions whereby the nucleotide incorporating biocatalyst extends the primer nucleic acid to produce an extended primer nucleic acid by incorporating the labeled nucleotides at a terminal end of the extended primer nucleic acid, wherein nucleotides that comprise different nucleobases comprise different labels, wherein the different labels produce detectable signals as or after the labeled nucleotides are incorporated at the terminal end of the extended primer nucleic acid, which detectable signals identify the labeled nucleotides incorporated at the terminal end of the extended primer nucleic acid and/or complementary nucleotides in the template nucleic acid, and wherein the detectable signals are detected as or after the labeled nucleotides are incorporated at the terminal end of the extended primer nucleic acid to thereby determine the base count of the subsequence of the template nucleic acid and/or the complement thereof. In other embodiments, the labels comprise different fluorescent labels and wherein the detectable signals are detected using a fluorescence microscope. In some embodiments, the at least one primer nucleic acid is a primer pair, wherein the primer pair is configured to hybridize with conserved regions of the two or more different bioagents and flanking variable regions of the two or more different bioagents.  

[0011] In particular embodiments, the terminal end of the extended primer nucleic acid is the 3′ terminal end. In further embodiments, the nucleotide incorporating biocatalyst comprises an enzyme selected from the group consisting of: a polymerase, a terminal transferase, a reverse transcriptase, a polynucleotide phosphorylase, and a telomerase. In additional embodiments, the nucleotide incorporating biocatalyst comprises one or more modifications. In some embodiments, the nucleotide incorporating biocatalyst is an enzyme derived from an organism that is selected from the group consisting of: Thermus anranikianii, Thermus aquaticus, Thermus caldophilus, Thermus chlorophilus, Thermus filiformis, Thermus flavus, Thermus igniorum, Thermus lacteus, Thermus ozolinai, Thermus ruber, Thermus rubens, Thermus scotoductus, Thermus silvanus, Thermus species Z05, Thermus species sps 17, Thermus species, Thermotoga maritima, Thermotoga neapolitana, Thermosipho africans, Anaerocellum thermophilum, Bacillus caldothermus, and Bacillus stearothermophilus.  

[0012] In certain embodiments, the nucleotide incorporating biocatalyst comprises a Φ29 DNA polymerase. In other embodiments, a label is attached to one of a heterocyclic base of a labeled nucleotide, a sugar moiety of a labeled nucleotide, and a phosphate group of a labeled nucleotide. In further embodiments, a linker attaches a label to a labeled nucleotide. In particular embodiments, the extended primer nucleic acid is complementary to a subsequence of the template nucleic acid. In further embodiments, the extended primer nucleic acid is complementary to a full-length sequence of the template nucleic acid. In certain embodiments, the primer nucleic acid comprises an intelligent primer. In other embodiments, the label comprises a fluorescent dye, a non-fluorescent label, a colorimetric label, a chemiluminescent label, a bioluminescent label, a radioisotope, an antibody, an antigen, biotin, a hapten, or an enzyme. In other embodiments, the label is a fluorescent dye selected from the group consisting of: a rhodamine dye, a fluorescein dye, a halo fluorescein dye, a dichlororhodamine dye, an energy transfer dye, a Lucifer dye, Oregon Green, and a cyanine dye. In some embodiments, the label is a fluorescent dye selected from the group consisting of: JOE, VIC, TET, HEX, PAM, R6G, R110, TAMRA, and ROX. In particular embodiments, the label is a radioisotope selected from the group consisting of: 3H, 14C, 32P, 35S, 42K, 45Ca, 59Fe, 125I, and 203Hg.  

[0013] In certain embodiments, the methods comprise determining at least partial base counts of subsequences of two or more template nucleic acids and/or complements thereof, using an approach that does not measure molecular mass of the template nucleic acids to produce the base count data. In other embodiments, the methods comprise querying the database to produce two or more matches of the base count data with two or more base count entries to thereby identify the template nucleic acid. In other embodiments, identifying the template nucleic acid comprises identifying an organismal source of the template nucleic acid and/or one or more properties of the template nucleic acid. In some embodiments, the properties of the template nucleic acid comprise a genotype, a virulence property, a pathogenic property, and/or a drug resistance property. In particular embodiments, the determining is performed in substantially real-time.  

[0014] In other embodiments, the organismal source comprises a mammal, a bacterium, a virus, a fungus, or a protozoan. In further embodiments, the organismal source is identified at one or more taxonomic rank levels selected from the group consisting of: a Domain, a Superphylum, a Superdivision, a Superclass, a Superorder, a Superfamily, a Super species, a Kingdom, a Phylum, a Division, a Class, a Legion, an Order, a Family, a Tribe, a Genus, a Species, a Subkingdom, a Sub phylum, a Subclass, a Suborder, a Subfamily, a Subtribe, a Subgenus, a Subspecies, an Infra kingdom, a Branch, an Infra phylum, an Infra class, an Infra order, an Alliance, an Infraspecies, a Microphylum, a Pan class, and a Parvorder.  

[0015] In some embodiments, the present invention provides systems comprising: at least one sequencing device; at least one primer pair configured to hybridize with conserved regions of two or more different bioagents and flanking variable regions of the two or more different bioagents; and a computer readable medium having one or more logic instructions for directing a system to: (a) determine at least a partial base count of at least a subsequence of at least one template nucleic acid and/or a complement thereof, using an approach that does not measure molecular mass of the template nucleic acid, to produce base count data; and (b) query a database comprising at least one base count entry corresponding to an identified nucleic acid to produce a match of the base count data with the base count entry to thereby identifying the template nucleic acid.  

[0016] In certain embodiments, the system comprise instructions for determining at least a partial base count of at least a subsequence of at least one template nucleic acid and/or a complement thereof, in substantially real-time without measuring the molecular mass of the template nucleic acid to produce base count data using the zero-mode waveguide and the primer nucleic acid and for identifying the template nucleic acid using the base count data. In certain
embodiments, the sequencing device comprises a zero-mode waveguide. In other embodiments, the systems comprise at least one nucleotide incorporating biocatalyst and/or labeled nucleotides. In further embodiments, the systems further comprise one or more containers for packaging the zero-mode waveguide, or the primer pair.

In certain embodiments, the present invention provides a computer program product, comprising a computer readable medium having one or more logic instructions for directing a system to: (a) determine at least a partial base count of at least a subsequence of at least one nucleic acid and/or a complement thereof (e.g., in substantially real-time), using an approach that does not measure molecular mass of the template nucleic acid, to produce base count data; and (b) query a database comprising at least one base count entry corresponding to an identified nucleic acid to produce a match of the base count data with the base count entry to thereby identifying the template nucleic acid.

In some embodiments, the present invention provides system comprising: a reaction vessel or substrate; a detector configured to detect detectable signals produced in or on the reaction vessel or substrate, which detectable signals correspond to at least some nucleobases incorporated into a nucleic acid to generate nucleobase incorporation data; a database of base count entries indexed to identified nucleic acids; and a controller configured to correlate the nucleobase incorporation data with detected base counts and query the database for a match between the detected base counts and the base count entries to thereby identify or detect the nucleic acids.

In particular embodiments, the reaction vessel or substrate comprises at least one zero-mode waveguide. In further embodiments, the detector comprises a fluorescence microscope. In other embodiments, the systems comprise at least one material transfer component that transfers material to and/or from the reaction vessel or substrate. In other embodiments, the systems comprise at least one thermal modulator configured to modulate temperature in the reaction vessel or substrate. In other embodiments, the reaction vessel or substrate comprises at least one primer nucleic acid is selected from the primer nucleic acids disclosed in one or more of the published documents incorporated by reference herein. In other embodiments, the systems comprise at least one nucleic acid amplification component and/or at least one nucleic acid amplification component.

In further embodiments, the present invention provides methods of identifying a target nucleic acid in a sample comprising: (a) at least partially sequencing the target nucleic acid, or amplified sequences thereof, to generate sequence data; (b) determining at least a partial base count from the sequence data to generate base count data; and (c) querying a database comprising at least one base count entry corresponding to the target nucleic acid to produce a match of the base count data with the base count entry, thereby identifying the target nucleic acid.

In some embodiments, the present invention provides methods of identifying a target nucleic acid in a sample comprising: (a) amplifying one or more segments of the nucleic acid in the sample using at least one purified oligonucleotide primer pair to generate amplified sequences, wherein the primer pair is configured to hybridize with conserved regions of the two or more different bioagents and flank variable regions of the two or more different bioagents; (b) at least partially sequencing the amplified sequences to generate sequence data; (c) determining at least a partial base count from the sequence data to generate base count data; and (d) querying a database comprising at least one base count entry corresponding to the target nucleic acid to produce a match of the base count data with the base count entry, thereby identifying the target nucleic acid.

In certain embodiments, the molecular mass of the target sequence or the amplified sequences is not determined in the identifying the target nucleic acid. In further embodiments, the sequencing is performed in real-time or substantially real-time. In other embodiments, the identifying the target nucleic acid is performed in substantially real-time. In further embodiments, the at least partially sequencing the target nucleic acid is performed with a device comprising at least one zero-mode waveguide. In some embodiments, the primer pair comprises forward and reverse primers that are about 20 to 35 nucleobases in length.

In other embodiments, the present invention provides systems comprising: (a) a sequencing device configured to generate nucleic acid sequence data corresponding to the nucleic acid sequence of one or more amplicons produced using at least one purified oligonucleotide primer pair that comprises forward and reverse primers, wherein the primer pair comprises nucleic acid sequences that are substantially complementary to nucleic acid sequences of two or more different bioagents; and (b) a controller operably connected to the sequencing device, the controller configured to: (i) determine at least a partial base count from the nucleic acid sequence data to generate base count data, and (ii) query a database with the base count data, wherein the database comprises at least one base count entry corresponding to a target nucleic acid.

In certain embodiments, the database comprises base composition information (entries) for at least three different bioagents. In other embodiments, the database comprises base composition information for at least 2 . . . 10 . . . 50 . . . 100 . . . 1000 . . . 10,000, or 100,000 different bioagents. In some embodiments, the base composition data comprises at least 10 . . . 50 . . . 100 . . . 500 . . . 1000 . . . 10,000 . . . or 100,000 unique base compositions. In further embodiments, the database is stored on a local computer. In particular embodiments, the database is accessed from a remote computer over a network. In further embodiments, the base composition information in the database is associated with bioagent identity. In certain embodiments, the base composition data in the database is associated with bioagent geographic origin.

In some embodiments, the present invention provides methods of identifying a target nucleic acid in a sample comprising: (a) amplifying one or more segments of the target nucleic acid in the sample using at least one purified oligonucleotide primer pair to generate a sample comprising amplified sequences, wherein said primer pair comprises a forward primer and a reverse primer and is configured to hybridize with conserved regions of two or more different bioagents and flank variable regions of the two or more different bioagents; (b) treating the sample with reagents such that the amplified sequences in the sample are digested into monodeoxynucleotides; (c) quantitating the monodeoxynucleotides using HPLC, paper chromatography, or similar non-mass spectrometry based technique such that base count data is generated; and (f) querying a database comprising at least one base count entry corresponding to the target nucleic acid to produce a match of the base count data with said base
count entry, thereby identifying the target nucleic acid. In certain embodiments, the methods further comprise a step after step (a) but before step (b) of treating the sample comprising amplified sequences such that either the sense strands corresponding to the forward primer or said anti-sense strands corresponding to the reverse primer of the amplified sequences are removed from the sample in order to generate a purified sample. In certain embodiments, step (c) employs a phosphodiesterase (e.g., such as snake venom phosphodiesterase I).

BRIEF DESCRIPTION OF THE DRAWINGS

[0026] The foregoing summary and detailed description is better understood when read in conjunction with the accompanying drawings which are included by way of example and not by way of limitation.

[0027] FIG. 1 shows one embodiment of a biogent identification process of the present invention.

[0028] FIG. 2 shows one embodiment of a biogent identification process of the present invention.

[0029] FIG. 3 shows one embodiment of a sequencing system composed of zero-move waveguides.

[0030] FIG. 4 shows a process diagram illustrating one embodiment of the primer pair selection process.

[0031] FIG. 5 shows a process diagram illustrating one embodiment of the primer pair validation process. Here select primers are shown meeting test criteria. Criteria include but are not limited to, the ability to amplify targeted biogent nucleic acid, the ability to exclude non-target biogents, the ability to produce unexpected amplions, the ability to dimerize, the ability to have analytical limits of detection of ±100 genomic copies/reaction, and the ability to differentiate amongst different target organisms.

[0032] FIG. 6 shows a block diagram showing a representative system.

DETAILED DESCRIPTION OF EMBODIMENTS

[0033] The present invention provides systems, methods, and compositions for nucleic acid detection based on non-mass determined base compositions. For example, in certain embodiments, base count data for a template nucleic acid is generated without measuring the molecular mass of the template nucleic acid (e.g., by sequencing the template nucleic acid) and a database comprising base count entries is queried to identify the target nucleic acid. In particular embodiments, sequencing is employed which is conducted in substantially real-time.

[0034] The present invention provides methods that are useful in detecting and characterizing nucleic acids. The methods typically involve determining at least partial base counts (i.e., partial or complete base counts) and/or the sequence of target or target nucleic acids in real-time without measuring the molecular mass of the template nucleic acid. Exemplary applications of the method include diagnosing infectious or genetic diseases, forensics, genotyping, and companion diagnostics, among many other applications. In addition, the invention also provides related zero-mode waveguide (e.g., arrays of zero-mode waveguides), kits, computer program products and systems.

[0035] The term “partial base composition” or “partial base count” refers to the number of each residue of at least one nucleobase type (e.g., a given purine nucleobase type, a given pyrimidine nucleobase type, a given nucleobase analog type, and/or the like), but not each residue comprised in an amplicon or other nucleic acid (e.g., for single or multiple strands of those nucleic acids), without consideration for the linear arrangement of these residues in the strand(s) of the amplicon. For example, if a given amplicon or other nucleic acid includes four nucleobase types (e.g., adenosine (A), guanosine (G), cytidine (C), and (deoxy)thymidine (T)), a partial base count for that amplicon or other nucleic acid would include the number of any one of those four nucleobase types (e.g., [A,G], [G,C], or [T,T]), any two of those four nucleobase types (e.g., [A,G], [G,C], [A,T], [G,C], or [C,T]), or at most any three of those four nucleobase types (e.g., [A,G,C], [A,C,T], [A,G,T], or [G,C,T]), in which w, x, y and/or z are each independently a whole number representing the number of said nucleoside residues in that amplicon or other nucleic acid. To further illustrate, if a nucleic acid has the following composition: ATTTGCTAAGGTAAACG, then partial base counts for that nucleic acid include [A,w], [G,x], [C,y], [T,z], [A,w,G], [A,w,C], [A,w,T], [G,x,C], [G,x,T], [C,y,T], [A,w,G,C], [A,w,G,T], [A,w,C,T], [G,x,C,T], or [G,x,C,T].

[0036] FIG. 1 schematically illustrates an exemplary process according to one embodiment of the invention. As shown, the process includes obtaining a sample with template nucleic acids. The sample is prepared, for example, by purifying or at partially isolating the template nucleic acids. As shown, the template nucleic acids are amplified, for example, in polymerase chain reactions or other amplification approaches to produce amplicons. In some embodiments, template nucleic acids are analyzed directly, for example, without being amplified. Amplicon base counts and/or sequences are determined, for example, using an array of zero-mode waveguides and a fluorescence microscope (and, e.g., primer nucleic acids (e.g., intelligent primers), polymerses (e.g., Φ29 DNA polymerses, etc.), phosphorylated nucleotides, etc, as also described in, for example, Kornreich et al. (2008) “Selective aluminum passivation for targeted immobilization of single DNA polymerase molecules in zero-mode waveguide nanostructures” Proc. Nat’l Acad. Sci. U.S.A. 105(4):1176-1181, herein incorporated by reference. The determined amplicon base counts and/or amplicon sequence information is used to query a base counts and/or sequence database to detect, identify, characterize, etc. the template nucleic acids. See also, FIG. 2 for additional exemplary variations.

[0037] Sequencing Technologies

[0038] As described above, embodiments of the present invention involve determining the base composition, or partial base composition, or a target sequence without determining the mass of the target sequence (e.g., without using mass spectrometry related methods). In certain embodiments, such methods of determining base compositions employ sequencing methods. The present invention is not limited by the type of sequencing method employed. Exemplary sequencing methods are described below.

[0039] Illustrative non-limiting examples of nucleic acid sequencing techniques include, but are not limited to, chain terminator (Sanger) sequencing and dye terminator sequencing. Those of ordinary skill in the art will recognize that because RNA is less stable in the cell and more prone to nucleic acid degradation, RNA is usually reverse transcribed to DNA before sequencing.

[0040] Chain terminator sequencing uses sequence-specific termination of a DNA synthesis reaction using modified nucleotide substrates. Extension is initiated at a specific site
on the template DNA by using a short radioactive, or other labeled, oligonucleotide primer complementary to the template at that region. The oligonucleotide primer is extended using a DNA polymerase, standard four deoxynucleotide bases, and a low concentration of one chain terminating nucleotide, most commonly a di-deoxynucleotide. This reaction is repeated in four separate tubes with each of the bases taking turns as the di-deoxynucleotide. Limited incorporation of the chain terminating nucleotide by the DNA polymerase results in a set of related DNA fragments that are terminated only at positions where that particular di-deoxynucleotide is used. For each reaction tube, the fragments are size-separated by electrophoresis in a slab polyacrylamide gel or a capillary tube filled with a viscous polymer. The sequence is determined by reading which lane produces a visualized band from the labeled primer as you scan from the top of the gel to the bottom.

Dye terminator sequencing alternatively labels the terminators. Complete sequencing can be performed in a single reaction by labeling each of the di-deoxynucleotide chain-terminators with a separate fluorescent dye, which fluoresces at a different wavelength.

A set of methods referred to as “next-generation sequencing” techniques have emerged as alternatives to Sanger and dye-terminator sequencing methods (Voelkerding et al., Clinical Chem., 55: 641-658, 2009; MacLean et al., Nature Rev. Microbiol., 7: 287-296; each herein incorporated by reference in their entirety). Next-generation sequencing (NGS) methods share the common feature of massively parallel, high-throughput strategies, with the goal of lowering costs in comparison to older sequencing methods. NGS methods can be broadly divided into those that require template amplification and those that do not. Amplification-requiring methods include pyrosequencing commercialized by Roche as the 454 technology platforms (e.g., GS 20 and GS FLX), the Solexa platform commercialized by Illumina, and the Supported Oligonucleotide Ligation and Detection (SOLiD) platform commercialized by Applied Biosystems. Non-amplification approaches, also known as single-molecule sequencing, are exemplified by the Heliscope platform commercialized by Helicos Biosciences, and emerging platforms commercialized by VisiGien, Oxford Nanopore Technologies Ltd., and Pacific Biosciences, respectively.

In pyrosequencing (Voelkerding et al., Clinical Chem., 55: 641-658, 2009; MacLean et al., Nature Rev. Microbiol., 7: 287-296; each herein incorporated by reference in their entirety), template DNA is fragmented, end-repaired, ligated to adaptors, and clonally amplified in situ by capturing single template molecules with beads bearing oligonucleotides complementary to the adaptors. Each bead bearing a single template type is compartmentalized into a water-in-oil microvesicle, and the template is clonally amplified using a technique referred to as emulsion PCR. The emulsion is disrupted after amplification and beads are deposited into individual wells of a picotitre plate functioning as a flow cell during the sequencing reactions. Ordered, iterative introduction of each of the four dNTP reagents occurs in the flow cell in the presence of sequencing enzymes and luminescent reporter such as luciferase. In the event that an appropriate dNTP is added to the 3’ end of the sequencing primer, the resulting production of ATP causes a burst of luminescence within the well, which is recorded using a CCD camera. It is possible to achieve read lengths greater than or equal to 400 bases, and 1 x 10^6 sequence reads can be achieved, resulting in up to 500 million base pairs (Mb) of sequence.

In the Solexa/Illumina platform (Voelkerding et al., Clinical Chem., 55: 641-658, 2009; MacLean et al., Nature Rev. Microbiol., 7: 287-296; U.S. Pat. No. 6,833,246; U.S. Pat. No. 7,115,400; U.S. Pat. No. 6,969,488; each herein incorporated by reference in its entirety), sequencing data are produced in the form of shorter-length reads. In this method, single-stranded fragmented DNA is end-repaired to generate 5’-phosphorylated blunt ends, followed by Klenow-mediated addition of a single A base to the 3’ end of the fragments. A-addition facilitates addition of T-overhang adaptor oligonucleotides, which are subsequently used to capture the template-adaptor molecules on the surface of a flow cell that is studded with oligonucleotide anchors. The anchor is used as a PCR primer, but because of the length of the template and its proximity to other nearby oligonucleotides, extension by PCR results in the “arching over” of the molecule to hybridize with an adjacent anchor oligonucleotide to form a bridge structure on the surface of the flow cell. These loops of DNA are denatured and cleaned. Forward strands are then sequenced with reversible dye terminators. The sequence of incorporated nucleotides is determined by detection of post-incorporation fluorescence, with each fluor and block removed prior to the next cycle of dNTP addition. Sequence read length ranges from 36 nucleotides to over 50 nucleotides, with overall output exceeding 1 billion nucleotide pairs per analytical run.

Sequencing nucleic acid molecules using SOLiD technology (Voelkerding et al., Clinical Chem., 55: 641-658, 2009; MacLean et al., Nature Rev. Microbiol., 7: 287-296; U.S. Pat. No. 5,912,148; U.S. Pat. No. 6,130,073; each herein incorporated by reference in their entirety) also involves fragmentation of the template, ligation to oligonucleotide adaptors, attachment to beads, and clonal amplification by emulsion PCR. Following this, beads bearing template are immobilized on a derivatized surface of a glass flow-cell, and a primer complementary to the adaptor oligonucleotide is annealed. However, rather than utilizing this primer for 3’ extension, it is instead used to provide a 5’ phosphate group for ligation to interrogation probes containing two probe-specific bases followed by 6 degenerate bases and one of four fluorescent labels. In the SOLiD system, interrogation probes have 16 possible combinations of the two bases at the 3’ end of each probe, and one of four fluor at the 5’ end. Fluor color and thus identity of each probe corresponds to specified color-space coding schemes. Multiple rounds (usually 7) of probe annealing, ligation, and fluor detection are followed by denaturation, and then a second round of sequencing using a primer that is offset by one base relative to the initial primer. In this manner, the template sequence can be computationally re-constructed, and template bases are interrogated twice, resulting in increased accuracy. Sequence read length averages 35 nucleotides, and overall output exceeds 4 billion bases per sequencing run.

In certain embodiments, nanopore sequencing in employed (see, e.g., Astier et al., J Am Chem Soc. 2006 Feb. 8; 128(5):1705-10, herein incorporated by reference). The theory behind nanopore sequencing has to do with what occurs when the nanopore is immersed in a conducting fluid and a potential (voltage) is applied across it: under these conditions a slight electric current due to conduction of ions through the nanopore can be observed, and the amount of current is exceedingly sensitive to the size of the nanopore. If
DNA molecules pass (or part of the DNA molecule passes) through the nanopore, this can create a change in the magnitude of the current through the nanopore, thereby allowing the sequences of the DNA molecule to be determined.

[0047] HeliScope by Helicos BioSciences (Voelkerding et al., Clinical Chem., 55: 641-658, 2009; Maclennan et al., Nature Rev. Microbiol., 7: 287-296; U.S. Pat. No. 7,169,560; U.S. Pat. No. 7,282,337; U.S. Pat. No. 7,482,120; U.S. Pat. No. 7,501,245; U.S. Pat. No. 6,818,395; U.S. Pat. No. 6,911,345; U.S. Pat. No. 7,501,245; each herein incorporated by reference in their entirety) is the first commercialized single-molecule sequencing platform. This method does not require clonal amplification. Template DNA is fragmented and polyadenylated at the 3’ end, with the final adenosine bearing a fluorescent label. Denatured polyadenylated template fragments are ligated to poly(dT) oligonucleotides on the surface of a flow cell. Initial physical locations of captured template molecules are recorded by a CCD camera, and then label is cleaved and washed away. Sequencing is achieved by addition of polymerase and serial addition of fluorescently-labeled dNTP reagents. Incorporation events result in fluorescent signal corresponding to the dNTP and signal is captured by a CCD camera before each round of dNTP addition. Sequence read length ranges from 25-50 nucleotides, with overall output exceeding 1 billion nucleotide pairs per analytical run.

[0048] Another exemplary nucleic acid sequencing approach developed by Stratas Genomics, Inc. that is also optionally adapted for use with the present invention involves the use of Xpanders. This sequencing process typically includes providing a daughter strand produced by a templatedirected synthesis. The daughter strand generally includes a plurality of subunits coupled in a sequence corresponding to a contiguous nucleotide sequence of all or a portion of a target nucleic acid in which the individual subunits comprise a tether, at least one probe or nucleobase residue, and at least one selectively cleavable bond. The selectively cleavable bond(s) is/are cleaved to yield an Xpander of a length longer than the plurality of the subunits of the daughter strand. The Xpander typically includes the tethers and reporter elements for parsing genetic information in a sequence corresponding to the contiguous nucleotide sequence of all or a portion of the target nucleic acid. Reporter elements of the Xpander are then detected. Additional details relating to Xpander-based approaches are described in, for example, U.S. Patent Publication No. 20000039777, entitled "HIGH THROUGHPUT NUCLEIC ACID SEQUENCING BY EXPANSION," that was filed Jun. 19, 2008, which is incorporated herein in its entirety.

[0049] Other emerging single molecule sequencing methods include real-time sequencing by synthesis using a VigiGen platform (Voelkerding et al., Clinical Chem., 55: 641-658, 2009; U.S. Pat. No. 7,329,492; U.S. patent application Ser. No. 11/671,956; U.S. patent application Ser. No. 11/781,166; each herein incorporated by reference in their entirety) in which immobilized, primed DNA template is subjected to strand extension using a fluorescently-modified polymerase and fluorescent acceptor molecules, resulting in detectable fluorescence resonance energy transfer (FRET) upon nucleotide addition.

[0050] Another real-time single molecule sequencing system developed by Pacific Biosciences (Voelkerding et al., Clinical Chem., 55: 641-658, 2009; Maclennan et al., Nature Rev. Microbiol., 7: 287-296; U.S. Pat. No. 7,170,050; U.S. Pat. No. 7,302,146; U.S. Pat. No. 7,313,308; U.S. Pat. No. 7,476,503; all of which are herein incorporated by reference) utilizes reaction wells 50-100 nm in diameter and encompassing a reaction volume of approximately 20 zeptoliters (10^-23 L). Sequencing reactions are performed using immobilized template, modified phi29 DNA polymerase, and high local concentrations of fluorescently labeled dNTPs. High local concentrations and continuous reaction conditions allow incorporation events to be captured in real time by fluorescent signal detection using laser excitation, an optical waveguide, and a CCD camera.

[0051] In certain embodiments, the single molecule real time (SMRT) DNA sequencing methods using zero-mode waveguides (ZMWs) developed by Pacific Biosciences, or similar methods, are employed. With this technology, DNA sequencing is performed on SMRT chips, each containing thousands of zero-mode waveguides (ZMWs). A ZMW is a hole, tens of nanometers in diameter, fabricated in a 100 nm metal film deposited on a silicon dioxide substrate. Each ZMW becomes a nanophotonic visualization chamber providing a detection volume of just 20 zeptoliters (10^-21 liters). At this volume, the activity of a single molecule can be detected amongst a background of thousands of labeled nucleotides.

[0052] The ZMW provides a window for watching DNA polymerase as it performs sequencing by synthesis. Within each chamber, a single DNA polymerase molecule is attached to the bottom surface such that it permanently resides within the detection volume. Phospholinked nucleotides, each type labeled with a different colored fluorophore, are then introduced into the reaction solution at high concentrations which promote enzyme speed, accuracy, and processivity. Due to the small size of the ZMW, even at these high, biologically relevant concentrations, the detection volume is occupied by nucleotides only a small fraction of the time. In addition, visits to the detection volume are fast, lasting only a few microseconds, due to the very small distance that diffusion has to carry the nucleotides. The result is a very low background.

[0053] As the DNA polymerase incorporates complementary nucleotides, each base is held within the detection volume for tens of milliseconds, which is orders of magnitude longer than the amount of time it takes a nucleotide to diffuse in and out of the detection volume. During this time, the engaged fluorophore emits fluorescent light whose color corresponds to the base identity. Then, as part of the natural incorporation cycle, the polymerase cleaves the bond holding the fluorophore in place and the dye diffuses out of the detection volume. Following incorporation, the signal immediately returns to baseline and the process repeats.

[0054] Unhampered and uninterrupted, the DNA polymerase continues incorporating bases at a speed of tens per second. In this way, a completely natural long chain of DNA is produced in minutes. Simultaneous and continuous detection occurs across all of the thousands of ZMWs on the SMRT chip in real time. This approach has the capability to produce reads thousands of nucleotides in length.

[0055] Some aspects the invention are generally directed to optical detection or monitoring systems (e.g., ZMWs discussed above), methods enabled by such systems, and components of such systems for monitoring, in real-time, optical signals that emanate from multiple discrete sources of those optical signals. In particular, the optical detection and monitoring systems are generally capable of monitoring discrete signals from potentially very large numbers of different sig-
nal sources, optionally separating and/or deconvolving such signals into constituent signal events, and doing so in real-time, despite that such signals may be changing rapidly, over time.

[0056] The systems typically include all or a portion of a collection of different functional elements. These elements include the multiple discrete sources that include the capability of generating optical signals. In some aspects, such sources include chemical, biochemical and/or biological reactants, or mimics of such reactants that are capable of generating optical signals that are indicative of their presence, reaction or conversion. While the sources may be capable of generating optical signals on their own, in certain cases, a source of excitation radiation is also provided to excite optical signals, e.g., fluorescence, within the sources.

[0057] The systems also typically include optical elements that direct, separate, and/or otherwise alter optical signals from these sources (as well as excitation radiation directed at such sources), in order to ultimately derive optimal amounts of information from such signals when they are ultimately detected. Consequently, the systems typically include an optical detection system for detecting the potentially large numbers of signals that were directed from the sources, and optionally separated and/or otherwise altered by the optical elements. Signals detected by the optical detection system are then recorded and processed by appropriate processing systems and data management processes to provide output of the system in user ready formats.

[0058] The systems are typically applied in the monitoring of arrays or collections of spatially discrete chemical, biochemical and/or biological reactions that generate optically detectable signals, such as chromogenic reactions, luminescent or luminogenic reactions, or fluorescent or fluorogenic reactions. A few examples of reactions include those that are regularly performed in the pharmaceutical, biotechnology and healthcare diagnostic fields, i.e., base count analyses, immunosassays, enzymatic assays, receptor assays, nucleic acid hybridization assays, nucleic acid synthesis reactions, cellular assays, and many others.

[0059] Typically, the progress of the reactions used in application of the systems described herein result in one or more of the consumption, production and/or conversion of a material that is capable of generating an optically detectable signal, either alone, or in response to an external stimulus, e.g., excitation radiation. By way of example, certain reactants may become fluorescent upon reaction with another reactant, or may have their fluorescence altered or reduced upon such reaction. As such, the fluorescence emitted from the reaction in response to an excitation radiation will change as the reaction progresses. The systems generally provide for the source of such signals, e.g., the area in which the reaction occurs, including optionally, the reactants and/or products, the optical elements for collecting, directing and optionally separating and/or altering such signals from such sources, and the ultimate detection of such signals, as well as the manipulation of the resulting data to yield optimal value and information for the user.

[0060] The systems typically include all or a subset of a substrate that includes all or a subset of the sources of optical signals, an optional excitation light source, an optical train that includes the various optical elements for collection, direction and/or manipulation of the optical signals and optional excitation light, optical detectors for receiving, detecting and recording (or putting into a form for recordation) the optical signals, as well as processors for processing data derived from the optical detectors.

[0061] A general schematic representation of the system as set forth above, is illustrated in FIG. 3. As shown, the system includes a substrate that includes a plurality of discrete sources of optical signals, e.g., reaction wells or optical confinements. An excitation light source, e.g., laser, is optionally provided in the system and is positioned to direct excitation radiation at the various signal sources. This is typically done by directing excitation radiation at or through appropriate optical components, e.g., dichroic and objective lenses, that direct the excitation radiation at the substrate, and particularly the signal sources. Emitters signals from source are then collected by the optical components, e.g., objective lenses, and passed through additional optical elements, e.g., dichroic, prism lens, and lens, until they are directed to and impinge upon an optical detection system, e.g., detector array. The signals are then detected by detector array, and the data from that detection is transmitted to an appropriate data processing unit, e.g., computer, where the data is subjected to interpretation, analysis, and ultimately presented in a user ready format, e.g., on display, or printed on, from printer.


DEFINITIONS AND FURTHER DESCRIPTION

[0063] It is to be understood that the terminology used herein is for thhe purpose of describing particular embodiments only, and is not intended to be limiting. Further, unless defined otherwise, all technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which this invention pertains. In describing and claiming the present invention, the following terminology and grammatical variants will be used in accordance with the definitions set forth below.

[0064] As used herein, the term “about” means encompassing plus or minus 10%. For example, about 200 nucleotides refers to a range encompassing between 180 and 220 nucleotides.

[0065] As used herein, the term “amplicon” or “bioagent identifying amplicon” refers to a nucleic acid generated using the primer pairs described herein. The amplicon is typically double stranded DNA; however, it may be RNA and/or DNA. In some embodiments, the amplicon comprises DNA complementary to HPV RNA, DNA, or cDNA. In some embodiments, the amplicon comprises sequences of conserved regions/primer pairs and intervening variable region. As discussed herein, primer pairs are configured to generate amplicons from bioagent nucleic acid. As such, the base composition of any given amplicon may include the primer pair, the complement of the primer pair, the conserved regions and the variable region from the bioagent that was amplified to generate the amplicon. One skilled in the art understands that the incorporation of the designed primer pair sequences into an amplicon may replace the native sequences at the primer binding site, and complement thereof. In certain embodiments, after amplification of the target region using the primers the resultant amplicons having the primer sequences are used to generate the base composition data. Bioagent identifying amplicons generate base compositions that are preferably unique to the identity of a bioagent.

[0066] The term “amplifying” or “amplification” in the context of nucleic acids refers to the production of multiple copies of a polynucleotide, or a portion of the polynucleotide, typically starting from a small amount of the polynucleotide (e.g., a single polynucleotide molecule), where the amplification products or amplicons are generally detectable. Amplification of polynucleotides encompasses a variety of chemical and enzymatic processes. The generation of multiple DNA copies from one or a few copies of a target or template DNA molecule during a polymerase chain reaction (PCR) or a ligase chain reaction (LCR) are forms of amplification. Amplification is not limited to the strict duplication of the starting molecule. For example, the generation of multiple cDNA molecules from a limited amount of RNA in a sample using reverse transcription (RT)-PCR is a form of amplification. Furthermore, the generation of multiple RNA molecules from a single DNA molecule during the process of transcription is also a form of amplification.

[0067] The term “attached” refers to interactions including, but not limited to, covalent bonding, ionic bonding, chemisorption, physisorption, and combinations thereof.

[0068] As used herein, “viral nucleic acid” includes, but is not limited to, DNA, RNA, or DNA that has been obtained from viral RNA, such as, for example, by performing a reverse transcription reaction. Viral RNA can either be single-stranded (of positive or negative polarity) or double-stranded.

[0069] As used herein, the term “base composition” or “base count” refers to the number of each residue comprised in an amplicon or other nucleic acid, without consideration for the linear arrangement of these residues in the strand(s) of the amplicon. The amplicon residues comprise, adenosine (A), guanosine (G), cytidine (C), (deoxy)thymidine (T), uracil (U), inosine (I), nitroindoles such as 5-nitroindole or...
As used herein, "broad range survey primers" are primers designed to identify an unknown bioagent as a member of a particular biological division (e.g., an order, family, class, Glade, or genus). However, in some cases the broad range survey primers are also able to identify unknown bioagents at the species or sub-species level. As used herein, "division-wide primers" are primers designed to identify a bioagent at the species level and "drill-down" primers are primers designed to identify a bioagent at the sub-species level. As used herein, the "sub-species" level of identification includes, but is not limited to, strains, subtypes, variants, and isolates. Drill-down primers are not always required for identification at the sub-species level because broad range survey intelligent primers may, in some cases provide sufficient identification resolution to accomplish this identification objective. Broad range survey primers may be used in the non-mass determined base compositions methods and systems of the present invention.

As used herein, the terms "complementary" or "complementarity" are used in reference to polynucleotides (i.e., a sequence of nucleotides) related by the base-pairing rules. For example, the sequence "5'-A-G-3'," is complementary to the sequence "3'-T-C-A-5'." Complementarity may be "partial," in which only some of the nucleic acids' bases are matched according to the base pairing rules. Or, there may be "complete" or "total" complementarity between the nucleic acids. The degree of complementarity between nucleic acid strands has significant effects on the efficiency and strength of hybridization between nucleic acid strands. This is of particular importance in amplification reactions, as well as detection methods that depend upon binding between nucleic acids.

The term "conserved region" in the context of nucleic acids refers to a nucleobase sequence (e.g., a subsequence of a nucleic acid, etc.) that is the same or similar in two or more different regions or segments of a given nucleic acid molecule (e.g., an intramolecular conserved region), or that is the same or similar in two or more different nucleic acid molecules (e.g., an intermolecular conserved region). To illustrate, a conserved region may be present in two or more different taxonomic ranks (e.g., two or more different genera, two or more different species, two or more different subspecies, and the like) or in two or more different nucleic acid molecules from the same organism. To further illustrate, in certain embodiments, nucleic acids comprising at least one conserved region typically have between about 70%-100%, between about 80-100%, between about 90-100%, between about 95-100%, or between about 99-100% sequence identity in that conserved region. A conserved region may also be selected or identified functionally as a region that permits generation of amplicons via primer extension through hybridization of a completely or partially complementary primer to the conserved region for each of the target sequences to which conserved region is conserved.

As used herein, in some embodiments the term "database" is used to refer to a collection of base composition and/or partial base composition data. The base composition data in the database is indexed to bioagents and to primer pairs. The base composition data reported in the database comprises the number of at least one type of nucleoside in an amplicon (e.g., A, T, G, or C) that would be generated for each bioagent using each primer. The database can be populated by empirical data. In this aspect of populating the database, a bioagent is selected and a primer pair is used to generate an
amplicon. Note that base composition entries in the database may be derived from sequencing data (i.e., known sequence information). An entry in the database is made to associate the base composition with the biocatalyst and the primer pair used. The database may also be populated using other databases comprising biocatalytic information. For example, using the GenBank database it is possible to perform electronic PCR using an electronic representation of a primer pair. This in silico method may provide the base composition for any or all selected biocatalyst(s) stored in the GenBank database. The information may then be used to populate the base composition database as described above. A base composition database can be in silico, a written table, a reference book, a spreadsheet or any format generally amenable to databases. Preferably, it is in silico on computer readable media.

[0078] The term “detect”, “detecting” or “detection” refers to an act of determining the existence or presence of one or more targets (e.g., biocatalyst nucleic acids, amplicons, etc.) in a sample.

[0079] As used herein, the term “etiology” refers to the causes or origins, of diseases or abnormal physiological conditions.

[0080] Nucleic acids are “extended” or “elongated” when additional nucleotides (or other analogous molecules) are incorporated into the nucleic acids. For example, a nucleic acid is optionally extended by a nucleotide incorporating biocatalyst, such as a polymerase that typically adds nucleotides at the 3′ terminal end of a nucleic acid.

[0081] An “extended primer nucleic acid” refers to a primer nucleic acid to which one or more additional nucleotides have been added or otherwise incorporated (e.g., covalently bonded to).

[0082] As used herein, the term “gene” refers to a nucleic acid (e.g., DNA) sequence that comprises coding sequences necessary for the production of a polypeptide, precursor, or RNA (e.g., rRNA, tRNA). The polypeptide can be encoded by a full length coding sequence or by any portion of the coding sequence so long as the desired activity or functional properties (e.g., enzymatic activity, ligand binding, signal transduction, immunogenicity, etc.) of the full-length sequence or fragment thereof are retained.

[0083] A “genotype” refers to all or part of the genetic constitution of a nucleic acid molecule, cell or subject, or group of nucleic acid molecules, cells or subjects. For example, a genotype includes the particular mutations and/or alleles (e.g., polymorphisms, such as single nucleotide polymorphism, SNPs or the like) present at a given locus or distributed in a genome.

[0084] A “heterocyclic ring” refers to a monocyclic or bicyclic ring that is either saturated, unsaturated, or aromatic, and which comprises one or more heteroatoms independently selected from nitrogen, oxygen and sulfur. A heterocyclic ring may be attached to the sugar moiety, or analog thereof, of a nucleotide of the invention via any heteroatom or carbon atom. Exemplary heterocyclic rings include morpholinline, pyrrolindinyl, pyrrolidinyl, piperidinyl, hydantoinyl, valerolactamyl, oxiranyl, oxetanyl, tetrahydrofuranyl, tetrahydropyranyl, tetrahydropyridinyl, tetrahydroprimidinyl, tetrahydrothiophenyl, tetrahydrothiopyranyl, tetrahydrothiopyridinyl, tetrahydrothiophenyl, tetrahydrothiopyranyl, 5-furyl, benzofuranyl, thiophenyl, benzothiophenyl, pyrydyl, indolyl, isoindolyl, azaindolyl, pyridyl, quinolinyl, isoquinolinyl, oxazolyl, isooxazolyl, benzoxazolyl, pyrazolyl, imidazolyl, benzimidazolyl, thiazolyl, benzothiazolyl, iso(thiazolyl), pyridazinyl, pyrimidinyl, pyrazinyl, quinolinyl, pthialdazinyl, quinazolinyl, and the like.

[0085] A “homocyclic ring” refers to a saturated (but not aromatic) carbocyclic ring, such as cyclopropyl, cyclobutyl, cyclopentyl, cyclohexane, cycloheptane, cyclohexene, and the like.

[0086] As used herein, the term “heterologous gene” refers to a gene that is not in its natural environment. For example, a heterologous gene includes a gene from one species introduced into another species. A heterologous gene also includes a gene native to an organism that has been altered in some way (e.g., mutated, added in multiple copies, linked to non-native regulatory sequences, etc). Heterologous genes are distinguished from endogenous genes in that the heterologous gene sequences are typically joined to nucleic acid sequences that are not found naturally associated with the gene sequences in the chromosome or are associated with portions of the chromosome not found in nature (e.g., genes expressed in loci where the gene is not normally expressed).

[0087] The terms “homology,” “homologous” and “sequence identity” refer to a degree of identity. There may be partial homology or complete homology. A partially homologous sequence is one that is less than 100% identical to another sequence. Determination of sequence identity is described in the following example: a primer 20 nucleobases in length which is otherwise identical to another 20 nucleobase primer but having two non-identical residues has 18 of 20 identical residues (18/20 = 0.9 or 90% sequence identity). In another example, a primer 15 nucleobases in length having all residues identical to a 15 nucleobase segment of a primer 20 nucleobases in length would have 15/20 = 0.75 or 75% sequence identity with the 20 nucleobase primer. In context of the present invention, sequence identity is meant to be properly determined when the query sequence and the subject sequence are both described and aligned in the 5′ to 3′ direction. Sequence alignment algorithms such as BLAST, will return results in two different alignment orientations. In the Plus/Plus orientation, both the query sequence and the subject sequence are aligned in the 5′ to 3′ direction. On the other hand, in the Plus/Minus orientation, the query sequence is in the 5′ to 3′ direction while the subject sequence is in the 3′ to 5′ direction. It should be understood that with respect to the primers of the present invention, sequence identity is properly determined when the alignment is designated as Plus/Plus. Sequence identity may also encompass alternate or “modified” nucleobases that perform in a functionally similar manner to the regular nucleobases adenine, thymine, guanine, and cytosine with respect to hybridization and primer extension in amplification reactions. In a non-limiting example, if the 5-propynyl pyrimidines propyne C and/or propyne T replace one or more C or T residues in one primer which is otherwise identical to another primer in sequence and length, the two primers will have 100% sequence identity with each other. In another non-limiting example, Inosine (I) may be used as a replacement for G or T and effectively hybridize to C, A or U (uracil). Thus, if inosine replaces one or more C, A or U residues in one primer which is otherwise identical to another primer in sequence and length, the two primers will have 100% sequence identity with each other. Other such modified or universal bases may exist which would perform in a functionally similar manner for hybridization and amplification reactions and will be understood to fall within this definition of sequence identity.
As used herein, “housekeeping gene” or “core viral gene” refers to a gene encoding a protein or RNA involved in basic functions required for survival and reproduction of a bioagent. Housekeeping genes include, but are not limited to, genes encoding RNA or proteins involved in translation, replication, recombination and repair, transcription, nucleotide metabolism, amino acid metabolism, lipid metabolism, energy generation, uptake, secretion and the like.

As used herein, the term “hybridization” or “hybridize” is used in reference to the pairing of complementary nucleic acids. Hybridization and the strength of hybridization (i.e., the strength of the association between the nucleic acids) is influenced by such factors as the degree of complementarity between the nucleic acids, stringency of the conditions involved, the melting temperature (Tm) of the formed hybrid, and the G:C ratio within the nucleic acids. A single molecule that contains pairing of complementary nucleic acids within its structure is said to be “self-hybridized.” An extensive guide to nucleic hybridization may be found in Tijssen, Laboratory Techniques in Biochemistry and Molecular Biology-Hybridization with Nucleic Acid Probes, part I, chapter 2, “Overview of principles of hybridization and the strategy of nucleic acid probe assays,” Elsevier (1993), which is incorporated by reference.

A “label” refers to a moiety attached (covalently or non-covalently), or capable of being attached, to a molecule, which moiety provides or is capable of providing information about the molecule (e.g., descriptive, identifying, etc. information about the molecule). Exemplary labels include fluorescent labels, non-fluorescent labels, colorimetric labels, chemiluminescent labels, bioluminescent labels, radioactive labels, mass-modifying groups, antibodies, antigens, biotin, haptons, and enzymes (including, e.g., peroxidase, phosphatase, etc.).

The term “kit” is used in reference to a combination of articles that facilitate a process, method, assay, analysis or manipulation of a sample. Kits can contain instructions describing how to use the kit (e.g., instructions describing the methods of the invention), zero-mode waveguides, primer nucleic acids, nucleotide incorporating biocatalysts, labeled nucleotides, chemical reagents, as well as any other components.

A “linker” or “spacer” refers to a chemical moiety that covalently or non-covalently (e.g., ionically, etc.) attaches a compound or substituent group to, e.g., a solid support, another compound or group, or the like. For example, a linker optionally attaches a label (e.g., a fluorescent dye, a radionuclide, etc.) to a nucleotide or the like. Linkers are typically bifunctional chemical moieties and in certain embodiments, they comprise cleavable attachments, which can be cleaved by, e.g., heat, an enzyme, a chemical agent, electromagnetic radiation, etc. to release materials or compounds from, e.g., a solid support, another compound, etc. A careful choice of linker allows cleavage to be performed under appropriate conditions compatible with the stability of the compound and assay method. Generally a linker has no specific biological activity other than to, e.g., join chemical species together or to preserve some minimum distance or other spatial relationship between such species. However, the constituents of a linker may be selected to influence some property of the linked chemical species such as three-dimensional conformation, net charge, hydrophobicity, etc. Additional description of linker molecules is provided in, e.g., Lyttle et al. (1996) Nucleic Acids Res. 24(14):2793, Shchepino et al. (2001) Nucleosides, Nucleotides, & Nucleic Acids 20:369, Doronina et al. (2001) Nucleosides, Nucleotides, & Nucleic Acids 20:1007, Trawick et al. (2001) Bioconjugate Chem. 12:900, Olejnik et al. (1998) Methods in Enzymology 291:135, Pjiljevic et al. (2003) J. Am. Chem. Soc. 125(12):3486, Ward, et al., U.S. Pat. No. 4,711,955, Stavrionopolos, U.S. Pat. No. 4,707,552, and Stavrionopolos, U.S. Pat. No. 4,707,440, which are each incorporated by reference.

A “mass modifying” group modifies the mass, typically measured in terms of molecular weight as daltons, of a nucleic acid that comprises the group. For example, mass modifying groups that increase the discrimination between at least two nucleic acids with single base differences in size or sequence can be used to facilitate sequencing using, e.g., molecular weight determinations.

A “mixture” refers to a combination of two or more different components. A “reaction mixture” refers a mixture that comprises molecules that can participate in and/or facilitate a given reaction.

The term “molecular mass” refers to the mass of a compound (e.g., a nucleic acid, etc.) as determined, for example, using mass spectrometry.

A “modified” enzyme refers to an enzyme comprising a monomer sequence in which at least one monomer of the sequence differs from a monomer in a reference sequence, such as a native or wild-type form of the enzyme or another modified form of the enzyme, e.g., when the two sequences are aligned for maximum identity. Exemplary modifications include monomer insertions, deletions, and substitutions. The modified enzymes (i.e., protein- or nucleic acid-based catalysts) of the invention have been or are optionally created by various diversity generating methods. Although essentially any method can be used to produce a modified enzyme, certain exemplary techniques include recombinig (e.g., via recursive recombination, synthetic recombination, or the like) two or more nucleic acids encoding one or more parental enzymes, or by mutating one or more nucleic acids that encode enzymes, e.g., using recursive ensemble mutagenesis, cassette mutagenesis, random mutagenesis, in vivo mutagenesis, site directed mutagenesis, or the like. A nucleic acid encoding a parental enzyme typically includes a gene that, through the mechanisms of transcription and translation, produces an amino acid sequence corresponding to a parental enzyme, e.g., a native form of the enzyme. Modified enzymes also include chimeric enzymes that have identifiable component sequences (e.g., structural and/or functional domains, etc.) derived from two or more parents. Also included within the definition of modified enzymes are those comprising chemical modifications (e.g., attached substituent groups, altered substituent groups, etc.) relative to a reference sequence.

A “moiety” or “group” refers to one of the portions into which something, such as a molecule, is divided (e.g., a functional group, substituent group, or the like). For example, a nucleotide typically comprises a basic group (e.g., adenine, thymine, cytosine, guanine, uracil, or an analog basic group), a sugar moiety (e.g., a moiety comprising a sugar ring or an analog thereof), and one or more phosphate groups.

As used herein, the term “primer” or “primer nucleic acid” refers to an oligonucleotide, whether occurring naturally as in a purified restriction digest or produced synthetically, that is capable of acting as a point of initiation of synthesis when placed under conditions in which synthesis of
a primer extension product that is complementary to a nucleic acid strand is induced (e.g., in the presence of nucleotides and an inducing agent such as a biocatalyst (e.g., a DNA polymerase or the like) and at a suitable temperature and pH). The primer is typically single stranded for maximum efficiency in amplification, but may alternatively be double stranded. If double stranded, the primer is generally first treated to separate its strands before being used to prepare extension products. In some embodiments, the primer is an oligodeoxyribonucleotide. The primer is sufficiently long to prime the synthesis of extension products in the presence of the inducing agent. The exact lengths of the primers will depend on many factors, including temperature, source of primer and the use of the method.

[0099] As used herein, “intelligent primers” or “primers” or “primer pairs,” in some embodiments, are oligonucleotides that are designed to bind to conserved sequence regions of one or more biocatalytic nucleic acids to generate biocatalyst identifying amplicons. In some embodiments, the bound primers flank an intervening variable region between the conserved binding sequences. Upon amplification, the primer pairs yield amplicons e.g., amplification products that provide base composition variability between the two or more biocatalysts. The variability of the base compositions allows for the identification of one or more individual biocatalysts from, e.g., two or more biocatalysts based on the base composition distinctions. In some embodiments, the primer pairs are also configured to generate amplicons amenable to molecular mass analysis. Further, the sequences of the primer members of the primer pairs are not necessarily fully complementary to the conserved region of the reference biocatalyst. For example, in some embodiments, the sequences are designed to be “best fit” amongst a plurality of biocatalysts at these conserved binding sequences. Therefore, the primer members of the primer pairs have substantial complementarity with the conserved regions of the biocatalysts, including the reference biocatalyst.

[0100] In some embodiments of the invention, the oligonucleotide primer pairs described herein can be purified. As used herein, “purified oligonucleotide primer pair,” “purified primer pair,” or “purified” means an oligonucleotide primer pair that is chemically synthesized to have a specific sequence and a specific number of linked nucleosides. This term is meant to explicitly exclude nucleosides that are generated at random to yield a mixture of several compounds of the same length each with randomly generated sequence. As used herein, the term “purified” or “to purify” refers to the removal of one or more components (e.g., contaminants) from a sample.

[0101] The term “nucleic acid” or “nucleic acid molecule” refers to any nucleic acid containing molecule, including but not limited to, DNA or RNA. The term encompasses sequences that include any of the known base analogs of DNA and RNA including, but not limited to, 4-acetylcytosine, 8-hydroxy-N9-methyladenosine, azidinylyctosine, pseudouridine, N4-acetylcytosine, N4-hydroxycytosine, N4-acetyluracil, 5-(carboxamidomethyl)-uracil, 5-fluorouracil, 5-bromouracil, 5-carboxamidomethyluracil, 2-thiouracil, 5-carboxamidomethyl-2-thiouracil, inosine, N9-isopentenyladenine, 1-methyladenine, 1-methyl-5-uracil, 1-methylguanine, 1-methylinosine, 2,2-dimethylguanine, 2-methyladenine, 2-methylguanine, 3-methylcytosine, 5-methylcytosine, N4-methyladenine, 7-methylguanine, 5-methylaminomethyluracil, 5-methoxyguanine, 7-methylguanine, 2-thiouracil, 4-thiouracil, 2-thiouracil, 5-uracil, 5-methyluracil, N-uracil-5-oxyacetic acid methylster, uracil-5-oxyacetic acid methylster, uracil-5-oxyacetic acid, pseudouracil, queosine, 2-thiocytosine, 5-methyl-2-thiouracil, 2-thiouracil, 4-thiouracil, 5-methyluracil, N-uracil-5-oxyacetic acid methylster, uracil-5-oxyacetic acid, pseudouracil, queosine, 2-thiocytosine, and 2,6-diaminopurine.

[0102] As used herein, the term “nucleoside” is synonymous with other terms in use in the art including “nucleoside,” “deoxynucleoside,” “nucleic acid residue,” “deoxyribonucleoside residue,” “nucleotide triphosphate (NTP),” or deoxynucleotide triphosphate (dNTP). As is used herein, a nucleoside includes natural and modified residues, as described herein.

[0103] A “nucleoside” refers to a nucleic acid component that comprises a base or basic group (e.g., comprising at least one homocyclic ring, at least one heterocyclic ring, at least one aryl group, and/or the like) covalently linked to a sugar moiety (e.g., a ribose sugar, etc.), a derivative of a sugar moiety, or a functional equivalent of a sugar moiety (e.g., an analog, such as carbocyclic ring). For example, when a nucleoside includes a sugar moiety, the base is typically linked to a 1'-position of that sugar moiety. As described above, a base can be naturally occurring (e.g., a purine base, such as adenine (A) or guanine (G), a pyrimidine base, such as thymine (T), cytosine (C), or uracil (U)), or non-naturally occurring (e.g., 7-deazapurine base, a pyrazolo[3,4-d]pyrimidine base, a propynyl-DN base, etc.). Exemplary nucleosides include ribonucleosides, deoxyribonucleosides, ribocytosinucleosides, carbocyclic nucleosides, etc.).

[0104] A “nucleotide” refers to an ester of a nucleoside, e.g., a phosphate ester of a nucleoside. For example, a nucleotide can include 1, 2, 3, or more phosphate groups covalently linked to a 5' position of a sugar moiety of the nucleoside.

[0105] A “nucleotide incorporating biocatalyst” refers to a catalyst that catalyzes the incorporation of nucleotides into a nucleic acid. Nucleotide incorporating biocatalysts are typically enzymes. An “enzyme” is a protein- and/or nucleic acid-based catalyst that acts to reduce the activation energy of a chemical reaction involving other compounds or “substrates.” A “nucleotide incorporating enzyme” refers to an enzyme that catalyzes the incorporation of nucleotides into a nucleic acid. Exemplary nucleotide incorporating enzymes include, e.g., DNA polymerases, RNA polymerases, terminal transferases, reverse transcriptases, telomerase, polynucleotide phosphorylases, and the like.

[0106] An “oligonucleotide” refers to a nucleic acid that includes at least two nucleic acid monomer units (e.g., nucleotides), typically more than three monomer units, and more typically greater than ten monomer units. The exact size of an oligonucleotide generally depends on various factors, including the ultimate function or use of the oligonucleotide. To further illustrate, oligonucleotides are typically less than 200 residues long (e.g., between 15 and 100), however, as used herein, the term is also intended to encompass longer polynucleotide chains. Oligonucleotides are often referred to by their length. For example a 24 residue oligonucleotide is referred to as a “24-mer”. Typically, the nucleoside monomers are linked by phosphodiester bonds or analogs thereof, including phosphorothioate, phosphorodithioate, phosphoroselenoate, phosphorodiselenoate, phosphorodithioate, phosphorothioate, and the like, including associated counterions, e.g., H+, NH4+, Na+, and the like, if such counterions are present. Further, oligonucleotides are typically single-stranded. Oligonucleotides are optionally prepared by any suitable method, including, but not limited
to, isolation of an existing or natural sequence, DNA replication or amplification, reverse transcription, cloning and restriction digestion of appropriate sequences, or direct chemical synthesis by a method such as the phosphodiester method of Narang et al. (1979) *Meth Enzymol.* 68: 90-99; the phosphodiester method of Brown et al. (1979) *Meth Enzymol.* 68: 109-151; the diethylphosphoramidite method of Beaucage et al. (1981) *Tetrahedron Lett.* 22: 1859-1862; the tri-ester method of Matteucci et al. (1981) *Am Chem Soc.* 103: 3185-3191; automated synthesis methods; or the solid support method of U.S. Pat. No. 4,458,066, entitled “PROCESS FOR PREPARING POLYNUCLEOTIDES,” issued Jul. 3, 1984 to Caruthers et al., or other methods known to those skilled in the art. All of these references are incorporated by reference.

[0107] As used herein a “sample” refers to anything capable of being analyzed by the methods provided herein. In some embodiments, the sample comprises or is suspected to comprise one or more nucleic acids capable of analysis by the methods. Preferably, the samples comprise nucleic acids (e.g., DNA, RNA, cDNAs, etc.) from one or more biogens. Samples can include, for example, blood, saliva, urine, feces, anorectal swabs, vaginal swabs, cervical swabs, and the like. In some embodiments, the samples are “mixture” samples, which comprise nucleic acids from more than one subject or individual. In some embodiments, the methods provided herein comprise purifying the sample or purifying the nucleic acid(s) from the sample. In some embodiments, the sample is purified nucleic acid.

[0108] A “sequence” of a biopolymer refers to the order and identity of monomer units (e.g., nucleotides, etc.) in the biopolymer. The sequence (e.g., base sequence) of a nucleic acid is typically read in the 5' to 3' direction.

[0109] As is used herein, the term “single primer pair identification” means that one or more biogens can be identified using a single primer pair. A base composition signature for an amplicon may singly identify one or more biogens.

[0110] A “solid support” refers to a solid material which can be derivatized with, or otherwise attached to, a chemical moiety, such as a primer nucleic acid, a template nucleic acid, or the like. Exemplary solid supports include a zero-mode waveguide array, a plate, a bead, a microbead, a fiber, a whisker, a comb, a hybridization chip, a membrane, a single crystal, a ceramic layer, a self-assembling monolayer, and the like.

[0111] As used herein, a “sub-species characteristic” is a genetic characteristic that provides the means to distinguish two members of the same biogen species. For example, one viral strain may be distinguished from another viral strain of the same species by possessing a genetic change (e.g., for example, a nucleotide deletion, addition or substitution) in one of the viral genes, such as the RNA-dependent RNA polymerase.

[0112] A “subsequence” or “fragment” refers to any portion of an entire nucleic acid sequence.

[0113] As used herein, in some embodiments the term “substantial complementarity” means that a primer member of a primer pair comprises between about 70%–100%, or between about 80-100%, or between about 90-100%, or between about 95-100%, or between about 99-100% complementarity with the conserved binding sequence of a nucleic acid from a given biogen. These ranges of complementarity and identity are inclusive of all whole or partial numbers embraced within the recited range numbers. For example, and not limitation, 75.667%, 82%, 91.2435% and 97% complementarity or sequence identity are all numbers that fall within the above recited range of 70% to 100%, therefore forming a part of this description.

[0114] A “system” in the context of analytical instrumentation refers to a group of objects and/or devices that form a network for performing a desired objective.

[0115] A “template nucleic acid” refers to a nucleic acid to which a primer nucleic acid can hybridize and be extended. Accordingly, template nucleic acids include subsequences that are at least partially complementary to the primer nucleic acids. Template nucleic acids can be derived from essentially any source. To illustrate, template nucleic acids are optionally derived or isolated from, e.g., cultured microorganisms, uncultured microorganisms, complex biological mixtures, tissues, sera, pooled sera or tissues, multispecies consortia, ancient, fossilized or other nonliving biological remains, environmental isolates, soils, groundwater, waste facilities, deep-sea environments, or the like. Further, template nucleic acids optionally include or are derived from, e.g., individual cDNA molecules, cloned sets of cDNAs, cDNA libraries, extracted RNAs, natural RNAs, in vitro transcribed RNAs, characterized or uncharacterized genomic DNAs, cloned genomic DNAs, genomic DNA libraries, enzymatically fragmented DNAs or RNAs, chemically fragmented DNAs or RNAs, physically fragmented DNAs or RNAs, or the like. Template nucleic acids can also be chemically synthesized using techniques known in the art. In addition, template nucleic acids optionally correspond to at least a portion of a gene or are complementary thereto.

[0116] As used herein, “triangulation identification” means the use of more than one primer pair to generate a corresponding amplicon for identification of a biogen. The more than one primer pair can be used in individual wells or vessels or in a multiplex PCR assay. Alternatively, PCR reactions may be carried out in single wells or vessels comprising a different primer pair in each well or vessel. Following amplification the amplicons are pooled into a single well or container which is then subjected to base composition analysis (e.g., which does not involve molecular mass analysis). The combination of pooled amplicons can be chosen such that the expected ranges of base compositions of individual amplicons are not overlapping and thus will not complicate identification of signals. Triangulation is a process of elimination, wherein a first primer pair identifies that an unknown biogen may be one of a group of biogens. Subsequent primer pairs are used in triangulation identification to further refine the identity of the biogen amongst the subset of possibilities generated with the earlier primer pair. Triangulation identification is complete when the identity of the biogen is determined. The triangulation identification process may also be used to reduce false negative and false positive signals, and enable reconstruction of the origin of hybrid or otherwise biogen engineered biogens. For example, identification of the three part toxin genes typical of *B. anthracis* (Bouwen et al., *J Appl Microbiol.*, 1999, 87, 270-278) in the absence of the expected compositions from the *B. anthracis* genome would suggest a genetic engineering event.

[0117] As used herein, the term “unknown biogen” can mean, for example: (i) a biogen whose existence is not known (for example, the SARS coronavirus was unknown prior to April 2003) and/or (ii) a biogen whose existence is known (such as the well known bacterial species *Staphylococcus aureus* for example) but which is not known to be in a
sample to be analyzed. For example, if the method for identification of coronaviruses disclosed in commonly owned U.S. patent Ser. No. 10/829,826 (incorporated herein by reference in its entirety) was to be employed prior to April 2003 to identify the SARS coronavirus in a clinical sample, both meanings of "unknown" biogent are applicable since the SARS coronavirus was unknown to science prior to April, 2003 and since it was not known what biogent (in this case a coronavirus) was present in the sample. On the other hand, if the method of U.S. patent Ser. No. 10/829,826 was to be employed subsequent to April 2003 to identify the SARS coronavirus in a clinical sample, the second meaning (ii) of "unknown" biogent would apply because the SARS coronavirus became known to science subsequent to April 2003 because it was not known what biogent was present in the sample.

[0118] As used herein, the term "variable region" is used to describe a region that falls between any one primer pair described herein. The region possesses distinct base compositions between at least two biogents, such that at least one biogent can be identified at, for example, the family, genus, species or sub-species level. The degree of variability between the at least two biogents need only be sufficient to allow for identification using mass spectrometry analysis, as described herein.

[0119] As used herein, a "wobble base" is a variation in a codon found at the third nucleotide position of a DNA triplet. Variations in conserved regions of sequence are often found at the third nucleotide position due to redundancy in the amino acid code.

[0120] In certain embodiments, provided herein are methods, compositions, kits, and related systems for the detection and identification of biogents (e.g., species of HPV) using biogent identifying amplicons. In some embodiments, primers are selected to hybridize to conserved sequence regions of nucleic acids derived from a biogent and which flank variable sequence regions to yield a biogent identifying amplicon which can be amplified and which is amenable to base composition analysis. In some embodiments, the corresponding base composition of one or more different amplicons is queried against a database of base compositions indexed to biogents and to the primer pair used to generate the amplicon. A match of the measured base composition to a database entry base composition associates the sample biogent to an indexed biogent in the database. Thus, the identity of the unknown biogent is determined. No prior knowledge of the unknown biogent is necessary to make an identification. In some instances, the measured base composition associates with more than one database entry base composition. Thus, a second/subsequent primer pair is generally used to generate an amplicon, and its measured base composition is similarly compared to the database to determine its identity in triangulation identification. Furthermore, the methods and other aspects of the invention can be applied to rapid parallel multiplex analyses, the results of which can be employed in a triangulation identification strategy. Thus, in some embodiments, the present invention provides rapid throughput and does not require nucleic acid sequencing or knowledge of the linear sequences of nucleobases of the amplified target sequence for biogent detection and identification.


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Summary (ed. Knobler S E, Mahmoud A, Lemon S.) The 

In some embodiments, amplicons corresponding to 
bioagent identifying amplicons are obtained using the poly-
merase chain reaction (PCR). Other amplification methods 
may be used such as ligase chain reaction (LCR), low-string-
genency single primer PCR, and multiple strand displacement 
amplification (MDA). (Michael, S. F., Biotechniques (1994), 

One embodiment of a process flow diagram used for 
primer selection and validation process is depicted in FIGS. 4 
and 5. For each group of organisms, candidate target 
sequences are identified (200) from which nucleotide 
sequence alignments are created (210) and analyzed (220). 
Primers are then configured by selecting priming regions 
(230) to facilitate the selection of candidate primer pairs 
(240). The primer pair sequence is typically a “best fit” 
amongst the aligned sequences, such that the primer pair 
sequence may or may not be fully complementary to the 
hybridization region on any one of the bioagents in the align-
ment. Thus, best fit primer pair sequences are those with 
sufficient complementarity with two or more bioagents to 
hybridize with the two or more bioagents and generate an 
amplicon. The primer pairs are then subjected to in silico 
analysis by electronic PCR (ePCR) (300) wherein bioagent 
identifying amplicons are obtained from sequence databases 
such as GenBank or other sequence collections (310) and 
tested for specificity in silico (320). Bioagent identifying 
amplicons obtained from ePCR of GenBank sequences (310) 
may also be analyzed by a probability model which predicts 
the capability of a given amplicon to identify unknown bio-
agents. Preferably, the base compositions of amplicons with 
favorable probability scores are then stored in a base composi-
tion database (325). Alternatively, base compositions of the 
bioagent identifying amplicons obtained from the primers 
and GenBank sequences are directly entered into the base 
composition database (330). Candidate primer pairs (240) are 
validated by in vitro amplification by a method such as PCR 
analysis (400) of nucleic acid from a collection of organisms 
(410). Amplicons thus obtained are analyzed to confirm the 
sensitivity, specificity and reproducibility of the primers used 
to obtain the amplicons (420).

Synthesis of primers is well known and routine in 
the art. The primers may be conveniently and routinely made 
through the well-known technique of solid phase synthesis. 
Equipment for such synthesis is sold by several vendors 
including, for example, Applied Biosystems (Foster City, 
Calif.). Any other means for such synthesis known in the art 
may additionally or alternatively be employed.

In some embodiments, primers are employed as 
compositions for use in methods for identification of bio-
agents as follows: a primer pair composition is contacted with 
nucleic acid of an unknown isolate suspected of comprising a 
target bioagent. The nucleic acid is then amplified by a 
nucleic acid amplification technique, such as PCR, for 
example, to obtain an amplicon that represents a bioagent 
identifying amplicon. The base composition of the double-
stranded amplicon, or single strand corresponding to only 
the forward or reverse primer, is determined by techniques 
such as sequencing, HPLC, and paper chromatography (see, 
e.g., Voelkerding et al., Clinical Chem., “Next-generation 
sequencing: from basic research to diagnostics.””; 55, 641-658, 
2009; MacLean et al., Nature Rev. Microbiol., 7: 287-296, 
and Manderville and Kropinski, “Approaches to the Composi-
tional Analysis of DNA.” Methods Mol Biol 2000; 502:11-
7, all of which are herein incorporated by reference). A mea-
sured base composition calculated therefrom is then 
compared with a database of base compositions indexed to 
primer pairs and to known bioagents. A match between the 
base composition of the amplicon and the database base com-
position for that indexed primer pair correlates the measured 
base composition with an indexed bioagent, thus identifying 
the unknown bioagent (e.g. the species of virus). In some 
embodiments, the method is repeated using a different primer 
pair to resolve possible ambiguities in the identification pro-
cess or to improve the confidence level for the identification 
assignment (trangulation identification). In some embodi-
ments, for example, where the unknown is a novel, previously 
uncharacterized organism, the base composition from an 
amplicon generated from the unknown is matched with one or 
more best match base compositions from a database to predict 
a family, genus, species, sub-type, etc. of the unknown. 
Such information may assist further characterization of the 
unknown or provide a physician treating a patient infected by 
the unknown with a therapeutic agent best calculated to treat 
the patient.

In certain embodiments, the bioagent is detected 
with the systems and methods of the present invention in 
combination with other bioagents, including viruses, bacte-
ria, fungi, or other bioagents. In particular embodiments, a 
panel is employed that includes a first bioagent and other 
related or un-related bioagents. Such panels may be specific 
for a particular type of bioagent, or specific for a specific type 
of test (e.g., for testing the safety of blood, one may include 
commonly present viral pathogens such as HCV, HIV, and 
bacteria that can be contracted via a blood transfusion).

In some embodiments, a bioagent identifying 
amplicon may be produced using only a single primer (either
the forward or reverse primer of any given primer pair), provided an appropriate amplification method is chosen, such as, for example, low stringency single primer PCR (LSSPP-PCR).

[0129] In some embodiments, the oligonucleotide primers are broad range survey primers which hybridize to conserved regions of nucleic acid. The broad range primer may identify the unknown bioagent depending on which bioagent is in the sample. In other cases, the base composition of an amplicon does not provide sufficient resolution to identify the unknown bioagent as any one bioagent or at below the species level. These cases generally benefit from further analysis of one or more amplicons generated from at least one additional broad range survey primer pair, or from at least one additional division-wide primer pair, or from at least one additional drill-down primer pair. Identification of sub-species characteristics may be required, for example, to determine a clinical treatment of patient, or in rapidly responding to an outbreak of a new species, sub-type, etc. of pathogen to prevent an epidemic or pandemic.

[0130] One with ordinary skill in the art of design of amplification primers will recognize that a given primer need not hybridize with 100% complementarity in order to effectively prime the synthesis of a complementary nucleic acid strand in an amplification reaction. Primer pair sequences may be a “best fit” amongst the aligned bioagent sequences, thus they need not be fully complementary to the hybridization region of any one of the bioagents in the alignment. Moreover, a primer may hybridize over one or more segments such that intervening or adjacent segments are not involved in the hybridization event (e.g., for example, a loop structure or a hairpin structure). Thus, in some embodiments, an extent of variation of 70% to 100%, or any range falling within, of the sequence identity is possible relative to the specific primer sequences disclosed herein. To illustrate, determination of sequence identity is described in the following example: a primer 20 nucleobases in length which is identical to another 20 nucleobase primer having two non-identical residues has 18 of 20 identical residues (18/20 = 0.9 or 90% sequence identity). In another example, a primer 15 nucleobases in length having all residues identical to a 15 nucleobase segment of another 20 nucleobases in length would have 15/20 = 0.75 or 75% sequence identity with the 20 nucleobase primer. Percent identity need not be a whole number, for example when a 28 consecutive nucleobase primer is completely identical to a 31 consecutive nucleobase primer (28/31 = 0.9032 or 90.3% identical).

[0131] Percent homology, sequence identity or complementarity, can be determined by, for example, the Gap program (Wisconsin Sequence Analysis Package, Version 8 for Unix, Genetics Computer Group, University Research Park, Madison Wis.), using default settings, which uses the algorithm of Smith and Waterman (Adv. Appl. Math., 1981, 2, 482-489). In some embodiments, complementarity of primers with respect to the conserved priming regions of viral nucleic acid, is between about 70% and about 80%. In other embodiments, homology, sequence identity or complementarity, is between about 80% and about 90%. In yet other embodiments, homology, sequence identity or complementarity, is at least 90%, at least 92%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99% or is 100%.

[0132] In some embodiments, the primers described herein comprise at least 70%, at least 75%, at least 80%, at least 85%, at least 90%, at least 92%, at least 94%, at least 95%, at least 96%, at least 98%, or at least 99%, or 100% (or any range falling within) sequence identity with the primer sequences specifically disclosed herein.

[0133] In some embodiments, the oligonucleotide primers are 13 to 35 nucleobases in length (13 to 35 linked nucleotide residues). These embodiments comprise oligonucleotide primers 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 34 or 35 nucleobases in length, or any range there within.

[0134] In some embodiments, any given primer comprises a modification comprising the addition of a non-templated T residue to the 5’ end of the primer (i.e., the added T residue does not necessarily hybridize to the nucleic acid being amplified). The addition of a non-templated T residue has an effect of minimizing the addition of non-templated A residues as a result of the non-specific enzyme activity of, e.g., Taq DNA polymerase (Magneon et al., Biotechniques, 1996, 21, 700-709), an occurrence which may lead to ambiguous results arising from molecular mass analysis.

[0135] Primers may contain one or more universal bases. Because any variation (due to codon wobble in the third position) in the conserved regions among species is likely to occur in the third position of a DNA (or RNA) triplet, oligonucleotide primers can be designed such that the nucleotide corresponding to this position is a base which can bind to more than one nucleotide, referred to herein as a “universal nucleobase.” For example, under this “wobble” base pairing, inosine (I) binds to U, C or A; guanine (G) binds to U or C, and uridine (U) binds to U or C. Other examples of universal nucleobases include nitroindoles such as 5-nitroindole or 3-nitropyrrrole (Loakes et al., Nucleosides and Nucleotides, 1995, 14, 1001-1003), the degenerate nucleotides dP or dK, an acyclic nucleoside analog containing 5-nitroindazole (Van Aerschot et al., Nucleosides and Nucleotides, 1995, 14, 1053-1056) or the purine analog 1-(2-deoxy-beta-D-ribofuranoxy)-imidazole-4-carboxamide (Sala et al., Nucl. Acids Res., 1996, 24, 3302-3306).

[0136] In some embodiments, to compensate for weaker binding by the wobble base, oligonucleotide primers are configured such that the first and second positions of each triplet are occupied by nucleotide analogs which bind with greater affinity than the unmodified nucleotide. Examples of these analogs include, but are not limited to, 2,6-diaminopurine which binds to thymine, 5-propynyluracil which binds to adenine and 5-propynylcytosine and phenoxazines, including G-clamp, which binds to G. Propynylated pyrimidines are described in U.S. Pat. Nos. 5,645,985, 5,830,653 and 5,484,908, each of which is commonly owned and incorporated herein by reference in its entirety. Propynylated primers are described in U.S Pre-Grant Publication No. 2003-0170682; also commonly owned and incorporated herein by reference in its entirety. Phenoxazines are described in U.S. Pat. Nos. 5,502,177, 5,763,588, and 6,005,096, each of which is incorporated herein by reference in its entirety. G-clamps are described in U.S. Pat. Nos. 6,007,992 and 6,028,183, each of which is incorporated herein by reference in its entirety.

[0137] In some embodiments, non-template primer tags are used to increase the melting temperature (Tm) of a primer-template duplex in order to improve amplification efficiency. A non-template tag is at least three consecutive A or T nucleotide residues on a primer which are not complementary to the template. In any given non-template tag, A can be replaced by C or G and T can also be replaced by C or G. Although
Watson-Crick hybridization is not expected to occur for a non-template tag relative to the template, the extra hydrogen bond in a G-C pair relative to an A-T pair confers increased stability of the primer-template duplex and improves amplification efficiency for subsequent cycles of amplification when the primers hybridize to strands synthesized in previous cycles.

In other embodiments, propynylated tags may be used in a manner similar to that of the non-template tag, wherein two or more 5-propynylcytidine or 5-propynyluridine residues replace template matching residues on a primer. In other embodiments, a primer contains a modified internucleoside linkage such as a phosphorothioate linkage, for example.

In some embodiments, the primers contain mass-modifying tags. Reducing the total number of possible base compositions of a nucleic acid of specific molecular weight provides a means of avoiding a possible source of ambiguity in the determination of base composition of amplicons.

In some embodiments, assignment of previously unobserved base compositions (also known as “true unknown base compositions”) to a given phylogeny can be accomplished via the use of pattern classifier model algorithms. Base compositions, like sequences, may vary slightly from strain to strain within species, for example. In some embodiments, the pattern classifier model is the mutational probability model. A polypeptide model is a polypeptide model that incorporates both the restrictions among strains and position dependence of a given nucleobase within a triplet. In certain embodiments, a polypeptide pattern classifier is used to classify a test or unknown organism according to its amplicon base composition.

In some embodiments, it is possible to manage this diversity by building “base composition probability clouds” around the composition constraints for each species. A “pseudo four-dimensional plot” may be used to visualize the concept of base composition probability clouds. Optimal primer design typically involves an optimal choice of biobagent identifying amplicons and maximizes the separation between the base composition signatures of individual bioagents. Areas where clouds overlap generally indicate regions that may result in a misclassification, a problem which is overcome by a triangulation identification process using biobagent identifying amplicons not affected by overlap of base composition probability clouds.

In some embodiments, base composition probability clouds provide the means for screening potential primer pairs in order to avoid potential misclassifications of base compositions. In other embodiments, base composition probability clouds provide the means for predicting the identity of an unknown biobagent whose assigned base composition has not been previously observed and/or indexed in a biobagent identifying amplicon base composition database due to evolutionary transitions in its nucleic acid sequence.

Provided herein is biobagent classifying information at a level sufficient to identify a given biobagent. Furthermore, the process of determining a previously unknown base composition for a given biobagent (for example, in a case where sequence information is unavailable) has utility by providing additional biobagent indexing information with which to populate base composition databases. The process of future biobagent identification is thus improved as additional base composition signature indexes become available in base composition databases.

In certain embodiments, a sample comprising an unknown biobagent is contacted with a primer pair which amplifies the nucleic acid from the biobagent, and a known quantity of a polynucleotide that comprises a calibration sequence. The amplification reaction then produces two amplicons: a biobagent identifying amplicon and a calibration amplicon. The biobagent identifying amplicon and the calibration amplicon are distinguishable by base composition while being amplified at essentially the same rate. Reflecting differential base compositions can be accomplished by choosing as a calibration sequence, a representative biobagent identifying amplicon (from a specific species of biobagent) and performing, for example, a 2-8 nucleobase deletion or insertion within the variable region between the two priming sites, a calibration sequence with a different base composition due to base substitutions. The amplified sample containing the biobagent identifying amplicon and the calibration amplicon is then subjected to base composition analysis (e.g., sequencing, HPLC, paper chromatography, etc.) without determining the molecular mass of the sequences. The resulting base composition analysis of the nucleic acid of the biobagent and of the calibration sequence provides base composition data and abundance data for the nucleic acid of the biobagent and of the calibration sequence. The base composition data obtained for the nucleic acid of the biobagent enables identification of the unknown biobagent by base composition analysis. The abundance data enables calculation of the quantity of the biobagent, based on the knowledge of the quantity of calibration polynucleotide contacted with the sample.

In some embodiments, construction of a standard curve in which the amount of calibration or calibrant polynucleotide spiked into the sample is varied provides additional resolution and improved confidence for the determination of the quantity of biobagent in the sample. Alternatively, the calibration polynucleotide can be amplified in its own reaction vessel or vessels under the same conditions as the biobagent. A standard curve may be prepared there from, and the relative abundance of the biobagent determined by methods such as linear regression. In some embodiments, multiplex amplification is performed where multiple biobagent identifying amplicons are amplified with multiple primer pairs which also amplify the corresponding standard calibration sequences. In this or other embodiments, the standard calibration sequences are optionally included within a single construct (preferably a vector) which functions as the calibration polynucleotide.

In some embodiments, the calibrant polynucleotide is used as an internal positive control to confirm that amplification conditions and subsequent analysis steps are successful in producing a measureable amplicon. Even in the absence of copies of the genome of a biobagent, the calibration polynucleotide gives rise to a calibration amplicon. Failure to produce a measurable calibration amplicon indicates a failure of amplification or subsequent analysis step such as amplicon purification or base composition determination. Reaching a conclusion that such failures have occurred is, in itself, a useful event. In some embodiments, the calibration sequence is comprised of DNA. In some embodiments, the calibration sequence is comprised of RNA.

In some embodiments, a calibration sequence is inserted into a vector which then functions as the calibration...
polynucleotide. In some embodiments, more than one calibration sequence is inserted into the vector that functions as the calibration polynucleotide. Such a calibration polynucleotide is herein termed a “combination calibration polynucleotide.” It should be recognized that the calibration method should not be limited to the embodiments described herein. The calibration method can be applied for determination of the quantity of any bioagent identifying amplicon when an appropriate standard calibrant polynucleotide sequence is designed and used.

[0148] In certain embodiments, primer pairs are configured to produce bioagent identifying amplicons within more conserved regions of a bioagent, while others produce bioagent identifying amplicons within regions that may evolve more quickly. Primer pairs that characterize amplicons in a conserved region with low probability that the region will evolve past the point of primer recognition are useful, e.g., as a broad range survey-type primer. Primer pairs that characterize amplicon regions corresponding to an evolving genomic region are useful, e.g., for distinguishing emerging bioagent strain variants.

[0149] The primer pairs described herein provide reagents, e.g., for identifying diseases caused by emerging types of bioagents. Base composition analysis eliminates the need for prior knowledge of bioagent sequence to generate hybridization probes. Thus, in another embodiment, there is provided a method for determining the etiology of a particular strain when the process of identification is carried out in a clinical setting, and even when a new strain is involved. This is possible because the methods may not be confounded by naturally occurring evolutionary variations.

[0150] Another embodiment provides a means of tracking the spread of any species or strain of particular bioagents when a plurality of samples obtained from different geographical locations are analyzed by methods described above in an epidemiological setting. For example, a plurality of samples from a plurality of different locations may be analyzed with primers which produce bioagent identifying amplicons, a subset of which identifies a specific strain. The corresponding locations of the members of the strain-containing subset indicate the spread of the specific strain to the corresponding locations.

[0151] Also provided are kits for carrying out the methods described herein. In some embodiments, the kit may comprise a sufficient quantity of one or more primer pairs to perform an amplification reaction on a target polynucleotide from a bioagent to form a bioagent identifying amplicon. In some embodiments, the kit may comprise from one to twenty primer pairs, from one to ten primer pairs, from one to eight pairs, from one to five primer pairs, from one to three primer pairs, or from one to two primer pairs.

[0152] In some embodiments, the kit may also comprise a sufficient quantity of reverse transcriptase, a DNA polymerase, suitable nucleoside triphosphates (including any of those described above), a DNA ligase, and/or reaction buffer, or any combination thereof, for the amplification processes described above. The kit may also comprise reagents necessary for performing sequencing methods, or HPLC or paper chromatography (see, e.g., Voelkerding et al., Clinical Chem., “Next-generation sequencing: from basic research to diagnostics,” 55: 641-658, 2009; MacLean et al., Nature Rev. Microbiol., 7: 287-296, and Manderville and Kropinski, “Approaches to the Compositional Analysis of DNA,” Methods Mol. Biol. 2009; 502:11-7, all of which are herein incorporated by reference).

[0153] A kit may further include instructions pertinent for the particular embodiment of the kit, such instructions describing the primer pairs and amplification conditions for operation of the method. In some embodiments, the kit further comprises instructions for analysis, interpretation and dissemination of data acquired by the kit. In other embodiments, instructions for the operation, analysis, interpretation and dissemination of the data of the kit are provided on computer readable media. A kit may also comprise amplification reaction containers such as microcentrifuge tubes, microtiter plates, and the like. A kit may also comprise reagents or other materials for isolating bioagent nucleic acid or bioagent identifying amplicons from amplification reactions, including, for example, detergents, solvents, or ion exchange resins which may be linked to magnetic beads. A kit may also comprise a table of measured or calculated base compositions of bioagents using the primer pairs of the kit.

[0154] The invention also provides systems that can be used to perform various assays relating to bioagent detection or identification. In certain embodiments, systems include sequencing devices (or HPLC equipment or paper chromatography equipment) configured to detect base compositions of amplicons produced using purified oligonucleotide primer pairs described herein. Other devices/equipment that are optionally adapted for use in the systems of the invention are described further below. In some embodiments, systems also include controllers operably connected to sequencing devices and/or other system components. In some of these embodiments, controllers are configured to correlate the base compositions of the amplicons with bioagents to effect detection or identification. As described herein, the base compositions generally correspond to the bioagent species identities. In certain embodiments, controllers include, or are operably connected to, databases of known base compositions of amplicons of known species of bioagents produced with the primer pairs described herein. Controllers are described further below.

[0155] In certain embodiments, the oligonucleotides are arrayed on solid supports, whereas in others, they are provided in one or more containers, e.g., for assays performed in solution. In certain embodiments, the systems also include at least one detector or detection component (e.g., a spectrometer) that is configured to detect detectable signals produced in the container or on the support. In addition, the systems also optionally include at least one thermal modulator (e.g., a thermal cycling device) operably connected to the containers or solid supports to modulate temperature in the containers or on the solid supports, and/or at least one fluid transfer component (e.g., an automated pipettor) that transfers fluid to and/or from the containers or solid supports, e.g., for performing one or more assays (e.g., nucleic acid amplification, real-time amplicon detection, etc.) in the containers or on the solid supports.

[0156] Examples of suitable thermocycling devices that are optionally utilized are available from many different commercial suppliers, including Mastercycler® devices (Eppendorf North America, Westbury, N.Y., U.S.A.), the COBAS® AMPLICOR Analyzer (Roche Molecular Systems, Inc., Pleasanton, Calif., U.S.A.), MyCycler and iCycler Thermal Cyclers (Bio-Rad Laboratories, Inc., Hercules, Calif., U.S.A.), and the SmartCycler System (Cepheid, Sunnyvale, Calif.)
U.S.A.), among many others. In other exemplary embodiments, sample preparation components, nucleic acid amplification components, and related fluid handling or material transfer components are integrated with the systems described herein, e.g., to fully automate a given nucleic acid amplification and analysis process. Instruments that can be adapted for this purpose include, for example, the m2000™ automated instrument system (Abbott Laboratories, Abbott Park, Ill., U.S.A.), the GeneXpert System (Cepheid, Sunnyvale, Calif., U.S.A.), and the COBAS® Amplicor® System (Roche Molecular Systems, Inc., Pleasanton, Calif., U.S.A.), and the like.

[0157] Detectors are typically structured to detect detectable signals produced, e.g., in or proximal to another component of the given assay system (e.g., in a container and/or on a solid support). Suitable signal detectors that are optionally utilized, or adapted for use, herein detect, e.g., fluorescence, phosphorescence, radioactivity, absorbance, refractive index, luminescence, or mass. Detectors optionally monitor one or a plurality of signals from upstream and/or downstream of the performance of, e.g., a given assay step. For example, detectors optionally monitor a plurality of optical signals, which correspond in position to "real-time" results. Example detectors or sensors include photomultiplier tubes, CCD arrays, optical sensors, temperature sensors, pressure sensors, pH sensors, conductivity sensors, or scanning detectors. Detectors are also described in, e.g., Skoog et al., Principles of Instrumental Analysis, 6th Ed., Brooks Cole (2006), Currell, Analytical Instrumentation: Performance Characteristics and Quality, John Wiley & Sons, Inc. (2000), Sharma et al., Introduction to Fluorescence Spectroscopy, John Wiley & Sons, Inc. (1999), Valure, Molecular Fluorescence: Principles and Applications, John Wiley & Sons, Inc. (2002), and Gore, Spectrophotometry and Spectrofluorimetry: A Practical Approach, 2nd Ed., Oxford University Press (2000), which are each incorporated by reference.

[0159] As mentioned above, the systems of the invention also typically include controllers that are operably connected to one or more components (e.g., detectors, databases, thermal modulators, fluid transfer components, robotic material handling devices, and the like) of the given system to control operation of the components. More specifically, controllers are generally included either as separate or integral system components that are utilized, e.g., to receive data from detectors to effect and/or regulate temperature in the containers, or to effect and/or regulate fluid flow to or from selected containers. Controllers and/or other system components are optionally coupled to an appropriately programmed processor, computer, digital device, information appliance, or other logic device (e.g., including an analog to digital or digital to analog converter as needed), which functions to instruct the operation of these instruments in accordance with preprogrammed or user input instructions, receive data and information from these instruments, and interpret, manipulate and report this information to the user. Suitable controllers are generally known in the art and are available from various commercial sources.

[0160] Any controller or computer optionally includes a monitor, which is often a cathode ray tube ("CRT") display, a flat panel display (e.g., active matrix liquid crystal display or liquid crystal display), or others. Computer circuitry is often placed in a box, which includes numerous integrated circuit chips, such as a microprocessor, memory, interface circuits, and others. The box also optionally includes a hard disk drive, a floppy disk drive, a high capacity removable drive such as a writeable CD-ROM, and other common peripheral elements. Inputting devices such as a keyboard or mouse optionally provide for input from a user. These components are illustrated further below.

[0161] The computer typically includes appropriate software for receiving user instructions, either in the form of user input into a set of parameter fields, e.g., in a graphic user interface (GUI), or in the form of preprogrammed instructions, e.g., preprogrammed for a variety of different specific operations. The software then converts these instructions to appropriate language for instructing the operation of one or more controllers to carry out the desired operation. The computer then receives the data from, e.g., sensors/detectors included within the system, and interprets the data, either provides it in a user understood format, or uses that data to initiate further controller instructions, in accordance with the programming.

[0162] FIG. 6 is a schematic showing a representative system that includes a logic device in which various aspects of the present invention may be embodied. As will be understood by practitioners in the art from the teachings provided herein, aspects of the invention are optionally implemented in hardware and/or software. In some embodiments, different aspects of the invention are implemented in either client-side logic or server-side logic. As will be understood in the art, the invention or components thereof may be embodied in a media program component (e.g., a fixed media component) containing logic instructions and/or data that, when loaded into an appropriately configured computing device, cause that device to perform as desired. As will also be understood in the art, a fixed media containing logic instructions may be delivered to a viewer on a fixed media for physically loading into a viewer's computer or a fixed media containing logic instructions may reside on a remote server that a viewer accesses through a communication medium in order to download a program component.

[0163] More specifically, FIG. 6 schematically illustrates computer 1000 to which sequencing device or system 1002 (e.g., SMRT detection array from Pacific Biosciences), fluid transfer component 1004 (e.g., a sample injection needle or the like), and database 1008 are operably connected. Optionally, one or more of these components are operably connected to computer 1000 via a server (not shown in FIG. 6). During operation, fluid transfer component 1004 typically transfers reaction mixtures or components thereof (e.g., aliquots comprising amplicons) from multi-well container 1006 to sequencing device. Sequencing device 1002 then detects the nucleic acid sequence of the amplicons. Computer 1000 then typically receives this sequence data, calculates base compositions from this data, and compares it with entries in database 1008 to identify species or strains of biocagents in a given sample. It will be apparent to one of skill in the art that one or more components of the system schematically depicted in FIG. 6 are optionally fabricated integral with one another (e.g., in the same housing).

[0164] While the present invention has been described with specificity in accordance with certain of its embodiments, the following examples serve only to illustrate the invention and are not intended to limit the same. In order that the invention disclosed herein may be more efficiently understood, examples are provided below. It should be understood that...
these examples are for illustrative purposes only and are not to be construed as limiting the invention in any manner.

Example 1

Non-Mass Determined Base Composition Analysis for Target Sequence Detection

This example illustrates that pathogens can be identified and distinguished from one another using complete or partial base composition or count information without the need for mass determination of bases. For example, Table I shows base composition (single strand) results for 16S-1100-1188 (16S rRNA) primer amplification reactions for different species of bacteria. Such base composition data can be generated by any type of sequencing methodology or system (see, e.g., Voelkerding et al., Clinical Chem., “Next-generation sequencing: from basic research to diagnostics,” 55: 641-658, 2009; and Macl.ean et al., Nature Rev. Microbiol., 7: 287-296). Species which are repeated in the table (e.g., Clostridium botulinum) are different strains which have different base compositions in the 16S-1100-1188 region. As shown in Table I, for example, a complete base composition using this primer amplification reaction will uniquely identify and distinguish Mycobacterium avium (i.e., A15G23C18T16) from the other organisms listed in Table I, as well as various partial base compositions using the same primer amplification reaction (e.g., A15G23C18T16, A15G23C18T16, C18T16, etc.).

### TABLE I-continued

<table>
<thead>
<tr>
<th>Organism name</th>
<th>Base comp.</th>
<th>Organism name</th>
<th>Base comp.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Acinetobacter</td>
<td>A23G26C21T19</td>
<td>Yersinia pseudotuberculosis</td>
<td>A26G3C16T13</td>
</tr>
<tr>
<td>calcoacetic</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Leptospira</td>
<td>A23G26C21T15</td>
<td>Clostridium botulinum</td>
<td>A26G2C16T24</td>
</tr>
<tr>
<td>borgonieri</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Leptospira</td>
<td>A23G26C21T15</td>
<td>Clostridium septic</td>
<td>A26G2C16T26</td>
</tr>
<tr>
<td>interrogans</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Clostridium</td>
<td>A23G27C19T19</td>
<td>Francisella tularensis</td>
<td>A26G2C16T19</td>
</tr>
<tr>
<td>perfringens</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Bacillus</td>
<td>A23G27C19T15</td>
<td>Acinetobacter calcoacetic</td>
<td>A26G2C16T19</td>
</tr>
<tr>
<td>anthracis</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Bacillus</td>
<td>A23G27C19T15</td>
<td>Bacteroides fragilis</td>
<td>A26G2C16T22</td>
</tr>
<tr>
<td>cereus</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Bacillus</td>
<td>A23G27C19T15</td>
<td>Chlamydia pneumonia</td>
<td>A26G2C16T16</td>
</tr>
<tr>
<td>psittaci</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>shigella</td>
<td>A23G27C19T15</td>
<td>Borelia burgdorferi</td>
<td>A26G2C16T19</td>
</tr>
<tr>
<td>hydrophila</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Escherichia</td>
<td>A23G27C19T15</td>
<td>Streptococcus miller</td>
<td>A26G2C16T19</td>
</tr>
<tr>
<td>coli</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Pseudomonas</td>
<td>A23G27C19T15</td>
<td>Mycoplasma arfasei</td>
<td>A26G2C16T19</td>
</tr>
<tr>
<td>putida</td>
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<td></td>
</tr>
<tr>
<td>Escherichia</td>
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<td>Mycoplasma arfasei</td>
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</tr>
<tr>
<td>coli</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Shigella</td>
<td>A23G27C19T15</td>
<td>Mycoplasma mycoplasma</td>
<td>A26G2C16T20</td>
</tr>
<tr>
<td>dysenteriae</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Some of these organisms can be distinguished using multiple primers. For example, Chlamydia trachomatis can be distinguished from Chlamydia pneumoniae AR39 and other organisms listed in Table II using the primer 16S_971-1062 and the primer 16S_1228-1310 in addition to 16S 1100-1188 based on a complete or partial base composition for the amplifiers.

### TABLE II

<table>
<thead>
<tr>
<th>Organism name</th>
<th>Base comp.</th>
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<th>Base comp.</th>
</tr>
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<tr>
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<tr>
<td>Leptospira</td>
<td>A23G26C21T15</td>
<td>Clostridium botulinum</td>
<td>A26G2C16T24</td>
</tr>
<tr>
<td>borgonieri</td>
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</tr>
<tr>
<td>Leptospira</td>
<td>A23G26C21T15</td>
<td>Clostridium septic</td>
<td>A26G2C16T26</td>
</tr>
<tr>
<td>interrogans</td>
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<td></td>
</tr>
<tr>
<td>Clostridium</td>
<td>A23G27C19T19</td>
<td>Francisella tularensis</td>
<td>A26G2C16T19</td>
</tr>
<tr>
<td>perfringens</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Bacillus</td>
<td>A23G27C19T15</td>
<td>Acinetobacter calcoacetic</td>
<td>A26G2C16T19</td>
</tr>
<tr>
<td>anthracis</td>
<td></td>
<td></td>
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</tr>
<tr>
<td>Bacillus</td>
<td>A23G27C19T15</td>
<td>Bacteroides fragilis</td>
<td>A26G2C16T22</td>
</tr>
<tr>
<td>cereus</td>
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<tr>
<td>Bacillus</td>
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<td>psittaci</td>
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</tr>
<tr>
<td>dysenteriae</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
Example 2

HPLC Base Composition Determination

Example 1 can be repeated and instead of sequencing methodologies, HPLC type methodologies can be used to determine base compositions of the various amplicons. Methods for determining base compositions with HPLC and UV-vis detection are described in Manderville and Kroppinski, “Approaches to the Compositional Analysis of DNA,” Methods Mol Biol. 2009; 502:11-7, which is herein incorporated by reference. Briefly, a given PCR primer-pair generated amplicon can be rendered single-stranded (e.g., by heat) and then the forward primer strand or reverse primer strand can be removed from the sample so only the sense strand or antisense strand is present. This can be done, for example, by subjecting the amplicon containing sample to a solid support that has the forward primer linked to the solid support. Sense strands can then hybridize to the solid support, while reverse strands are washed away. This purified sample can then be digested (e.g., to completion) to generate mononucleo- seides (e.g., the sample can be expose to alkaline phosphatase and a phosphodiesterase, see Huang et al., Free Radical Biology & Medicine, 31:1341-1351, 2001, herein incorporated by reference). This sample may then be filtered to remove enzymes. This filtered sample is then injected onto a HPLC column (e.g., C18 column, such as 5 um Agilent ZORBAX Eclipse XDB C18 column, 4.6 mm×150 mm) with 0.1M triethylammonium acetate (pH 6.5) containing 5% CH3CN (buffer A) and 0.1M triethylammonium acetate (pH 6.5) containing 65% CH3CN (buffer B) operated at a flow rate of 1 ml/min and ambient temperature. This system may employ isocratic elution with 95% buffer A and 5% buffer B. The order of elution for these conditions is dC, dG, dT, dA. Standards for the four deoxynucleosides should be used for comparison and to generate standard curves that can be used for quantification. The amount of each deoxynucleoside in the single stranded amplicon is determined by integration to give the area under each peak in the HPLC trace. These areas must be divided by the following extinction coefficients at 254 nm to take into account the different absorptions of the deoxynucleosides at the detection wavelength (Connolly, 1991, “Oligonucleotides Containing Modified Bases, In F. Eckstein (Ed.), Oligonucleotides and Analogues A Practical Approach, Oxford University Press, New York, herein incorporated by reference). Levels of the normal nucleosides are quantified using the standard curves derived from standards. The extinction coefficients at 254 nm: dC (6×10^4), dG (13.5×10^4), T (7×10^4), and dA (14.3×10^4). Once the base compositions are determined in this manner, they can be compared to database entries and unknown biogentic amplicons can be identified.

What is claimed is:

1. A method of identifying a template nucleic acid, the method comprising:
   (a) determining at least a partial base count of at least a subsequence of at least one template nucleic acid and/or a complement thereof, using an approach that does not measure molecular mass of the template nucleic acid, to produce base count data; and
   (b) querying a database comprising at least one base count entry corresponding to an identified nucleic acid to produce a match of the base count data with the base count entry, thereby identifying the template nucleic acid.

* * * * *