Title: NON-INVASIVE ANALYTE MEASUREMENT DEVICE FOR MEASURING TEARS AND OTHER OCULAR ELEMENTS USING ELECTROMAGNETIC RADIATION AND METHOD OF USING THE SAME

Abstract: A method of non-invasively measuring the presence, absence or concentration of one or more analytes in an ocular element of a subject, the subject including an eye with an ocular surface and a tear layer, includes exposing at least a portion of the tear layer and/or other ocular elements of the subject to electromagnetic radiation without contact with the ocular surface; detecting electromagnetic radiation reflected from the tear layer and/or other ocular elements without contact with the ocular surface; and determining a radiation signature of the reflected electromagnetic radiation to determine the presence, absence or concentration of the one or more analytes in the tear layer and/or other ocular elements of the subject.
For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.
NON-INVASIVE ANALYTE MEASUREMENT DEVICE FOR MEASURING TEARS AND OTHER OCULAR ELEMENTS USING ELECTROMAGNETIC RADIATION AND METHOD OF USING THE SAME

FIELD OF THE INVENTION

[01] The present invention relates to the non-invasive measurement of glucose and other medically important analytes through the use of infrared radiation measurements on tears and other ocular elements.

BACKGROUND OF THE INVENTION

[02] Diabetes remains one of the most serious and under-treated diseases facing the worldwide healthcare system. Diabetes is a chronic disease where the body fails to maintain normal levels of glucose in the bloodstream. It is now the fifth leading cause of death from disease in the U.S. today and accounts for about 15% of the entire healthcare budget. People with diabetes are classified into two groups: Type 1 (formerly known as "juvenile onset" or "insulin dependent" diabetes, that are required to take insulin to maintain life) and Type 2 (formerly known as "adult onset" or "non-insulin dependent," that may require insulin but may sometimes be treated by diet and oral hypoglycemic drugs). In both cases, without dedicated and regular blood glucose measurement, all patients face the possibility of the complications of diabetes that include cardiovascular disease, kidney failure, blindness, amputation of limbs and premature death.

[03] The number of cases of diabetes in the U.S. has jumped 40% in the last decade. This high rate of growth is believed to be due to a combination of genetic and lifestyle origins that appear to be a long-term trend, including obesity and poor diet. The American Diabetes Association (ADA) and others estimate that about 17 million Americans and over 150 million people worldwide have diabetes, and it is estimated that up to 40% of these people are currently undiagnosed [American Diabetes Association, "Facts & Figures"].

[04] Diabetes must be "controlled" in order to delay the onset of the disease complications. Therefore, it is essential for people with diabetes to measure their blood glucose levels several times per day in an attempt to keep their glucose levels within the normal range (80 to 126 mg/dl). These glucose measurements are used to determine the amount of insulin or alternative treatments necessary to bring the glucose level to within
-2-
target limits. Self-Monitoring of Blood Glucose (SMBG) is an ongoing process repeated multiple times per day for the rest of the patient's lifetime.

[05] All currently FDA approved invasive or "less-invasive" (blood taken from the arm or other non-fingertip site) glucose monitoring products currently on the market require the drawing of blood in order to make a quantitative measurement of blood glucose. The ongoing and frequent measurement requirements (1 to possibly 10 times per day) presents all diabetic patients with pain, skin trauma, inconvenience, and infection risk resulting in a general reluctance to frequently perform the critical measurements necessary for selecting the appropriate insulin dose or other therapy.

[06] These current product drawbacks have led to a poor rate of patient compliance. Among Type 1 diabetics, 39% measure their glucose levels less than once per day and 21% do not monitor their glucose at all. Among Type 2 diabetics who take insulin, only 26% monitor at least once per day and 47% do not monitor at all. Over 75% of non-insulin-taking Type 2 diabetics never monitor their glucose levels. Roper Starch Worldwide Survey. Of 1,186 diabetics surveyed, 91% showed interest in a non-invasive glucose monitor [www.childrenwithdiabetes.com]. As such, there is both a tremendous interest and clinical need for a non-invasive glucose sensor.

[07] The present invention seeks to replace the currently used blood glucose measurement methods, devices and instruments, including invasive measures and the use of glucose test strips, with an optical non-invasive instrument.

[08] Various methods have been developed related to non-invasive glucose sensing using a dermal testing site such as the finger or earlobe. These methods primarily employ instruments which measure blood-glucose concentration by generating and measuring light only in the near-infrared radiation spectrum. For example, U.S. Patent No. 4,882,492 (the '492 patent), expressly incorporated by reference herein, is directed to an instrument which transmits near-infrared radiation through a sample to be tested on the skin surface of a human. In the '492 patent, the near-infrared radiation that passes through the sample is split into two beams, wherein one beam is directed through a negative correlation filter and the second through a neutral density filter. The differential light intensity measured through the filters of the two light beams is proportional to glucose concentration according to the '492 patent.

[09] U.S. Patent No. 5,086,229 (the '229 patent), expressly incorporated by reference herein, is directed to an instrument which generates near-infrared radiation within the
spectrum of about 600 to about 1100 nanometers. According to the '229 patent, a person places their finger in between the generated near-infrared radiation source and a detector, which correlates the blood-glucose concentration based on the detected near-infrared radiation. Similarly, U.S. Patent No. 5,321,265 (the '265 patent), expressly incorporated by reference herein, also measures blood-glucose level using both near-infrared radiation and the fingertip as a testing site. The detectors disclosed in the '265 patent further comprise silicon photocells and broad bandpass filters.

[10] U.S. Patent No. 5,361,758 (the '758 patent), expressly incorporated by reference herein, is directed to an instrument which measures near-infrared radiation that is either transmitted through or is reflected from the finger or earlobe of a human. In the '758 patent, the transmitted or reflected light is separated by a grating or prism, and the near-infrared radiation is detected and correlated with blood-glucose concentration. This instrument of the '758 patent also comprises an additional timing and control program wherein the device takes measurements specifically in between heartbeats and can also adjust for temperature.

[11] U.S. Patent No. 5,910,109 (the '109 patent), expressly incorporated by reference herein, is also directed to an instrument for measuring blood-glucose concentration using near-infrared radiation and the earlobe as the testing site. The instrument of the '109 patent comprises four light sources of a very specific near-infrared emission spectrum, and four detectors having specific near-infrared detection spectra corresponding to the wavelength of the light sources. The signals detected by the four distinct detectors are averaged, and these averages are analyzed to determine blood-glucose concentration according to the '109 patent.

[12] The technique of using near-infrared radiation, wherein the near-infrared radiation is transmitted through or reflected from a dermal testing site and monitored for measuring glucose in vivo, is known to be inaccurate. The glucose concentration of interest is in the blood or the interstitial fluid, not on the surface of the dermis. Therefore these methods must penetrate down into the layers beneath the top layers of dermis. There are a number of substances in the dermis that can interfere with the near-infrared glucose signal. Additionally, there is a wide variation in the human dermis, both between individuals and within a given individual. Moreover, glucose simply lacks a satisfactory distinguishable "fingerprint" in the near-infrared radiation spectrum. Because near-infrared radiation is not sufficiently adsorbed by glucose and because of the level of tissue interferences found
in the dermis, this technique is substantially less desirable for the accurate measurement of blood-glucose concentrations.

[13] U.S. Patent No. 6,362,144 (the '144 patent), expressly incorporated by reference herein, discloses using the fingertip as a testing site, however, the described instrument uses attenuated total reflection (ATR) infrared spectroscopy. According to the '144 patent, a selected skin surface, preferably the finger, is contacted with an ATR plate while ideally maintaining the pressure of contact. In the '144 patent, the skin is then irradiated with a mid-infrared beam, wherein the infrared radiation is detected and quantified to measure blood-glucose levels. This technique is not ideal, however, if the surface of tissue from which the measurement is taken is very dense in the wavelength region of interest or is not amenable to direct contact with the ATR plate, such as an eye, conjunctiva, nose, mouth, or other orifice, cavity or piercing tract.

[14] The minimal depth of peripheral capillaries in epithelial tissues is typically about 40 microns. Again, there are physical characteristics as well as a number of substances present in the skin that can interfere with the desired glucose-specific signal. While useful in the laboratory, both the near-infrared transmission methods and the ATR method mentioned above are not practical, or may not be adequate for use in monitoring blood glucose concentration in patients.

[15] Methods have also been developed related to non-invasive glucose sensing using the eye as a testing site. For example, in both U.S. Patent Nos. 3,958,560 (the '560 patent) and 4,014,321 (the '321 patent), both expressly incorporated by reference herein, a device utilizing the optical rotation of polarized light is described. In the '560 and the '321 patents, the light source and light detector are incorporated into a contact lens which is placed in contact with the surface of the eye whereby the eye is scanned using a dual source of polarized radiation, each source transmitting in a different absorption spectrum at one side of the cornea or aqueous humor. The optical rotation of the radiation that passes through the cornea correlates with the glucose concentration in the cornea according to the '560 and '321 patents. While this method would be termed, "non-invasive" because the withdrawal of blood is not required, it may still cause significant discomfort or distort vision of the user because of the need to place the sensor directly in contact with the eye.

[16] U.S. Patent No. 5,009,230 (the '230 patent), expressly incorporated by reference herein, uses a polarized light beam of near-infrared radiation within the range of 940 to
1000 nm. In the '230 patent, the amount of rotation imparted by glucose present in the bloodstream of the eye on the polarized light beam is measured to determine glucose concentration. Again, the accuracy is limited because glucose simply lacks a sufficiently distinguishable “fingerprint” in this near-infrared radiation spectrum.

[17] Both U.S. Patent No. 5,209,231 (the '231 patent), and International Publication No. WO 92/07511 (the '511 application), both expressly incorporated by reference herein, similarly disclose the use of polarized light, which is initially split by a beam splitter into a reference beam and a detector beam, and then transmitted through a specimen, preferably the aqueous humor of the eye. The amount of phase shift as compared between the transmitted reference and detector beams are correlated to determine glucose concentration in the '231 patent and '511 application. U.S. Patent No. 5,535,743 (the '743 patent), expressly incorporated by reference herein, measures diffusely reflected light provided by the surface of the iris as opposed to the aqueous humor of the eye. According to the '743 patent, the measurement of optical absorption is possible whereas measurement of the optical rotation through the aqueous humor is not possible. In the '743 patent, the intensity of the diffusely reflected light, however, may be analyzed to obtain useful information on the optical properties of the aqueous humor, including blood-glucose concentration.

[18] U.S. Patent No. 5,687,721 (the '721 patent), expressly incorporated by reference herein, also discloses a method of measuring blood-glucose concentration by generating both a measurement and reference polarized light beam, and comparing the beams to determine the angle of rotation, which is attributable to the blood-glucose concentration. The preferable testing site disclosed, however, is the finger or other suitable appendage according to the '721 patent. The '721 patent further discloses and requires the use of a monochromatic laser and/or semi-conductor as a light source.

[19] U.S. Patent No. 5,788,632 (the '632 patent), expressly incorporated by reference herein, discloses a non-invasive instrument for determining blood-glucose concentration by transmitting a first beam of light through a first polarizer and a first retarder, then directing the light through the sample to be measured, transmitting the light through a second polarizer or retarder, and lastly detecting the light from the second detector. The rotation of measured polarized light is correlated to the blood-glucose concentration of the sample measured according to the '632 patent.
[20] U.S. Patent No. 5,433,197 (the '197 patent), expressly incorporated by reference herein, discloses a non-invasive instrument for determining blood-glucose concentration using a broad-band of near-infrared radiation which illuminates the eye in such a manner that the energy passes through the aqueous humor in the anterior chamber of the eye and is then reflected from the iris. The reflected energy then passes back through the aqueous humor and the cornea and is collected for spectral analysis. According to the '197 patent, the electrical signals representative of the reflected energy are analyzed by univariate and/or multivariate signal processing techniques to correct for any errors in the glucose determination. Again, the accuracy of the instrument in the '197 patent is limited because glucose simply lacks a sufficiently distinguishable "fingerprint" in this near-infrared radiation spectrum.

[21] Instruments and methods of using the body's naturally emitted radiation to measure blood-glucose concentration using the human body, and in particular, the tympanic membrane as a testing site have also been disclosed. U.S. Patent Nos. 4,790,324; 4,797,840; 4,932,789; 5,024,533; 5,167,235; 5,169,235; and 5,178,464, expressly incorporated by reference herein, describe various designs, stabilization techniques and calibration techniques for tympanic non-contact thermometers. In addition, U.S. Patent No. 5,666,956 (the '956 patent), expressly incorporated by reference herein, discloses an instrument which measures electromagnetic radiation from the tympanic membrane and computes monochromatic emissivity using Plank's law by measuring the radiation intensity, spectral distribution, and blackbody temperature. According to the '956 patent, the resultant monochromatic emissivity is variable depending on the spectral characteristics of the site measured, namely the blood-glucose concentration measured from the tympanic membrane. It should be noted, however, that the '956 patent equates skin surfaces of the body to a "gray-body" rather than a black-body with respect to its monochromatic emissivity. Therefore, according to the '956 patent, the accuracy of such skin surface-based methods utilizing natural black-body emitted radiation is not useful for analyte measurements, as compared to a method of subsurface analysis utilizing natural black-body radiation emitted from the tympanic membrane.

[22] The human body naturally emits from its surfaces infrared radiation whose spectrum, or radiation signature, is modified by the presence, absence or concentration of analytes in the body tissues. The eye is particularly well suited as a testing site to detect
this infrared radiation. For example, certain analytes, such as glucose, exhibit a minimal
time delay in glucose concentration changes between the eye and the blood, and the eye
provides a body surface with few interferences [Cameron et al, (3)2 DIABETES TECHNOL.
THER., 202-207 (2001)]. There is, therefore, in the field of non-invasive blood analyte
monitoring, an unmet need for a suitable instrument, and methodologies for using it, to
accurately measure analyte concentrations, such as blood glucose concentration, as well as
concentrations of other desired analytes, in subjects requiring this type of blood analyte
measurement.

SUMMARY OF THE INVENTION

[23] The present invention seeks to replace the currently used invasive blood glucose
measurement instruments and methods, including the use of glucose test strips, with a
hand-held, non-invasive measurement device that shines infrared radiation onto the tear
layer covering the eye, without physical contact with the ocular surface, and the reflected
signal can be used to determine the presence, absence or concentration of glucose and
other medically important analytes. In another embodiment, the device measures the
infrared radiation radiating from the tear layer covering the eye, without physical contact
with the ocular surface, and the naturally emitted signal can be used to determine the
presence, absence or concentration of glucose and other medically important analytes.
The non-invasive nature of glucose level measurement with the device makes glucose
level monitoring painless and simple.

[24] Another aspect of the invention involves a method of non-invasively measuring the
presence, absence or concentration of one or more analytes in an ocular element of a
subject, the subject including an eye with an ocular surface and a tear layer. The method
includes exposing at least a portion of the tear layer of the subject to electromagnetic
radiation without contact with the ocular surface; detecting electromagnetic radiation
reflected from the tear layer without contact with the ocular surface; and determining a
radiation signature of the reflected electromagnetic radiation to determine the presence,
absence or concentration of the one or more analytes in the tear layer of the subject.

[25] An additional aspect of the invention involves a method of non-invasively
measuring the presence, absence or concentration of one or more analytes in an ocular
element of a subject, the subject including an eye with an ocular surface and multiple
ocular elements. The method includes measuring the presence, absence or concentration
of the one or more analytes from the eye by profiling more than one different ocular element to determine an ideal ocular element for measuring the presence, absence or concentration of the one or more analytes; measuring the presence, absence or concentration of one or more analytes from the ideal ocular element by exposing at least a portion of the ideal ocular element to electromagnetic radiation without contact with the ocular element; detecting electromagnetic radiation reflected from the ocular element without contact with the ocular element; and determining a radiation signature of the reflected electromagnetic radiation to determine the presence, absence or concentration of the one or more analytes in the ocular element of the subject.

[26] A further aspect of the invention involves a non-invasive analyte measurement instrument for determining a concentration of one or more analytes in an ocular element of a subject, the subject including an eye with an ocular surface and a tear layer. The instrument includes means for exposing at least a portion of the tear layer of the subject to electromagnetic radiation without contact with the ocular surface; means for detecting electromagnetic radiation reflected from the tear layer without contact with the ocular surface; and means for determining a radiation signature of the reflected electromagnetic radiation to determine the presence, absence or concentration of the one or more analytes in the ocular element of the subject.

[27] A still further aspect of the invention involves a non-invasive analyte measurement instrument for determining a concentration of one or more analytes in an ocular element of a subject, the subject including an eye with an ocular surface and multiple ocular elements. The instrument includes means for profiling more than one different ocular element to determine an ideal ocular element for measuring analyte concentration; and means for measuring the presence, absence or concentration of one or more analytes from the ideal ocular element. The measuring means includes means for exposing at least a portion of the ideal ocular element to electromagnetic radiation without contact with the ocular element; means for detecting electromagnetic radiation reflected from the ocular element without contact with the ocular element; and means for determining a radiation signature of the reflected electromagnetic radiation to determine an analyte concentration in the ocular element of the subject.

[28] Another aspect of the invention involves a method of non-invasively measuring the presence, absence or concentration of one or more analytes in one or more ocular elements of a subject, the subject including an eye with an ocular surface. The method includes
exposing at least a portion of the one or more ocular elements of the subject to electromagnetic radiation without contact with the ocular surface; detecting electromagnetic radiation reflected from the one or more ocular elements without contact with the ocular surface; and determining a radiation signature of the reflected electromagnetic radiation to determine the presence, absence or concentration of the one or more analytes in the one or more ocular elements of the subject.

[29] A still further aspect of the invention involves a non-invasive analyte measurement instrument for determining the presence, absence or concentration of one or more analytes in one or more ocular elements of a subject, the subject including an eye with an ocular surface. The non-invasive analyte measurement instrument includes means for exposing at least a portion of the one or more ocular elements of the subject to electromagnetic radiation without contact with the ocular surface; means for detecting electromagnetic radiation reflected from the one or more ocular elements without contact with the ocular surface; and means for determining a radiation signature of the reflected electromagnetic radiation to determine the presence, absence or concentration of the one or more analytes in the one or more ocular elements of the subject.

[30] Other objectives, features and advantages of the present invention will become apparent from the following detailed description. The detailed description and the specific examples, although indicating specific embodiments of the invention, are provided by way of illustration only. Accordingly, the present invention also includes those various changes and modifications within the spirit and scope of the invention that may become apparent to those skilled in the art from this detailed description.

BRIEF DESCRIPTION OF THE DRAWINGS

[31] Figure 1, Panel A provides a graphical illustration of the human eye. Panel B shows the high degree of vascularization in the conjunctiva, with veins (V) and arterioles (A).

[32] Figure 2 provides a graphical illustration of one embodiment of the present invention, wherein analyte concentration is measured from the mid-infrared radiation reflected back from the eye.

[33] Figure 3 provides a flowchart of one embodiment of the present invention, comprising a method wherein a remote access user can receive a subject's measured analyte concentrations which have been downloaded and stored in a computer system.
[34] Figure 4 provides a graph of multiple dose response measurements using detection of varying concentrations of glucose using polyethylene membranes as the measurement surface.

[35] Figure 5 shows a plot of the glucose concentration versus mid-infrared absorption using polyethylene membranes as the measurement surface.

[36] Figure 6 shows a plot of the results obtained from mid-infrared measurements of glucose concentration using a rabbit eye as the surface from which the measurements were made.

[37] Figure 7 shows a plot of human data obtained from the conjunctiva of the patient's eye measured using mid-infrared absorption to determine blood glucose concentration of the patient.

[38] Figure 8 shows a plot of the data obtained from a human diabetic patient in a glucose tracking study demonstrating a correlation of glucose concentration with mid-infrared absorption measured from the human eye surface.

[39] Figure 9 shows the correlation between glucose measurements taken from the eye according to the methods of the present invention (squares) and SMBG measurements (diamonds).

[40] Figure 10 shows a schematic of an embodiment of an optical, non-invasive glucose monitor with depth profiling/adjustable focus to choose an ideal ocular element for measuring analyte concentrations.

DETAILED DESCRIPTION OF THE INVENTION

[41] It is understood that the present invention is not limited to the particular methodologies, protocols, instruments, and systems, etc., described herein, as these may vary. It is also to be understood that the terminology used herein is used for the purpose of describing particular embodiments only, and is not intended to limit the scope of the present invention. It must be noted that as used herein and in the appended claims, the singular forms "a," "an," and "the" include plural reference unless the context clearly dictates otherwise. Thus, for example, a reference to "a mid-infrared filter" is a reference to one or more filters and includes equivalents thereof known to those skilled in the art, and so forth. Further, for example, a reference to an instrument/monitor for non-invasively measuring the presence, absence or concentration of one or more analytes in an ocular element of a subject is a reference to the instrument/monitor and includes devices
(i.e., combination devices) that may integrate the instrument/monitor with one or more additional mechanisms. For example, but not by way of limitation, the instrument/monitor may be integrated with a wireless communication device to wirelessly transmit/receive information.

[42] Unless defined otherwise, all technical and scientific terms used herein have the same meanings as commonly understood by one of ordinary skill in the art to which this invention belongs. Preferred methods, devices, and materials are described, although any methods and materials similar or equivalent to those described herein can be used in the practice or testing of the present invention. All references cited herein are incorporated by reference herein in their entirety.

DEFINITIONS

[43] **Analyte**: As used herein describes any particular substance or chemical constituent to be measured. Analyte may also include any substance in the tissue of a subject, in a biological fluid (for example, blood, interstitial fluid, cerebrospinal fluid, lymph fluid or urine), or is present in air that was in contact with or exhaled by a subject, which demonstrates an infrared radiation signature. Analyte may also include any substance which is foreign to or not normally present in the body of the subject. Analytes can include naturally occurring substances, artificial substances, metabolites, and/or reaction products. In some embodiments, the analyte for measurement by the devices and methods described herein is glucose. However, other analytes are contemplated as well, including, but not limited to, metabolic compounds or substances, carbohydrates such as sugars including glucose, proteins, glycated proteins, fructosamine, hemoglobin A1c, peptides, amino acids, fats, fatty acids, triglycerides, polysaccharides, alcohols including ethanol, toxins, hormones, vitamins, bacteria-related substances, fungus-related substances, virus-related substances, parasite-related substances, pharmaceutical or non-pharmaceutical compounds, substances, pro-drugs or drugs, and any precursor, metabolite, degradation product or surrogate marker of any of the foregoing. Other analytes are contemplated as well, including, but not limited to, acarboxyprothrombin; acylcarnitine; adenine phosphoribosyl transferase; adenosine deaminase; albumin; alpha-fetoprotein; amino acid profiles (arginine (Krebs cycle), histidine/urocanic acid, homocysteine, phenylalanine/tyrosine, tryptophan); andrenostenedione; antipyrine; arabininol enantiomers; arginase; benzoylecgonine (cocaine); biotinidase; bioterin; c-reactive
protein; carnitine; carnosinase; CD4; ceruloplasmin; chenodeoxycholic acid; chloroquine; cholesterol; cholinesterase; conjugated 1-hydroxy-cholic acid; Cortisol; creatine kinase; creatine kinase MM isoenzyme; cyclosporin A; d-penicillamine; de-ethylchloroquine; dehydroepiandrosterone sulfate; nucleic acids (deoxyribonucleic acids and ribonucleic acids including native and variant sequences related to acetylator polymorphism, alcohol dehydrogenase, alpha 1-antitrypsin, cystic fibrosis, Down's syndrome, Duchenne/Becker muscular dystrophy, glucose-6-phosphate dehydrogenase, hemoglobin A, hemoglobin S, hemoglobin C, hemoglobin D, hemoglobin E, hemoglobin F, D-Punjab, beta-thalassemia, hepatitis B virus, HCMV, HIV-I, HTLV-I, Leber hereditary optic neuropathy, MCAD, PKU, Plasmodium vivax, sexual differentiation, 21-hydroxylase); 21-deoxycortisol; desbutylhalofantrine; dihydropteridine reductase; diptheria/tetanus antitoxin; erythrocyte arginase; erythrocyte protoporphyrin; esterase D; fatty acids/acylglycines; free -human chorionic gonadotropin; free erythrocyte porphyrin; free thyroxine (FT4); free triiodothyronine (FT3); fumarylacetoacetase; galactose/gal-1-phosphate; galactose-1-phosphate uridytransferase; gentamicin; glucose-6-phosphate dehydrogenase; glutathione; glutathione peroxidase; glycocholic acid; glycosylated hemoglobin; halofantrine; hemoglobin variants; hexosaminidase A; human erythrocyte carbonic anhydrase I; 17-alpha-hydroxyprogesterone; hypoxanthine phosphoribosyl transferase; immunoreactive trypsin; lactate; lead; lipoproteins ((a), B/A-I, ); lysozyme; mefloquine; netilmicin; phenobarbitone; phenytoin; phytic/pristanic acid; progesterone; prolactin; prolidase; purine nucleoside phosphorylase; quinine; reverse triiodothyronine (rT3); selenium; serum pancreatic lipase; sissomicin; somatomedin C; specific antibodies (adenovirus, antinuclear antibody, anti-zeta antibody, arbovirus, Aujeszky's disease virus, dengue virus, Dracunculus medinensis, Echinococcus granulosus, Entamoeba histolytica, enterovirus, Giardia duodenalis, Helicobacter pylori, hepatitis B virus, herpes virus, HIV-I, IgE (atopic disease), influenza virus, Leishmania donovani, leptospira, measles/mumps/rubella, Mycobacterium leprae, Mycoplasma pneumoniae, Myoglobin, Onchocerca volvulus, parainfluenza virus, Plasmodium falciparum, poliovirus, Pseudomonas aeruginosa, respiratory syncytial virus, rickettsia (scrub typhus), Schistosoma mansoni, Toxoplasma gondii, Treponema pallidium, Trypanosoma cruzi/rangeli, vesicular stomatis virus, Wuchereria bancrofti, yellow fever virus); specific antigens (hepatitis B virus, HIV-I); neurotransmitters (such as glutamate, GABA, dopamine, serotonin), opioid neurotransmitters (such as endorphins, and dynorphins),
neurokinins (such as substance P); succinylacetone; sulfadoxine; theophylline; thyrotropin (TSH); thyroxine (T4); thyroxine-binding globulin; trace elements; transferrin; UDP-galactose-4-epimerase; urea; prokaryotic and eukaryotic cell-surface antigens; peptidoglycans; lipopolysaccharide; uroporphyrinogen I synthase; vitamin A; white blood cells; and zinc protoporphyrin. Salts naturally occurring in blood or interstitial fluids can also constitute analytes in certain embodiments. The analyte can be naturally present in the biological fluid, for example, a metabolic product, an antigen, an antibody, and the like. Alternatively, the analyte can be introduced into the body, for example, a contrast agent for imaging, a radioisotope, a chemical agent, a fluorocarbon-based synthetic blood, or pharmaceutical composition, including but not limited to insulin; ethanol; cannabis (marijuana, tetrahydrocannabinol, hashish); inhalants (nitrous oxide, amyl nitrite, butyl nitrite, chlorohydrocarbons, hydrocarbons); cocaine (crack cocaine); stimulants (amphetamines, methamphetamine, Ritalin, Cylert, Preludin, Didrex, PreState, Voranil, Sandrex, Plegine); depressants (barbiturates, methaqualone, tranquilizers such as Valium, Librium, Miltown, Serax, Equanil, Tranxene); tricyclic antidepressants, benzodiazepines, acetaminophen (paracetamol, APAP), aspirin, methadone, hallucinogens (phencyclidine, lysergic acid, mescaline, peyote, psilocybin); narcotics (heroin, codeine, morphine, opium, meperidine, Percocet, Percodan, Tussionex, Fentanyl, Darvon, Talwin, Lomotil); designer drugs (analogs of fentanyl, meperidine, amphetamines, methamphetamine, and phencyclidine, for example, Ecstasy); anabolic steroids; and nicotine. The metabolic products of drugs and pharmaceutical compositions are also contemplated analytes.

Analytes such as neurochemicals and other chemicals generated within the body can also be analyzed, such as, for example, ascorbic acid, uric acid, dopamine, noradrenaline, 3-methoxytyramine (3MT), 3,4-dihydroxyphenylacetic acid (DOPAC), homovanillic acid (HVA), 5-hydroxytryptamine (5HT), and 5-hydroxyindoleacetic acid (5HIAA).

[44] **Conjunctiva**: As used herein describes the membranous tissue that covers the exposed surface of the eye and the inner surface of the eyelids.

[45] **Electromagnetic Radiation**: As used herein refers to any radiation energy, either generated from any source or naturally emitted, in the electromagnetic spectrum, namely, radiation energy having a frequency within the range of approximately $10^{23}$ hertz to 0 hertz and a wavelength within the range of approximately $10^{13}$ centimeter to infinity and...
including, in order of decreasing frequency, cosmic-ray photons, gamma rays, x-rays, ultraviolet radiation, visible light, infrared radiation, microwaves, and radio waves.

[46]   **Far-Infrared Radiation**: As used herein refers to any radiation, either generated from any source or naturally emitted, having wavelengths of about 50.00 to about 1000.00 microns.

[47]   **Flooding**: As used herein refers to broadly applying relatively widely diffused or spread-out rays of light onto a surface.

[48]   **Focused**: As used herein means mostly parallel rays of light that are caused to converge on a specific predetermined point.

[49]   **Infrared Radiation**: As used herein refers to any radiation, either generated from any source or naturally emitted, having wavelengths of about 0.78 to about 1000.00 microns.

[50]   **Mid-Infrared Radiation**: As used herein refers to any radiation, either generated from any source or naturally emitted, having wavelengths of about 2.50 microns to about 50.00 microns.

[51]   **Mid-Infrared Radiation Detector**: As used herein refers to any detector or sensor capable of registering infrared radiation. Examples of a suitable infrared radiation detectors include, but are not limited to, a thermocouple, a thermistor, a microbolometer, and a liquid nitrogen cooled Mercury Cadmium Telluride (MCT) detector. The combined detected infrared radiation may be correlated with wavelengths corresponding to analyte concentrations using means such as the Fourier transform to produce high resolution spectra.

[52]   **Near-Infrared Radiation**: As used herein refers to any radiation, either generated or naturally emitted, having wavelengths of about 0.78 to about 2.50 microns.
Non-invasive: As used herein refers to a method or instrument that does not break a subject's skin nor any other tissue barriers.

Ocular element: As used herein refers to an element of or relating to the eye such as, but not limited to the eyelid(s), the epithelial cells, the aqueous humor, the vitreous humor, various layers of the cornea, lens, various layers of the sclera, conjunctiva, interstitial fluid in the conjunctiva, tears, the tear layer, and blood vessels.

Surface: As used herein refers to any part of a subject's body that may be exposed to the external environment, including but not limited to, skin, the eye, ear, mouth, nose or any other orifice, body cavities, piercing tracts or other surface whether naturally occurring or artificial such as a surgically created surface. Also includes samples such as urine, tears and saliva, which do not require that the skin be punctured in order to obtain a sample for measurement.

Tears: The fluid secreted by the lacrimal gland and diffused between the eye and eyelids to moisten the parts and facilitate their motion.

Tear layer: The layer of fluid on the eye created by the tears.

Tissue: As used herein includes any tissue or component of a subject, including, but not limited to, skin, blood, body fluids, the eye, the tear layer of the eye, interstitial fluid, ocular fluid, bone, muscle, epithelium, fat, hair, fascia, organs, cartilage, tendons, ligaments, and any mucous membrane.

NON-INVASIVE GLUCOSE MEASUREMENT

In one aspect of the present invention, electromagnetic radiation, and more preferably, infrared radiation, and even more preferably, mid-infrared radiation, is flooded onto the eye surface using a radiation source. This flooded mid-infrared radiation is reflected from the eye to a detector. The reflected radiation is detected by a mid-infrared detection instrument placed before the eye. The radiation signature of the reflected mid-infrared radiation is affected by the presence or concentration of analytes. This provides a non-invasive method employing an instrument of the present invention to measure the
presence, absence, or concentration of one or more analytes, such as, but not limited to, glucose, from a tissue such as, but not limited to, the conjunctiva of a subject (Figure 2). [60] The use of tear fluids for chemical analysis holds great potential as a non-invasive approach for clinical diagnosis. (Chen, R. et al, J Cap. Elec. 003:5, 243-248, 1996). Gasset et.al. reported that the glucose concentration in tears is approximately 5% of the blood glucose concentration. (Gasset, et. al., Amer. J Ophthal. 65:414-209, 1968). Furthermore, the glucose in tears varied in proportion to the blood glucose concentration. These investigators and others have used a variety of techniques to collect the tear sample followed by a chemical method to measure the glucose concentration in the tear sample. Thus, the tear layer is an especially ideal ocular element for non-invasive measurement of the presence, absence or concentration of analytes in the tissue of a subject. The instruments/devices and methods below will be generally described in conjunction with the non-invasive measurement of the presence, absence or concentration of analytes in the tear layer of a subject.

[61] However, in alternative embodiments, one or more other and/or additional ocular elements including, but not limited to, the eyelid(s), the epithelial cells, the aqueous humor, the vitreous humor, various layers of the cornea, lens, various layers of the sclera, conjunctiva, interstitial fluid in the conjunctiva, tears, the tear layer, and blood vessels are the ocular element(s) for non-invasive measurement of the presence, absence or concentration of analytes in the tissue of a subject.

[62] Further, there is substantial evidence that fluctuations in blood glucose levels are well correlated with glucose levels in the aqueous humor of the eye [Steffes, 1(2) DIABETES TECHNOL. THER., 129-133 (1999)]. In fact, it is estimated that the time delay between the blood and aqueous humor glucose concentration averages only about five minutes [Cameron et al., 3(2) DIABETES TECHNOL. THER., 201-207 (2001)]. The aqueous humor is a watery liquid that lies between the lens and cornea, which bathes and supplies the nutrients to the cornea, lens and iris. The glucose in the eye is located throughout the various components and compartments of the eye, including, but not limited to, epithelial cells, the aqueous humor, the vitreous humor, various layers of the cornea, lens, various layers of the sclera, conjunctiva, tears, the tear layer, and blood vessels. The eye, including, but not limited to, the tear layer and the conjunctiva, is both an ideal and suitable body surface for non-invasive measurement of the presence, absence or concentration of analytes in the tissue of a subject.
MEASURING MID-INFRARED RADIATION

[63] When electromagnetic radiation is passed through a substance, it can either be absorbed or transmitted, depending upon its frequency and the structure of the molecules it encounters. Electromagnetic radiation is energy and hence when a molecule absorbs radiation it gains energy as it undergoes a quantum transition from one energy state \(E_{\text{in}}\) to another \(E_{\text{fin}}\). The frequency of the absorbed radiation is related to the energy of the transition by Planck's law: \(E_{\text{fin}} - E_{\text{in}} = E = h\nu = h\nu/\lambda\). Thus, if a transition exists which is related to the frequency of the incident radiation by Planck's constant, then the radiation can be absorbed. Conversely, if the frequency does not satisfy the Planck expression, then the radiation will be transmitted. A plot of the frequency of the incident radiation vs. some measure of the percent radiation absorbed by the sample is the radiation signature of the compound. The absorption of some amount of the radiation that is applied to a substance, or body surface containing substances, that absorbs radiation may result in a measurable decrease in the amount of radiation energy that actually passes through, or is affected by, the radiation absorbing substances. Such a decrease in the amount of radiation that passes through, or is affected by, the radiation absorbing substances may provide a measurable signal that may be utilized to measure the presence, absence or the concentration of an analyte.

[64] One embodiment of the present invention provides a method for non-invasively measuring the blood-analyte concentration in a subject comprising the steps of generating electromagnetic radiation which is flooded onto the tear layer of the subject, detecting the reflected electromagnetic radiation, correlating the spectral characteristics of the detected electromagnetic radiation with a radiation signature that corresponds to the analyte concentration, and analyzing the detected electromagnetic radiation signature to give an analyte concentration measurement. In another embodiment, the method includes a filtering step before detection, by filtering the electromagnetic radiation reflected back from a body surface so that only wavelengths of about 8.00 microns to about 11.00 pass through the filter. In this embodiment, the filtering step may be accomplished using absorption filters, interference filters, monochromators, linear or circular variable filters, prisms or any other functional equivalent known in the art. The detecting step may be accomplished using any electromagnetic radiation sensor such as a thermocouple, thermistor, microbolometer, liquid nitrogen cooled MCT, or any other functional...
equivalent known in the art. In alternative embodiments, the detector includes specular reflection optics for surface reflective measurements, and diffuse reflection optics for deeper ocular element reflective measurements. Correlating the spectral characteristics of the detected electromagnetic radiation may comprise the use of a microprocessor to correlate the detected electromagnetic radiation signature with a radiation signature of an analyte. If the analyte being measured is glucose, then the radiation signature generated may be within the wavelength range within about 8.0 to about 11.0 microns. The analyzing step further comprises a microprocessor using algorithms based on Plank's law to correlate the absorption spectrum with a glucose concentration. In another embodiment of the present invention, the analyzing step may comprise the use of a transform, such as, but not limited to, Kramers-Kronig transform or other classical transform known in the art, to transform the detected electromagnetic radiation signal to the analyte spectra for correlation.

Although the emitted and reflected electromagnetic radiation is described as mid-infrared radiation, in alternative embodiments, the electromagnetic radiation is infrared radiation or other types of electromagnetic radiation.

In another embodiment of the present invention, where glucose is the analyte of interest, an instrument comprising a electromagnetic radiation detector and a display may be held up to the tear layer of a subject. The electromagnetic radiation from the tear layer may optionally be filtered so that only wavelengths of about 8.0 microns to about 11.0 microns reach the electromagnetic radiation detector. The radiation signature of the electromagnetic radiation detected by the detector may then be correlated with a radiation signature that corresponds to a glucose concentration. The radiation signature may then be analyzed to give an accurate glucose concentration measurement. The measured glucose concentration may be displayed.

In another embodiment of the present invention, an instrument comprising an electromagnetic radiation generator, an electromagnetic radiation detector and a display may be held up to the eye of a subject. Electromagnetic radiation may be generated by the instrument and used for flooding or alternatively aiming a focused beam onto the eye of a subject. The electromagnetic radiation generated may be broad band or narrow band radiation, and may also be filtered to allow only desired wavelengths of radiation to reach the body surface. Any analyte, such as glucose, present in any constituent of the tear layer may absorb some of the generated radiation. The electromagnetic radiation that is not
absorbed by the tear layer may be reflected back to the instrument. The reflected electromagnetic radiation may optionally be filtered so that only wavelengths of about 8.0 microns to about 11.0 microns reach the electromagnetic radiation detector. The radiation signature of the electromagnetic radiation detected by the detector may then be correlated with a radiation signature that corresponds to analyte, such as glucose, concentration. The radiation signature may be analyzed to give an analyte, such as glucose, concentration. The measured analyte, such as glucose, concentration may be displayed by the instrument.

[68] Infrared radiation may be generated by the instrument of the present invention. Such infrared radiation may be generated by any suitable generator including, but not limited to, a narrow band wavelength generator or a broadband wavelength generator. In one embodiment of the present invention, an instrument may comprise a mid-infrared radiation generator. In another embodiment of the present invention, the instrument comprises a light source with one or more filters to restrict the wavelengths of the light reaching the tear layer. The mid-infrared generator may further comprise a heating element. The heating element of this embodiment may be a Nernst glower (zirconium oxide/yttrium oxides), a NiChrome wire (nickel-chromium wire), and a Globar (silicon-carbon rod), narrow band or broadband light emitting diodes, or any other functional equivalent known in the art. Mid-infrared radiation has wavelengths in the range of about 2.5 microns to about 50.0 microns. Analytes typically have a characteristic "fingerprint" or "signature" or "radiation signature" with respect to their mid-infrared radiation spectrum that results from the analyte’s affect on the mid-infrared radiation, such as absorption. Glucose in particular has a distinct spectral "fingerprint" or "signature" in the mid-infrared radiation spectrum, at wavelengths between about 8.0 microns to about 11.0 microns. This radiation signature of glucose may be readily generated for a wide variety of glucose concentrations utilizing a wide variety of body surfaces, such as the tear layer, for taking radiation signature data. In one embodiment of the present invention, an instrument may comprise a mid-infrared radiation filter, for filtering out all mid-infrared radiation not within a range of wavelengths from about 8.0 to about 11.0 microns. In other embodiments the filter is selected to filter out all mid-infrared radiation other than other than the wavelengths that provide the radiation signature of the desired analyte, such as glucose. Filtering mid-infrared radiation may be accomplished using absorption filters, interference filters, monochromators, linear or circular variable filters, prisms or any other functional equivalent known in the art.
In one embodiment of the present invention, the instrument may also comprise a mid-infrared radiation detector for detecting mid-infrared radiation. The mid-infrared radiation detector can measure the naturally emitted or reflected mid-infrared radiation in any form, including in the form of heat energy. Detecting the naturally emitted or reflected mid-infrared radiation may be accomplished using thermocouples, thermistsors, microbolometers, liquid nitrogen cooled MCT, or any other functional equivalent known in the art. Both thermocouples and thermistsors are well known in the art and are commercially available. For example, thermocouples are commonly used temperature sensors because they are relatively inexpensive, interchangeable, have standard connectors and can measure a wide range of temperatures [http://www.picoieh.com]. In addition, Thermometries' product portfolio comprises a wide range of thermistsors (thermally sensitive resistors) which have, according to type, a negative (NTC), or positive (PTC) resistance/temperature coefficient [http://www.tlicrniometric.coni].

The instrument of the present invention may also comprise a microprocessor. The microprocessor of this embodiment correlates the detected electromagnetic radiation with a radiation signature whose spectral characteristics provide information to the microprocessor about the analyte concentration being measured. The microprocessor of this embodiment analyzes the resultant radiation signature using suitable algorithms such as those based on Plank's law, to translate the radiation signature into an accurate analyte concentration measurement in the sample being measured.

It is readily apparent to those skilled in the art that a broad band light source may be modulated by an interferometer, such as in Fourier transform spectroscopy, or by an electro-optical or moving mask, as in Hadamard transform spectroscopy, to encode wavelength information in the time domain. A discrete wavelength band may be selected and scanned in center wavelength using, for example, an acousto-optical tuned filter. The instrument of the present invention having a radiation source, comprises one or more electromagnetic radiation sources, which provide radiation at many wavelengths, and also comprises one or more electromagnetic radiation detectors. The instrument may further comprise one or more filter or wavelength selector to remove, distinguish or select radiation of a desired wavelength, before or after detection by the detector.
CLINICAL APPLICATIONS

[72] It may be required for diabetes patients and subjects at risk for diabetes to measure their blood glucose levels regularly in an attempt to keep their blood glucose levels within an acceptable range, and to make an accurate recordation of blood-glucose levels for both personal and medical records. In one aspect of the present invention, the instrument may also comprise an alphanumeric display for displaying the measured blood-glucose concentration. The alphanumeric display of this embodiment may comprise a visual display and an audio display. The visual display may be a liquid crystal display (LCD), a plasma display panel (PDP), and a field emission display (FED) or any other functional equivalent known in the art. An audio display, capable of transmitting alphanumeric data and converting this alphanumeric data to an audio display, may be provided with an audio source comprising recorded audio clips, speech synthesizers and voice emulation algorithms or any other functional equivalent known in the art.

[73] Self-Monitoring of Blood Glucose (SMBG) is an ongoing process repeated multiple times per day for the rest of the diabetic patient's lifetime. Accurate recordation of these measurements are crucial for diagnostic purposes. A facile storage and access system for this data is also contemplated in this invention. In one aspect of the present invention, an instrument for non-invasively measuring blood-glucose concentration further comprises a microprocessor and a memory which is operatively linked to the microprocessor for storing the blood glucose measurements. The instrument of this embodiment further comprises a communications interface adapted to transmit data from the instrument to a computer system. In this embodiment the communications interface selected may include, for example, serial, parallel, universal serial bus (USB), FireWire, Ethernet, fiber optic, co-axial, twisted pair cables, a wireless communication link (e.g., WLAN, WIFI, Bluetooth, infrared) or any other functional equivalent known in the art. The communications interfaces (250, 450) may include, for example, serial, parallel, universal serial bus (USB), FireWire, Ethernet, fiber optic, co-axial, twisted pair cables, and/or a wireless communication link (e.g., WLAN, WIFI, Bluetooth, infrared).

[74] In addition to storing blood-glucose measurement data within an instrument, the present invention includes a computer system for downloading and storing these measurement data to facilitate storage and access to this information. The present invention further contemplates a computer processor, a memory which is operatively linked to the computer processor, a communications interface adapted to receive and send
data within the computer processor, and a computer program stored in the memory which executes in the computer processor. The computer program of this embodiment further comprises a database, wherein data received by the database may be sorted into predetermined fields, and the database may be capable of graphical representations of the downloaded analyte concentrations. The graphical representations of this embodiment may include, but are not limited to, column, line, bar, pie, XY scatter, area, radar, and surface representations.

[75] The computer system contemplated by the present invention should be accessible to a remote access user via an analogous communications interface for use as a diagnostic, research, or other medically related tool. Physicians, for example, could logon to the computer system via their analogous communications interface and upload a patient's blood-glucose measurements over any period of time. This information could provide a physician with an accurate record to use as a patient monitoring or diagnostic tool such as, for example, adjusting medication levels or recommending dietary changes. Other remote access users contemplated may include research institutes, clinical trial centers, specialists, nurses, hospice service providers, insurance carriers, and any other health care provider.

[76] The present invention has demonstrated that glucose can be non-invasively measured using a mid-infrared signal from an ocular element. Studies have been performed in a variety of systems, in vitro studies using glucose solutions in a gelatin matrix, and human studies including a diabetic human volunteer with varying blood glucose concentrations.

[77] All studies, including the human studies, clearly demonstrate the dose-response of blood glucose concentrations using mid-infrared measurement techniques compared to standard SMBG monitoring test strips.

EXAMPLES

[78] The following examples are provided to describe and illustrate the present invention. As such, they should not be construed to limit the scope of the invention. Those in the art will well appreciate that many other embodiments also fall within the scope of the invention, as it is described hereinabove and in the claims.
EXAMPLE 1
Experimental In-Vitro Model to Test Precision and Accuracy of the Instrument

Instrumentation

The instrument used for the mid-infrared measurements was the SOC 400 portable FTIR. The SOC 400 portable FTIR is based on an interferometer and was originally designed for the U.S. Army to detect battlefield gases. This instrument has been modified to allow measurements on in vitro models using glucose solutions in a gelatin matrix and also on human eyes. These modifications have included the installation of a filter to allow only energy in the 7 to 13 micron region to be measured and also the modification of the faceplate to permit easier placement of the instrument for human studies.

In Vitro Studies

Studies were performed to demonstrate that solutions with varying concentrations of glucose would give a mid-infrared dose-response. Hydrophilic polyethylene membranes from Millipore Corporation were saturated with glucose solutions with concentrations at 2000 mg/dl and lower. The series of curves generated in this experiment are shown in Figure 4. For this plot, the following equation was used: Absorption = -ln (sample spectrum/gold reference spectrum). When the glucose concentration is plotted against the absorption at 9.75 microns, the plot shown in Figure 5 was observed. These studies confirmed that glucose concentration can be measured in an aqueous environment in the mid-infrared wavelength range.

EXAMPLE 2
Experimental Rabbit Model

Ketamine Anesthesitized Rabbit studies

As noted in the scientific literature (Cameron et al., DIABETES TECH. THER., (1999) 1(2): 135-143), rabbits anesthetized with Ketamine experience a rapid and marked increase in blood glucose concentration, due to the release of glucose from the liver. We have confirmed this in a series of experiments and observed that the rabbit blood sugar can change from -125 mg/dl to -325 mg/dl in 60 minutes, as measured with a LNX ExpressView blood glucose meter. These experiments require a preliminary use of gas anesthesia (Isoflurane) prior to the use of Ketamine. The rabbit was immobilized such that after anesthesia, the eyeball was available for measurements with the SOC 400 portable
FTIR. Once the animal was unconscious, a drop of blood from a vein was taken and
tested on a blood glucose test strip with the LXN ExpressView blood glucose meter. Such
samples were taken every fifteen minutes throughout the study. The gas must be
discontinued in order for the Ketamine effect to fully manifest itself. The drying out of the
eye may be prevented by suturing the eyelids and using the sutures to open the eye for the
measurement and then allowing them to close after the measurement to moisten the
eyeball.

[82] The data from the rabbit study measuring glucose concentration from an ocular
element yielded the results with a regression coefficient (R squared) of 0.86, shown in
Figure 6.

EXAMPLE 3
Human Clinical Study

Human Studies

[83] Several studies were performed with non-diabetic and diabetic human volunteers.
Prior to performing these studies it was confirmed that the infrared radiation being used
poses no health hazard.

[84] Several experiments with a diabetic volunteer were performed. The subject was
asked to adjust his food intake and insulin administration in order to have his glucose
levels move from approximately 100 to 300 mg/dl over a three to four hour timeframe.
During the study, the patient took duplicate fingerstick glucose measurements and was
scanned with the SOC 400 approximately every five minutes. Prior to collecting the
infrared scan, the instrument operator aligned the SOC 400 with the subjects' eye to
attempt to collect the strongest signal being reflected off of the eye.

[85] In one study performed on the patient using the SOC 400 measuring off of the
surface of the patient's eyeball, the following correlation was observed, as shown in
Figure 7. As seen, the correlation of the signal with the glucose concentration is clear.

Human Study using the SOC 400

[86] A glucose tracking study was performed using the diffuse detector for the SOC
400. A glucose tracking study was performed with a diabetic volunteer and the results
shown in Figure 8 demonstrate that the glucose concentration changes were clearly
detected and measured using an instrument and method of the present invention. The
-25-
correlation between the measurements taken with the instrument of the present invention using the methods of the present invention is shown in Figure 9. Measurements using the instruments and methods of the present invention showed very close correlation to SMBG measurements (squares and diamonds respectively).

EXAMPLE 4
A Method Wherein a Remote Access User Can Receive a Subject's Measured Analyte Concentrations That Have Been Downloaded and Stored in a Computer System

[87] One aspect of the present invention relates to a method of downloading and storing a subject's measured analyte concentrations (Figure 3). A subject first measures the analyte concentration from a body surface such as their eye 100, whereby reflected mid-infrared radiation 150 is measured using a non-invasive instrument/monitor 200. The non-invasive instrument 200 further comprises a communications interface 250 which is capable of connecting 300 the non-invasive instrument 200 through the communications interface 250 to a computer system 400. The communications interface 250 is specifically adapted to transmit data from the instrument to the computer system 400. The computer system 400 comprises a computer processor, a computer program which executes in the computer processor, and an analogous communications interface 450. The measured analyte concentrations from the non-invasive instrument 200 are downloaded via the communications interface 250 to the computer system 400. A remote access user 500, having a computer system with an analogous communications interface 450 is capable of retrieving the downloaded measured analyte concentrations from the computer system 400. The communications interfaces 250, 450 may include, for example, serial, parallel, universal serial bus (USB), FireWire, Ethernet, fiber optic, coaxial, twisted pair cables, and/or a wireless communication link (e.g., WLAN, WIFI, Bluetooth, infrared). This information is used, for example, to provide data, warnings, advice or assistance to the patient or physician, and to track a patient's progress throughout the course of the disease.

[88] With reference to Figure 10, an embodiment of an optical, non-invasive glucose instrument/monitor 200a with depth profiling/adjustable focus to choose the best ocular element (and/or depth/layer) 430 for non-invasive measurement of the presence, absence or concentration of analytes in the ocular element of a subject will be described. The monitor 200a may be used to measure one or more of the following ocular elements 430 (and/or one or more depths/layers therein): eyelid(s), epithelial cells, the aqueous humor,
the vitreous humor, various layers of the cornea, lens, various layers of the sclera, conjunctiva, interstitial fluid in the conjunctiva, tears, the tear layer, and blood vessels.

[89] In the embodiment shown, the monitor 200a includes a movable lens 428, a mid-infrared generator 438, a sensor 432, and a controller 434. The lens 428 is longitudinally aligned with the sensor 432 and is movable in a longitudinal direction towards and away from the sensor 432. When the monitor 32 and the user's eye are located in close proximity, the mid-infrared radiation generator 438 emits mid-infrared radiation 50 from the monitor 32 and the lens 48 focuses the mid-infrared radiation 50 on the ocular element (and/or depth/layer) 430 of the eye 40. The sensor 432 receives reflected mid-infrared radiation from the ocular element (and/or depth/layer) 430 of the eye 40 through the lens 428 (alternatively, the mid-infrared radiation may be naturally emitted from the ocular element). The lens 428 focuses the reflected light onto the sensor 432. The controller 434 reads the information obtained by the sensor 432. If an ideal reading is not obtained with the ocular element (and/or depth/layer) 430, or if additional readings are desired, the controller 434 controls the movement of lens 428 to profile one or more different ocular elements (and/or depths/layers) 430. The controller 434 determines the ideal ocular element (and/or depth/layer) 430 and provides measurement information on the presence, absence or concentration of analytes (e.g., glucose concentration) to the user (or other entities, locations, devices).

[90] In alternative embodiments, the monitor 200a includes specular reflection optics for surface reflective measurements, or diffuse reflection optics for deeper ocular element (and/or depth/layer) 430 reflective measurements.

[91] In alternative embodiments, the monitor 200a may include one or more of a movable device 200a, a movable housing (or sub housing), a movable lens 428, a movable mid-infrared radiation generator 438, and/or a movable sensor 432 to adjust the focus of the monitor 200a for profiling different ocular elements (and/or depth/layers) 430.

[92] In one or more embodiments, the monitor 200a may include a polarizer to profile different ocular elements (and/or depth/layers) 430 and/or the monitor 200a may employ optics that vary the angle of optic components to profile different ocular elements (and/or depth/layers) 430.

[93] The monitor 200a is advantageous in that it can determine the ideal ocular element (and/or depth layer) 430 from which to obtain a reading for non-invasive measurement of the presence, absence or concentration of analytes in the tissue of a subject. If an ocular
element (and/or depth layer) 430 is not providing a sufficient reading, one or more additional ocular elements (and/or depth layers) 430 may be measured for determining an ideal ocular element (and/or depth layer) 430 for determining the presence, absence or concentration of analytes in the tissue of a subject. The measurements from the ideal ocular element (and/or depth layer) 430 are then provided to the user (or other entities, locations, devices).

[94] In an alternative embodiment, the monitor 200a may be configured such that the mid-infrared generator 438 interrogates the ocular region with mid-infrared radiation. The sensor 432 is configured to operate in cooperation with the mid-infrared generator 438 to receive reflected mid-infrared radiation at a certain time interval that corresponds with the desired depth level in the ocular region at which the measurement is to be taken. In one embodiment, the lens 428 may be employed (e.g., by focusing on the surface of the eye) to determine when the monitor 200a is in the correct range of proximity to the ocular region so that the emission of mid-infrared radiation and the receipt of reflected mid-infrared radiation can be optimally coordinated in time. Other techniques for determining the correct proximity of the monitor 200a to the ocular region may also be employed. For example, the mid-infrared generator may take an initial reading to gauge the distance of the monitor from the ocular surface based on the timing of the received reflected mid-infrared radiation.

[95] The above description of the disclosed embodiments is provided to enable any person skilled in the art to make or use the invention. Various modifications to these embodiments will be readily apparent to those skilled in the art, and the generic principles described herein can be applied to other embodiments without departing from the spirit or scope of the invention. Thus, it is to be understood that the description and drawings presented herein represent a presently preferred embodiment of the invention and are therefore representative of the subject matter which is broadly contemplated by the present invention. It is further understood that the scope of the present invention fully encompasses other embodiments that may become obvious to those skilled in the art and that the scope of the present invention is accordingly limited by nothing other than the appended claims.
WHAT IS CLAIMED IS:

1. A method of non-invasively measuring the presence, absence or concentration of one or more analytes in an ocular element of a subject, the subject including an eye with an ocular surface and a tear layer, comprising:
   exposing at least a portion of the tear layer of the subject to electromagnetic radiation without contact with the ocular surface;
   detecting electromagnetic radiation reflected from the tear layer without contact with the ocular surface; and
   determining a radiation signature of the reflected electromagnetic radiation to determine the presence, absence or concentration of the one or more analytes in the tear layer of the subject.

2. The method of claim 1, wherein the method further includes profiling one or more ocular elements in addition to the tear layer to determine an ideal ocular element for measuring the presence, absence or concentration of the one or more analytes.

3. The method of claim 2, wherein the ocular element is at least one of: the eyelid, epithelial cells, the aqueous humor, the vitreous humor, various layers of the cornea, lens, various layers of the sclera, conjunctiva, interstitial fluid in the conjunctiva, tears, the tear layer, and blood vessels.

4. The method of claim 1, wherein the method further includes profiling one or more different depths in the tear layer to determine an ideal depth in the tear layer for measuring at least one of the concentration, presence, and absence of the one or more analytes.

5. The method of claim 1, wherein the electromagnetic energy is infrared radiation.

6. The method of claim 1, wherein detecting includes detecting electromagnetic radiation with specular reflection optics.
7. The method of claim 1, wherein detecting includes detecting electromagnetic radiation with diffuse reflection optics.

8. A method of non-invasively measuring the presence, absence or concentration of one or more analytes in an ocular element of a subject, the subject including an eye with an ocular surface and multiple ocular elements, comprising:
   measuring the presence, absence or concentration of the one or more analytes from the eye by profiling more than one different ocular element to determine an ideal ocular element for measuring the presence, absence or concentration of the one or more analytes;
   measuring the presence, absence or concentration of one or more analytes from the ideal ocular element by
      exposing at least a portion of the ideal ocular element to electromagnetic radiation without contact with the ocular element;
      detecting electromagnetic radiation reflected from the ocular element without contact with the ocular element; and
      determining a radiation signature of the reflected electromagnetic radiation to determine the presence, absence or concentration of the one or more analytes in the ocular element of the subject.

9. The method of claim 8, wherein the ocular element is at least one of: the eyelid, the epithelial cells, the aqueous humor, the vitreous humor, various layers of the cornea, lens, various layers of the sclera, conjunctiva, interstitial fluid in the conjunctiva, tears, the tear layer, and blood vessels.

10. The method of claim 8, wherein profiling more than one different ocular element further includes profiling one or more different depths in the ocular element to determine an ideal depth in the ocular element for measuring the presence, absence or concentration of the one or more analytes.

11. The method of claim 8, wherein the electromagnetic energy is infrared radiation.
12. The method of claim 8, wherein detecting includes detecting electromagnetic radiation with specular reflection optics.

13. The method of claim 8, wherein detecting includes detecting electromagnetic radiation with diffuse reflection optics.

14. A non-invasive analyte measurement instrument for determining the presence, absence or concentration of one or more analytes in an ocular element of a subject, the subject including an eye with an ocular surface and a tear layer, comprising:

   means for exposing at least a portion of the tear layer of the subject to electromagnetic radiation without contact with the ocular surface;

   means for detecting electromagnetic radiation reflected from the tear layer without contact with the ocular surface; and

   means for determining a radiation signature of the reflected electromagnetic radiation to determine the presence, absence or concentration of the one or more analytes in the ocular element of the subject.

15. The instrument of claim 14, further including means for profiling one or more ocular elements in addition to the tear layer to determine an ideal ocular element for measuring at least one of the concentration, presence, and absence of the one or more analytes.

16. The instrument of claim 14, further including means for profiling one or more different depths in the tear layer to determine an ideal depth in the tear layer for measuring at least one of the concentration, presence, and absence of the one or more analytes.

17. The instrument of claim 14, wherein the detecting means includes specular reflection optics.

18. The instrument of claim 14, wherein the detecting means includes diffuse reflection optics.
19. A non-invasive analyte measurement instrument for determining the presence, absence or concentration of one or more analytes in an ocular element of a subject, the subject including an eye with an ocular surface and multiple ocular elements, comprising:

means for profiling more than one different ocular element to determine an ideal ocular element for measuring analyte concentration;

means for measuring the presence, absence or concentration of one or more analytes from the ideal ocular element, including

means for exposing at least a portion of the ideal ocular element to electromagnetic radiation without contact with the ocular element;

means for detecting electromagnetic radiation reflected from the ocular element without contact with the ocular element; and

means for determining a radiation signature of the reflected electromagnetic radiation to determine the presence, absence or concentration of the one or more analytes in the ocular element of the subject.

20. The instrument of claim 19, wherein the measuring means further includes means for profiling one or more different depths in the ocular element to determine an ideal depth in the ocular element for measuring analyte concentration.

20. The instrument of claim 19, wherein the detecting means includes specular reflection optics.

21. The instrument of claim 19, wherein the detecting means includes diffuse reflection optics.
FIG. 4

FIG. 5
RABBIT RESULTS AT 8.7 MICRONS WITH SPECULAR DETECTOR

FIG. 6

HUMAN STUDY AT 8.6 MICRONS WITH SPECULAR DETECTOR

FIG. 7