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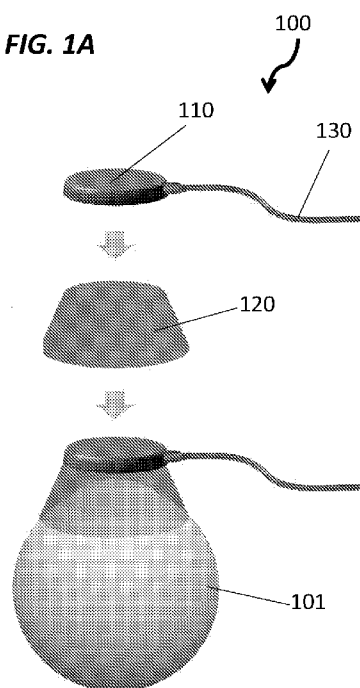
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[Continued on next page]

(54) Title: OCULAR CRANIAL NERVE MONITORING SYSTEM

FIG. 1A



(57) Abstract: An ocular cranial nerve monitoring system (OCNMS) that provides continuous measurement of pupillary reactivity in the eye. The system features a sensor component that produces a stimulating light and a recording light and records reflected recording light so as to calculate pupillary diameter and response to stimulation as measured by latency, velocity, and/or amplitude. This system of the present invention may be used for indirect assessment of the 2nd and 3rd cranial nerve (CN 2, CN3) pathways and/or intracranial pressure in patients and may be used intraoperatively. The data obtained from this system may allow for immediate corrective actions, which may help prevent permanent deficits and improve patient safety and surgical outcomes. In some instances, the system may help avoid unnecessary or invasive procedures (e.g., catheters inserted for intracranial pressure monitoring).



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OCULAR CRANIAL NERVE MONITORING SYSTEM

CROSS REFERENCE

[0001] This application claims priority to U.S. Patent Application No. 62/286,052, filed January 22, 2016, the specification(s) of which is/are incorporated herein in their entirety by reference.

FIELD OF THE INVENTION

[0002] The present invention relates to methods and systems for measuring pupillary light reflex responses (pupillometry), more particularly to a device or system featuring a solid-state camera and light source (e.g., visible, infrared) mounted onto an extraocular lens/shield for measuring pupillary light reflex responses, e.g., bilaterally, in real-time or near real-time. The system may be used, e.g., intraoperatively, to determine the integrity of one or more cranial nerves, e.g., 2nd and/or 3rd cranial nerve, as well as to infer intracranial pressure and measure ocular muscle activity.

BACKGROUND OF THE INVENTION

[0003] Electrophysiological monitoring of motor, sensory, and cranial nerve function is routine for many neurosurgical procedures. These techniques quantify changes in stimulatory current and resulting amplitude of EEG or muscular activity. With cranial nerves in particular, somatic motor nerves (cranial nerves (CNs) 12, 11, 10/9, 7, 5) and some special sensory nerves (CN 8) are readily monitored for feedback during complex surgical procedures in those respective regions. The upper cranial nerves, however, remain more difficult to monitor. Cranial nerves 3 and 4 do have motor tracts but placement of a needle lead into their respective extraocular muscles can be difficult. Assessment of the optic nerves (CN 2) remains even more problematic because the most often used modality, visual evoked potentials (VEP), measures not only nerve integrity, but also that of the optic tracts and visual cortex. In practice it has not been found to be entirely reliable. The eyes are a vital window to the function and health of the central nervous system and brain. Pupillary reflexes, extraocular motor function, and direct retinal inspection can all provide invaluable information ranging from cranial nerve, tract, and nuclei integrity to an indirect measure of intracranial pressure. Indeed, cranial nerve function has been used to assess intracranial pressure in critical ICU patients and attempts have been made to

correlate brainstem auditory evoked responses (BAERs, e.g., CN 8) and intermittent/periodic reflex pupillometry (CN 3) with intracranial pressure. The current state-of-the-art in reflex pupillometry involves hand-held devices based on 40 year-old technology. A brief light source illuminates the retina, and subsequent pupillary response (including latency, velocity, and amplitude) is recorded with an infrared light-source and camera. These measurements are intermittent and require logistically difficult manipulations including manual eyelid opening and measurement of the contralateral response. In the clinic setting with awake patients, these impediments are easily overcome; however, in the ICU or OR with critical and comatose patients or anesthetized patients, such interventions are not often feasible.

[0004] The present invention features an ocular cranial nerve monitoring system (OCNMS), e.g., an on-eye or near-eye (e.g. contact lens or shield) reflex pupillometry device that can provide continuous measurement (e.g., real-time or near real-time measurement) of the eye (a pair of systems is used for monitoring both eyes. Briefly, the system features the use of a stimulating light (e.g., white light) and a recording light (e.g., infrared light) as well as a camera design for continuously measuring pupillary diameter and response to stimulation as measured by latency, velocity, and/or amplitude, and/or any other appropriate parameters. This monitoring modality can record both ipsilateral and contralateral (consensual) pupillary responses simultaneously when the device is employed over both eyes; this is an important distinction compared to existing technology. This may allow for an indirect assessment of the 2nd and 3rd cranial nerve (CN 2, CN3) pathways and indirectly measure intracranial pressure. The system of the present invention (OCNMS) may be used intraoperatively, e.g., in anesthetized patients, or in comatose and/or neurologically compromised patients (e.g., patients in the ICU). (The system may also be able to be used on awake patients.) As such, the system can be used to provide immediate feedback concerning manipulations of particular nerves, e.g., CN 2, CN3) and/or the brainstem. The system or the present invention (OCNMS) may also be used to provide feedback concerning intracranial pressure. This feedback may allow for immediate corrective actions, which may help prevent permanent deficits and improve patient safety and surgical outcomes. In some instances, this feedback may help avoid unnecessary or invasive procedures (e.g., Ventriculostomy catheters inserted for intracranial pressure monitoring).

[0005] The present invention also features applications and methods of use of said ocular cranial nerve monitoring system. For example, in some embodiments, the OCNMS is used during anterior skull base procedures (e.g., transnasal, transcranial) so as to provide immediate feedback as particular cranial nerves (e.g., CN 2, CN3) were manipulated, potentially limiting damage by alerting the surgeon to the impending insult. In some embodiments, the OCNMS may also allow for a more reliable stimulation and measurement of visual evoked potentials (VEP) to assess the integrity of the entire visual pathway (retina to cortex).

[0006] In some embodiments, the system of the present invention is used for monitoring intracranial pressure in patients with severe traumatic brain injury (TBI) or malignant cerebral edema (or other appropriate conditions), e.g., in the ICU. For example, patients with severe TBI or malignant cerebral edema are at risk for severe disability or death from progressive intracranial hypertension. Historical observations ("blown pupil") and recent pupillometry literature suggest that subtle pupillary dysfunction (increased latency, decreased velocity) presages rising intracranial pressure. Continuously recording these parameters using the system of the present invention may allow earlier interventions (e.g., diuretics, barbiturate coma, decompressive craniectomy) before irreversible damage has occurred.

[0007] The system of the present invention may also be used as a predictor of elevated intracranial pressure, e.g., to predict the patients that would require invasive monitoring and/or intervention in the first place. This may help provide appropriate intervention in a timely and effective manner so as to help avoid poor or worsening outcomes. This may also help avoid the use of said invasive procedures (e.g. Ventriculostomy) in individuals that did not need them, thereby limiting primarily hemorrhagic and infectious risks. The system may also help function to prognosticate a patient's outcome, e.g., a traumatic brain injury patient's outcome.

SUMMARY OF THE INVENTION

[0008] The present invention features ocular cranial nerve monitoring systems for monitoring pupillary reactivity. In some embodiments, the system comprises a sensor component comprising a plurality of image sensors and a plurality of light emitters disposed on a sensor component surface. The light emitters may be adapted to emit a stimulating light and a recording light, wherein the stimulating light

has a wavelength that is effective for triggering pupil reactivity. The image sensors may be adapted to detect recording light reflected from the iris. The system may (or may not) further comprise an intermediate plate having a sensor surface connected to the sensor surface of the sensor component and an eye surface for placing atop the eye, the eye surface being curved, wherein the intermediate plate allows or promotes transmission of recording light reflected from the iris to the image sensors. The system may further comprise a computer system operatively connected to the image sensors, wherein the computer system is adapted to calculate a pupillary reactivity parameter based on recording light detected by the image sensors. In some embodiments, the sensor component is curved to accommodate the curvature of the eye. In some embodiments, the system continuously activates the sensor component so as to obtain continuous measurements of the pupillary reactivity parameter.

[0009] In some embodiments, the system is used to monitor intracranial pressure. In some embodiments, the system is used to monitor intracranial pressure intraoperatively. In some embodiments, the system is used to monitor cranial nerve (CN) 2, CN 3, or both CN 2 and CN 3 integrity or function. In some embodiments, the image sensor (112) comprises a complementary metal-oxide-semiconductor (CMOS) sensor, a wafer-layer optic (WLO) sensor, or a wafer-level packaging (WLP) sensor. In some embodiments, the light emitter, the image sensors, or both the light emitter and the image sensors comprise OLED-on-silicon technology. In some embodiments, the recording light is infrared light or ambient light or visible light. In some embodiments, the wavelength of the recording light is from 400 to 700 nm. In some embodiments, the pupillary reactivity parameter is velocity and/or amplitude and/or latency.

[0010] In some embodiments, the image sensors and light emitters are arranged on the sensor surface in an alternating pattern. In some embodiments, the alternating pattern is a checkerboard pattern, a radial pattern, or a diagonal pattern. In some embodiments, the image sensors and light emitters are arranged on the sensor surface such that they do not obstruct a line of sight extending through an apex of the sensor component. In some embodiments, the image sensors and light emitters are arranged on the sensor surface at locations that do not obstruct central vision, paracentral vision, or near peripheral vision.

[0011] In some embodiments, the intermediate plate comprises a bundle of optical fibers. In some embodiments, the intermediate plate comprises a zero-depth optical window. In some embodiments, the zero-depth optical window comprises a bundle of optical fibers. In some embodiments, the optical fibers comprise a multi-fiber plate, a fiber optic plate, a fiber optic faceplate, or a fiber optic taper. In some embodiments, the intermediate plate is hollow. In some embodiments, the intermediate plate matches curvature of an eye to allow surface tension to hold the system in place when placed on the eye. In some embodiments, the intermediate plate is disposable.

[0012] In some embodiments, the system further comprises extensions extending from one or more sides of the sensor component, wherein electrodes are disposed on ends of the extensions, the electrodes function to stimulate muscles around the eye. In some embodiments, the electrodes are operatively connected to the computer processing system. In some embodiments, the computer system is operatively connected to the sensors via a wireless mechanism. In some embodiments, the wireless mechanism comprises a radio-frequency identification (RFID) system. In some embodiments, the wireless mechanism is for collecting measurements of the pupillary reactivity. In some embodiments, the wireless mechanism is for providing power to the image sensor or the light emitters. In some embodiments, the sensor surface comprises a rigid support material. In some embodiments, the rigid support material comprises polymethyl methacrylate (PMMA), polystyrene (PS), or a combination thereof. In some embodiments, the sensor surface comprises a flexible support material. In some embodiments, the flexible support material comprises a polymer adapted for flexible electrodes. In some embodiments, the flexible support material comprises polyimide (PI), polyethylene terephthalate (PET), polyether ether ketone (PEEK), or a combination thereof. In some embodiments, the sensor surface comprises a combination of a rigid support material and a flexible support material. In some embodiments, the sensor surface matches curvature of an eye to allow surface tension to hold the system in place when placed on the eye.

[0013] The present invention also features methods of detecting pupillary reactivity. In some embodiments, the method comprises activating on a patient an ocular cranial nerve monitoring system of the present invention. In some embodiments, the

light emitters emit stimulating light and recording light to the iris and pupil, wherein the stimulating light triggers pupil reaction, the sensors detect recording light reflected from the iris, and the computer system calculates a pupillary reactivity parameter based on recording light detected by the sensors.

[0014] The present invention also features methods of monitoring of intracranial pressure. In some embodiments, the method comprises activating on a patient an ocular cranial nerve monitoring system according to the present invention. In some embodiments, the light emitters emit stimulating light and recording light to the iris and pupil, the stimulating light triggers pupil reaction, the sensors detect recording light reflected from the iris, and the computer system calculates a pupillary reactivity parameter based on recording light detected by the sensors, wherein the pupillary reactivity parameter is indicative of intracranial pressure.

[0015] The present invention also features methods of monitoring of integrity or function of cranial nerve (CN) 2, CN 3, or CN 2 and CN3. In some embodiments, the method comprises activating on a patient an ocular cranial nerve monitoring system according to the present invention. In some embodiments, the light emitters emit stimulating light and recording light to the iris and pupil, the stimulating light triggers pupil reaction, the sensors detect recording light reflected from the iris, and the computer system calculates a pupillary reactivity parameter based on recording light detected by the sensors, wherein the pupillary reactivity parameter is indicative of CN 2, CN 3, or CN 2 and CN3 integrity or function.

[0016] Any feature or combination of features described herein are included within the scope of the present invention provided that the features included in any such combination are not mutually inconsistent as will be apparent from the context, this specification, and the knowledge of one of ordinary skill in the art. Additional advantages and aspects of the present invention are apparent in the following detailed description and claims.

BRIEF DESCRIPTION OF THE DRAWINGS

[0017] FIG. 1A shows a perspective view of an embodiment of the ocular cranial nerve monitoring system (OCNMS) of the present invention. For example, the system may comprise sensor (e.g., an OLED-on-silicon sensor) that sends light and captures images. The system may further comprise an intermediate plate (e.g., an

optical fiber plate, optionally disposable) that may match the eyeball's curvature (or similar) and size to the sensor. In some embodiments, the system is assembled using adhesives, such as PVA adhesives. The system may be placed on the eyeball as shown in FIG. 1A.

[0018] FIG. 1B shows a schematic view of the sensor, e.g., an OLED-on-silicon sensor (a bottom view, e.g., as viewed from the intermediate plate or optical fiber plate). The black squares represent CMOS sensors, the white squares represent OLED emitter (e.g., visible or NIR, etc.). The pattern of sensors and emitters may be checkerboard or any other appropriate pattern not limited to those patterns disclosed herein.

[0019] FIG. 1C shows the processing of an intermediate plate (e.g., an optical fiber plate): an unprocessed optical fiber plate (flat).

[0020] FIG. 1D shows the processing of an intermediate plate (e.g., an optical fiber plate): an optical fiber plate processed to conform to the eyeball curvature or similar.

[0021] FIG. 1E shows the processing of an intermediate plate (e.g., an optical fiber plate): a processed optical fiber plate with the sensor (e.g., OLED-on-silicon sensor) attached.

[0022] FIG. 2A shows a perspective view of an embodiment of the ocular cranial nerve monitoring system (OCNMS) of the present invention. For example, the system may comprise a meniscus lens incorporating OLED-on-silicon pixels that can send light and capture images. The system may have a generally concave curvature that conforms to the shape of the eye or similar.

[0023] FIG. 2B shows a schematic view of the sensor, e.g., an OLED-on-silicon sensor (a bottom view, e.g., as viewed from the optical plate). The black squares represent CMOS sensors, and the white squares represent OLED emitter (e.g., visible or NIR, etc.). The pattern of sensors and emitters may be checkerboard or any other appropriate pattern not limited to those patterns disclosed herein.

[0024] FIG. 2C shows the processing of a lens (e.g., PMMA, etc.): an unprocessed lens.

[0025] FIG. 2D shows the processing of a lens (e.g., PMMA, etc.): a lens processed with the sensor (e.g., OLED-on-silicon sensor) attached.

[0026] FIG. 3 shows an alternative embodiment (a top view) of the system of the present invention. The system comprises extensions that may feature electrodes for contacting muscle fibers around the eye to facilitate EMG monitoring modalities.

[0027] FIG. 4A shows a perspective view of an embodiment of the ocular cranial nerve monitoring system (OCNMS) of the present invention.

[0028] FIG. 4B shows a wired image sensor (as viewed from the bottom). The black circle is a wafer-level optics (WLO) technology image sensor, and the white circle is an LED emitter (e.g., visible, NIR, etc.). The pattern of white and dark circles is shown as an example. The present invention is not limited to this configuration.

[0029] FIG. 4C shows an unprocessed lens (e.g., PMMA, etc.).

[0030] FIG. 4D shows a processed lens with embedded wafer-level optics (WLO) sensors (black) and LED emitters (white).

[0031] FIG. 5A shows a wired image sensor (as viewed from the bottom). The black circle is a wafer-level optics (WLO) technology image sensor, and the white circle is an LED emitter (e.g., visible, NIR, etc.). The pattern shown is not obstructing central vision. The pattern of white and dark circles is shown as an example. The present invention is not limited to this configuration.

[0032] FIG. 5B shows an unprocessed lens (e.g., PMMA, etc.).

[0033] FIG. 5C shows a processed lens with embedded wafer-level optics (WLO) sensors (black) and LED emitters (white).

[0034] FIG. 6A shows a wired image sensor (as viewed from the bottom). The black circle is a wafer-level optics (WLO) technology image sensor, and the white circle is an LED emitter (e.g., visible, NIR, etc.). The dotted ring is an RFID antenna. The pattern shown is not obstructing central vision. The pattern of white and dark circles is shown as an example. The present invention is not limited to this configuration.

[0035] FIG. 6B shows an unprocessed lens (e.g., PMMA, etc.).

[0036] FIG. 6C shows a processed lens with embedded wafer-level optics (WLO) sensors (black), LED emitters (white), and an antenna added (dotted).

DETAILED DESCRIPTION OF THE INVENTION

[0037] The present invention features an ocular cranial nerve monitoring system (OCNMS), e.g., an on-eye or near-eye reflex pupillometry device that can provide continuous measurement (e.g., real-time or near real-time measurement) of the eye (a pair of systems is used for monitoring both eyes). As previously discussed, the system features the use of a stimulating light (e.g., a pulse of light) and a recording light (e.g., infrared light) as well as a sensor/camera design for continuously measuring pupillary diameter and response to stimulation (e.g., as measured by

latency, velocity, and/or amplitude, and/or any other appropriate parameters). This may allow for an indirect assessment of the 2nd and 3rd cranial nerve (CN 2, CN3) pathways and monitoring of intracranial pressure.

Embodiment 1

[0038] Referring to FIG. 1A, FIG. 1B, FIG. 1C, FIG. 1D, and FIG. 1E, in some embodiments, the ocular cranial nerve monitoring system (OCNMS) (100) comprises a sensor component (110). The sensor component (110) is adapted to send a pulse of light, e.g., the stimulating light (e.g., white/visible light) and sense or capture images via a recording light (e.g., infrared light). For example, in some embodiments, the sensor component (110) comprises image sensors (112) and light emitters (114). The image sensors (112) may comprise complementary metal-oxide-semiconductor (CMOS) sensors; however, the image sensors (112) are not limited to CMOS sensors. In some embodiments, the light emitters (114) comprise organic light emitting diodes (OLEDs); however, the light emitters (114) are not limited to OLEDs. In some embodiments, the sensor component (110) comprises an OLED-on-silicon sensor. In some embodiments, the light emitters (114) can emit light of one or more wavelengths, e.g., visible, infrared (IR), near infrared (NIR), etc. The light emitters (114) and image sensors (112) may be arranged in any appropriate configuration such as but not limited to a checkerboard configuration shown in FIG. 1B. The present invention is not limited to this configuration.

[0039] As previously discussed, the light emitters (114) provide a stimulating light and a recording light. In some embodiments, the stimulating light comprises white light or any particular visible light. For example, in some embodiments, the stimulating light has a wavelength between 400 nm and 700 nm. In some embodiments, the stimulating light has a wavelength between 400 and 800 nm. In some embodiments, the stimulating light has a wavelength between 380 and 700 nm. In some embodiments, the stimulating light has a wavelength between 380 and 800 nm. In some embodiments, the recording light comprises infrared light. For example, in some embodiments, the recording light has a wavelength between 700 nm and 15 μ m. In some embodiments, the recording light has a wavelength between 800 nm and 15 μ m. In some embodiments, the recording light has a wavelength between 700 nm and 1400 nm. In some embodiments, the recording light has a wavelength between 750 nm and 1400 nm. In some embodiments, the recording

light has a wavelength between 1400 nm and 3 μ m. In some embodiments, the recording light has a wavelength between 3 μ m and 8 μ m. In some embodiments, the recording light has a wavelength between 8 μ m and 15 μ m. In some embodiments, the recording light has a wavelength between 15 μ m and 30 μ m. In some embodiments, the recording light is visible light.

[0040] The sensor component (110) may provide the stimulating light, which triggers pupil reaction. The sensor component (110) may simultaneously function as a camera or sensor for detecting the recording light (e.g., IR light). The sensor component (110) may be adapted to measure pupil area or diameter, and may detect the velocity and latency (e.g., lag or wait time). For example, the IR light that bounces off the eye may be detected, whereas the IR light that continues through the pupil may not be detected. Based on what is detected, the system may calculate the appropriate pupil reactivity parameters. The system may measure this pupillary response simultaneously in both pupils (e.g., ipsilateral and consensual) if the monitoring devices are employed over both eyes. As previously discussed, this is an important distinction compared with existing technologies.

[0041] The system (100) of the present invention may feature components or systems that can filter out or differentiate between recording light that is reflected off the retina from that which is reflected from the iris. The system (100) may further comprise an intermediate plate (120) for adapting the sensor component (110) to the eye (101). The intermediate plate (120) provides a means of adapting the sensor component (110) to the curvatures and/or sizes of the eye. The intermediate plate (120) can also help to block any interfering light from accidentally interfering with the sensor component (110), e.g., the intermediate plate (120) provides a dark environment for the light pulses and recordings.

[0042] The intermediate plate (120) has a sensor surface that attaches or connects to the sensor component (110). The intermediate plate also has an eye surface (a curved surface) that is for placing atop the eye. In some embodiments, the intermediate plate comprises optical fibers (e.g., a bundle of optical fibers) that transmit light, e.g., the intermediate plate is an optical fiber plate. In some embodiments, the intermediate plate is hollow. The intermediate plate (120) is not limited to the aforementioned configurations and may comprise any appropriate

components that allow for the image capturing of the eye via the sensor component (110). In some embodiments, the optical fibers may be narrowed (or tapered) so as to change the size of the field or image. In some embodiments, the intermediate plate (120) is disposable. In some embodiments, the sensor surface of the intermediate plate (120) is attached to the sensor component (110) via any appropriate adhesive or lubricant. In some embodiments, the intermediate plate (120) matches the curvature of the eye to allow surface tension to hold the system in place when placed on the eye, e.g., like a contact lens.

[0043] In some embodiments, the system (100), e.g., the sensor component (110), is operatively connected to a computer processing system and/or a power source via a wire (130) (e.g., for cost-effectiveness). In some embodiments, the system (100), e.g., the sensor component (110), is wirelessly operatively connected to a computer processing system. In some embodiments, the sensor component (110) comprises a battery system. In some embodiments, the sensor component (110) functions as a camera. In some embodiments, the sensor component (110) is operatively connected to a camera. In some embodiments, the sensor component (110) and/or camera is operatively connected to a computer system that is adapted to measure and records the pupillary constriction (latency, velocity, amplitude) and process the parameters appropriately to provide information usable to surgeons or other medical professionals.

[0044] Because the system of the present invention is adapted to measure pupillary reactivity on a continuous basis, the data obtained from the system of the present invention may be used to show trends wherein a normal reactivity changes to an abnormal reactivity. This may provide an indicator to the surgeon or medical professional that action should be taken, e.g., action related to the cranial nerve affected, action related to intracranial pressure, etc.

Embodiment 2

[0045] Referring to FIG. 2A, FIG. 2B, FIG. 2C, and FIG. 2D, in some embodiments, the ocular cranial nerve monitoring system (OCNMS) (200) comprises a sensor component (210) that is curved and adapted to contact the eye (101). For example, the system (200) may comprise a meniscus lens incorporating OLED-on-silicon pixels that can send light and capture images. The system (200) may have a

generally concave curvature that conforms to the shape of the eye or similar. In some embodiments, the system (200), e.g., the sensor component (210) or other appropriate component, matches the curvature of the eye to allow surface tension to hold the system in place when placed on the eye (e.g., like a contact lens). In some embodiments, the system (200) comprises multiple chips and wires, creating an array configuration so as to accommodate the light emitters and sensors into a curved lens-like configuration.

[0046] As previously discussed, the sensor component (210) is adapted to send a pulse of light, e.g., the stimulating light (e.g., white/visible light) and sense or capture images via a recording light (e.g., infrared light). For example, in some embodiments, the sensor component (210) comprises image sensors (212) and light emitters (214). The image sensors (212) may comprise complementary metal-oxide-semiconductor (CMOS) sensors; however, the image sensors (212) are not limited to CMOS sensors. In some embodiments, the light emitters (214) comprise organic light emitting diodes (OLEDs); however, the light emitters (214) are not limited to OLEDs. In some embodiments, the sensor component (210) comprises an OLED-on-silicon sensor. In some embodiments, the light emitters (214) can emit light of one or more wavelengths, e.g., visible, infrared (IR), near infrared (NIR), etc. The light emitters (214) and image sensors (212) may be arranged in any appropriate configuration such as a checkerboard configuration shown in FIG. 1B. The present invention is not limited to this configuration.

[0047] As previously discussed, the light emitters provide a stimulating light and a recording light. In some embodiments, the stimulating light comprises white light or any particular visible light. For example, in some embodiments, the stimulating light has a wavelength between 400 nm and 700 nm. In some embodiments, the stimulating light has a wavelength between 400 and 800 nm. In some embodiments, the stimulating light has a wavelength between 380 and 700 nm. In some embodiments, the stimulating light has a wavelength between 380 and 800 nm. In some embodiments, the recording light comprises infrared light. In some embodiments, the recording light comprises visible light, e.g., 400 to 700 nm or similar. In some embodiments, the recording light comprises ambient light (e.g., a camera may be able to capture images with ambient light, e.g., without any added light). In some embodiments, the recording light has a wavelength between 700 nm

and 1 mm. In some embodiments, the recording light has a wavelength between 800 nm and 1 mm. In some embodiments, the recording light has a wavelength between 700 nm and 1400 nm. In some embodiments, the recording light has a wavelength between 750 nm and 1400 nm. In some embodiments, the recording light has a wavelength between 1400 nm and 3 μ m. In some embodiments, the recording light has a wavelength between 3 μ m and 8 μ m. In some embodiments, the recording light has a wavelength between 8 μ m and 15 μ m. In some embodiments, the recording light has a wavelength between 15 μ m and 1 mm.

[0048] The sensor component (210) may provide the stimulating light, which triggers pupil reaction. The sensor component (210) may simultaneously function as a camera or sensor for detecting the recording light (e.g., IR light). The sensor component (210) may be adapted to measure pupil area or diameter, and may detect the velocity and latency (e.g., lag or wait time). For example, the IR light that bounces off the eye may be detected, whereas the IR light that continues through the pupil may not be detected. Based on what is detected, the system may calculate the appropriate pupil reactivity parameters. The system (200) of the present invention may feature components or systems that can filter out or differentiate between recording light that is reflected off the retina from that which is reflected from the iris.

[0049] As previously discussed, in some embodiments, the system (200), e.g., the sensor component (210), is operatively connected to a computer processing system and/or a power source via a wire (230) (e.g., for cost-effectiveness). In some embodiments, the system (200), e.g., the sensor component (210), is wirelessly operatively connected to a computer processing system. In some embodiments, the sensor component (210) comprises a battery system. As previously discussed, in some embodiments, the sensor component (210) functions as a camera. In some embodiments, the sensor component (210) is operatively connected to a camera. In some embodiments, the sensor component (210) and/or camera is operatively connected to a computer system that is adapted to measure and records the pupillary constriction (latency, velocity, amplitude) and process the parameters appropriately to provide information usable to surgeons or other medical professionals.

[0050] Because the system of the present invention is adapted to measure pupillary reactivity on a continuous basis, the data obtained from the system of the present invention may be used to show trends wherein a normal reactivity changes to an abnormal reactivity. This may provide an indicator to the surgeon or medical professional that action should be taken, e.g., action related to the cranial nerve affected, action related to intracranial pressure, etc.

Methods and Applications for Use

[0051] As previously discussed, the ocular cranial nerve monitoring system (OCNMS) of the present invention may be used intraoperatively, e.g., in anesthetized patients, or in comatose and/or neurologically compromised patients (e.g., patients in the ICU). (The system may also be able to be used on awake patients.) The system may be used to provide feedback regarding manipulations of particular nerves, e.g., CN 2, CN 3, etc., and/or intracranial pressure. As such, the present invention features methods of monitoring CN 2 and/or CN 3 and/or brainstem integrity and/or function. The present invention also features methods of monitoring intracranial pressure. The feedback obtained from the system of the present invention may allow for immediate corrective actions, which may help prevent permanent deficits and improve patient safety and surgical outcomes. In some instances, this feedback may help avoid unnecessary or invasive procedures (e.g., catheters inserted for intracranial pressure monitoring).

[0052] More specifically, in some embodiments, the system is used during anterior skull base procedures (e.g., transnasal, transcranial) so as to provide immediate feedback as particular cranial nerves (e.g., CN 2, CN3) were manipulated, potentially limiting damage by alerting the surgeon to the impending insult. In some embodiments, the OCNMS may also allow for a more reliable stimulation and measurement of visual evoked potentials (VEP) to assess the integrity of the entire visual pathway (retina to cortex).

[0053] In some embodiments, the system of the present invention is used for monitoring intracranial pressure in patients with severe traumatic brain injury (TBI) or malignant cerebral edema (or other appropriate conditions), e.g., in the ICU. For example, patients with severe TBI or malignant cerebral edema are at risk for severe disability or death from progressive intracranial hypertension. Historical observations

("blown pupil") and recent pupillometry literature suggest that subtle pupillary dysfunction (increased latency, decreased velocity) presages rising intracranial pressure. Continuously recording these parameters using the system of the present invention may allow earlier interventions (e.g., diuretics, barbiturate coma, decompressive craniectomy) before irreversible damage has occurred.

[0054] The system of the present invention may also be used as a predictor of elevated intracranial pressure, e.g., to predict the patients that would require invasive monitoring and/or intervention. This may help provide appropriate intervention in a timely and effective manner so as to help avoid poor or worsening outcomes. This may also help avoid the use of said invasive procedures in individuals that did not need them. The system of the present invention may also be used in other applications, for example in epilepsy or seizure monitoring applications. In some embodiments, the system of the present invention may be used for multiple sclerosis applications. In some embodiments, the system may be used to monitor any condition wherein a change in reactivity of the pupils is indicative of a particular outcome.

Additional Features

[0055] The system of the present invention is not limited to the configurations shown and described herein. In some embodiments, the system further comprises an irrigation port to help moisten or lubricate the cornea. In some embodiments, the system may also be fitted with additional neuro-physiologic monitoring modalities such as the capability to display light patterns for use with visual evoked potential (VEP) monitoring. In some embodiments, the system comprises additional electrodes around the periphery, which might also allow sensing and stimulation of electrical current in the extraocular muscles so that cranial nerves 3, 4, and 6 might also be assessed for motor integrity. For example, referring to FIG. 3, in some embodiments, the ocular cranial nerve monitoring system (OCNMS) (300) comprises extensions (320) extending from one or more sides of the sensor component (310). The extensions (320) may comprise electrodes (340) that can contact muscle fibers around the eye. The electrodes (340) may be operatively connected to the sensor component (310) and/or a computer processing system. The electrodes (340) may allow for monitoring CN 4 and/or CN6 integrity and/or function.

Additional Embodiments

[0056] Referring to FIG. 4A, FIG. 4B, FIG. 4C, and FIG. 4D, in some embodiments, the ocular cranial nerve monitoring system (OCNMS) (400) comprises a sensor component (410) that is curved and adapted to contact the eye (101). For example, the system (400) may comprise a meniscus lens incorporating components that can send light and capture images. The system (400) may have a generally concave curvature that conforms to the shape of the eye or similar. In some embodiments, the system (400), e.g., the sensor component (410) or other appropriate component, matches the curvature of the eye to allow surface tension to hold the system in place when placed on the eye (e.g., like a contact lens). In some embodiments, the system (400) comprises multiple chips and wires, creating an array configuration so as to accommodate the light emitters and sensors into a curved lens-like configuration.

[0057] As previously discussed, the sensor component (410) is adapted to send a pulse of light, e.g., the stimulating light (e.g., white/visible light) and sense or capture images via a recording light (e.g., infrared light). For example, in some embodiments, the sensor component (410) comprises image sensors (412) and light emitters (414). The image sensors (412) may comprise wafer-level optics (WLO) technology sensors; however, the image sensors (412) are not limited to WLO sensors. In some embodiments, the light emitters (414) comprise light emitting diodes (LEDs); however, the light emitters (414) are not limited to LEDs. The light emitters (414) and image sensors (412) may be arranged in any appropriate configuration. The sensor component (410) may provide the stimulating light, which triggers pupil reaction. The sensor component (410) may simultaneously function as a camera or sensor for detecting the recording light (e.g., IR light). The sensor component (410) may be adapted to measure pupil area or diameter, and may detect the velocity and latency (e.g., lag or wait time). For example, the IR light that bounces off the eye may be detected, whereas the IR light that continues through the pupil may not be detected. Based on what is detected, the system may calculate the appropriate pupil reactivity parameters. The system (400) of the present invention may feature components or systems that can filter out or differentiate between recording light that is reflected off the retina from that which is reflected from the iris.

[0058] As previously discussed, in some embodiments, the system (400), e.g., the

sensor component (410), is operatively connected to a computer processing system and/or a power source via a wire (430) (e.g., for cost-effectiveness). In some embodiments, the system (400), e.g., the sensor component (410), is wirelessly operatively connected to a computer processing system. In some embodiments, the sensor component (410) comprises a battery system. As previously discussed, in some embodiments, the sensor component (410) functions as a camera. In some embodiments, the sensor component (410) is operatively connected to a camera. In some embodiments, the sensor component (410) and/or camera is operatively connected to a computer system that is adapted to measure and records the pupillary constriction (latency, velocity, amplitude) and process the parameters appropriately to provide information usable to surgeons or other medical professionals.

[0059] Referring to FIG. 5A, FIG. 5B, and FIG. 5C, in some embodiments, the sensor component (510) comprises image sensors (512) and light emitters (514). The image sensors (512) may comprise wafer-level optics (WLO) technology sensors; however, the image sensors (512) are not limited to WLO sensors. In some embodiments, the light emitters (514) comprise light emitting diodes (LEDs); however, the light emitters (514) are not limited to LEDs. The light emitters (514) and image sensors (512) may be arranged in any appropriate configuration. Note the configuration shown in FIG. 5A, FIG. 5B, and FIG. 5C is not obstructing central vision (vertical dotted line shown in FIG. 5C). As previously discussed, in some embodiments, the system, e.g., the sensor component (510), is operatively connected to a computer processing system and/or a power source via a wire (530) (e.g., for cost-effectiveness). In some embodiments, the system, e.g., the sensor component (510), is wirelessly operatively connected to a computer processing system.

[0060] Referring to FIG. 6A, FIG. 6B, and FIG. 6C, in some embodiments, the sensor component (610) comprises image sensors (612), light emitters (614), and a radio-frequency identification (RFID) antenna (616) for wireless connectivity. The image sensors (612) may comprise wafer-level optics (WLO) technology sensors; however, the image sensors (612) are not limited to WLO sensors. In some embodiments, the light emitters (614) comprise light emitting diodes (LEDs); however, the light emitters (614) are not limited to LEDs. The light emitters (614) and

image sensors (612) may be arranged in any appropriate configuration. Note the configuration shown in FIG. 6A, FIG. 6B, and FIG. 6C is not obstructing central vision (vertical dotted line in FIG. 6C).

[0061] As previously discussed, the system, e.g., the sensor component, etc., may be operatively connected to a computer system, e.g., via a wireless mechanism. In some embodiments, the wireless mechanism comprises a radio-frequency identification (RFID) system, e.g., an RFID antenna (616). In some embodiments, the wireless mechanism is for collecting measurements of the pupillary reactivity. In some embodiments, the wireless mechanism is for providing power to the image sensor or the light emitters. As previously discussed, the image sensors and light emitters may be arranged on the sensor surface such that they do not obstruct a central line of sight, e.g., a line of sight extending through an apex of the sensor component. In some embodiments, the image sensors and light emitters may be arranged on the sensor surface such that they do not obscure central, paracentral, or near peripheral vision. In some embodiments, wireless connectivity and/or sensor and light emitter configurations that do not obstruct vision may be used to allow the system of the present invention to be used in a patient that is awake. For example, the sensor and light emitter configuration may provide minimal obstruction to vision. The wireless connectivity may allow for the patient to be mobile while using the system. The use of wafer-level optic sensors and/or wafer-level packaging sensors may provide for a low-cost and/or very small technology.

[0062] The disclosures of the following documents are incorporated in their entirety by reference herein: U.S. Pat. No. 5,297,554; WO 1990/012534; Chateau et al., 1996, *Optom Vis Sci* 73(12):733-41; Fison et al., 1979, *Br J Ophthal* 63:195-199; Ellis CJK, 1981, *Br J Ophthal* 65:754-759; Sautter et al, 1991, *Brain Res* 565:23-33; Klöcker N et al., 2001, *Journal of Neuroscience Methods* 110:147-153; Gellrich et al., 2002, *Br J Ophthalmol* 86:233-237; Bergamin and Kardon, 2003, *Invest Ophthalmol Vis Sci* 44:1546-1554; Taylor et al., 2003, *J Neurosurg* 98(1):205-213; Wilhelm and Wilhelm B, 2003, *J Neuro-Ophthalmol* 23:42-49; Bremner FD, 2004, *Eye* 18:1175-1181; Lee et al., 2005, *Korean Journal of Ophthalmology* 19(2):149-152; Swanson et al., 2005, *Invest Ophthalmol Vis Sci* 46:3737-3741; Thompson HS, 2005, *J Neuro-Ophthalmol* 25(1):44-49; Fountas et al., 2006, *Neurocrit. Care* 5:55-60; Franzen et al., 2009, *Biol Psychol* 80(3):300-305; Kim SJ, Kim YJ, Park KH:

Eye 23:727–733, 2009; Papageorgiou et al, 2009, J Neuro-Ophthalmol 29:33–36; Wargha et al., 2009, Vision Research 49:295–300; Yan et al., 2009, Liver Transplantation 15:1718-1727; Ma et al., 2010, Neuroscience Letters 476:3–8; Chen et al., 2011, Surg Neurol Int 2:82; You et al., 2012, PLoS ONE 7(12):e52061; Ortube et al, 2013, Investigative Ophthalmology & Visual Science 54(1):9-18; Zhu et al, 2013, PLoS ONE 8(7):e68935; Chen et al, 2014, SpringerPlus 3:548; Koehler and Wijdicks, 2015, J Neurosurg 122(2):453-463.

[0063] Various modifications of the invention, in addition to those described herein, will be apparent to those skilled in the art from the foregoing description. Such modifications are also intended to fall within the scope of the appended claims. Each reference cited in the present application is incorporated herein by reference in its entirety.

[0064] Although there has been shown and described the preferred embodiment of the present invention, it will be readily apparent to those skilled in the art that modifications may be made thereto which do not exceed the scope of the appended claims. Therefore, the scope of the invention is only to be limited by the following claims. Reference numbers recited in the claims are exemplary and for ease of review by the patent office only, and are not limiting in any way. In some embodiments, the figures presented in this patent application are drawn to scale, including the angles, ratios of dimensions, etc. In some embodiments, the figures are representative only and the claims are not limited by the dimensions of the figures. In some embodiments, descriptions of the inventions described herein using the phrase “comprising” includes embodiments that could be described as “consisting of”, and as such the written description requirement for claiming one or more embodiments of the present invention using the phrase “consisting of” is met.

[0065] The reference numbers recited in the below claims are solely for ease of examination of this patent application, and are exemplary, and are not intended in any way to limit the scope of the claims to the particular features having the corresponding reference numbers in the drawings.

WHAT IS CLAIMED IS:

1. An ocular cranial nerve monitoring system (100) for monitoring pupillary reactivity, said system (100) comprising:
 - a. sensor component (110) comprising a plurality of image sensors (112) and a plurality of light emitters (114) disposed on a sensor component surface, the light emitters (114) are adapted to emit a stimulating light and a recording light, wherein the stimulating light has a wavelength that is effective for triggering pupil reactivity, the image sensors (112) are adapted to detect recording light reflected from the iris;
 - b. an intermediate plate (120) having a sensor surface connected to the sensor surface of the sensor component (110) and an eye surface for placing atop the eye, the eye surface being curved, wherein the intermediate plate allows or promotes transmission of recording light reflected from the iris to the image sensors (112); and
 - c. a computer system operatively connected to the image sensors (112), the computer system is adapted to calculate a pupillary reactivity parameter based on recording light detected by the image sensors; wherein the system continuously activates the sensor component (110) so as to obtain continuous measurements of the pupillary reactivity parameter.
2. The system (100) of claim 1, wherein the system is used to monitor intracranial pressure.
3. The system (100) of claim 2, wherein the system is used to monitor intracranial pressure intraoperatively.
4. The system (100) of claim 1, wherein the system is used to monitor cranial nerve (CN) 2, CN 3, or both CN 2 and CN 3 integrity or function.
5. The system (100) of claim 1, wherein the image sensor (112) comprises a complementary metal-oxide-semiconductor (CMOS) sensor, a wafer-layer optic (WLO) sensor, or a wafer-level packaging (WLP) sensor.
6. The system (100) of claim 1, wherein the light emitter (114) comprises an organic light emitting diode (OLED).
7. The system of claim 1, wherein the light emitter (114), the image sensors (112), or both the light emitter (114) and the image sensors (112) comprise OLED-on-silicon technology.

8. The system (100) of claim 1, wherein the wavelength of the stimulating light is between 380 nm and 700 nm.
9. The system (100) of claim 1, wherein the recording light is infrared light.
10. The system (100) of claim 1, wherein the recording light is ambient light or visible light.
11. The system (100) of claim 1, wherein the wavelength of the recording light is from 400 to 700 nm.
12. The system (100) of claim 1, wherein the pupillary reactivity parameter is velocity.
13. The system (100) of claim 1, wherein the pupillary reactivity parameter is amplitude.
14. The system (100) of claim 1, wherein the pupillary reactivity parameter is latency.
15. The system (100) of claim 1, wherein the image sensors (112) and light emitters (114) are arranged on the sensor surface in an alternating pattern.
16. The system (100) of claim 15, wherein the alternating pattern is a checkerboard pattern, a radial pattern, or a diagonal pattern.
17. The system (100) of claim 1, wherein the image sensors (112) and light emitters (114) are arranged on the sensor surface such that they do not obstruct a line of sight extending through an apex of the sensor component.
18. The system (200) of claim 1, wherein the image sensors (212) and light emitters (214) are arranged on the sensor surface at locations that do not obstruct central vision, paracentral vision, or near peripheral vision.
19. The system (100) of claim 1, wherein the intermediate plate (120) comprises a bundle of optical fibers.
20. The system (100) of claim 1, wherein the intermediate plate (120) comprises a zero-depth optical window.
21. The system of claim 20, wherein the zero-depth optical window comprises a bundle of optical fibers.
22. The system of claim 21, wherein the optical fibers comprise a multi-fiber plate, a fiber optic plate, a fiber optic faceplate, or a fiber optic taper.
23. The system (100) of claim 1, wherein the intermediate plate (120) is hollow.
24. The system (100) of claim 1, wherein the intermediate plate (120) matches curvature of an eye to allow surface tension to hold the system in place when

placed on the eye.

25. The system (100) of claim 1, wherein the intermediate plate (120) is disposable.
26. The system (100) of claim 1 further comprising extensions (320) extending from one or more sides of the sensor component, wherein electrodes (340) are disposed on ends of the extensions (320), the electrodes (340) function to stimulate muscles around the eye.
27. The system (100) of claim 26, wherein the electrodes (340) are operatively connected to the computer processing system.
28. The system (100) of claim 26, wherein the system is used for monitoring CN 4, CN 6, or both CN 4 and CN6 integrity or function.
29. The system of claim 1, wherein the computer system is operatively connected to the sensors via a wireless mechanism.
30. The system of claim 29, wherein the wireless mechanism comprises a radio-frequency identification (RFID) system.
31. The system of claim 29, wherein the wireless mechanism is for collecting measurements of the pupillary reactivity.
32. The system of claim 29, wherein the wireless mechanism is for providing power to the image sensor (112) or the light emitters (114).
33. An ocular cranial nerve monitoring system (200) for monitoring pupillary reactivity, said system (200) comprising:
 - a. sensor component (210) comprising a plurality of image sensors (212) and a plurality of light emitters (214) disposed on a sensor surface, the sensor component (210) is curved to accommodate the curvature of the eye, wherein the light emitters (214) are adapted to emit a stimulating light and a recording light, the stimulating light has a wavelength that is effective for triggering pupil reactivity, wherein the image sensors (212) are adapted to detect recording light reflected from the iris; and
 - b. a computer system operatively connected to the image sensors (212), the computer system is adapted to calculate a pupillary reactivity parameter based on recording light detected by the image sensors; wherein the system continuously activates the sensor component (210) so as to obtain continuous measurements of the pupillary reactivity

parameter.

34. The system (200) of claim 33, wherein the system is used to monitor intracranial pressure.
35. The system (200) of claim 34, wherein the system is used to monitor intracranial pressure intraoperatively.
36. The system (200) of claim 33, wherein the system is used to monitor cranial nerve (CN) 2, CN 3, or both CN 2 and CN 3 integrity or function.
37. The system (200) of claim 33, wherein the image sensor (212) comprises a complementary metal-oxide-semiconductor (CMOS) sensor, a wafer-layer optic (WLO) sensor, or a wafer-level packaging (WLP) sensor.
38. The system (200) of claim 33, wherein the light emitter (214) comprises an organic light emitting diode (OLED).
39. The system (200) of claim 33, wherein the sensor surface comprises a rigid support material.
40. The system (200) of claim 39, wherein the rigid support material comprises polymethyl methacrylate (PMMA), polystyrene (PS), or a combination thereof.
41. The system (200) of claim 33, wherein the sensor surface comprises a flexible support material.
42. The system (200) of claim 41, wherein the flexible support material comprises a polymer adapted for flexible electrodes.
43. The system (200) of claim 41, wherein the flexible support material comprises polyimide (PI), polyethylene terephthalate (PET), polyether ether ketone (PEEK), or a combination thereof.
44. The system (200) of claim 33, wherein the sensor surface comprises a combination of a rigid support material and a flexible support material.
45. The system (200) of claim 33, wherein the sensor surface matches curvature of an eye to allow surface tension to hold the system in place when placed on the eye.
46. The system (200) of claim 33, wherein the wavelength of the stimulating light is between 380 nm and 700 nm.
47. The system (200) of claim 33, wherein the recording light is infrared light.
48. The system (200) of claim 33, wherein the recording light is visible or ambient light.
49. The system (200) of claim 33, wherein the pupillary reactivity parameter is

velocity.

50. The system (200) of claim 33, wherein the pupillary reactivity parameter is amplitude.
51. The system (200) of claim 33, wherein the pupillary reactivity parameter is latency.
52. The system (200) of claim 33, wherein the image sensors (212) and light emitters (214) are arranged on the sensor surface in an alternating pattern.
53. The system (200) of claim 52, wherein the alternating pattern is a checkerboard pattern, a radial pattern, or a diagonal pattern.
54. The system (200) of claim 33, wherein the image sensors (212) and light emitters (214) are arranged on the sensor surface such that they do not obstruct a line of sight extending through an apex of the sensor component.
55. The system (200) of claim 33, wherein the image sensors (212) and light emitters (214) are arranged on the sensor surface at locations that do not obstruct central vision, paracentral vision, or near peripheral vision.
56. The system (200) of claim 33 further comprising extensions (320) extending from one or more sides of the sensor component, wherein electrodes (340) are disposed on ends of the extensions (320), the electrodes (340) function to stimulate muscles around the eye.
57. The system (200) of claim 56, wherein the electrodes (340) are operatively connected to the computer processing system.
58. The system (200) of claim 56, wherein the system is used for monitoring CN 4, CN 6, or both CN 4 and CN6 integrity or function.
59. The system of claim 1, wherein the computer system is operatively connected to the sensors via a wireless mechanism.
60. The system of claim 59, wherein the wireless mechanism comprises a radio-frequency identification (RFID) system.
61. The system of claim 59, wherein the wireless mechanism is for collecting measurements of the pupillary reactivity.
62. The system of claim 59, wherein the wireless mechanism is for providing power to the image sensor (212) or the light emitters (214).
63. A method of detecting pupillary reactivity, said method comprising:
activating on a patient an ocular cranial nerve monitoring system according to any of claims 1-62, wherein the light emitters emit stimulating light and

recording light to the iris and pupil, the stimulating light triggers pupil reaction, the sensors detect recording light reflected from the iris, and the computer system calculates a pupillary reactivity parameter based on recording light detected by the sensors.

64. A method of monitoring of intracranial pressure, said method comprising: activating on a patient an ocular cranial nerve monitoring system according to any of claims 1-62, wherein the light emitters emit stimulating light and recording light to the iris and pupil, the stimulating light triggers pupil reaction, the sensors detect recording light reflected from the iris, and the computer system calculates a pupillary reactivity parameter based on recording light detected by the sensors, wherein the pupillary reactivity parameter is indicative of intracranial pressure.
65. A method of monitoring of integrity or function of cranial nerve (CN) 2, CN 2, or CN 2 and CN3, said method comprising: activating on a patient an ocular cranial nerve monitoring system according to any of claims 1-62, wherein the light emitters emit stimulating light and recording light to the iris and pupil, the stimulating light triggers pupil reaction, the sensors detect recording light reflected from the iris, and the computer system calculates a pupillary reactivity parameter based on recording light detected by the sensors, wherein the pupillary reactivity parameter is indicative of CN 2, CN 3, or CN 2 and CN3 integrity or function.
66. The method of any of claims 63-65, wherein the method is performed on an anesthetized patient.
67. The method of any of claims 63-65, wherein the method is performed intraoperatively.
68. The method of any of claims 63-65, wherein the pupillary reactivity parameter is amplitude.
69. The method of any of claims 63-65, wherein the pupillary reactivity parameter is velocity.
70. The method of any of claims 63-65, wherein the pupillary reactivity parameter is latency.

FIG. 1A

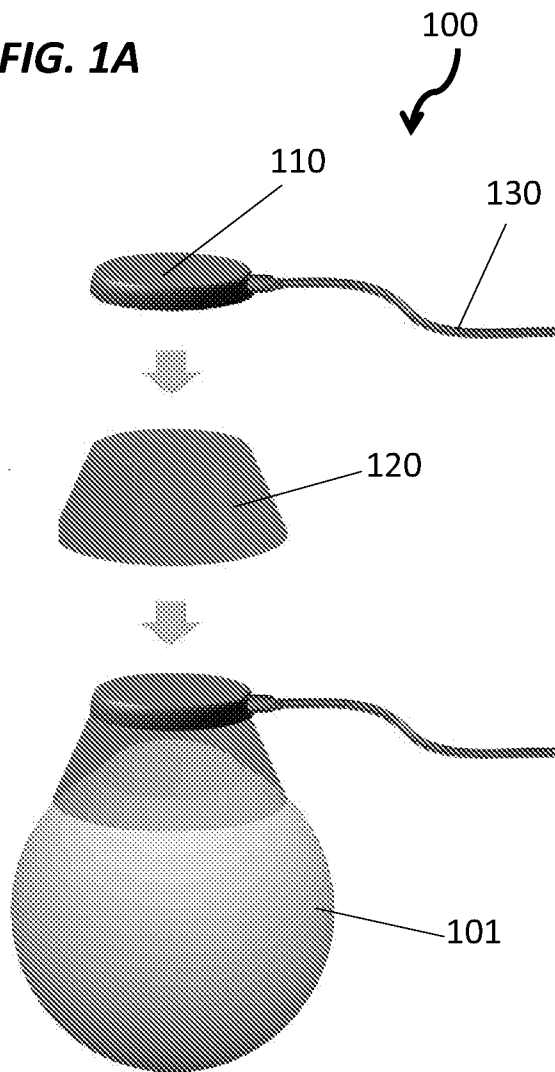


FIG. 1B

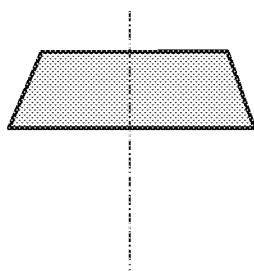
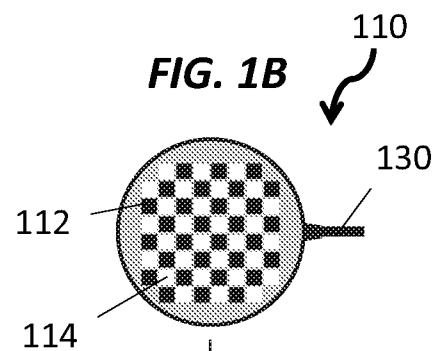


FIG. 1C

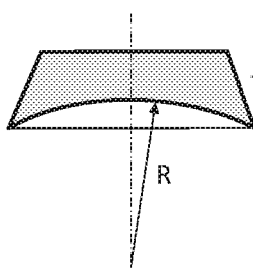


FIG. 1D

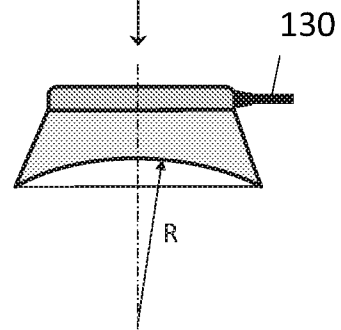


FIG. 1E

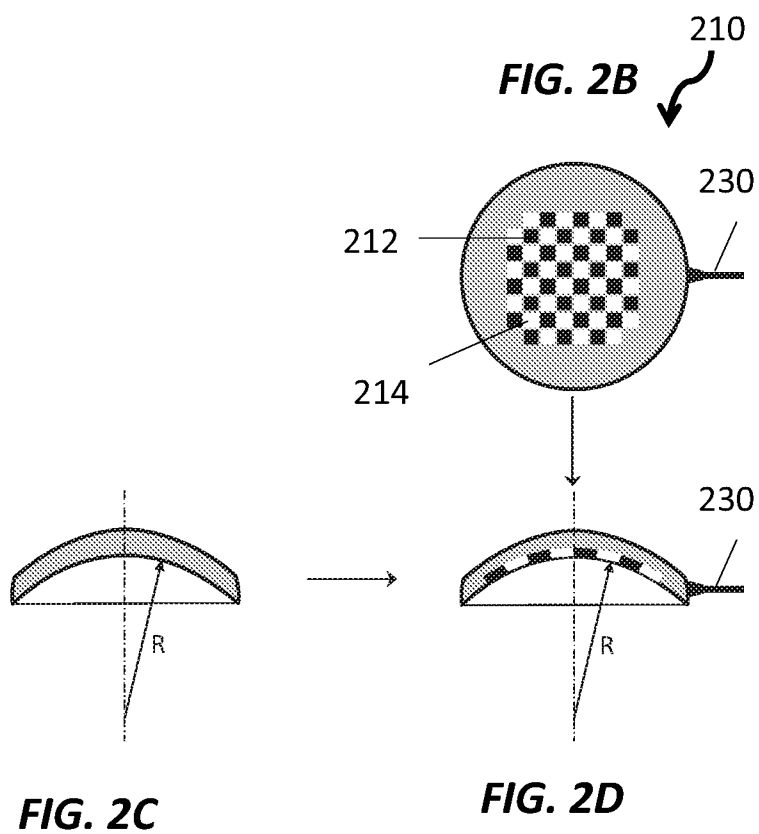
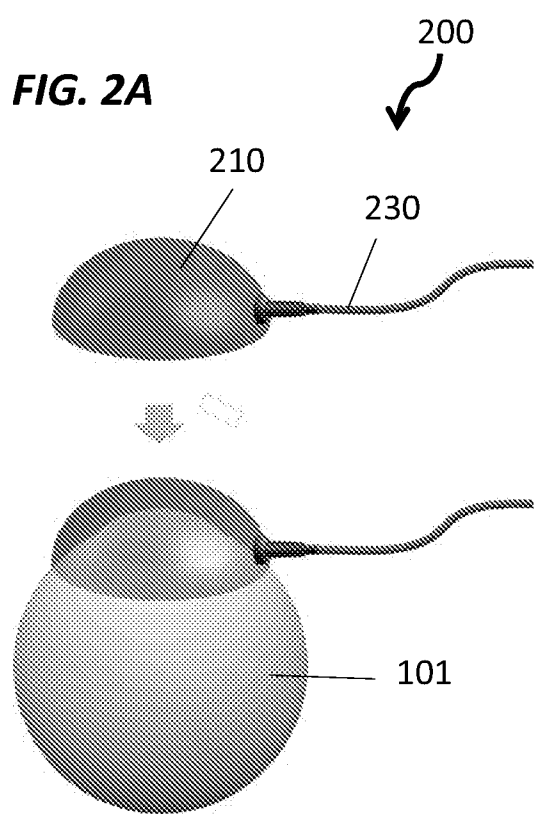


FIG. 3

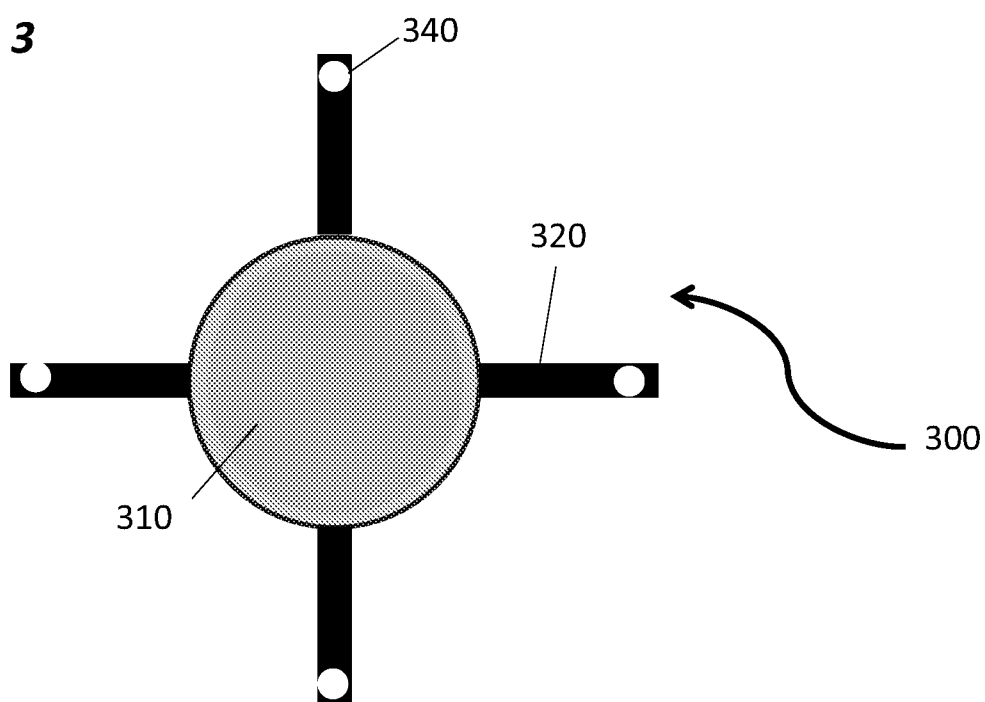


FIG. 4A

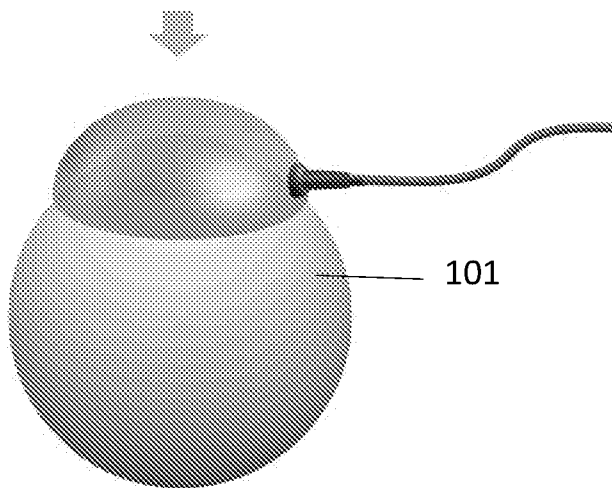
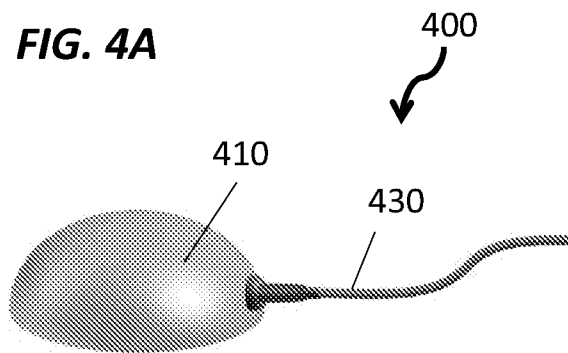


FIG. 4B

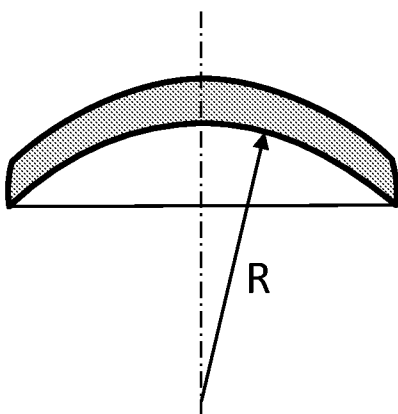
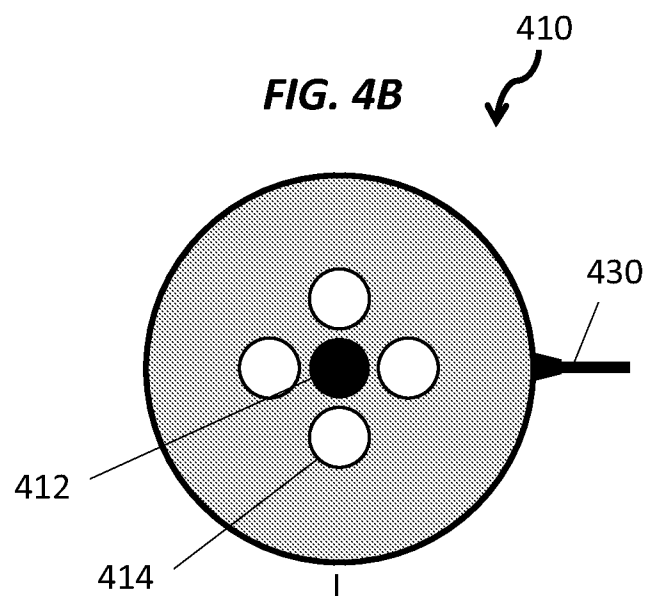


FIG. 4C

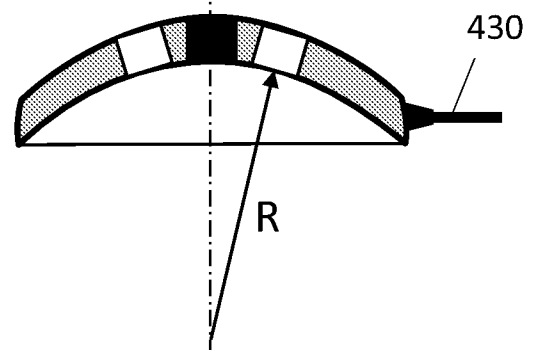
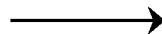
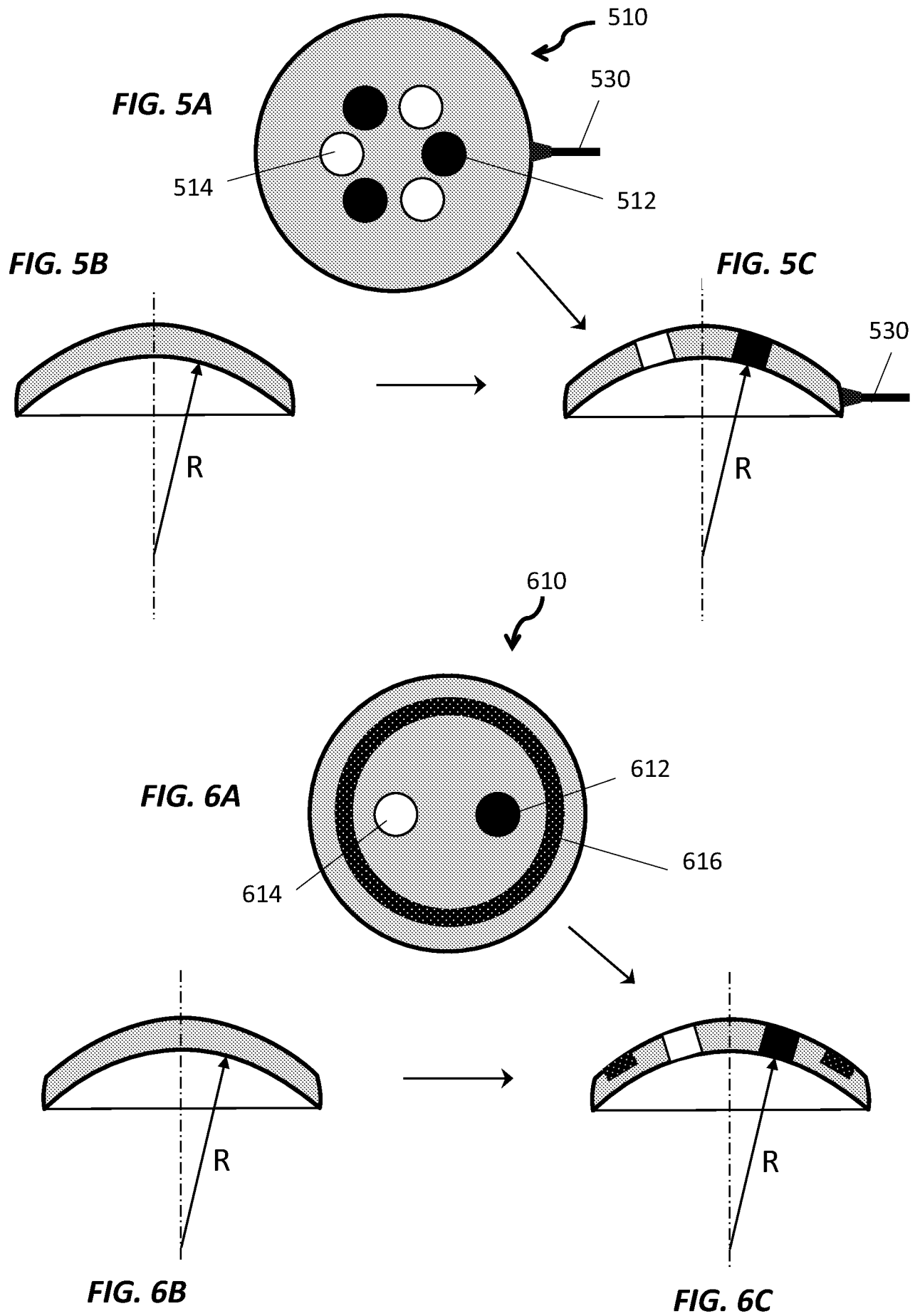


FIG. 4D



INTERNATIONAL SEARCH REPORT

International application No.

PCT/US2017/014552

A. CLASSIFICATION OF SUBJECT MATTER

IPC(8) - A61B 3/00; A61B 3/14; A61B 3/16; A61B 13/00; A61N 5/00; A61N 5/10 (2017.01)

CPC - A61B 3/00; A61B 3/14; A61B 3/16; A61B 5/00; A61B 13/00; A61N 5/00; A61N 5/10 (2017.02)

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

See Search History document

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

USPC - 351/206; 351/211; 351/221; 351/246; 378/65; 600/558; 606/4 (keyword delimited)

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

See Search History document

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 6,820,979 B1 (STARK et al) 23 November 2004 (23.11.2004) entire document	33-53
A	US 7,801,271 B2 (GERTNER et al) 21 September 2010 (21.09.2010) entire document	1-65
A	US 2008/0212026 A1 (MOLNAR et al) 04 September 2008 (04.09.2008) entire document	1-65
A	US 2011/0077548 A1 (TORCH) 31 March 2011 (31.03.2011) entire document	1-65

☐ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier application or patent but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

10 March 2017

Date of mailing of the international search report

31 MAR 2017

Name and mailing address of the ISA/US

Mail Stop PCT, Attn: ISA/US, Commissioner for Patents
P.O. Box 1450, Alexandria, VA 22313-1450

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Authorized officer

Blaine R. Copenheaver

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PCT OSP: 571-272-7774

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US2017/014552

Box No. II Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:

2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

3. ☒ Claims Nos.: 66-70
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box No. III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying additional fees, this Authority did not invite payment of additional fees.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:

4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest and, where applicable, the payment of a protest fee.
- ☐ The additional search fees were accompanied by the applicant's protest but the applicable protest fee was not paid within the time limit specified in the invitation.
- ☐ No protest accompanied the payment of additional search fees.