



(43) International Publication Date  
19 May 2016 (19.05.2016)

(10) International Publication Number  
**WO 2016/077741 A1**

- (51) International Patent Classification:  
*G01J 3/02* (2006.01) *G01N 21/64* (2006.01)
- (21) International Application Number:  
PCT/US2015/060658
- (22) International Filing Date:  
13 November 2015 (13.11.2015)
- (25) Filing Language: English
- (26) Publication Language: English
- (30) Priority Data:  
62/079,273 13 November 2014 (13.11.2014) US  
62/186,449 30 June 2015 (30.06.2015) US
- (71) Applicant: MARQUETTE UNIVERSITY [US/US]; PO  
Box 1881, Milwaukee, WI 53201-1881 (US).
- (72) Inventor; and
- (71) Applicant : KOVRIGUINE, Evgueni [RU/US]; 17975  
Continental Drive, Brookfield, WI 53045 (US).
- (74) Agents: MCBRIDE, M. Scott et al.; Andrus Intellectual  
Property Law, LLP, 100 East Wisconsin Avenue, Suite  
1100, Milwaukee, WI 53202 (US).
- (81) Designated States (unless otherwise indicated, for every  
kind of national protection available): AE, AG, AL, AM,

AO, AT, AU, AZ, BA, BB, BG, BH, BN, BR, BW, BY,  
BZ, CA, CH, CL, CN, CO, CR, CU, CZ, DE, DK, DM,  
DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT,  
HN, HR, HU, ID, IL, IN, IR, IS, JP, KE, KG, KN, KP, KR,  
KZ, LA, LC, LK, LR, LS, LU, LY, MA, MD, ME, MG,  
MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM,  
PA, PE, PG, PH, PL, PT, QA, RO, RS, RU, RW, SA, SC,  
SD, SE, SG, SK, SL, SM, ST, SV, SY, TH, TJ, TM, TN,  
TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW.

- (84) Designated States (unless otherwise indicated, for every  
kind of regional protection available): ARIPO (BW, GH,  
GM, KE, LR, LS, MW, MZ, NA, RW, SD, SL, ST, SZ,  
TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, RU,  
TJ, TM), European (AL, AT, BE, BG, CH, CY, CZ, DE,  
DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU,  
LV, MC, MK, MT, NL, NO, PL, PT, RO, RS, SE, SI, SK,  
SM, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ,  
GW, KM, ML, MR, NE, SN, TD, TG).

**Published:**

- with international search report (Art. 21(3))
- before the expiration of the time limit for amending the  
claims and to be republished in the event of receipt of  
amendments (Rule 48.2(h))

(54) Title: ADAPTER FOR A CELL HOLDER OF A SPECTROFLUOROMETER

(57) Abstract: Disclosed herein are spectrofluorometric adaptors for cell holders in spectrofluorometers and spectrofluorometric methods that enable fluorescence detection from surfaces of a microcell in conventional horizontal-beam spectrofluorometers equipped with standard cell holders and having multiple excitation channels.



WO 2016/077741 A1

## ADAPTER FOR A CELL HOLDER OF A SPECTROFLUOROMETER

### CROSS-REFERENCE TO RELATED PATENT APPLICATIONS

[0001] The present application claims the benefit of prior under 35 U.S.C. § 119(e) to U.S. Provisional Application No. 62/186,449, filed on June 30, 2015, and U.S. Provisional Patent Application No. 62/079,273, filed on November 13, 2014, the contents of which are incorporated herein by reference in their entireties.

### BACKGROUND

[0002] The field of the invention relates to spectrofluorometry. In particular, the field of the invention relates to adapters for cell holders of spectrofluorometers, wherein the adapter is configured to hold and position a microcell in the cell holder.

[0003] Fluorescence spectroscopy is one of the most sensitive methods for detecting biologically important molecules such as proteins, metabolites, and signaling molecules. In fluorescence spectroscopy, a light photon hits a molecule in an excitation process where the energy of the photon may be absorbed and re-emitted by the molecule through fluorescence. Often, the wavelength of the emitted photon is significantly longer than the wavelength of the excitation photon, which manifests itself as fluorescence in a different color. For example, excitation with blue light may cause green fluorescence. This phenomenon allows for sensitive detection of molecules capable of fluorescence emission (*i.e.*, “fluorophores”) because the emitted light may be readily differentiated from the background, scattered excitation light on the basis of the wavelength of the fluorescence emission. Fluorescence measurements are used extensively in many laboratory and clinical assays. The instruments for fluorescence detection range from small benchtop devices specialized for specific analysis to large versatile setups that enable detection of multiple fluorophores in samples and analysis of the multiple fluorophores’ properties.

[0004] Fluorescence measurement typically is performed in solution. In a typical fluorometric measurement, a liquid sample is placed in a quartz or glass cell, and the cell then is placed in a

cell holder of a fluorometer. The light beam from an excitation source then is directed into the quartz or glass cell through a side wall to illuminate the liquid sample inside the cell, which fluoresces. Fluorescence is emitted approximately equally in all directions and, generally, is much weaker than the excitation light, which may reflect and scatter upon illuminating the liquid sample. Therefore, in order to acquire accurate fluorescence measurements, it is important to separate the fluorescence signal from the reflected and scattered light of the excitation source.

**[0005]** To achieve the best separation, the most common design of the high-sensitivity spectrofluorometer involves detection of fluorescence at a 90° angle relative to the direction of the excitation light beam. (*See* Figure 1). As illustrated in Figure 1A, an excitation beam **6** enters through an excitation aperture **8** in an excitation screen **9a** and contacts the illuminated area **4** of a cell **2**. Fluorophores in a sample solution **16** contained in the cell **2** are excited and emit fluorescence **14** which is detected through the observed area **10** of the cell **2** as the fluorescence passes through the emission aperture **12** in an emission screen **9b** to a detector. As illustrated in Figure 1, to reduce contribution from scattered light, the excitation light or beam **6** typically is “trimmed” using an excitation screen **9a** having a narrow, rectangular or linear, excitation aperture **8** which permits the excitation beam **6** to illuminate only a middle area of the cell **4** and to prevent the excitation beam **6** from illuminating the corners of the cell **2**. The same technique is used for detecting fluorescence where an emission screen **9b** having a narrow, rectangular or linear, excitation aperture **12** permits the emission beam **14** to exit the middle of the cell **2** as an observed area **10** and enter the emission channel of a detector for detection. As a result, the fluorescence signal is detected exclusively from the middle of the sample cell, which is referred to as the “working volume” (see Figure 1.B), thus avoiding detecting any light (fluorescence or scattered) originating from the cell walls. As indicated in Figure 1B, a “working volume” **18** of the sample solution **16** is contacted by the excitation light **6** as it enters through an excitation aperture in the cell holder **11**. Fluorescence emission **14** from the working volume can pass only through the emission aperture in the cell holder **11**. This approach to measuring fluorescence from soluble molecules is utilized in the majority of highly sensitive commercial spectrofluorometers.

[0006] Difficulties in detecting fluorescence may arise due to high optical density and sample turbidity. For example, difficulties regarding high optical density may result when the concentration of the fluorophore in the solution is high enough that the solution absorbs all of the excitation light before it has a chance to reach the “working volume” in the middle of the cell, a phenomenon termed “inner filtering effect.” The same difficulty may arise when turbidity in the sample due to suspended particles blocks the excitation beam. To enable observation of fluorescence in these situations, a conventional workaround involves illuminating the front surface of the cell and exposing the same surface to the emission channel. This workaround permits collection of the fluorescence signal from the portion of solution near the inner cell wall provided that the emission signal can be distinguished from the reflected and scattered excitation beam.

[0007] In recent years, some research has focused on the observation of fluorescence from solution-glass interfaces in order to investigate molecular structures absorbed on the glass surface (*e.g.*, phospholipid bilayers). Most of these studies have been performed using horizontal glass slides where the solution drops are placed on top of the slide and observation are made from the bottom of the slide using fluorescence microscopes. The advantage of this setup is in its microscopic resolution of the surface details. However, the drawback to this setup is its poor accuracy in measurements of fluorescence intensity and, generally, its lack of spectral resolution in excitation and emission channels because of limitations of standard fluorescence microscopes. While it is possible to couple a fluorescence microscope with more sophisticated multi-wavelength excitation and emission channels, this is not a routine type of a fluorescence microscope and is extremely expensive to build.

[0008] One commercially available solution to enable observation of molecular layers absorbed on the inner glass surface of a sample cell using a standard horizontal-beam spectrofluorometer involves the use of triangular cells originally designed for measurements of samples with high optical density and/or turbidity. (*See* Figure 2). Such cells are offered by the optical cell manufactures such as Starna Cells Inc. (*See* Figure 3).

[0009] However, all of the cells that are currently manufactured have the front surface

oriented at a 45 degree angle to the excitation channel, which creates a strong reflection of the excitation beam directly into the sensitive emission channel, effectively destroying signal-to-noise ratio of the measurement. Therefore, these triangular cells are not utilized for observation of molecular layers and are employed only in the analysis of turbid solutions when the fluorescence signal has a relatively high intensity.

**[0010]** Although unconventional cells having a non-45 degree angle relative to the excitation channel are not commercially available, such unconventional cells could be designed. However, an additional problem in using such unconventional cells would arise from the fact that the non-45-degree angle relative to the excitation beam would make the cells asymmetric such that the cell may be used only with the excitation beam coming from one specific side of the cell. This is problematic because many commercial spectrofluorometers have two excitation channels that illuminate the cell from opposite sides, for example, spectrofluorometers having a left channel equipped with a steady-state light source, and having a right channel equipped with a pulsed light source for time-domain measurements. Therefore, it would be impossible to perform steady-state and time-domain measurements on the same molecular sample using an unconventional cell having a non-45 degree angle. In this situation, the user would need to have two cells each made for the specific direction of the excitation light and create two identical samples in the two cells, which would complicate research design and increase costs.

**[0011]** Another important requirement for optical cells in biochemical research arises from the fact that sample quantities may be scarce for typical measurements. Therefore, there is a demand for so-called “microcells” such that the volume of the sample is within a microliter range (*e.g.*, < 100  $\mu$ l). In order to utilize microcells in spectrofluorometers having 1-cm cell holders, adapters have been designed which center the microcell in the cell holder and provide reduced-size apertures for illumination and observation of the center of the sample solution while avoiding the cell corners (*i.e.*, similar to illumination and emission screens).

**[0012]** Therefore, new microcell adapters for cell holders of spectrofluorometers are desirable. The new adapters preferably should receive and hold a microcell at a variable

position (*e.g.*, a variable skewed position) for detecting fluorescence of a sample at the inner surface of the microcell and permit use of the same microcell in spectrofluorometers that utilize multiple excitation channels for illuminating the microcell from opposite sides (*e.g.*, from a right channel and from a left channel). The new adapters preferably should be configured for use in a commercially available spectrofluorometer having a 1-cm cell holder without having to modify the spectrofluorometer.

### SUMMARY

**[0013]** Disclosed herein are spectrofluorometric adapters for cell holders in spectrofluorometers and spectrofluorometric methods that enable fluorescence detection from surfaces of a microcell in conventional horizontal-beam spectrofluorometers equipped with standard cell holders and having multiple excitation channels.

**[0014]** The presently disclosed adapters are configured for a cell holder of a spectrofluorometer in which the cell holder is substantially square or rectangular in horizontal cross-section and the adapter includes an adapter body that fits into the cell holder. The cell holder and adapter body have centered, vertical axes which are substantially aligned.

**[0015]** The adapter body includes a cavity for receiving a microcell, which is a rectangular prism that is substantially square or rectangular in horizontal cross-section. The microcell has a centered, vertical axis such that when the microcell is positioned in the cavity of the adapter body, the vertical axis of the microcell may be offset relative to the vertical axes of the cell holder and adapter body and/or the microcell may be rotated horizontally about its vertical axis relative to the cell holder and adapter body. As such, the microcell may be turned about its vertical axis and/or horizontally shifted relative to the vertical axes of the cell holder and adapter body such that the excitation beam contacts the front wall and/or rear wall of the microcell and is observed by the emission channel from an orthogonal direction. In some embodiments, the excitation beam may contact the front wall and/or rear wall at an angle other than 90 degrees and at a position other than the center of the front wall and/or other than the center of the back wall of the microcell. In further embodiments, the microcell is placed in

a horizontally skewed position relative to the cell holder and, generally, off-center to achieve illumination and observation of the cell wall instead the center of the cell volume. In even further embodiments, the axis of the microcell is aligned horizontally relative to the emission beam but shifted horizontally backward such that an inner wall of an emission side of the microcell is aligned in parallel with the excitation beam.

**[0016]** In some embodiments, the adapter body includes a substantially square or rectangular bottom having a bottom cavity for receiving the microcell and a substantially square or rectangular top having a top cavity for receiving the microcell. The bottom and top of the adapter body typically are connected by at least one vertical side between the bottom and the top that holds the bottom and top in square alignment. Preferably, the bottom and top of the adapter body are connected by at least two vertical sides, which optionally are two adjacent vertical sides that form a 90 degree angle, where the two vertical sides are between the bottom and the top of the adapter body and hold the bottom and top of the adapter body in square or rectangular alignment. The vertical sides of the adapter body typically include side windows that permit entrance of excitation light or exit of emission light. Optionally, the vertical sides include vertical slits for receiving an excitation window screen or an emission window screen that include a variable rectangular or linear slit therethrough permitting entrance of an excitation beam or exit of an emission signal.

**[0017]** In some embodiments, the bottom of the adapter body includes horizontal slits for receiving an insert where the insert, after being inserted, closes the bottom cavity for receiving the microcell, where the microcell sits on the insert and the insert prevents the microcell from passing through the bottom of the adapter body. Preferably, the adapter is vertically invertible in the cell holder where the top of the adapter body includes a slit for receiving the insert, similar to the bottom of the adapter body, such that the insert in the slit of the bottom is removable and replaceable to the slit of the top to open the bottom cavity and close the top cavity. The adapter then may be vertically inverted in the cell holder such that the top is inverted to the bottom and the bottom is inverted to the top. As such, in one vertical position, the adapter may be utilized to hold a microcell, illuminate the microcell from a left-hand

excitation channel, and read the fluorescence at an emission channel. Subsequently, the microcell may be removed from the adapter, the adapter may be inverted to the opposite vertical position (*i.e.*, by removing the insert at the bottom of the adapter body and replacing the insert to the top of the adapter body and inverting the adapter), the microcell may be replaced into the inverted adapter, the microcell may be illuminated from a right-hand excitation channel (*e.g.*, at the identical incident angle and relative position to excitation beam as with the left-hand excitation channel), and the fluorescence may be read from an emission channel.

**[0018]** Also disclosed are methods for performing spectrofluorometric analysis. The methods optionally may utilize the presently disclosed adapters. In one method for performing a spectrofluorometric analysis on a sample as contemplated herein, the method includes: (a) placing the sample in a microcell, (b) placing the microcell in the adapter of as disclosed herein, (c) placing the adapter in the cell holder of the spectrofluorometer, and (d) performing the spectrofluorometric analysis. Suitable samples for this method may include fluorescent lipid bilayers that form on an interior surface of the microcell in which performing the spectrofluorometric analysis comprises detecting fluorescence at the interior surface of the microcell.

**[0019]** In another method for performing a spectrofluorometric analysis on a sample as contemplated herein, the method comprises: (a) placing the sample in a microcell, (b) placing the microcell in the invertible adapter as disclosed herein, (c) placing the adapter in the cell holder of the spectrofluorometer, (d) performing the spectrofluorometric analysis, (e) removing the microcell from the adapter, (f) inverting the adapter, (g) replacing the microcell into the adapter, and (h) performing the spectrofluorometric analysis. Suitable samples for this method may include fluorescent lipid bilayers that form on an interior surface of the microcell in which performing the spectrofluorometric analysis comprises detecting fluorescence at the interior surface of the microcell.

**[0020]** In another method for performing a spectrofluorometric analysis on a sample as contemplated herein, the method comprises: (a) placing a sample comprising lamellar vesicles



in a microcell that is substantially square or rectangular in horizontal cross-section (*i.e.*, not a microcell that is triangular in horizontal cross-section) and forming supported, fluorescent lipid bilayers on an interior surface of the microcell, (b) placing the microcell in an adapter, (c) placing the adapter in the cell holder of a spectrofluorometer, (d) performing the spectrofluorometric analysis by detecting fluorescence from the supported lipid bilayers at the interior surface of the microcell. In this method, the supported lipid bilayers may be formed by placing a solution or suspension comprising fluorescent lamellar vesicles in the microcell, allowing the fluorescent lamellar vesicles to contact the interior wall of the microcell and form supported, fluorescent lipid bilayers, flushing out the solution or suspension and excess fluorescent lamellar vesicles that have not attached to the interior wall of the microcell with a flushing solution that does not comprise the fluorescent lamellar vesicles, and refilling the microcell with the flushing solution. Flushing may be performed by introducing flushing solution in a manner that keeps the microcell filled with flushing solution during the entire flushing procedure.

#### BRIEF DESCRIPTION OF THE FIGURES

[0021] Figure 1. Convention horizontal-beam fluorescence configuration of the prior art. A. Illustration of fluorescence detection from a sample via excitation light and emission. B. Exemplary illumination schematic.

[0022] Figure 2. Top view of a triangular cell having a 45 degree incident surface of the prior art.

[0023] Figure 3. Triangular cells manufactured by Starna Cells, Inc. and having a 45 degree incident surface of the prior art.

[0024] Figure 4. Illustration of one embodiment of an adaptor as contemplated herein. Shown is an adaptor with two horizontal sections at the top and the bottom with a square cavity to hold a microcell (see Figure 4.A.). The microcell position with respect to the excitation window and emission window in this particular embodiment is shown in Figure 4.B. The adaptor may have an emission side with an aperture to expose a particular portion of

the cell to the detector (see Figure 4.C.) Variable apertures may be present in excitation screens and/or emission screens (Figure 4.C) inserted into vertical slits in the top section of the adapter and slid into the excitation window and/or emission window, respectively. The top section and the bottom section of the adapter may include horizontal slits across the square cell cavity for positioning of a sliding insert (Figure 1.D.) upon which the microcell may sit. Figure 4. E. illustrates two views of a single working prototype of the 1-cm adaptor with the installed emission and excitation window screens, the sliding insert, and a standard Starna 3-mm microcell.

[0025] Figure 5. Exemplary configurations regarding microcell placement and angle relative to excitation and emission channels.

[0026] Figure 6. Embodiments of orientations of a narrow-rectangle cell in an adaptor.

[0027] Figure 7. Embodiment of the adapter to allow for adjustment of cell position and orientation directly in the fluorometer. (A) Rotating plate assembled with a microcell. (B) Stationary plate. Directions of the excitation beam and emission window are indicated with the arrows on the plate. (C and D) Stationary and rotating plates assembled with different orientation of the cell with respect to the excitation and emission beams.

[0028] Figure 8. Confocal microscopy imaging of supported lipid bilayers formed on the interior surface of the fluorometer cell. (a) The rectangular quartz cell (3 mm path) on a sample platform to observe the inner cell wall facing the objective. (b) Vertical cross-section of the microcell. The microscope was focused at three different vertical coordinates to image the slices above, at, and below the inner cell wall (positions I, II, and III). The cross section of the cell is schematically shown as a black outline. The lipid bilayer location is indicated by the interior line forming a square. The objective was centered at the vertical inner wall boundary to create strong contrast between the fluorescent cell content on the left and non-fluorescent optical material of the cell wall on the right of the image. The drawing is not to scale. (c) Images of the 3 mm cell containing supported lipid bilayer including 1% (mol) of Rhodamine-DPPE taken with the 40x objective. The slices were imaged at three vertical positions (I, II,

and III) of the confocal plane corresponding to the relative Z-coordinate of 25  $\mu\text{m}$ , 0  $\mu\text{m}$ , and -200  $\mu\text{m}$ , respectively. Thickness of the optical section was 4.3  $\mu\text{m}$ .

**[0029]** Figure 9. Detection of fluorescence from supported lipid bilayers and evaluation of sensitivity of the method. (a) Positioning of the rectangular 3 mm microcell inside the custom-designed cell adaptor. The excitation and emission channels are indicated by arrows, the light paths focused by optics and trimmed by custom slits are schematically depicted by dotted and solid lines. (b) For comparison, schematic of the microcell placement in the standard 3 mm cell adaptor (Starna, Cat# FCA3) avoiding direct illumination of the front and rear inner cell walls. (c) Solid line, the rhodamine emission spectrum of the supported lipid bilayer in the 3 mm microcell recorded using the custom adaptor shown in Panel a. The bilayer was deposited utilizing 100  $\mu\text{M}$  LUV containing 1% rhodamine-DPPE. Dotted line, the same sample was placed in the centered position of the FCA3 adaptor (Panel b) showing bilayer fluorescence excited by the light scattered in solution. Dashed line, a reference 5  $\mu\text{M}$  LUV solution with 1% rhodamine recorded in FCA3 adaptor (Panel b).

#### DETAILED DESCRIPTION

**[0030]** The disclosed subject matter further may be described utilizing terms as defined below.

**[0031]** Unless otherwise specified or indicated by context, the terms “a”, “an”, and “the” mean “one or more.”

**[0032]** As used herein, “about”, “approximately,” “substantially,” and “significantly” will be understood by persons of ordinary skill in the art and will vary to some extent on the context in which they are used. If there are uses of the term which are not clear to persons of ordinary skill in the art given the context in which it is used, “about” and “approximately” will mean plus or minus  $\leq 10\%$  of the particular term and “substantially” and “significantly” will mean plus or minus  $> 10\%$  of the particular term.

**[0033]** As used herein, the terms “include” and “including” have the same meaning as the

terms “comprise” and “comprising.” The terms “comprise” and “comprising” should be interpreted as being “open” transitional terms that permit the inclusion of additional components further to those components recited in the claims. The terms “consist” and “consisting of” should be interpreted as being “closed” transitional terms that do not permit the inclusion of additional components other than the components recited in the claims. The term “consisting essentially of” should be interpreted to be partially closed and allowing the inclusion only of additional components that do not fundamentally alter the nature of the claimed subject matter.

**[0034]** The presently disclosed adapters are configured for use in a standard horizontal-beam spectrofluorometer with a typical cell holder (*e.g.*, a 1-cm cell holder or another size) without having to modify the spectrofluorometer. The adapters are specifically configured for permitting observation of surface fluorescence in a microcell that is placed in the adapter. In some embodiments, the adapter comprises or consists of an adapter body with a cavity for receiving and holding a microcell, window screens with variable slits permitting entrance of excitation light and/or exit of emission light, and a removable bottom slit that is replaceable to the top of the adapter body such that the adapter is vertically invertible.

**[0035]** The adapter body positions a standard quartz or glass microcell at a selected angle relative to excitation and emission windows in the adapter body and relative to the cell holder. The adapter body may include an excitation window screen and/or an emission window screen that include a variable rectangular or linear apertures therethrough permitting entrance of an excitation beam or exit of an emission signal. The variable rectangular or linear aperture limits entrance of the excitation beam to the selected surfaces of the microcell and limits exposure of selected portions of the illuminated surfaces of the microcell to the emission window, while minimizing scattering and reflection of the excitation light into the emission window. The angle and position of the microcell inside the body of the adaptor as well as the widths and positions of the rectangular or linear apertures may be adjusted to maximize intensity of the fluorescence signal and to reduce excitation light captured by the emission window.

[0036] To enable illumination and observation of fluorescence at the surface of the microcell, the microcell, generally, is positioned askew or off-center in the cell holder and at a variable angle (*e.g.*, at an angle other than 0, 45, or 90 degrees) relative to the incident excitation beam, relative to the exit emission beam, and relative to the cell holder. In some embodiments, the microcell can be considered to be turned about a vertical axis relative to the incident excitation beam, relative to the exit emission beam, and relative to the cell holder. This configuration is inherently asymmetric and requires the excitation light to always enter from one specified direction. In other words, this configuration is impermissive of multiple excitation channels coming from different directions. Because standard fluorometers may have two independent excitation channels illuminating the microcell from opposite sides (*i.e.*, left-hand excitation and right-hand excitation), the presently disclosed adaptor is designed to be “vertically invertible” (*i.e.*, from top-to-bottom) with the microcell being inserted in one vertical end of the adaptor for use with a first excitation channel and the microcell being inserted in the opposite vertical end of the adaptor for use with the second excitation channel. To allow for the invertible design, an insert placed in horizontal slits at the bottom of the adaptor body is removable and may be re-positioned in horizontal slits at the top of the adaptor body (*i.e.*, the insert may be positioned at either vertical end of the adaptor body to create a “top” or a “bottom” at either end for holding the microcell).

[0037] Referring now to the figures, Figure 4 illustrates one embodiment of an adaptor **15** as contemplated herein. Shown is an adaptor **15** with two horizontal sections at the top **30a** and the bottom **30b** with a square cavity **22** to hold a microcell **2** (see Figure 4.A and Figure 4.B.) and spaced vertically apart by an excitation side **50a** and an emission side **50b** at a 90 degree angle to the excitation side **50a**. The adaptor **15** has an excitation side **50a** with an excitation window **38** for receiving an excitation beam **6** which will contact a portion of a cell **2** within the adaptor **15**. The emission side **50b** has an emission window **42** through which a emission beam **14** can be emitted and detected.

[0038] The amount of the microcell **2** exposed to the excitation beam **6** may be varied or adjusted by inserting an excitation screen **40a** into a vertical slit **26** in the top section **30a** of

the adapter **15** (see Figure 4.A. and 4.C.) As illustrated in Figure 4.C., the excitation screen **40a** is slid into the vertical slit **26** and covers a portion of the excitation window **38**. The portion of the cell exposed to the excitation beam **6** can be varied by varying the size of the aperture **44** in the excitation screen **40a**. The adapter **15** also may include a vertical slit (not shown) in the top section **30a** for receiving an emission screen **40b**. The emission screen **40b** may be slid into the emission window **42** such that only a portion of the emission beam **14** is allowed to pass through the emission window **42** to the detector.

[0039] As illustrated in Figure 4.B., the microcell **2** is held by the adapter in a position that is askew, off-center, and/or turned about a vertical axis relative to the adapter **15**. As such, the adapter **15**, when placed into a square cell holder will position the microcell **2** in the cell holder at a position that is askew, off-center, and/or turned about a vertical axis and/or at a variable angle (*e.g.*, at an angle other than 0, 45, or 90 degrees) relative to the incident excitation beam **6**, relative to the exit emission beam **14**, and relative to the cell holder.

[0040] As illustrated in Figure 4.D., the adapter **15** may be vertically reversible. As illustrated, the top section **30a** and the bottom **30b** of the adapter **15** may include horizontal slits **60** across the square cavity **22** for inserting a sliding insert **70** in to the bottom section upon which the microcell **2** may sit. Subsequently, the microcell **2** may be removed, the adapter **15** may be inverted vertically, the sliding insert **70** may be replaced to the bottom section **30b** of the adapter **15**, and the microcell **2** may be replaced into the square cavity **22** of the adapter **15** where the microcell **2** sits on the sliding insert **70**. Figure 4.E. illustrates two different views of a single prototype having a microcell **2** inserted into an adapter **15**.

[0041] The disclosed adapters may be configured to position a microcell at any selected position within a cell holder. Figure 5 illustrates exemplary configurations of adapter and positioning of the microcell relative to the excitation beam and emission beam. A. Microcell **2** turned and vertical axis off-center relative to adapter **15** (and cell holder); B. Microcell **2** turned and vertical axis off-center relative to adapter **15** (and cell holder); C. Microcell **2** horizontally aligned with emission channel **14** but shifted vertically downward with respect to

excitation beam **6**, effectively aligning the inside cell wall of the emission side of the microcell **2** in parallel with the excitation beam; D. Microcell **2** turned and vertical axis off-center relative to adapter **15** (and cell holder).

[0042] The disclosed adapters may be configured to receive and position a rectangular microcell at any selected position within a cell holder. Figure 6 illustrates embodiments of configuration of adapters for rectangular microcells and positioning of the microcells relative to the excitation beam and emission beam. A. Microcell **2** aligned with excitation beam **6** and emission channel **14**; B. Microcell **2** horizontally aligned with emission channel **14** but shifted horizontally backward with respect to excitation beam **6**, effectively aligning the inside cell wall of the emission side of the microcell **2** in parallel with the excitation beam; C. Microcell **2** having vertical axis aligned with adapter **15** but vertically turned counter-clockwise relative to adapter **15**; D. Microcell **2** turned and vertical axis off-center relative to adapter **15**.

[0043] The adapter may be configured to rotate the microcell adjustably while the adapter holding the microcell is position in the cell holder. As illustrated in Figure 7, the adapter includes a rotating plate (Figure 7.A.) having an opening for inserting a microcell, and a stationary plate (Figure 7.B.). The rotating plate may be placed into the stationary plate (Figure 7.C.) and the stationary plate may be inserted into a cell holder. The rotating plate of the adapter may be rotated adjustably in order to rotate the inserted microcell and vary the rotatable orientation of the microcell with respect to the excitation beam and emission beam (Figure 7.D.), while the adapter is positioned in the cell holder.

[0044] Figure 8. Confocal microscopy imaging of supported lipid bilayers formed on the interior surface of the fluorometer cell. (a) The rectangular quartz cell (3 mm path) on a sample platform to observe the inner cell wall facing the objective. (b) Vertical cross-section of the microcell. The microscope was focused at three different vertical coordinates to image the slices above, at, and below the inner cell wall (positions I, II, and III). The cross section of the cell is schematically shown as a black outline. The lipid bilayer location is indicated by the interior line forming a square. The objective was centered at the vertical inner wall boundary to create strong contrast between the fluorescent cell content on the left and non-fluorescent

optical material of the cell wall on the right of the image. The drawing is not to scale. (c) Images of the 3 mm cell containing supported lipid bilayer including 1% (mol) of Rhodamine-DPPE taken with the 40x objective. The slices were imaged at three vertical positions (I, II, and III) of the confocal plane corresponding to the relative Z-coordinate of 25  $\mu\text{m}$ , 0  $\mu\text{m}$ , and -200  $\mu\text{m}$ , respectively. Thickness of the optical section was 4.3  $\mu\text{m}$ .

## EXAMPLES

[0045] The following examples are illustrative and are not intended to limit the scope of the claimed subject matter.

### [0046] Example 1

[0047] The presently disclosed adapter may be utilized in methods for detecting fluorescence from molecular layers absorbed on the inner surfaces of a fluorometer microcell using a standard horizontal beam fluorometer equipped with a standard 1-cm cell holder. The detection of fluorescence on the inner surfaces of the microcell is enabled by (1) placing the microcell at a specific position and angle with respect to the excitation and emission windows with the aid of the disclosed adaptor inserted in the cell holder, and (2) utilizing customized screens having variable apertures at specific positions in windows of the adapter body that control excitation and emission paths through the adaptor and microcell to optimize the ratio of fluorescence-to-scatter light intensity. The adapter may be used to position the fluorometer microcell inside the fluorometer cell holder off-center and/or at an angle in order to appropriately expose the surfaces of the microcell to the excitation and emission channels. The desired positioning of the microcell is achieved by the adapter body and square or rectangular cavity therein which may be askew or angled (*i.e.*, turned about a vertical axis) relative to the incident excitation beam, relative to the exit emission beam, and relative to the cell holder. As such, when the microcell is placed in the cavity, the microcell and the surfaces of the microcell are positioned askew or angled (*i.e.*, turned about a vertical axis) relative to the incident excitation beam, relative to the exit emission beam, and relative to the cell holder. Figure 4.A illustrates one embodiment of the adaptor showing two horizontal sections at the



top and the bottom with a square cavity to hold the microcell. The microcell position with respect to the excitation and emission windows in this particular embodiment is shown in Figure 4.B.

**[0048]** In Figure 4.B, the excitation light illuminates the inner surface of the microcell and causes fluorescence that is observed through the emission window. The width of the aperture in the window screen and the position of the window screen in the excitation path are adjustable to restrict illumination to specific cell surface(s). Black bars in Figure 4.B indicate window screen portions blocking the light, and the opening between them is an aperture that allows the excitation light to illuminate the specified portion of the cell. Likewise, the side of the adaptor facing the emission channel (*i.e.* light detector) has a variable aperture to expose a particular portion of the cell to the detector. The variable apertures may be present in window screens (Figure 4.C) inserted into the vertical slits in the emission and excitation sides of the adaptor as in Figure 4.A or the variable apertures may be an integral part of the vertical sides of the adaptor body provided the variable aperture allow for illumination and/or observation of specific cell areas.

**[0049]** Particular combinations of aperture widths, aperture positions, positioning of the microcell inside the holder, and the angle of the microcell relative to the excitation beam may be selected to increase intensity of the observed fluorescence signal from inner surfaces while reducing contributions from the scattered and reflected light of the excitation beam.

**[0050]** The top and the bottom of the adaptor feature narrow horizontal slits across the cell cavity for positioning of the sliding insert (Figure 4.D) upon which the microcell may sit. As such, the sliding insert prevents the microcell from passing through the bottom of the adaptor. The insert may be placed at either end of the adaptor (*i.e.*, the top end or the bottom end) to enable use of the adaptor in spectrofluorometers having multiple excitation channels (*e.g.*, in spectrofluorometers having right- and left-sided excitation channels). Therefore, only one adaptor is required for use in a two-channel spectrofluorometer. Figure 4.E illustrates a working prototype of the 1-cm adaptor with the installed emission and excitation window screens, the sliding insert, and a standard Starna 3-mm microcell.

[0051] The configuration of the adapter and the window screens may vary depending on the geometry of the excitation beam. For example, when the excitation beam is a well-focused steady-state source, the adapter may utilize a window screen having a relatively narrow aperture on the excitation side to avoid strong reflections from the edges of the microcell. On the other hand, when the excitation beam is a pulsed LED, the adapter may utilize a window screen having a relatively wide aperture or the adapter may not utilize a window screen at all to facilitate illumination of the cell surfaces. Some exemplary configurations are illustrated in Figure 5.

[0052] In some embodiments, the disclosed adapters may be utilized in experiments that include analyzing fluorescence from bilayer molecular structures that are immobilized on the inner surface of a microcell. The bilayer molecular structures are immobilized on the inner surface thus enabling analysis of interactions of different ligands with the bilayer and its protein components in variable conditions. Multiple experiments are possible by repeatedly exposing the same bilayer to a series of biochemical agents. The proposed adaptor is a low-cost solution to advance academic research on membrane proteins in supported lipid bilayers, and to facilitate development of clinical tests for membrane protein function.

[0053] References.

[0054] 1. Galush *et al.* (2008), "Quantitative fluorescence microscopy using support lipid bilayer standards," *Biophys. J.* 95, 2512-9.

[0055] 2. Crane *et al.* (2007), "Fluorescence Microscopy to Study Domains in Supported Lipid Bilayers." In *Methods in Membrane Lipids*, Vol. 400, pp. 481-488.

[0056] 3. Lakowicz, J.R. (2010) *Principles of Fluorescence Spectroscopy*. 3<sup>rd</sup> edit. Springer.

[0057] 4. U.S. Patent No. 8,115,922.

**[0058]** Example 2

**[0059]** Another embodiment of the disclosed adaptor is designed to utilize a microcell that has a rectangular cross-section with one narrow and another long dimension. Figure 6 demonstrates the horizontal cross-section of this embodiment of the adaptor with a narrow-rectangle cell positioned in different orientations in the adaptor with respect to the excitation beam. In Figure 6, the horizontal cross-section of the square cell adaptor is shown with a black outline. The outer and inner boundaries of the microcell are shown with thick and thin black outlines, respectively. The excitation light beam focused by optics of the excitation channel and trimmed by adaptor slits is schematically shown with solid outline. Horizontal dashed line indicates center of the excitation beam and serves as a reference for the cell position. The portion of the cell exposed to the emission channel is indicated with slanted dashed lines.

**[0060]** As illustrated in Figure 6, Position A allows the inner cell walls be illuminated at a very sharp angle due to the fact that optics is adjusted to focus the beam in the middle of the cell holder. The sharp angle of the incident excitation light leads to a very small amount of scattering into the emission channel thus increasing sensitivity. In addition, due to elongated geometry of the cell, the cell corners are located outside of the area observed by the emission channel. In this position both inner cell walls are illuminated to a similar degree because the cell is centered with respect to the cell holder. Position B improves illumination of one inner cell wall (*i.e.*, the front cell wall as illustrated) by shifting the cell backward in the adaptor to increase efficiency of excitation of fluorophores absorbed at the front, inner cell wall. Position C further increases intensity of light hitting the inner cell wall by rotating the cell in the adaptor where the cell is rotated about a center axis. Position D reduces scattering from the left cell corner by rotating the cell in the adaptor about a skewed axis (*i.e.*, where the cell is moved from the center position of the adaptor and rotated).

**[0061]** The disclosed narrow-rectangle microcell of this embodiment may provide advantages over a square microcell. First, the useful surface that may be exposed to the emission channel in a narrow-rectangle microcell is wider. For example, the cross-section cell of the narrow-

rectangle microcell is 10 mm wide while the cross-section of the square microcell is only 3 mm. Also, the corners of the narrow-rectangle microcell are located farther away from the center of the emission window, and thus are less likely to contribute to scattering. Slanted dashed lines in Figure 6 indicate an approximately useful width for an emission window using the narrow-rectangle microcell. Further, the inner front and back walls of the narrow-rectangle microcell are separated by a relatively small distance (*e.g.*, 1 mm) such that both inner cell walls (*i.e.*, front inner cell wall and back inner cell wall) will be exposed to the excitation beam in most orientations of the microcell in the adapter. This effectively doubles the amount of useful fluorescence signal emitted from the sample layers absorbed on these inner walls, therefore increasing sensitivity of detection when the narrow-rectangle microcell is used.

#### [0062] Example 3

[0063] In some embodiments of the disclosed adapters, the position and orientation of the cell in the adapter may be adjustable, for example, to correct for small variation in tuning of the focusing optics of some fluorometers. Accurate illumination of the inner cell surfaces is sensitive to such variability and may be addressed by either providing a set of adapters with different cell position, which the end user will be able to choose from to obtain maximum signal, or by using a direct position-adjustment mechanism. One prototype of such a mechanism of the adapter to allow for adjustment of cell position and orientation directly in the fluorometer is shown in Figure 7, including (A) rotating plate assembled with a microcell; (B) stationary plate, directions of the excitation beam and emission window are indicated with the arrows on the plate; and (C and D) stationary and rotating plates assembled with different orientation of the cell with respect to the excitation and emission beams.

[0064] The stationary plate is mounted on top of the fluorometer cell holder and has a square opening aligned with the cavity of the cell holder. The stationary plate has flaps extending into the holder cavity down from the plate surface to keep the stationary plate position fixed at all

times. The rotating plate with a microcell is inserted into the opening of the stationary plate until the cell reaches the bottom of the cell holder and the rotating plate rests on top of the stationary plate. In this assembly the microcell is positioned vertically in the cell holder (C). The horizontal position of the microcell and its angle relative to the excitation and emission beams are easily adjusted by shifting the rotating plate with the microcell relatively to the stationary plate. The optimal position (*e.g.*, as evaluated by the maximum useful fluorescence signal and minimum scattering) may be marked on the stationary plate, which allows this position to be reproduced next time with necessary precision. Using the recorded angle and position parameters, the adaptor body may be manufactured with the matching location of the cell cavity resulting in non-adjustable optimized adaptor. Alternatively, the cell-position adjustment may be performed for every particular experiment using the described mechanism for adjusting the position of the microcell or a similar mechanism.

**[0065]** Example 4

**[0066]** Reference is made to the attached manuscript entitled “Fluorescence of supported phospholipid bilayers recorded in a conventional horizontal-beam spectrofluorometer,” E.A. Kovrigina, and E.L. Kovrigin, submitted to the Journal of Fluorescence, the contents of which are incorporated herein by reference in its entirety.

**[0067]** Title

**[0068]** Fluorescence of supported phospholipid bilayers recorded in a conventional horizontal-beam spectrofluorometer

**[0069]** Abstract

**[0070]** Supported phospholipid bilayers are a convenient model of cellular membranes in studies of membrane biophysics and protein-lipid interactions. Traditionally, supported lipid bilayers are formed on a flat surface of a glass slide to be observed through fluorescence

microscopes. This paper describes a method to enable fluorescence detection from the supported lipid bilayers using standard horizontal-beam spectrofluorometers instead of the microscopes. In the proposed approach, we form the supported lipid bilayers on the inner optical surfaces of the standard fluorescence microcell. To enable observation of the bilayer absorbed on the cell wall, the microcell is placed in a standard fluorometer cell holder and specifically oriented to expose the inner cell walls to both excitation and emission channels with a help of the custom cell adaptor. In our measurements, the signal intensity from supported bilayers doped with 1% (mol) of rhodamine-labeled lipid in the standard 3-mm optical microcell was equivalent to fluorescence of the 70-80 nM reference solution of rhodamine recorded in a commercial microcell adaptor. Because no modifications to the instruments are required in this method, a variety of steady-state and time-domain fluorescence measurements may be performed with supported phospholipid bilayers using standard horizontal-beam spectrofluorometers.

#### [0071] Introduction

[0072] Supported phospholipid bilayers were long established as a model of cellular phospholipid membranes [1]. The phospholipid bilayers attached to the glass surface (with and without cross-linking to the glass) are physically very similar to other membrane mimics in their biophysical properties [2]. The main approach to recording fluorescence signals from such supported bilayers is through the microscopes with the bilayer deposited on the horizontal surface of the glass slide and covered by a layer of the aqueous buffer [3,4]. Fluorescence microscopy of supported bilayers allows analysis of spontaneous phase separation [5,6] as well as detection of molecular events at the bilayer with a single-molecule resolution [7,8]. One highly useful feature of the supported lipid bilayer (relatively to liposomes) is that it is *immobilized* allowing for dynamic replacement of the solution in contact with the bilayer. Supported lipid bilayers are routinely studied in the flow-cell mode with the microscopy setups to reveal interactions of membrane proteins with lipids and with soluble ligands (for example, [9,10]). However, if one is particularly interested in *spectral* properties of the lipid-associated fluorophores and their changes due to ligand interactions, the

microscopes must use multi-wavelength lasers and include detectors with spectral resolution, which are expensive and not generally available (for the example of such setup see [11]). The best practical approach to record fluorescence of lipid bilayers in standard spectrofluorometers was to utilize solutions (suspensions) of liposomes or protein-liposome complexes [12-15]. Yet, liposomes are not very stable and do not allow for easy replacement of the surrounding buffer as they are freely diffusing particles. Supported lipid bilayers represent a convenient alternative because they are easily prepared by spontaneous fusion of lipid vesicles with the glass surfaces [1,2] yet require use of sophisticated fluorescence microscopes for their analysis. This paper reports a proof-of-principle experiment where a supported lipid bilayer is created in a standard fluorometer microcell and fluorescence measurements are performed in a typical horizontal-beam spectrofluorometer. We believe that this inexpensive approach may serve a number of needs in studies membrane protein interactions with lipids and soluble ligands.

#### [0073] Material and Methods

[0074] Preparation of liposomes. Lipid mixtures contained DOPC:DOPG:fluorophore in the molar ratio of 89:10:1. Rhod-DPPE was used as a fluorescent marker of the lipid bilayer. All lipids were obtained from Avanti Polar Lipids Inc. and used without further purification. Large unilamellar vesicles (LUV) were prepared by extrusion through 0.2  $\mu\text{m}$  polycarbonate membrane using Avanti Mini-Extruder. The total concentration of lipids during extrusion was 1 mM. The lipids were diluted to 0.1 mM for deposition of the supported lipid bilayers. Use of LUV versus multi-lamellar vesicles (MLV) or small unilamellar vesicles (SUV) does not appear to be a strict requirement in this protocol; Brian and McConnell [1] successfully utilized MLV in their pioneer work while SUV were employed by Cremer and Boxer [16] for the same purpose.

[0075] Preparation of supported lipid bilayers. Supported lipid bilayers were formed on the inner surface of a rectangular 3-mm fluorometer microcell (Starna #3-3.45-Q-3) by completely filling the cell with the LUV solution of 0.1 mM total lipid (a minimum concentration allowing for the stable bilayer formation [1,5,16]). Bilayers with negatively

charged lipids were reported to form very fast at physiological pH and ionic strength [16]. To keep the bilayer adhered to the walls, the cell must stay filled at all times because air-water boundary destroys the attached bilayers [16]). Therefore, to remove the initial LUV solution, we flushed the cell with a continuous flow of the phosphate saline buffer. To perform the flush, the cell was positioned horizontally and the buffer was introduced towards the bottom of the cell with the needle; the displaced solution was allowed to exit the cell and drip into the waste container. Capillary forces were sufficient to keep cells completely filled during the entire procedure. Relatively wide spacing between walls in the microcell (3 mm) made the buffer exchange process somewhat inefficient— 50 to 100 ml of buffer at a flow rate (approx.) 2 ml/min was required to ensure complete removal of residual LUV from the bulk solution inside the microcell. Quality of the LUV removal was checked by residual fluorescence of the bulk solution in the microcell.

**[0076]** Confocal fluorescence microscopy. Images of supported lipid bilayers were taken using Nikon Perfect Focus Ti-E inverted research microscope. To support the 3 mm cell in horizontal position in front of the objective, the custom cell holder was utilized.

**[0077]** Fluorescence spectroscopy. Photon Technologies International QuantaMaster 40 spectrofluorometer (Horiba) was used for recording emission spectra from supported lipid bilayers. The fluorometer was equipped with a standard 1 cm sample holder, emission and excitation monochromators, and Xenon steady-state excitation source. A custom cell adaptor was designed to hold the standard 3 mm microcell from Starna (Cat# 3-3.45-Q-3). The adaptor included internal slits to trim the excitation beam and to expose the illuminated cell surface to the emission channel. Details of the adaptor design are described in the U.S. Provisional Patent Applications No. 62/079,273 and No. 62/186,449 filed with U.S. Patent Office on November 13, 2014 and June 30, 2015, respectively.

## **[0078]** Results

**[0079]** Visualization of the supported lipid bilayers. Prior to spectroscopic investigation, formation of supported lipid bilayers on the inner surfaces of the fluorometer cell may be



confirmed using fluorescence microscopy. Figure 8 shows representative rhodamine fluorescence images of the inner volume of the 3-mm rectangular microcell. The sample platform was adjusted such that left-hand-side of the image corresponds to interior of the optical cell, while the right-hand-side is always inside the material of the vertical cell wall (Panels a and b). To demonstrate that the fluorescent lipids are found exclusively on the inner walls for the optical cell and not in the solution, we imaged confocal planes at three vertical coordinates (Panels b and c). The position III is 200  $\mu\text{m}$  below the inner wall and shows no fluorescence as the confocal plane is entirely localized inside the horizontal cell wall. The position II is aligned with the inner surface of the horizontal cell wall and reveals bright fluorescence (left of the cell wall junction) corresponding to rhodamine in the phospholipid bilayer coating the inner cell surface. The uppermost position I is 25  $\mu\text{m}$  above the inner surface of horizontal cell wall intersecting the vertical inner cell surface in the middle of the image. Left-hand-side of the image revealed no significant fluorescence from solution inside the cell indicating that all suspended LUV were successfully removed by flushing procedure. The bright straight line in the middle originates from fluorescence of the supported lipid bilayer formed on the vertical inner cell wall. Real thickness of the lipid bilayer cannot be estimated from these images due to inherent resolution limit of optical spectroscopy of about 400 nm at this wavelength. Therefore, we relied on literature reports that the absorbed lipids form a bilayer with the expected thickness of about 5 nm as was previously established [5,16]. Black area on the right of the image corresponds to the non-fluorescent interior of the vertical cell wall.

**[0080]** Supported lipid bilayers in a horizontal-beam spectrofluorometer. For observation of fluorescence from phospholipid bilayers we set the microcell inside the standard 1-cm cell holder to expose the surface with the bilayer to both excitation and emission channels. Figure 9.a. shows one of the possible positions of the microcell for bilayer observation compared to the standard centered position (Panel b). Panel c gives emission spectra recorded from the supported lipid bilayers and the diluted reference LUV solution. In this measurement, we used 560 nm excitation light with identical 5-nm slits in the excitation and emission monochromators. Illumination of the surface of the cell (Panel a) generates both the useful

fluorescence from the surface and scattered light from the corners of the microcell; therefore, one has to use additional adjustable slits trimming the excitation beam as well as limiting exposure of the emission channel to cell edges. Solid line in Panel c shows fluorescence signal from the most optimal combination of cell position and adjustable slits. This optimal combination is likely to slightly vary from one instrument to another due to differences in tuning of the focusing optics.

**[0081]** To confirm that the recorded fluorescence comes from the supported bilayer rather than from any residual LUV in the bulk solution, we tested the solution fluorescence from the same sample using a standard microcell adaptor with a centered cell location (FCA3, Starna) shown in Panel b. The FCA3 adaptor is manufactured to enable observation of fluorescence from internal volume of the solution; its narrow windows shield corners of the cell from exposure to excitation and emission channels. In our supported bilayer preparation we flushed the microcell with a buffer in several stages, every time recording residual LUV fluorescence from the bulk solution in the microcell using FCA3 adaptor. After passing about 50 ml of buffer through the cell, we observed the rhodamine fluorescence drop to a minimum value shown as a dash-dot-dashed line in Figure 9.c. Further flushing with another 50 ml did not reduce it any more indicating that this residual fluorescence signal likely comes from the supported bilayers on the front and back walls (facing the emission channel) excited indirectly by the light scattered in solution.

**[0082]** Relative sensitivity of measurements from the supported lipid bilayer sample may be evaluated by comparison of its fluorescence to the signal intensity of a reference LUV solution in a centered adaptor FCA3. The dashed black line in Figure 9.c is a spectrum of the initial LUV solution used for preparation of supported lipid bilayers diluted to 5  $\mu\text{M}$  total lipid concentration prior to the measurement (resulting in 50 nM concentration of rhodamine). Thus, the signal from supported lipid bilayers with 1% (mol) rhodamine in a 3-mm microcell was equivalent to 70-80 nM of rhodamine in LUV solution (recorded with 5 nm excitation and emission slits in the 3-mm cell).

**[0083]** Discussion

[0084] Typical applications of supported lipid bilayers utilized glass slides or flow-cells positioned horizontally to observe the bilayers from below or above in some kind of a microscopy setup. Supported lipid bilayers formed on the inner surfaces of the fluorometer cells may be investigated while oriented vertically because van der Waals forces holding the bilayers in place are stronger than gravity while the optical cell ensures that the bilayer is covered with solution at all times. In our experiments, supported lipid bilayer samples were stable for more than a week at 4°C. Vertical orientation of lipid bilayers in our method has another advantage over the horizontal orientation: biomolecular aggregates (or dust) sediment to the bottom of the microcell, thus leaving the observation area. With the supported lipid bilayers located horizontally in the microscopy setups, all aggregates settle directly onto the bilayer under study thus potentially biasing measurement results.

[0085] Observation of fluorescence from the solution near the cell surface dates back as far as to Sir Frederick William Herschel who described fluorescence of optically dense quinine solutions [17]. Since then the solutions of high optical density or/and significant turbidity were routinely studied with illumination of the cell surface at an angle such that the incident light penetrates a thin layer of solution inside the cell, which fluoresces. These applications typically utilize triangular cells with the cell surface oriented at the 45° angle to both emission and excitation channels. This arrangement, however, is not suitable for fluorescence detection from supported lipid bilayers because the weak fluorescence signal is swamped with the excitation light reflected into the monochromator thus destroying the signal/noise ratio. Lakowicz recommended smaller 20-30° angles to reduce scattered light with such triangular cells [18]—this is approximately the angle we are using in Figure 9.a. However, the optical cells that would expose the front cell surfaces to the excitation beam at these sharp angles are not generally available. Utilizing the commercial microcell that is smaller than the size of the standard 1-cm cell holder allowed us to explore multiple positions and orientations of the cell achieving illumination of the cell surfaces at different angles.

[0086] Another requirement in biochemical applications is a small size of the sample. The triangular 45° cells manufactured by Starna or Hellma require 1.7 ml of solutions to fill. Here

we utilized a 3-mm Starna microcell, which nominally holds 315  $\mu\text{L}$  of a sample bringing it to the volume range of flow-cells in microscopy applications. By attaching a cell cap with the needle and inlet-outlet tubing one might construct a flow cell to be able to replace solution in the cell directly in the fluorometer (in this study, for the sake of simplicity, we were flushing the cell outside of the instrument).

**[0087]** It has been demonstrated that proteoliposomes may be used to create supported lipid bilayers incorporating membrane proteins [1,2]. In our method, by adding the ligand to the bulk solution one may perform titration studies of receptor-ligand interactions. Subsequent flushing out the ligand regenerates the unbound form of the membrane-associated receptor thus allowing for multiple repeated experiments with the same bilayer sample. Many spectrofluorometers are equipped with temperature-controlled cell holders opening an avenue for studies of temperature-dependent thermodynamic and kinetic properties of the supported bilayer samples. Our approach to observation of the supported bilayer fluorescence is also agnostic to the type of the excitation light source, therefore time-domain studies are equally possible on the same samples used for spectral recording. In summary, a combination of a straightforward bilayer-forming technique, commercial optical cells coupled with the simple custom adaptors, and the standard spectrofluorometer hardware allows for an inexpensive way of recording fluorescence from supported lipid bilayers thus enabling detailed spectroscopic studies of lipid membranes, membrane proteins, and their interactions.

**[0088]** References

**[0089]** 1. Brian AA, McConnell HM (1984) Allogeneic stimulation of cytotoxic T cells by supported planar membranes. *Proc Natl Acad Sci U S A* 81: 6159-6163.

**[0090]** 2. Tamm LK, McConnell HM (1985) Supported phospholipid bilayers. *Biophysical Journal* 47: 105-113.

**[0091]** 3. Bagatolli LA (2007) Membranes and Fluorescence Microscopy. In: C.D. Geddes (ed.), editor. *Reviews in Fluorescence*. pp. 33-51.

- [0092] 4. Galush WJ, Nye JA, Groves JT (2008) Quantitative fluorescence microscopy using supported lipid bilayer standards. *Biophys J* 95: 2512-2519.
- [0093] 5. Crane JM, Tamm LK (2007) Fluorescence Microscopy to Study Domains in Supported Lipid Bilayers. *Methods in Membrane Lipids*. pp. 481-488.
- [0094] 6. Pinto SN, Fernandes F, Fedorov A, Futerman AH, Silva LC, et al. (2013) A combined fluorescence spectroscopy, confocal and 2-photon microscopy approach to re-evaluate the properties of sphingolipid domains. *Biochimica Et Biophysica Acta-Biomembranes* 1828: 2099-2110.
- [0095] 7. Hell SW, Wichmann J (1994) Breaking the diffraction resolution limit by stimulated emission: stimulated-emission-depletion fluorescence microscopy. *Optics Letters* 19: 780-782.
- [0096] 8. Kastantin M, Walder R, Schwartz DK (2012) Identifying Mechanisms of Interfacial Dynamics Using Single-Molecule Tracking. *Langmuir* 28: 12443-12456.
- [0097] 9. Lin W-C, Iversen L, Tu H-L, Rhodes C, Christensen SM, et al. (2014) H-Ras forms dimers on membrane surfaces via a protein–protein interface. *Proceedings of the National Academy of Sciences* 111: 2996-3001.
- [0098] 10. Iversen L, Tu H-L, Lin W-C, Christensen SM, Abel SM, et al. (2014) Ras activation by SOS: Allosteric regulation by altered fluctuation dynamics. *Science* 345: 50-54.
- [0099] 11. Raicu V, Singh DR (2013) FRET Spectrometry: A New Tool for the Determination of Protein Quaternary Structure in Living Cells. *Biophysical Journal* 105: 1937-1945.
- [00100] 12. Kinnunen P, Alakoskela J-M, Laggner P, Nejat D (2003) Phase Behavior of Liposomes. *Methods in Enzymology*: Academic Press. pp. 129-147.
- [00101] 13. Janoff AS (1999) *Liposomes: Rational Design*: Marcel Dekker.

**[00102]** 14. Walden P (1994) Liposomes as tools for the reconstitution of biological systems. In: Philippot J.R. SF, editor. Liposomes as tools in basic research and industry. Boca Raton: CRC Press. pp. 71-88.

**[00103]** 15. Woodle MC, Papahadjopoulos D (1989) Liposome preparation and size characterization. *Methods Enzymol* 171: 193-217.

**[00104]** 16. Cremer PS, Boxer SG (1999) Formation and Spreading of Lipid Bilayers on Planar Glass Supports. *The Journal of Physical Chemistry B* 103: 2554-2559.

**[00105]** 17. Herschel FW (1845) On a case of superficial colour presented by a homogeneous liquid internally colorless. *Philosophical Transactions of the Royal Society of London* 135: 143-145.

**[00106]** 18. Lakowicz JR (2010) *Principles of Fluorescence Spectroscopy*: Springer. 954 p.

**[00107]** In the foregoing description, it will be readily apparent to one skilled in the art that varying substitutions and modifications may be made to the invention disclosed herein without departing from the scope and spirit of the invention. The invention illustratively described herein suitably may be practiced in the absence of any element or elements, limitation or limitations which is not specifically disclosed herein. The terms and expressions which have been employed are used as terms of description and not of limitation, and there is no intention that in the use of such terms and expressions of excluding any equivalents of the features shown and described or portions thereof, but it is recognized that various modifications are possible within the scope of the invention. Thus, it should be understood that although the present invention has been illustrated by specific embodiments and optional features, modification and/or variation of the concepts herein disclosed may be resorted to by those skilled in the art, and that such modifications and variations are considered to be within the scope of this invention.

**[00108]** Citations to a number of patent and non-patent references are made herein. The cited references are incorporated by reference herein in their entireties. In the event that there is an inconsistency between a definition of a term in the specification as compared to a definition of the term in a cited reference, the term should be interpreted based on the definition in the specification.

## CLAIMS

I claim:

1. An adapter for a cell holder of a spectrofluorometer, the cell holder being substantially square or rectangular in horizontal cross-section, the adapter comprising a body that fits into the cell holder, the cell holder and adapter body having centered, vertical axes that are substantially aligned, the adapter body comprising a cavity for receiving a microcell, the microcell being a rectangular prism and being substantially square or rectangular in horizontal cross-section, the microcell having a centered, vertical axis such that the microcell may be positioned in the cavity of the adapter body such that at least one of the following conditions are met: (1) the vertical axis of the microcell is offset relative to the vertical axes of the cell holder and adapter body; (2) the microcell is rotated horizontally about its vertical axis relative to the cell holder and adapter body; and (3) the axis of the microcell is aligned horizontally relative to the emission beam but shifted horizontally backward such that an inner wall of an emission side of the microcell is aligned in parallel with the excitation beam.

2. The adapter of claim 1, wherein the microcell is positioned in the cavity of the adapter body such that both of condition (1) and condition (2) are met.

3. The adapter of claim 1 or 2, wherein the microcell is rotated horizontally about its vertical axis relative to the cell holder and adapter body at an angle relative to the cell holder that is not 45 degrees.

4. The adapter of any of the foregoing claims, wherein the adapter body comprises a substantially square or rectangular bottom having a bottom cavity for receiving the microcell, a substantially square or rectangular top having a top cavity for receiving the microcell, and at least one vertical side between the bottom and the top and holding the bottom and top in square or rectangular alignment.

5. The adapter of claim 4, comprising at least two vertical sides between



the substantially square or rectangular bottom and the substantially square or rectangular top and holding the bottom and top in square or rectangular alignment.

6. The adapter of claim 5, wherein the two vertical sides are adjacent and form a 90 degree angle.

7. The adapter of any of claims 4-6, wherein the vertical side comprises a side window permitting entrance of excitation light or exit of emission light.

8. The adapter of claim 7, wherein the vertical sides comprise vertical slits for receiving an excitation window screen or an emission window screen, the window screens including apertures for permitting entrance or exit of light.

9. The adapter of any of claims 4-8, wherein the bottom comprises horizontal slits for receiving an insert, the insert closing the bottom cavity for receiving the microcell and preventing the microcell from passing through the bottom of the adapter.

10. The adapter of claim 9, wherein the adapter is vertically invertible in the cell holder, wherein the top comprises a slit for receiving the insert such that the insert in the slit of the bottom is removable and replaceable to the slit of the top to open the bottom cavity and close the top cavity, and the adapter then is vertically inverted in the cell holder such that the top is inverted to the bottom and the bottom is inverted to the top.

11. The adapter of any of the foregoing claims, wherein the adapter comprises a rotating plate into which the microcell is inserted.

12. The adapter of claim 11, wherein the adapter further comprises a stationary plate into which the rotating plate is inserted, and the stationary plate is attached to the body of the adapter or is an integral part of the adapter body.

13. A method for performing a spectrofluorometric analysis on a sample, the method comprising: (a) placing the sample in a microcell, (b) placing the microcell in the adapter of any of the foregoing claims, (c) placing the adapter in the cell holder of the

spectrofluorometer, and (d) performing the spectrofluorometric analysis.

14. The method of claim 13, wherein the sample comprises a fluorophore-labelled lipid bilayer that forms on an interior surface of the microcell and performing the spectrofluorometric analysis comprises detecting fluorescence at the interior surface of the microcell.

15. A method for performing a spectrofluorometric analysis on a sample, the method comprising: (a) placing the sample in a microcell, (b) placing the microcell in the invertible adapter of claim 10, (c) placing the adapter in the cell holder of the spectrofluorometer, (d) performing the spectrofluorometric analysis, (e) removing the microcell from the adapter, (f) inverting the adapter, (g) replacing the microcell into the adapter, and (h) performing the spectrofluorometric analysis.

16. The method of claim 15, wherein the sample comprises a fluorophore-labelled lipid bilayer that forms on an interior surface of the microcell and performing the spectrofluorometric analysis comprises detecting fluorescence at the interior surface of the microcell.

17. A method for performing a spectrofluorometric analysis on a sample comprising supported lipid bilayers, the method comprising: (a) placing a sample comprising fluorophore-labelled lamellar vesicles in a microcell that is substantially square or rectangular in cross-section and forming supported, fluorophore-labelled lipid bilayers on an interior surface of the microcell, and (b) detecting fluorescence from the supported fluorophore-labelled lipid bilayers at the interior surface of the microcell.

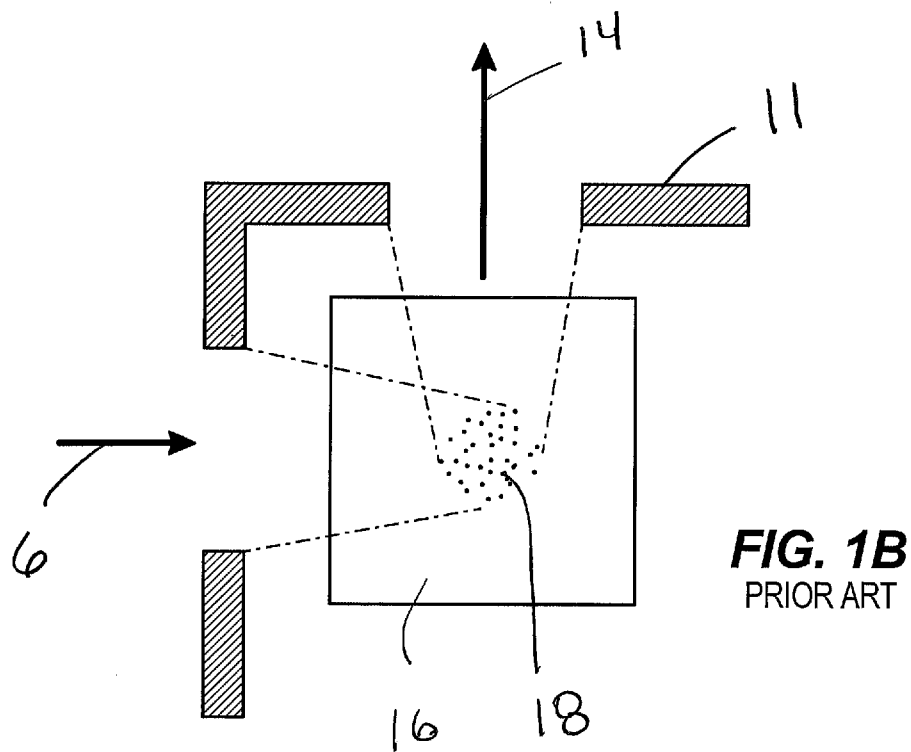
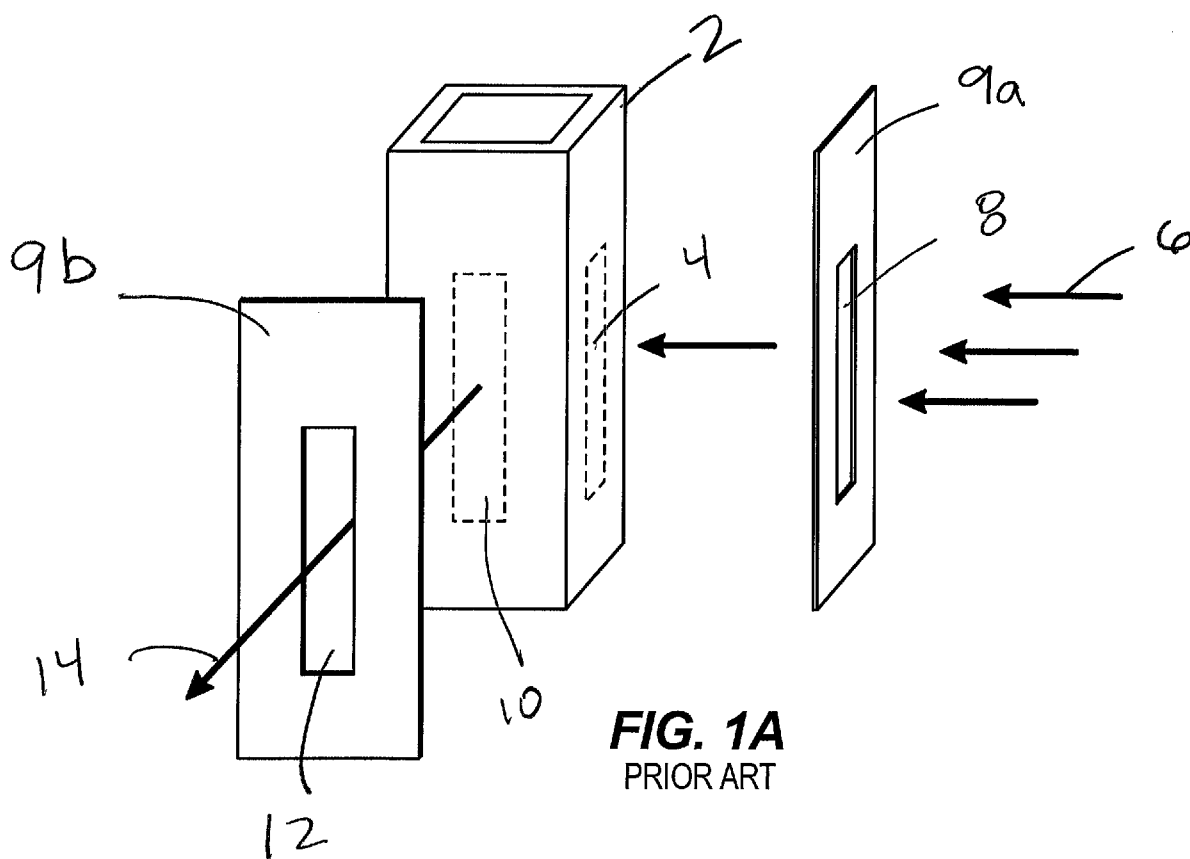
18. The method of claim 17, wherein the supported lipid bilayers are formed by placing a solution or suspension comprising fluorescent lamellar vesicles in the microcell, allowing the fluorescent lamellar vesicles to attach to the interior wall of the microcell and form supported, fluorescent lipid bilayers, flushing out the solution or suspension and excess fluorescent lamellar vesicles that have not attached to the interior wall of the microcell with a flushing solution that does not comprise the fluorescent lamellar

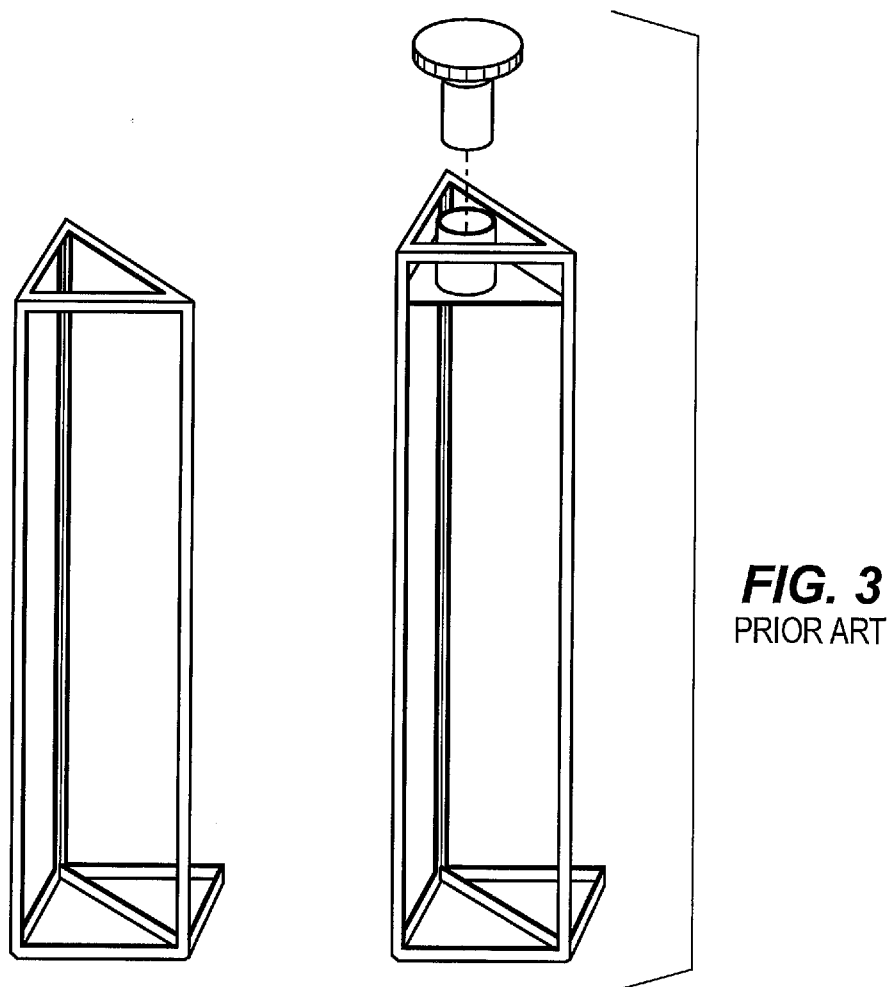
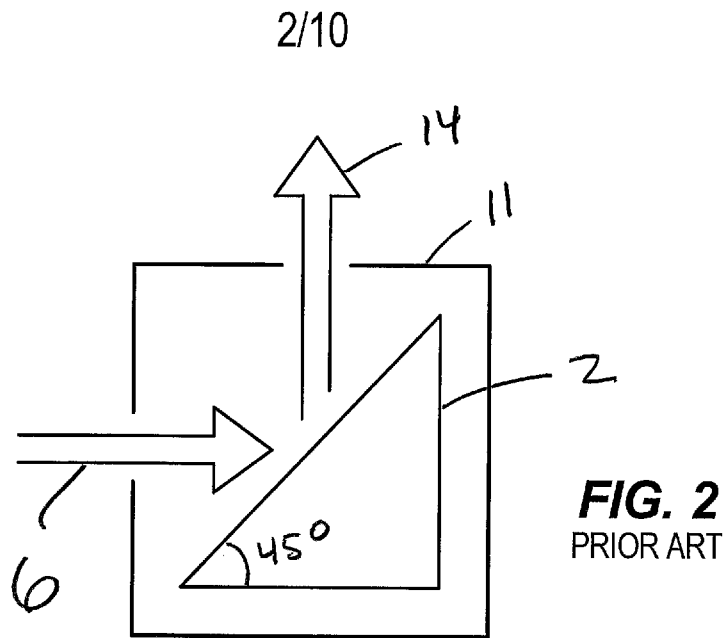
vesicles in a manner that keeps the microcell filled with flushing solution during the entire flushing procedure.

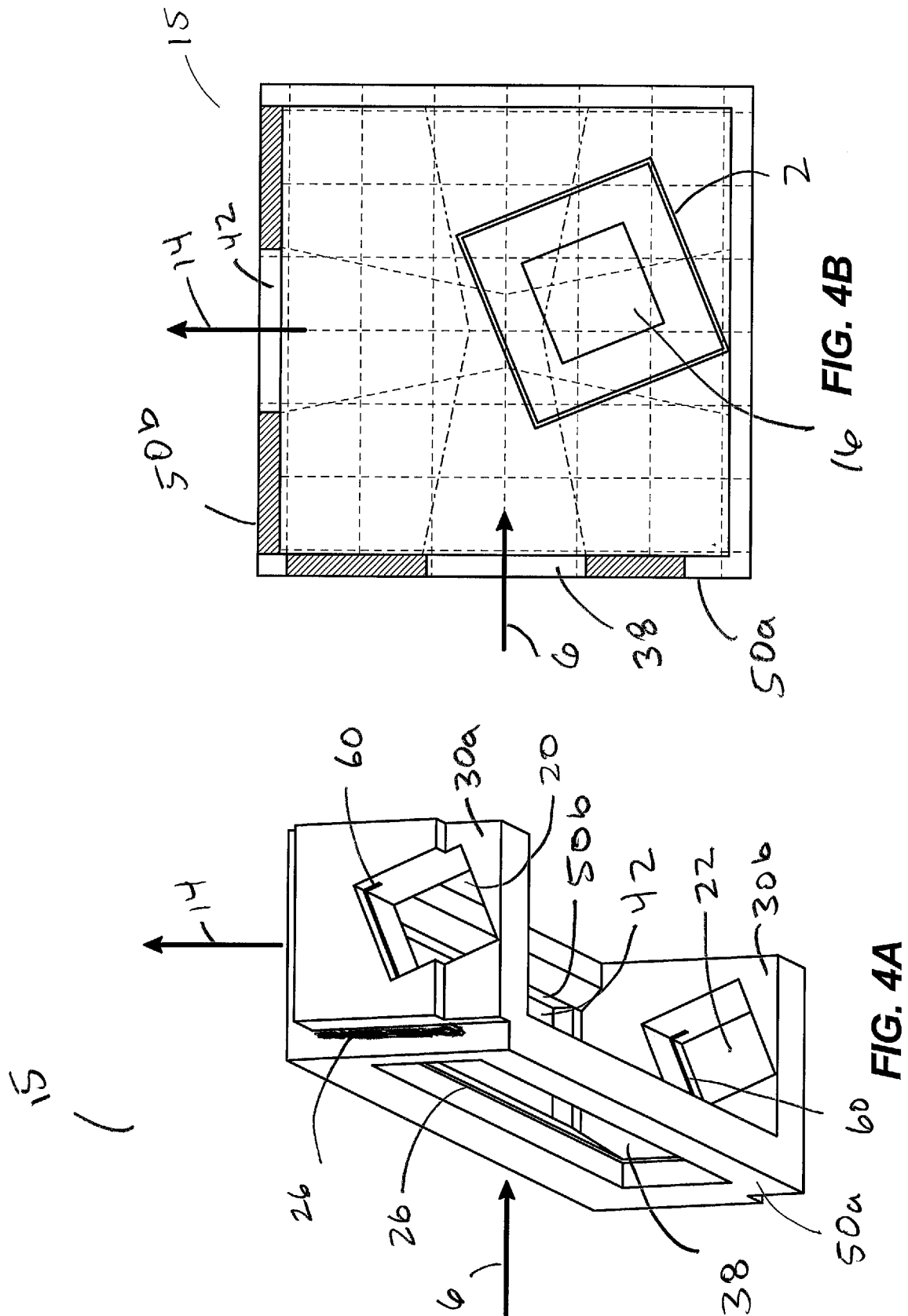
19. The method of claim 17 or 18, wherein detecting fluorescence comprises contacting the interior surface of the microcell with an excitation beam of the spectrofluorometer and exposing the interior surface of the microcell to the emission channel of the spectrofluorometer.

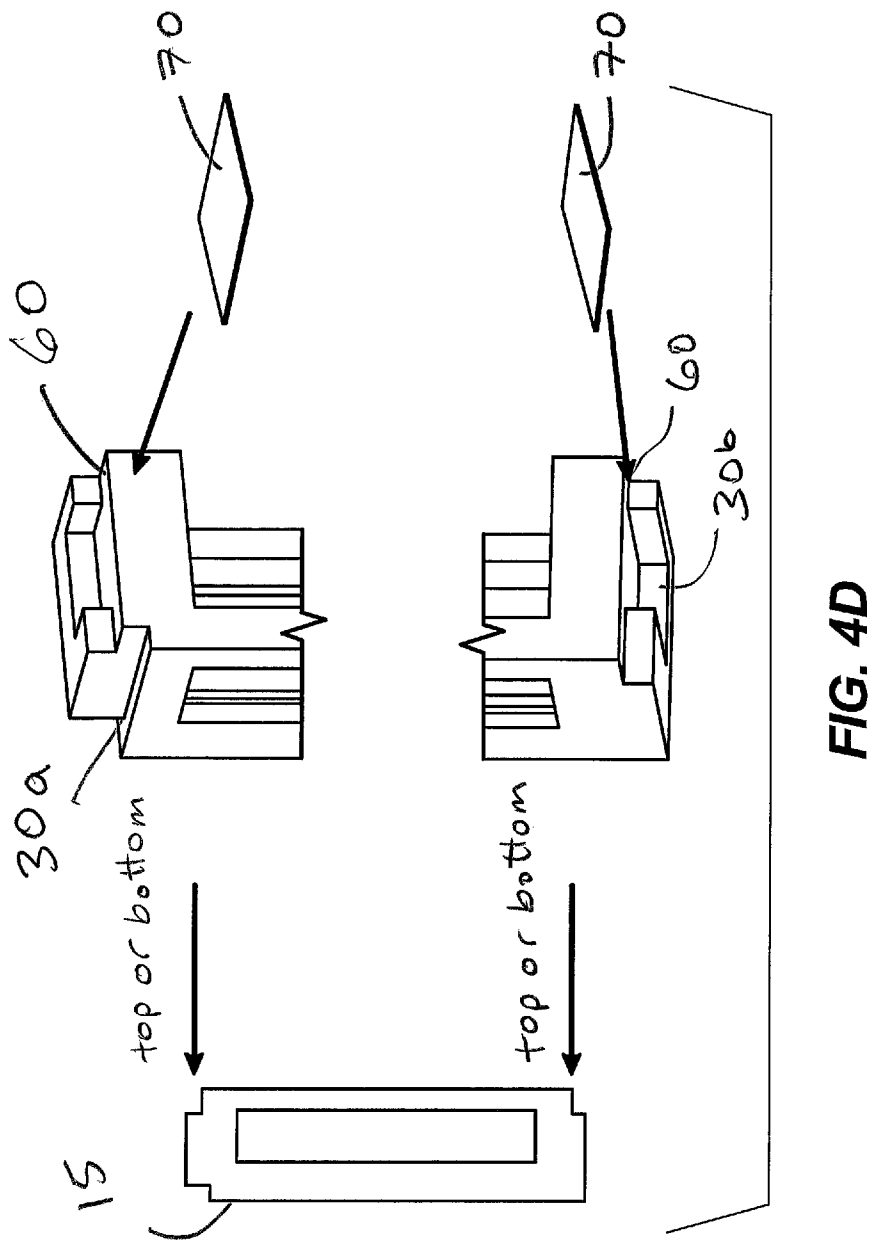
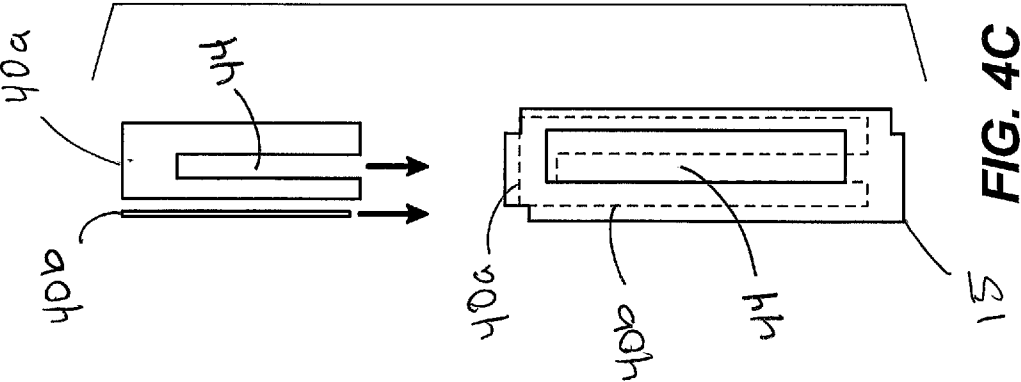
20. The method of claim 19, wherein the microcell is placed in an adapter and inserted into a spectrofluorometer prior to detecting fluorescence from the supported lipid bilayers at the interior surface of the microcell, and the adapter comprises internal slits that trim the excitation beam of the spectrofluorometer and expose the interior surface of the microcell to the emission channel.

1/10

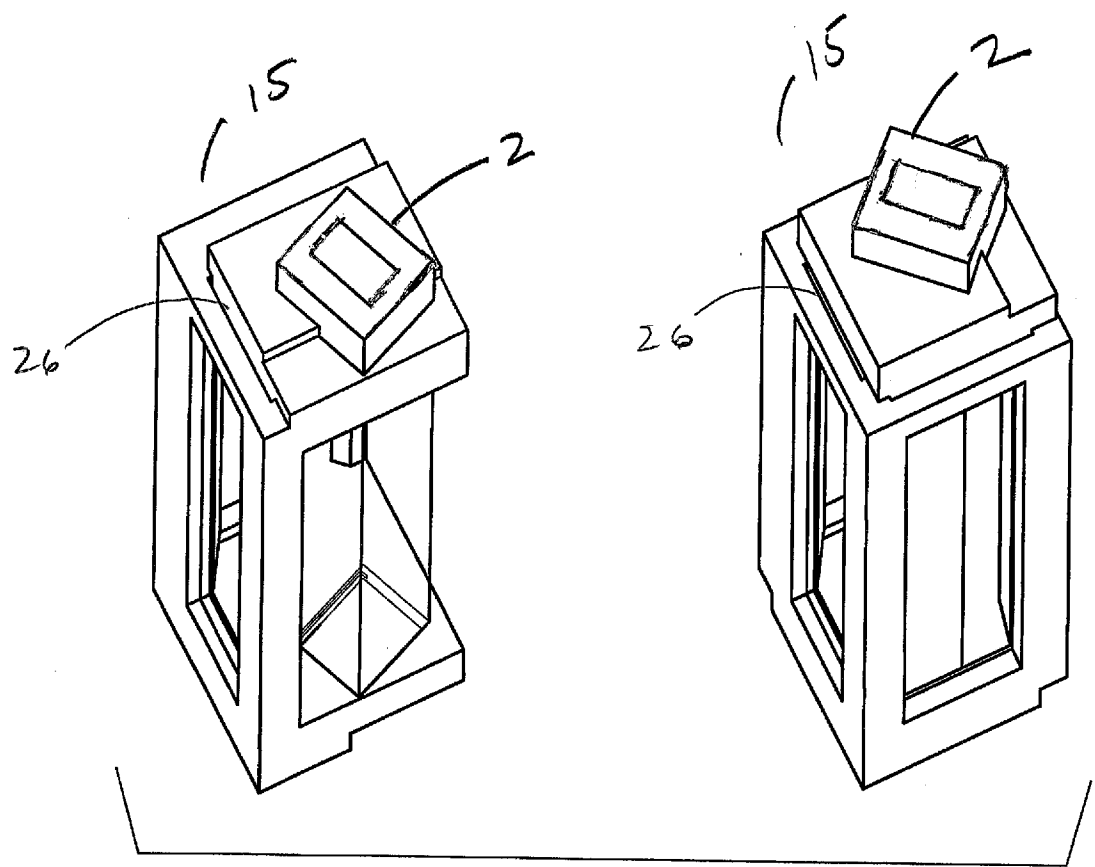








5/10



**FIG. 4E**



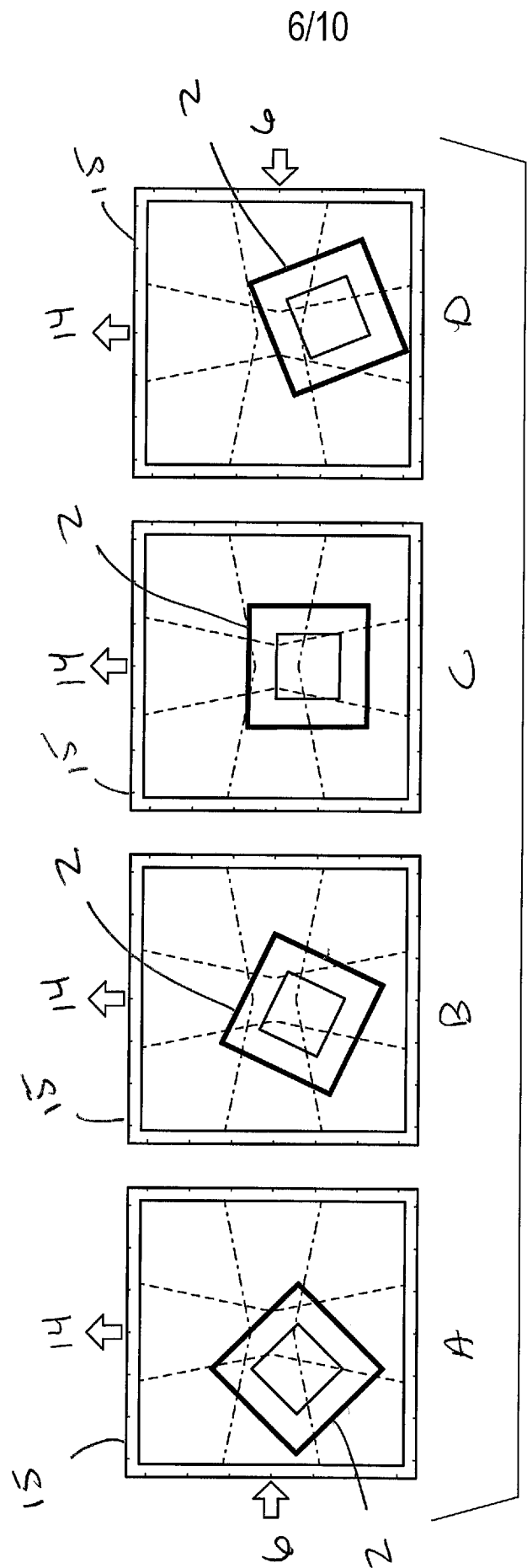
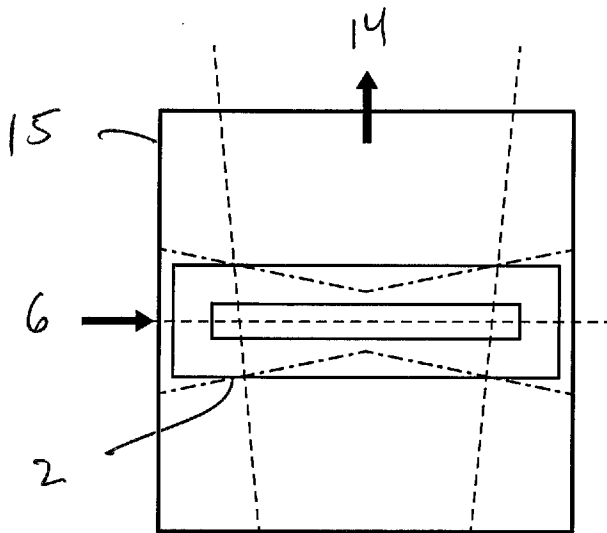
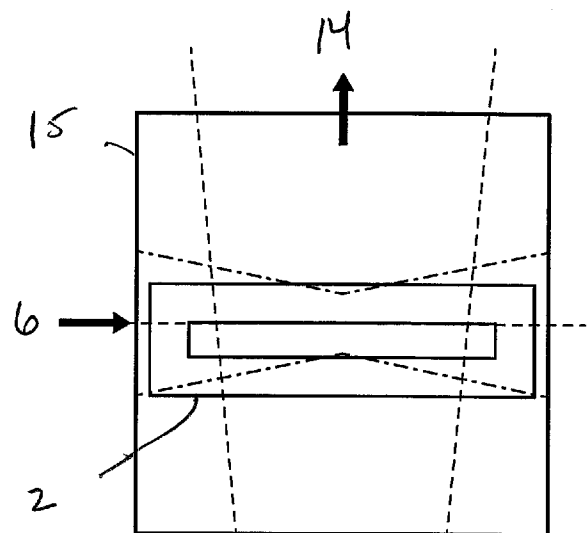


FIG. 5

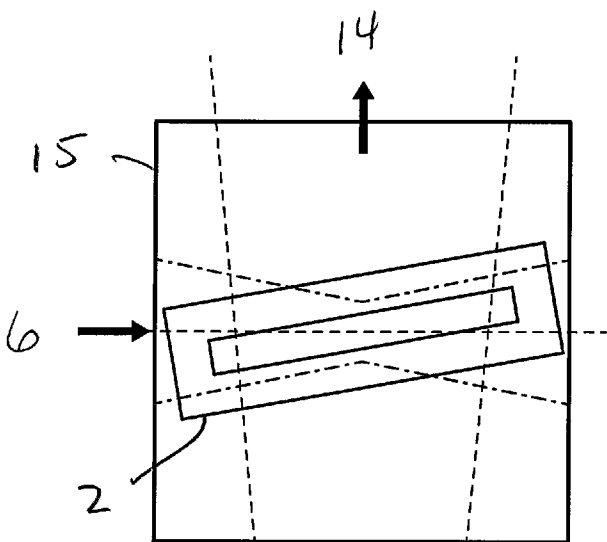
7/10



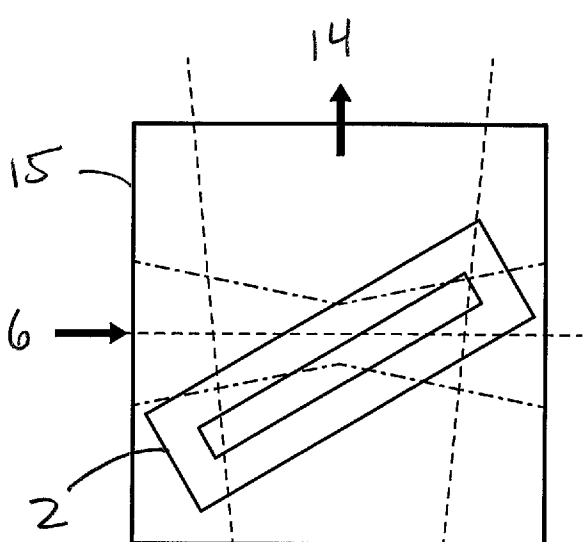
**FIG. 6A**



**FIG. 6B**

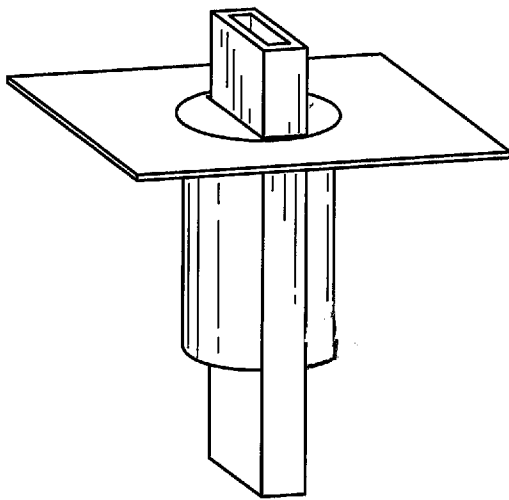


**FIG. 6C**

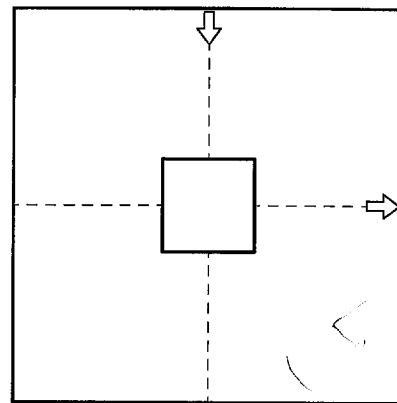


**FIG. 6D**

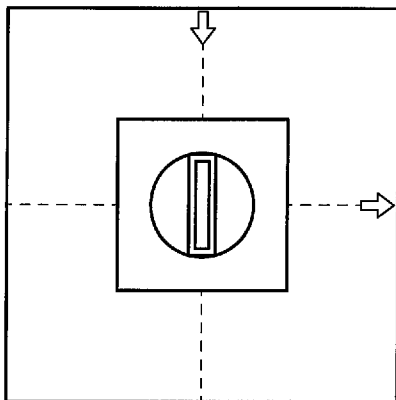
8/10



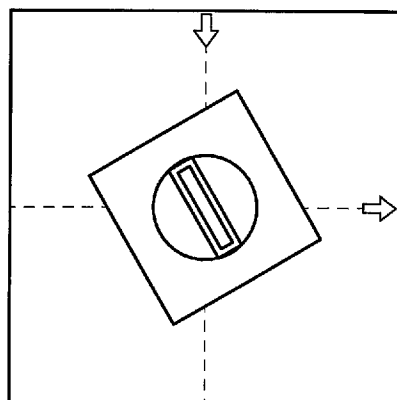
**FIG. 7A**



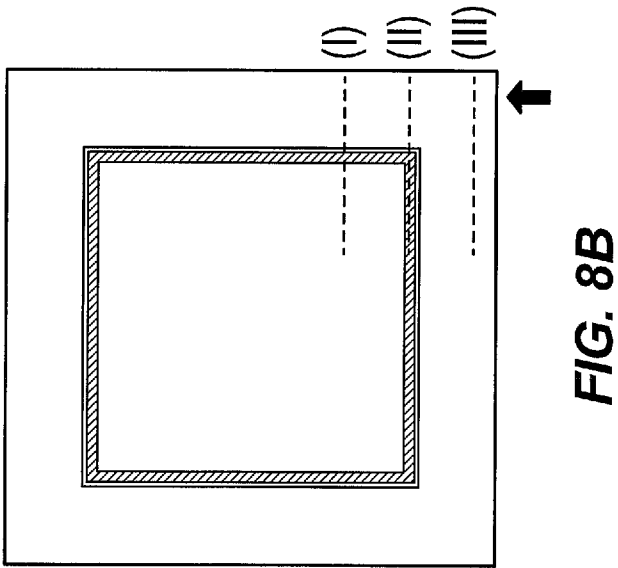
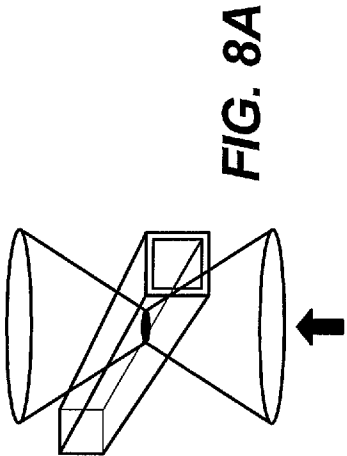
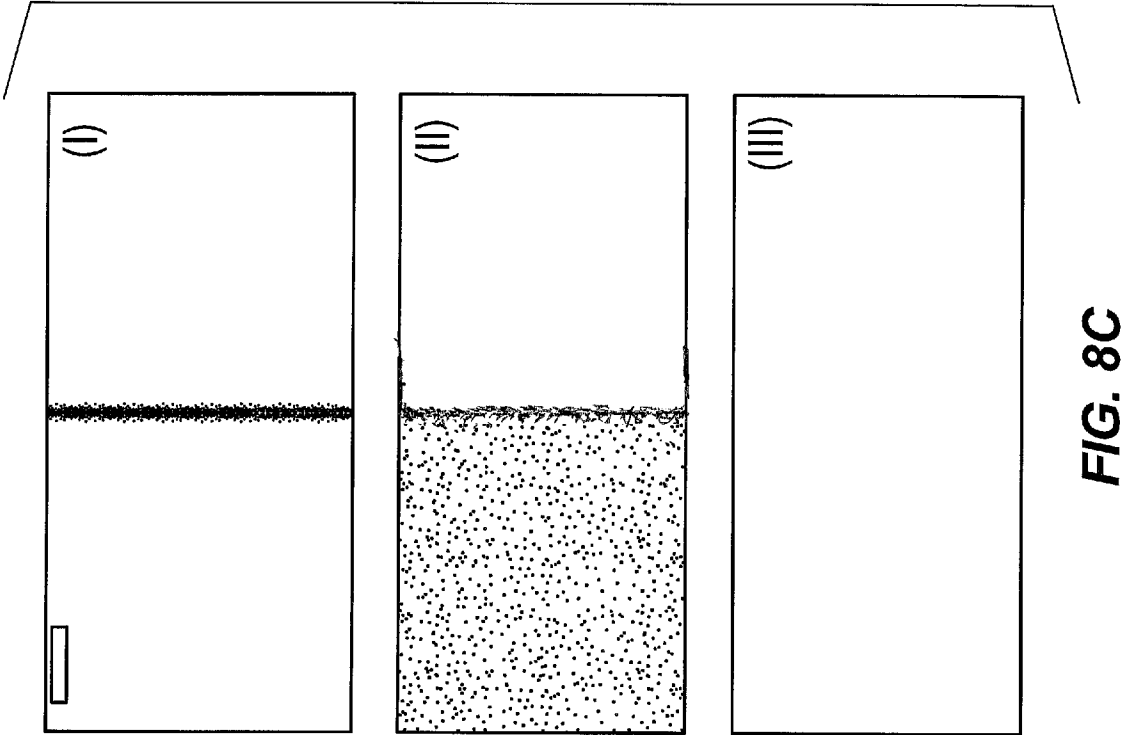
**FIG. 7B**



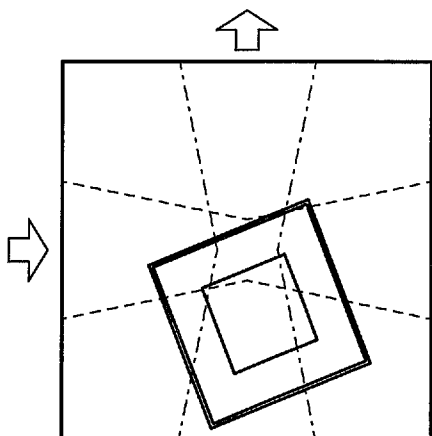
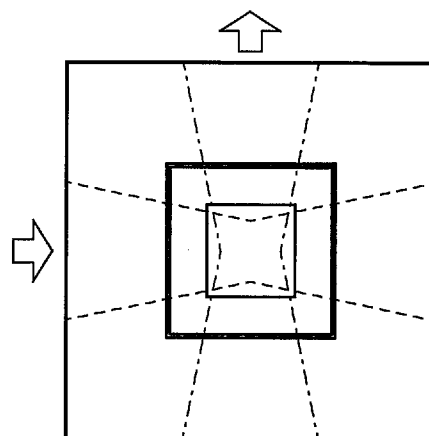
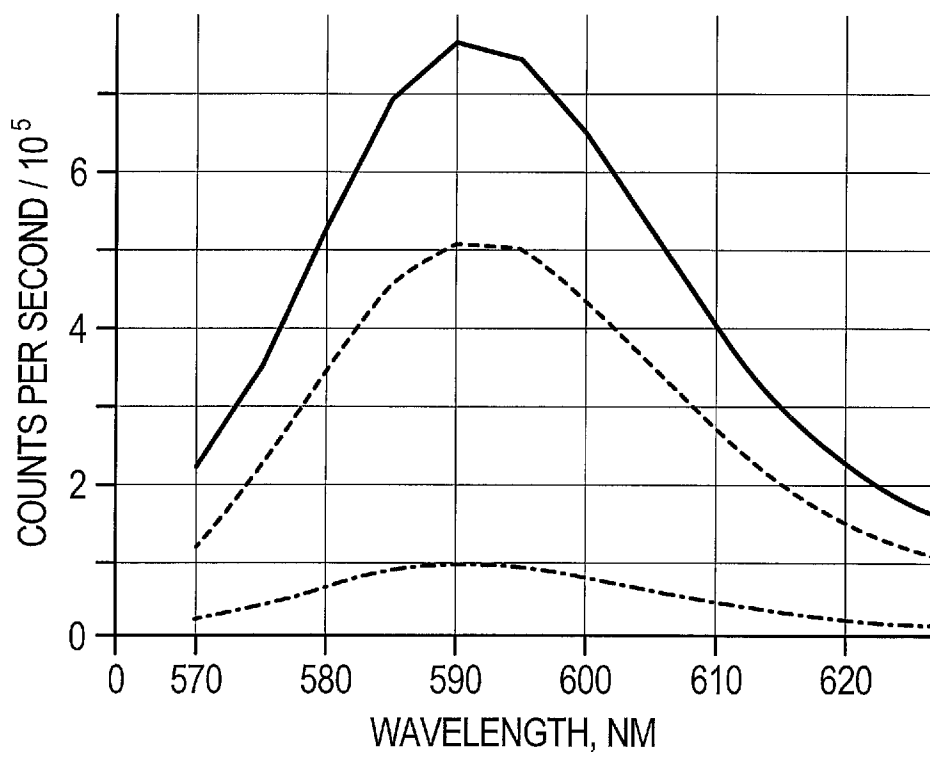
**FIG. 7C**



**FIG. 7D**



10/10

**FIG. 9A****FIG. 9B****FIG. 9C**

## INTERNATIONAL SEARCH REPORT

International application No.

PCT/US 2015/060658

## A. CLASSIFICATION OF SUBJECT MATTER

**G01J 3/02 (2006.01)**  
**G01N 21/64 (2006.01)**

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)

G01J 3/00, G01J 3/02, G01J 3/447, G01J 3/46, G01N 21/01, G01N 21/64, G01N 21/78

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

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

PatSearch (RUPTO internal), USPTO, PAJ, Esp@cenet, DWPI, EAPATIS, PATENTSCOPE, Information Retrieval System of FIPS

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	CN 102072880 A (BEIJING QINGBOHUA TECHNOLOGY CO., LTD) 25.05.2011	1-20
A	US 2010221838 A1 (UNIV CONNECTICUT) 02.09.2010	1-20
A	JPH 05126722 A (SHIMADZU CORP) 21.05.1993	1-20
A	RU 79181 U1 (FGUP NIEM) 20.12.2008	1-20
A	WO 2009/055940 A1 (VERANO FINANCIAL LTD et al.) 07.05.2009	1-20



Further documents are listed in the continuation of Box C.



See patent family annex.

* Special categories of cited documents:	"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
"A" document defining the general state of the art which is not considered to be of particular relevance	"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
"E" earlier document but published on or after the international filing date	"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
"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)	"&" document member of the same patent family
"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	

Date of the actual completion of the international search

15 March 2016 (15.03.2016)

Date of mailing of the international search report

31 March 2016 (31.03.2016)

Name and mailing address of the ISA/RU:

Federal Institute of Industrial Property,  
 Berezhkovskaya nab., 30-1, Moscow, G-59,  
 GSP-3, Russia, 125993  
 Facsimile No: (8-495) 531-63-18, (8-499) 243-33-37

Authorized officer

I. Zhestovskaya

Telephone No. (499) 240-25-91