



(43) International Publication Date
13 October 2016 (13.10.2016)

(51) International Patent Classification:
G01N 21/64 (2006.01) *G01Q 70/14* (2010.01)
G01Q 60/54 (2010.01)

(21) International Application Number:
PCT/US2016/026791

(22) International Filing Date:
8 April 2016 (08.04.2016)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
62/145,466 9 April 2015 (09.04.2015) US

(71) Applicant: **BIKANTA CORPORATION** [US/US]; 6694
Cedar Boulevard, Newark, California 94560 (US).

(72) Inventors: **BUMB, Ambika**; C/O Bikanta Corporation,
6694 Cedar Boulevard, Newark, California 94560 (US).
SARKAR, Susanta Kumar; C/O Bikanta Corporation,
6694 Cedar Boulevard, Newark, California 94560 (US).
NEUMAN, Keir Cajal; C/O Bikanta Corporation, 6694
Cedar Boulevard, Newark, California 94560 (US).

(74) Agents: **TRIMBLE, Alexander R.** et al.; Kilpatrick
Townsend & Stockton Llp, Two Embarcadero Center,
Eighth Floor, San Francisco, California 94111 (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,

[Continued on next page]

(54) Title: IMAGING SYSTEMS AND METHODS USING FLUORESCENT NANODIAMONDS

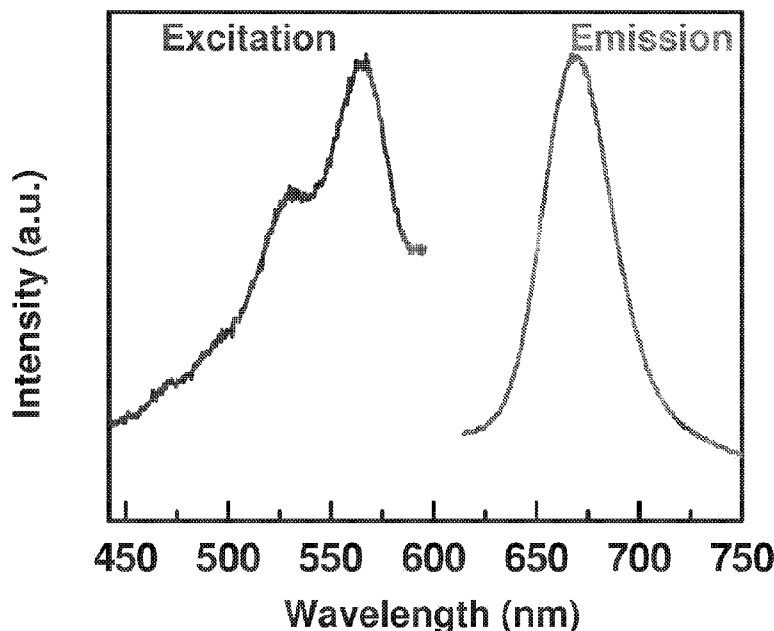


FIG. 1

(57) Abstract: Imaging systems and methods using fluorescent nanodiamonds are disclosed. The imaging systems and methods including applying a time-varying magnetic field to a specimen containing fluorescent nanodiamonds and comparing the fluorescence obtained with different magnetic fields to provide an image of the specimen.



SM, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, **Published:**
GW, KM, ML, MR, NE, SN, TD, TG).

— *with international search report (Art. 21(3))*

**IMAGING SYSTEMS AND METHODS USING
FLUORESCENT NANODIAMONDS**

CROSS-REFERENCES TO RELATED APPLICATIONS

[0001] This application claims priority to U.S. Provisional Application No. 62/145,466, filed April 9, 2015, which is incorporated in its entirety herein for all purposes.

**STATEMENT AS TO RIGHTS TO INVENTIONS MADE UNDER
FEDERALLY SPONSORED RESEARCH AND DEVELOPMENT**

[0002] The present subject matter was made with U.S. government support. The U.S. government has certain rights in this subject matter. This work was supported through grant number HL006087-02 BBC from the National Heart, Lung, and Blood Institute, National Institutes of Health.

BACKGROUND

[0003] Autofluorescence from naturally-occurring fluorescent biomolecules and fixative agents make it difficult to separate useful fluorescence from unwanted fluorescence due to overlapping emission spectra. Therefore, autofluorescence limits the capabilities of tissue and animal imaging. Even sophisticated spectral unmixing techniques cannot always reliably and accurately separate useful signal from background fluorescence.

BRIEF SUMMARY

[0004] One aspect of the invention provides an imaging method including: (a) acquiring a first fluorescent image of an object of interest impregnated with fluorescent nanodiamonds; (b) applying a magnetic field to the fluorescent nanodiamonds in order to decrease fluorescence of the fluorescent nanodiamonds; (c) acquiring a second fluorescent image of the object of interest; and (d) subtracting the second fluorescent image from the first fluorescent image to produce a resulting image.

[0005] This aspect can have a variety of embodiments. The object of interest can be a biological target. The biological target can be selected from the group consisting of: a cell, a plurality of cells, a tissue, an organ, and an organism.

[0006] The magnetic field can be generated by a permanent magnet. The magnetic field can be generated by an electromagnet.

[0007] The second fluorescent image can be acquired during application of the magnetic field.

[0008] Step (d) can be performed on a pixel-by-pixel basis.

[0009] The method can further include an additional step (e) of repeating steps (a)-(d) a plurality of times and averaging the resulting images. The plurality of times can be greater than 10.

[0010] Steps (a) and (c) can include applying an absorption wavelength to the object of interest. Another aspect of the invention provides a non-transitory computer readable medium containing program instructions executable by a processor. The computer readable medium includes: (a) program instructions that acquire a first fluorescent image of an object of interest impregnated with fluorescent nanodiamonds; (b) program instructions that apply a magnetic field to the fluorescent nanodiamonds in order to decrease fluorescence of the fluorescent nanodiamonds; (c) program instructions that acquire a second fluorescent image of the object of interest; and (d) program instructions that subtract the second fluorescent image from the first fluorescent image to produce a resulting image.

[0011] This aspect can have a variety of embodiments. The second fluorescent image can be acquired during application of the magnetic field.

[0012] Another aspect of the invention provides an imaging method including: (a) applying a time-varying magnetic field to an object of interest impregnated with fluorescent nanodiamonds to modulate the fluorescence of the fluorescent nanodiamonds; (b) acquiring a plurality of fluorescent images of the object of interest; and (c) for each corresponding pixel in the plurality of fluorescent images, calculating a fluorescence intensity using a lock-in technique.

[0013] This aspect can have a variety of embodiments. The object of interest can be a biological target. The biological target can be selected from the group consisting of: a cell, a plurality of cells, a tissue, an organ, and an organism.

[0014] The magnetic field can be generated by a permanent magnet. The magnetic field can be generated by an electromagnet.

[0015] Step (b) can include applying an absorption wavelength to the object of interest. The plurality of fluorescent images can be acquired by a wide-field camera. The plurality of fluorescent images can be acquired by a confocal microscope.

[0016] Another aspect of the invention provides a non-transitory computer readable medium containing program instructions executable by a processor. The computer readable medium includes: (a) program instructions that apply a time-varying magnetic field to an object of interest impregnated with fluorescent nanodiamonds to modulate the fluorescence of the fluorescent nanodiamonds; (b) program instructions that acquire a plurality of fluorescent images of the object of interest; and (c) program instructions that, for each corresponding pixel in the plurality of fluorescent images, calculate a fluorescence intensity using a lock-in technique.

[0017] Another aspect of the invention provides an imaging method including: (a) applying an absorption wavelength to an object of interest impregnated with fluorescent nanodiamonds; and (b) after a delay of at least at least about 6 nanoseconds, acquiring a fluorescent image of the object of interest.

[0018] This aspect can have a variety of embodiments. The object of interest can be a biological target. The biological target can be selected from the group consisting of: a cell, a plurality of cells, a tissue, an organ, and an organism.

[0019] The delay can be greater than about 10 nanoseconds. The delay can be greater than between about 10 nanoseconds and about 20 nanoseconds.

[0020] The absorption wavelength can be generated by a pulsed laser. The absorption wavelength can be between about 450 nm and about 650 nm or between about 900 nm and about 1300 nm. The absorption wavelength can be between about 450 nm and about 650 nm or between about 850 nm and about 1350 nm.

[0021] Another aspect of the invention provides an imaging method including: (a) applying an absorption wavelength to an object of interest impregnated with fluorescent nanodiamonds; and (b) after a delay of at least at least about 4 nanoseconds, acquiring a fluorescent image of the object of interest.

[0022] This aspect can have a variety of embodiments. The object of interest can be a biological target. The biological target can be selected from the group consisting of: a cell, a plurality of cells, a tissue, an organ, and an organism.

[0023] The delay can be greater than about 10 nanoseconds. The delay can be greater than between about 10 nanoseconds and about 20 nanoseconds.

[0024] The absorption wavelength can be generated by a pulsed laser. The absorption wavelength can be between about 450 nm and about 650 nm or between about 900 nm and about

1300 nm. The absorption wavelength can be between about 450 nm and about 650 nm or between about 850 nm and about 1350 nm.

[0025] Another aspect of the invention provides a non-transitory computer readable medium containing program instructions executable by a processor. The computer readable medium includes: (a) program instructions that apply an absorption wavelength to an object of interest impregnated with fluorescent nanodiamonds; and (b) program instructions that, after a delay of at least at least about 6 nanoseconds, acquire a fluorescent image of the object of interest.

[0026] In certain aspects, imaging systems are disclosed, comprising: an imaging stage for mounting a specimen; a radiation source configured to excite the triplet excited state of fluorescent nanodiamonds and directed on specimen; a fluorescence detector configured to detect triplet-triplet fluorescent nanodiamond fluorescence and configured to detect fluorescence from the specimen; and an apparatus configured to apply a time-varying magnetic field to the specimen.

BRIEF DESCRIPTION OF THE DRAWINGS

[0027] For a fuller understanding of the nature and desired objects of the present invention, reference is made to the following detailed description taken in conjunction with the accompanying drawing figures wherein like reference characters denote corresponding parts throughout the following views.

[0028] FIG. 1 depicts the absorption (excitation) and emission spectrums of diamond nitrogen-vacancy centers.

[0029] FIG. 2 depicts an imaging method according to an embodiment of the present subject matter.

[0030] FIG. 3 depicts an example of pixel-by-pixel subtraction of a second fluorescent image from a first fluorescent image according to an embodiment of present subject matter.

[0031] FIG. 4 depicts an imaging method according to another embodiment of the present subject matter.

[0032] FIG. 5 depicts an imaging method according to another embodiment of the present subject matter.

[0033] FIG. 6 depicts an imaging system according to an embodiment of the present subject matter.

[0034] FIG. 7A depicts the energy level diagram of negatively-charged nitrogen vacancy (NV) centers. FIG. 7B depicts a field of view containing FNDs. FIG. 7C depicts the intensity modulation of the FND depicted in FIG. 7A upon application of a modulating magnetic field with 0.1 Hz frequency and 100 Gauss amplitude.

[0035] FIG. 8A is a scanning confocal image of FNDs on the surface of a slide. FIG. 8B is a scanning confocal lineage taken under identical imaging conditions of the same field of view after the addition of $\sim 1 \mu\text{M}$ ALEXA FLUOR® 647 dye, which has a comparable emission spectrum to that of the FNDs. The fluorescence from the high concentration of ALEXA FLUOR® 647 dye completely obscures the fluorescence of the FNDs. FIG. 8C is a background-free lineage of the same field of view after processing images in the presence of the high ALEXA FLUOR® 647 dye background (as in FIG. 8B). The difference between pairs of images collected with and without the magnetic field was computed and 1000 of these difference images were averaged together to generate the processed image. Through this processing, images of the diamonds shown in FIG. 8A are recovered from lineages with high background like FIG. 8B.

[0036] FIG. 9A is a frame of a movie taken by a scanning confocal microscope. FIG. 9B shows the wide field background-free image after processing the movie pixel-by-pixel using a lock-in algorithm. FIG. 9C depicts the application of the lock-in algorithm to a bright pixel (horizontal coordinate, $x=109$ and vertical coordinate, $y=111$ from the top left corner of the image in FIG. 9A) corresponding to a FND. FIG. 9D illustrates the same lock-in algorithm applied to a dark pixel (horizontal coordinate, $x=137$ and vertical coordinate, $y=107$ from the top left corner of the image in FIG. 9A) corresponding to the background.

[0037] FIG. 10A depicts the fluorescence image acquired with conventional spectral unmixing techniques of the forequarters of a mouse after injection of FNDs. The image is an overlay of the background channel (top left) and the FND channel (top right). The site of injection of the FNDs is visible in the composite but the lymph node cannot be distinguished. FIG. 10B shows the same mouse imaged with the pairwise image subtraction technique. The lymph node is clearly visible in the processed image (top right) that was produced by pairwise subtraction of fluorescence images (top left) with the magnetic field on and off. FIG. 10C depicts the same images as FIG. 10B obtained with the wide-field lock-in technique. FIG. 10D depicts an image of the opened mouse chest cavity obtained with pairwise subtraction background free detection. The lymph node and the point of initial injection (white arrows) are

clearly visible. FIG. 10B depicts the same image as FIG. 10D obtained with wide-field lock in background free detection. FIG. 10F depicts the intensity as a function of time for points in the images where there are FNDs (1 and 3) and two background spots (2 and 4). The location of the four points are indicated on the images in FIGS. 10B and 10D.

[0038] FIG. 11A is an image of a lymph node (LN) injected with silica-coated FNDs without applied magnetic field. FIG. 11B is an image of the same field of view during application of a magnetic field of ~ 100 Gauss. FIG. 11C depicts the sum of 20 subtracted images of FIG. 11B from FIG. 11A. The FNDs are observed as bright spots in the otherwise dark field. FIGS. 11D and 11E depict the sums of 20 such subtractions when magnetic field was always OFF and ON, respectively.

[0039] FIG. 12A is a two-photon FLIM image of a different region of the same LN used in FIGS. 11A-11E. Longer lifetimes (indicated by red color) are believed to be due to FNDs, for which the reported lifetimes range from ~ 10 -20 ns. FIG. 12B is a background-free image of the same field of view obtained by subtracting images without and with a magnetic field, and adding 10 such subtractions.

[0040] FIG. 13 shows an imaging system including an alternating magnetic field apparatus and a total internal reflection fluorescence microscope according to certain embodiments.

[0041] FIG. 14 is a schematic showing the general components of a modulated magnetic field fluorescence imaging apparatus.

[0042] FIG. 15 shows various configurations for applying a modulated magnetic field according to certain embodiments.

[0043] FIG. 16 is a schematic showing elements of a small animal imaging instrument according to certain embodiments.

[0044] FIG. 17 is a schematic showing elements of a small animal imaging instrument modified for magnetic field modulation according to certain embodiments.

[0045] FIG. 18 is a schematic showing elements of a small animal imaging instrument for magnetic field modulation according to certain embodiments.

[0046] FIG. 19 shows a modulated magnetic field fluorescence imaging apparatus according to certain embodiments.

[0047] FIG. 20 shows a modulated magnetic field fluorescence imaging apparatus according to certain embodiments.

[0048] FIG. 21 shows a modulated magnetic field fluorescence imaging apparatus according to certain embodiments.

[0049] FIG 22 is a schematic showing elements of a hand held modulated magnetic field fluorescence imaging apparatus according to certain embodiments.

DEFINITIONS

[0050] The present subject matter is most clearly understood with reference to the following definitions:

[0051] As used herein, the singular form "a," "an," and "the" include plural references unless the context clearly dictates otherwise.

[0052] Unless specifically stated or obvious from context, as used herein, the term "about" is understood as within a range of normal tolerance in the art, for example within 2 standard deviations of the mean. "About" can be understood as within 10%, 9%, 8%, 7%, 6%, 5%, 4%, 3%, 2%, 1%, 0.5%, 0.1%, 0.05%, or 0.01% of the stated value. Unless otherwise clear from context, all numerical values provided herein are modified by the term about.

[0053] As used herein, the term "background-free image" refers to an image having a sufficiently high signal-to-noise ratio (SNR) such that the fluorescence of fluorescent nanodiamonds can be distinguished from the fluorescence of background elements such as endogenous proteins. Suitable SNRs include those greater than about 1:1, about 1.5:1, about 2:1, about 3:1, about 4:1, about 5:1, about 6:1, about 7:1, about 8:1, about 9:1, and about 10:1.

[0054] As used herein, the terms "comprises," "comprising," "containing," "having," and the like can have the meaning ascribed to them in U.S. patent law and can mean "includes," "including," and the like.

[0055] As used herein, the term "fluorescent image" refers to an image representing one or more fluorescent emissions. For example, the fluorescent emissions can be between about 625 nm and about 825 nm. Fluorescent images are typically obtained typically obtained by applying an absorption wavelength to an object of interest and simultaneously or after a delay, capturing an image of fluorescent emissions. Fluorescent images can be obtained using a variety of devices including fluorescent microscopes. The fluorescent image can, in some embodiments, be a two-dimensional image consisting of a plurality of pixels, each of which can be a numerical representation of the intensity of fluorescence as at particular location.

[0056] As used herein, the term "fluorescent nanodiamond" (abbreviated as "FND") refers to nanodiamonds that exhibit fluorescence when exposed to an appropriate absorption (excitation) spectrum. This fluorescence can be caused by the presence of nitrogen-vacancy (NV) centers, where a nitrogen atom is located next to a vacancy in the nanodiamond. The absorption (excitation) and emission spectrums of silica-coated FNDs having a diameter of about 100 nm are depicted in FIG. 1. The absorption (excitation) spectrum generally lies between about 450 nm and about 600 nm with an excitation peak at about 565 nm. (FNDs can also be excited by a two-photon process in the wavelength region between about 900 nm and about 1300 nm.) The emission spectrum generally lies between about 625 nm and about 750 nm, with an emission peak at about 700 nm. (In the working examples described herein, the samples were excited at 575 nm and a PTI fluorometer was used to obtain the excitation spectra.)

[0057] As used herein, the term "nanodiamond" refers to diamonds having a largest dimension of less than about 100 nm. For example, the largest dimension of a nanodiamond can be less than: about 100 nm, about 90 nm, about 80 nm, about 70 nm, about 60 nm, about 50 nm, about 40 nm, about 30 nm, about 20 nm, about 20 nm, about 10 nm, and the like.

[0058] As used herein, the term "object of interest" refers to any object of which a fluorescent image is desired. An object of interest can be a biological target, for example, a living organism or a sample of an organism. The object of interest can be an organism, one or more organs, one or more tissues, and/or one or more cells. Although the examples described herein are largely directed toward biological imaging, FNDs and the methods described herein can be used to increase the signal to noise in any imaging application in which a modulated magnetic field can be applied to the sample. For example, FNDs and the methods described herein can be used to image and study flow or morphology in a high background environment.

[0059] Unless specifically stated or obvious from context, the term "or," as used herein, is understood to be inclusive.

[0060] By "specifically binds" is meant recognition and binding to a target (*e.g.*, polypeptide, cell, surface antigen, and the like), but which does not substantially recognize and bind other molecules in a sample, for example, a biological sample.

[0061] The term "subject" as used herein, refers to any organism that is suitable for being imaged by the methods described herein. Such organisms include, but are not limited to,

human, dog, cat, horse, cow, sheep, goat, mouse, rat, guinea pig, monkey, avian, reptiles, bacteria, fungi, viruses, and the like.

[0062] The term "tissue" as used herein, refers to a subject's body. Nonlimiting examples of tissues include tissues from organs such as brain, heart, lung, liver, stomach, pancreas, colon, rectum, intestines, blood vessels, arteries, and the like.

[0063] Ranges provided herein are understood to be shorthand for all of the values within the range. For example, a range of 1 to 50 is understood to include any number, combination of numbers, or sub-range from the group consisting 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38, 39, 40, 41, 42, 43, 44, 45, 46, 47, 48, 49, or 50 (as well as fractions thereof unless the context clearly dictates otherwise).

DETAILED DESCRIPTION

[0064] Nitrogen vacancy centers in nanodiamonds are unique fluorescent sources that do not photobleach or blink. Remarkably, the fluorescence intensity of these fluorescent nanodiamonds can be modulated by a magnetic field of moderate strength (~ 0.01 T). Furthermore the fluorescence lifetime of nanodiamonds (~ 10 - 20 ns) is longer than the lifetime (≤ 5 - 6 ns) of most fluorophores that contribute to auto fluorescence. Both of these properties of nanodiamonds can be used to achieve background-free imaging. The modulation of the fluorescence intensity with applied magnetic field is a unique feature of the fluorescent nanodiamonds, which in combination with their other features enables a number of novel imaging applications. Background-free imaging is one of these applications.

[0065] The discovery allows background-free imaging of fluorescent nanodiamonds in tissue samples and *in vivo*, where conventional imaging is difficult due to background fluorescence. We present several techniques to reduce or eliminate background fluorescence by exploiting properties of the fluorescent nanodiamonds. In particular, magnetic field modulation of the fluorescence intensity offers a simple, robust, and easily adaptable method to obtain background-free imaging in a variety of imaging modalities, i.e., fluorescence microscopy, confocal fluorescence microscopy, and wide-field fluorescence animal imaging.

[0066] In one embodiment of the present subject matter, subtracting an image acquired with the magnetic field from one without the field collected under otherwise identical conditions

eliminates constant background fluorescence while highlighting the diamond fluorescence that is specifically reduced in one image.

[0067] In another embodiment, the field is modulated sinusoidally while images are acquired. Phase-sensitive detection of the modulated intensity can then be achieved by post-processing for camera-based imaging or through lock-in techniques in confocal-based imaging. This technique could be adapted for use in wide-field and confocal imaging systems. Importantly, this technique makes use of conventional continuous wave illumination.

[0068] Yet another embodiment makes use of the long excited state lifetime of the fluorescent nanodiamonds to reject shorter-lived background fluorescence. This technique relies on a pulsed laser and time-gated or lifetime imaging. Fluorescent nanodiamonds can be imaged with two-photon approaches, facilitating these lifetime-based background rejection techniques.

[0069] Referring now to FIG. 2, an imaging method 200 is provided.

[0070] In step S202, a first fluorescent image is acquired of an object of interest impregnated with fluorescent nanodiamonds. The fluorescent image can be acquired using conventional fluorescent imaging devices as described herein.

[0071] In step S204, a magnetic field is applied to the fluorescent nanodiamonds in order to decrease fluorescence of the fluorescent nanodiamonds. The magnetic field can be applied, for example, with a permanent magnet or an electromagnet. The magnetic field modulates the fluorescence intensity of the fluorescent nanodiamonds such that the fluorescence intensity of the fluorescent nanodiamonds is less than in the first fluorescent image.

[0072] In step S206, a second fluorescent image is acquired of the object of interest. Preferably, the second fluorescent image has identical parameters to the first fluorescent image other than the decreased fluorescence intensity of fluorescent nanodiamonds as a result of the application of the magnetic field. The second fluorescent image is acquired while the magnetic field is applied.

[0073] In step S208, the second fluorescent image is subtracted from the first fluorescent image to produce a resulting image. The subtraction can be performed on a pixel-by-pixel basis. For example, as shown conceptually in FIG. 3, a second 10x10 pixel fluorescent image can be subtracted from a first 10x10 pixel fluorescent image to produce a resulting image. Because the only difference between the first fluorescent image and the second fluorescent image is the

modulation of fluorescent nanodiamonds, the background fluorescence (*e.g.*, from endogenous proteins) will be canceled out by the subtraction to produce a background-free fluorescent image.

[0074] Pixel-by-pixel subtraction can be performed manually or can be automated. A variety of commercially-available computer programs can perform image subtraction including, for example, MATLAB® software available from The MathWorks, Inc. of Natick, Massachusetts and IMAGEJ software available from the National Institutes of Health at <http://rsbweb.nih.gov/ij/>.

[0075] In step S210, steps S202-S208 are repeated a plurality of times (*e.g.*, greater than 10) and the resulting images are averaged on a pixel-by-pixel basis to improve the imaging quality.

[0076] Referring now to FIG. 4, another imaging method 400 is provided.

[0077] In step S402, time-varying magnetic field is applied to an object of interest impregnated with fluorescent nanodiamonds to modulate the fluorescence of the fluorescent nanodiamonds. The time-varying magnetic field can vary cyclically. For example, the magnitude of the magnetic field can vary sinusoidally. The magnetic field can be varied by modulating the current applied to an electromagnet or by modulating the distance between the magnet (either a permanent magnet or an electromagnet) and the object of interest.

[0078] In step S404, a plurality of fluorescent images of the object of interest are acquired. Preferably, the plurality of fluorescent images are captured at a plurality of points within a cycle. The sampling can occur at regular or irregular intervals and can, but need not, be matched to the frequency of the phase-modified magnetic field. The plurality of fluorescent images can be acquired pixel-by-pixel using a point imager such as a confocal microscope or multiple pixels at a time using a wide-field imager.

[0079] In step S406, the fluorescence intensity of each pixel is calculated using a lock-in technique. Lock-in techniques multiply the fluorescence intensity in each pixel of the plurality of fluorescent images by the amplitude of the magnetic field at the time of each respective fluorescent image and then calculate the intensity of the appropriately filtered or processed resulting products. All background fluorescence will not fluctuate in phase with the magnetic field modulation over all images and will therefore average to zero. The fluorescence intensity of the fluorescent nanodiamonds will vary in phase with the magnetic intensity and, therefore, average to half of the amplitude of the magnetic intensity.

[0080] Lock-in techniques are described in publications such as Richard Burdett, "Amplitude Modulated Signals - The Lock-in Amplifier," in *Handbook of Measuring System Design* (2005) and Stanford Research Systems, Inc., "About Lock-In Amplifiers: Application Note #3," available at <http://www.thinksrs.com/downloads/PDFs/ApplicationNotes/AboutLIAs.pdf>.

Generally speaking, single-pixel inputs obtained from point detectors can be processed directly by conventional lock-in amplifiers available from suppliers such as Stanford Research Systems, Inc. of Sunnyvale, California, while algorithms can be written using software such as MATLAB® to step through each set of corresponding pixels in a series of fluorescent images and perform a lock-in technique to produce a resulting background-free image.

[0081] If the fluorescent images are acquired pixel-by-pixel, a lock-in technique can be applied for that particular pixel before acquiring another pixel. If the images are acquired using a wide-field imager, a lock-in technique can be applied on a pixel-by-pixel basis after all images are acquired.

[0082] Referring now to FIG. 5, another imaging method 500 is provided.

[0083] In step S502, an absorption wavelength (*e.g.*, a brief pulse of 5 ns or less) is applied to an object of interest impregnated with fluorescent nanodiamonds. This absorption wavelength will excite FNDs, but may also excite various background elements such as endogenous proteins.

[0084] In step S504, after a delay of at least at least about 6 nanoseconds, acquiring a fluorescent image of the object of interest. After 6 nanoseconds, most (if not all) background fluorescence will have dissipated, while the FNDs continue to emit photons. Thus, a background-free image can be captured without the need for the image processing algorithms described above.

[0085] Referring now to FIG. 6, the methods described herein can be implemented in hardware and/or software. For example, FIG. 6 depicts a system 600 including a fluorescent imaging device 602 such as a fluorescent microscope and a computer 604. The fluorescent imaging device 602 can include an excitation light source, an image sensor (*e.g.*, including a charge-coupled device (CCD), complementary metal-oxide-semiconductor (CMOS) chip, photomultiplier tube (PMT), or avalanche photodiode (APD)), one or more lenses, and a filter adapted to block undesired wavelengths. The computer 604 can be a special-purpose or general-purpose computer can be communicatively coupled with the fluorescent imaging device 602 via

communication standards such as parallel or serial ports, Universal Serial Bus (USB), USB 2.0, Firewire, Ethernet, Gigabit Ethernet, and the like.

[0086] As understood by those of skill in the art, computer 604 can include various components such as a display device, a processor, and/or a storage device.

[0087] Display device can be any device capable of displaying graphics and/or text. Examples of display devices include a cathode ray tube (CRT), a plasma display, a liquid crystal display (LCD), an organic light-emitting diode display (OLEO), a light-emitting diode (LED) display, an electroluminescent display (ELD), a surface-conduction electron-emitter display (SED), a field emission display (FED), a nano-emissive display (NED), an electrophoretic display, a bichromal ball display, an interferometric modulator display, a bistable nematic liquid crystal display, and the like.

[0088] Processor is an electronic device (also known as a central processing unit or microprocessor) capable of executing instructions stored as hardware and/or software. Suitable processors are available from manufacturers such as Intel Corporation of Santa Clara, California or Advanced Micro Devices (AMD) of Sunnyvale, California.

[0089] Storage device can include persistent storage devices such as magnetic media (*e.g.* tapes, disks), optical media (*e.g.* CD-ROM, CD-R, CD-RW, DVD, HD DVD, BLU-RAY DISK®, Laserdisk), punch cards, and the like. Storage device can also include temporary storage devices known as memory (*e.g.*, random access memory).

[0090] In some embodiments, motion correction is applied to compensate for motion in the camera and/or the subject. For example, motion correction can be applied to a time series of images to correct for in-plane motion at the sub-pixel level.

[0091] In one embodiment, a non-rigid deformation map is calculated pairwise between each image and a common reference image using an optical flow method, which iteratively maximizes the local cross-correlation image subsets at different resolutions. The initial time frame can be chosen as a common reference for both images series acquired with external field ON and OFF in order so that the ON and OFF images are co-registered. A subpixel spline based interpolation can be used for application of the non-rigid deformation to minimize loss of spatial resolution. Following motion correction, the time series of images can be averaged to reduce random fluctuation due to noise thereby creating an average image for the ON and OFF. A difference image between ON and OFF motion corrected averages can be used to analyze the contrast of the

probe signal by subtraction of the background tissue. This signal from the nanodiamond can then be overlaid on the starting image.

[0092] As described herein, the present invention relates to use of fluorescent nanodiamonds to image an object of interest. Fluorescent nanodiamonds suitable for use in the present invention and methods for making such fluorescent molecules are well-known in the art. *see, e.g., Nanodiamonds: Applications in Biology and Nanoscale Medicine* (D. Ho ed., Springer 2009); *Molecular Imaging* (R. Weissleder *et al.* eds., 2010); *Medical Nanotechnology and Nanomedicine (Perspectives in Nanotechnology)* (H. F. Tibbals ed., CRC Press 2010); *Molecular Fluorescence* (B. Valeur ed., Wiley-VCH 2012); *Introduction to Nanomedicine and Nanobioengineering (Wiley Series in Biomedical Engineering and Multi-Disciplinary Integrated Systems)* (P. N. Prasad ed., Wiley 2012); Yu *et al.*, *J. Am. Chem. Soc.* 127:17604-17605 (2005); Mochalin *et al.*, *Nature Nanotechnology* 7:11-23 (2012); Epstein *et al.*, *Nature Physics* 1:94-98 (2005); Chow *et al.*, *Sci. Transl. Med.* 3:73ra21 (2011); Awschalom *et al.*, *Sci. Am.* 297:84-91 (2007); and Wilson, *Phys. Today* 64:17-18 (2011).

[0093] The fluorescent nanodiamonds can be provided as a solution, emulsion, suspension, microsphere, particle, microparticle, nanoparticle, liposomes, and the like.

[0094] In aspects of the invention, the fluorescent nanodiamonds are directly contacted with a sample (*e.g.*, when impregnating the fluorescent nanodiamond in a sample *in vitro*, *ex vivo*, *in situ*, etc.). For example, fluorescent nanodiamonds can be impregnated in a sample by injection (*e.g.*, microinjection) or use of a delivery vehicle. Suitable delivery vehicles are well-known in the art. Nonlimiting examples include lipid vesicles or other polymer carrier materials, lipoplexes (*see, e.g.*, U.S. Patent Application Publication No. 2003/0203865; and Zhang *et al.*, *J. Control Release*, 100:165-180 (2004)), pH-sensitive lipoplexes (*see, e.g.*, U.S. Patent Application Publication No. 2002/0192275), reversibly-masked lipoplexes (*see, e.g.*, U.S. Patent Application Publication Nos. 2003/0180950), cationic lipid-based compositions (*see, e.g.*, U.S. Patent No. 6,756,054; and U.S. Patent Application Publication No. 2005/0234232), cationic liposomes (*see, e.g.*, U.S. Patent Application Publication Nos. 2003/0229040, 2002/0160038, and 2002/0012998; U.S. Patent No. 5,908,635; and International Publication No. WO 01/72283), anionic liposomes (*see, e.g.*, U.S. Patent Application Publication No. 2003/0026831), pH-sensitive liposomes (*see, e.g.*, U.S. Patent Application Publication No. 2002/0192274; and Australian Publication No. 2003/210303), antibody-coated liposomes (*see, e.g.*, U.S. Patent

Application Publication No. 2003/0108597; and International Publication No. WO 00/50008), cell-type-specific liposomes (*see, e.g.*, U.S. Patent Application Publication No. 2003/0198664), liposomes containing nucleic acid and peptides (*see, e.g.*, U.S. Patent No. 6,207,456), liposomes containing lipids derivatized with releasable hydrophilic polymers (*see, e.g.*, U.S. Patent Application Publication No. 2003/0031704), lipid-entrapped molecules (*see, e.g.*, International Publication Nos. WO 03/057190 and WO 03/059322), lipid-encapsulated molecules (*see, e.g.*, U.S. Patent Application Publication No. 2003/0129221; and U.S. Patent No. 5,756,122), other liposomal compositions (*see, e.g.*, U.S. Patent Application Publication Nos. 2003/0035829 and 2003/0072794; and U.S. Patent No. 6,200,599), stabilized mixtures of liposomes and emulsions (*see, e.g.*, European Publication No. EP 1 304 160), emulsion compositions (*see, e.g.*, U.S. Patent No. 6,747,014), and micro-emulsions (*see, e.g.*, U.S. Patent Application Publication No. 2005/0037086).

[0095] In some aspects of the invention, the fluorescent nanodiamonds are administered *in vivo* to a subject (*e.g.*, delivering the fluorescent nanodiamond to a target object within the subject, including, but not limited to, a target cell, tissue, organ, organism, infectious agent, a virus, a bacteria, fungus, a parasite, and the like). When administered to a subject, the fluorescent nanodiamonds can be provided as a pharmaceutical composition including a pharmaceutically acceptable carrier. The term "pharmaceutically acceptable" means approved by a regulatory agency or listed in the U.S. Pharmacopeia or other generally recognized pharmacopeia for use in animals, and more particularly in humans. The term "carrier" refers to a diluent, adjuvant, excipient, or vehicle with which the therapeutic is administered. Such pharmaceutical carriers can be sterile liquids, such as water and oils, including those of petroleum, animal, vegetable or synthetic origin, such as peanut oil, soybean oil, mineral oil, sesame oil, olive oil, gel (*e.g.*, hydrogel), and the like. Saline is an exemplary carrier when the pharmaceutical composition is administered intravenously. Saline solutions and aqueous dextrose and glycerol solutions can also be employed as liquid carriers, particularly for injectable solutions.

[0096] Suitable pharmaceutical excipients include starch, glucose, lactose, sucrose, gelatin, malt, rice, flour, chalk, silica gel, sodium stearate, glycerol monostearate, talc, sodium chloride, dried skim milk, glycerol, propylene, glycol, water, ethanol, and the like. The composition, if desired, can also contain minor amounts of wetting or emulsifying agents, or pH buffering

agents. These compositions can take the form of solutions, suspensions, emulsion, tablets, pills, capsules, powders, sustained-release formulations and the like. Oral formulation can include standard carriers such as pharmaceutical grades of mannitol, lactose, starch, magnesium stearate, sodium saccharine, cellulose, magnesium carbonate, and the like. Examples of suitable pharmaceutical carriers are described in "Remington's Pharmaceutical Sciences" by E. W. Martin, the contents of which are hereby incorporated by reference in its entirety.

[0097] In embodiments, the imaging agent may be administered through different routes, including, but not limited to, oral, parenteral, buccal and sublingual, rectal, aerosol, nasal, intramuscular, subcutaneous, intradermal, and topical. The term parenteral as used herein includes, for example, intraocular, subcutaneous, intrapleural, intracutaneous, intravenous, intramuscular, intraarticular, intraarterial, intrasynovial, intrasternal, intrathecal, intralésional, and intracranial injection, or other infusion techniques. One of ordinary skill in the art would readily understand that the formulation should suit the mode of administration.

[0098] Formulations suitable for administration include aqueous and non-aqueous sterile solutions, which may contain anti-oxidants, buffers, bacteriostats and solutes which render the formulation isotonic with the blood of the intended recipient, and aqueous and non-aqueous sterile suspensions which may include suspending agents and thickening agents. The formulations may be presented in unit-dose or multi-dose containers, for example, sealed ampoules and vials, and may be stored in a freeze-dried (lyophilized) condition requiring only the addition of the sterile liquid carrier, for example, water, immediately prior to use. Extemporaneous solutions and suspensions may be prepared from sterile powders, granules and tablets commonly used by one of ordinary skill in the art.

[0099] For oral administration in the form of a tablet or capsule, the fluorescent nanodiamonds can be combined with an oral, non-toxic, pharmaceutically acceptable, inert carrier such as lactose, starch, sucrose, glucose, methyl cellulose, magnesium stearate, dicalcium phosphate, calcium sulfate, mannitol, sorbitol and the like. For oral administration in liquid form, the fluorescent nanodiamonds can be combined with any oral, non-toxic, pharmaceutically acceptable inert carrier such as ethanol, glycerol, water, and the like. Moreover, when desired or necessary, suitable binders, lubricants, disintegrating agents, and coloring agents can also be incorporated into the mixture. Suitable binders include starch, gelatin, natural sugars such as glucose or β -lactose, corn sweeteners, natural and synthetic gums such as acacia, tragacanth, or

sodium alginate, carboxymethylcellulose, polyethylene glycol, waxes, and the like. Lubricants used in these dosage forms include sodium oleate, sodium stearate, magnesium stearate, sodium benzoate, sodium acetate, sodium chloride, and the like. Disintegrators include, without limitation, starch, methyl cellulose, agar, bentonite, xanthan gum, and the like.

[0100] The fluorescent nanodiamonds used in the methods of the present invention can also be administered in the form of liposome delivery systems. Such delivery systems are well known in the art, and include but are not limited to, unilamellar vesicles, large unilamellar vesicles, and multilamellar vesicles. Liposomes can be formed from a variety of phospholipids, such as cholesterol, stearylamine, or phosphatidylcholines.

[0101] The administration of the fluorescent nanodiamonds to a subject can be by a general or local administration route. For example, the fluorescent nanodiamonds may be administered to the subject such that it is delivered throughout the body. Alternatively, the fluorescent nanodiamonds can be administered to a specific organ or tissue of interest.

[0102] The fluorescent nanodiamonds should have sufficient emission to assure reliable diagnosis. The amount of fluorescent nanodiamonds to be contacted with a sample or introduced into a subject in order to provide for detection can readily be determined by those skilled in the art. For example, increasing amounts of the fluorescent nanodiamonds can be applied or given to a subject until the fluorescent nanodiamonds are detected by the detection method of choice. In addition, those skilled in the art are also familiar with determining the amount of time sufficient for the fluorescent nanodiamonds to become associated with a target object. The amount of time necessary can easily be determined by introducing a detectable amount of the fluorescent nanodiamonds into a subject and then detecting the fluorescent nanodiamonds at various times after administration.

[0103] In some aspects, the fluorescent nanodiamonds can be associated with a molecule that preferentially binds to the target object. Such molecules are well-known in the art, and include but are not limited to target-binding agents having one or more target recognition moieties for the selective binding of the target-binding agents (and fluorescent nanodiamond) to a target molecule. The target recognition moiety is configured to specifically bind to a target molecule of a particular cell, tissue, organ, receptor, surface antigen, organism/infectious agent, and the like.

[0104] Examples of target recognition moieties include, but are not limited to, an antigen, ligand, receptor, one member of a specific binding pair, polyamide, peptide, carbohydrate,

oligosaccharide, polysaccharide, low density lipoprotein (LDL) or an apoprotein of LDL, steroid, steroid derivative, hormone, hormone-mimic, lectin, drug, antibiotic, aptamer, DNA, RNA, lipid, an antibody, an antibody-related polypeptide, and the like. In embodiments, the targeting moieties are polypeptides, carbohydrates, or lipids. The targeting moieties can also be antibodies, antibody fragments, or nanobodies. In other embodiments, the target recognition moiety can be a molecule or a macromolecular structure (*e.g.*, a liposome, a micelle, a lipid vesicle, or the like) that preferentially associates or binds to a particular tissue, receptor, organism/infectious agent, and the like.

[0105] One of ordinary skill in the art would readily understand how to make the fluorescent nanodiamond conjugates contemplated herein. For example, the fluorescent nanodiamonds can be covalently or non-covalently associated with the target binding agent/moiety. See Vijayanthimalal *et al.*, *Nanomedicine* 4:47-55 (2009); Vijayanthimalal *et al.*, *Biomaterials* 33: 7794-7802 (2012); Hartmann *et al.*, *Chemistry - A European Journal* 18:21, 6485-6492 (2012); Mochalin *et al.*, *Nat. Nanotechnology* 7:11-23 (2012); Weng *et al.*, *Diamond and Related Materials* 22:96-104 (2012); Alhaddad *et al.*, *Small* 7:3087-3095 (2011); Krueger, *J Mater. Chem.* 21:12571-12578 (2011); and Liu *et al.*, *Nanoscale Research Letters* 5:1045-1050 (2010); Rurack, *Supramolecular Chemistry Meets Hybrid (Nano)Materials: A Brief Look Ahead*, pages 689-700 of *The Supramolecular Chemistry of Organic-Inorganic Hybrid Materials* (Wiley, 2010).

[0106] In some embodiments, the FNDs are silica-coated FNDs. Methods for creating silica-coated nanodiamonds are described in A. Bumb *et al.*, "Silica encapsulation of fluorescent nanodiamonds for colloidal stability and facile surface functionalization," 135 *Journal of the American Chemical Society* 7815-18 (2013).

Working Example # 1 – Magnetic Modulation of FND Emission

[0107] FIG. 7A depicts the energy level diagram of NV⁻ centers in diamond showing spin-triplet ($m_s = 0$ and $m_s = \pm 1$) ground and excited states as well as the singlet metastable state. NV⁻ centers can be optically excited over a broad range of wavelengths (450-650 nm) (green arrows). NV⁻ centers in the $m_s = \pm 1$ sublevels of the excited states have a higher probability to decay via the metastable state (grey dashed arrows) than to the $m_s = \pm 1$ sublevels of the ground state. From the metastable state, NV⁻ centers predominantly transition to the $m_s = 0$ sublevel of the ground state without emitting visible light. Therefore, in the absence of a magnetic field, NV⁻

centers are rapidly pumped into the $m_s = 0$ sublevel of the ground state when excited. This results in an initial increase in fluorescence emission intensity as steady state is reached. In the presence of a magnetic field, the $m_s = 0$ and $m_s = \pm 1$ states are mixed, making the decay pathway through the metastable singlet state accessible and therefore decreasing the fluorescence emission intensity.

[0108] FIG. 7A depicts a field of view containing FNDs. FIG. 7B depicts the intensity modulation of the FND depicted in FIG. 7A upon application of a modulating magnetic field with 0.1 Hz frequency and 100 Gauss amplitude.

[0109] To show magnetic modulation of FND emission, a coverslip was prepared with FNDs. 500 μ l of 1 mg/ml poly-L-lysine (PLL) in PBS buffer was mixed with silica-coated FNDs, deposited on a #1 coverslip, and incubated overnight. A flow cell was made using double-sided tape to attach the coverslip to a glass slide.

[0110] Movies were obtained using a CARL ZEISS® LSM5 LIVE microscope. Each frame of the movie was a scanning confocal image with 250 ms time resolution. A 10X ZEISS® objective with 0.3 NA (EC Plan-Neofluar) was used to introduce the excitation light to stimulate the FNDs and to collect the FND emission. Samples were excited at 532 nm, emission was filtered using a long pass filter LP650 and detected using a photomultiplier tube (PMT).

[0111] An electromagnet (APW Company, Item# EM400-12-212, 4.0" Diameter Round Electromagnet) was powered by square wave voltage signal with an amplitude of 0 or 12V and zero offset. The sample was placed ~13 mm away from the face of the magnet, where the magnetic field strength was ~100 Gauss.

Working Example #2 – Background-Free Imaging by Pairwise Subtraction of Frames With and Without Magnetic Field

[0112] FIG. 8A is an image of a field of view with ~40 nm FNDs containing ~15 NV-imaged as in FIG. 7B. FIG. 8B is an image of the same field of view after introducing ~1 μ M ALEXA FLUOR® 647 dye solution into the flow cell. This dye has similar emission characteristics as the FNDs so the fluorescence from the FNDs is masked by the background fluorescence of high concentration of the ALEXA FLUOR® 647 dye. FIG 8C is an image of the same field of view after processing images in the presence of the high ALEXA FLUOR® 647 dye background (as in FIG 8B). The difference between pairs of images collected with and without the magnetic field was computed and 1,000 of these difference images were averaged

together to generate the processed image. Through this processing, images of the diamonds shown in FIG. 8A are recovered from images with high background like FIG. 8B.

Working Example #3 – Background-Free Imaging Using Wide-Field Lock-In Detection

[0113] FIG. 9A shows a frame of a movie (1000 frames with time resolution 0.25 s) taken by a scanning confocal microscope. The details of the imaging are identical to those in FIG. 7B above. A modulating magnetic field with 0.1 Hz frequency and 100 Gauss amplitude was applied during the acquisition of the movie.

[0114] FIG. 9B shows the wide field background-free image after processing the movie pixel-by-pixel using a lock-in algorithm.

[0115] FIG. 9C depicts the application of the lock-in algorithm to a bright pixel (horizontal coordinate, $x=109$ and vertical coordinate, $y=111$ from the top left corner of the image in FIG. 9A) corresponding to a FND. The top left panel of FIG. 9C illustrates the pixel values as a function of time. The top right panel of FIG. 9C illustrates the fast Fourier transform (FFT) as a function of frequency. The middle left panel of FIG. 9C depicts the pixel values from the top left panel multiplied by a reference sine wave $1+\sin(2\pi \times 0.1 \times t)$. The middle right panel of FIG. 9C depicts the corresponding FFT. The bottom left panel of FIG. 9C shows the pixel values multiplied with a reference cosine wave $1+\cos(2\pi \times 0.1 \times t)$. The bottom right panel of FIG. 9C depicts the corresponding FFT.

[0116] FIG. 9D illustrates the same lock-in algorithm applied to a dark pixel (horizontal coordinate, $x=137$ and vertical coordinate, $y=107$ from the top left corner of the image in FIG. 9A) corresponding to the background.

[0117] Referring to both FIGS. 9C and 9D, vertical dashed lines at 0.2 Hz are guides to indicate the values at twice the reference frequency. Means of three points around 0.2 Hz in the sine and cosine FFTs were calculated for each pixel. The background-free image in FIG. 9B corresponds to the sine and cosine means added in quadrature, *i.e.*, $i=\sqrt{(i\cos^2+i\sin^2)}$, where i is the mean intensity and $i\cos$ and $i\sin$ correspond to the means calculated from the FFT of the signal multiplied by the cosine and sine functions respectively. Means of the pixel values over 1000 frames were 173 and 18 for the pixels at (109,111) and (137,107) before applying the lock-in algorithm giving a signal-to-noise ratio of ~ 10 . Means of the FFT amplitudes for three points around 0.2 Hz are 21.42 and 0.28 for the pixels at (109,111) and (137,107) after applying the lock-in algorithm giving a signal-to-noise ratio of ~ 77 . Thus the lock-in algorithm increased the

signal to noise ratio by a factor of ~ 8 . Computations were implemented in MATLAB® software.

Working Example# 4 – Background-Free Imaging of Sentinel Lymph Node In Vivo Using Wide Field Pairwise Image Subtraction and Lock-In Detection

[0118] FIG. 10A depicts an image of the forequarters of a mouse obtained using conventional spectral unmixing methods to separate the emission of the FNDs (top right inset, red in overlay) from background fluorescence (top left inset, white in overlay). FNDs were not detected through the skin in the draining axillary lymph node. FIG. 10B depicts an image of the same mouse obtained by averaging 475 pairwise-subtracted images with and without the magnetic field. The processed image (top right inset, red in overlay) was overlaid on an unprocessed image obtained with the magnetic field off (top left inset, white in overlay). The white arrows point to the injection site in the footpad and the location of the auxiliary (sentinel) lymph node. Signal from the FNDs in the lymph node is clearly detected. FIG. 10C depicts an image obtained by lock-in detection of emission from FNDs from the same images used to generate FIG. 10B using the algorithm described in FIG. 9 (top right inset, red in overlay) overlaid on the unprocessed image obtained with the magnetic field off (top left inset, white in overlay). FIG. 10D depicts an image of the same mouse's open chest cavity by averaging of images obtained by pairwise subtracting images with and without the magnetic field. The processed image (bottom inset, red in overlay) was overlaid on an unprocessed image obtained with the magnetic field off (top inset, white in overlay). The white arrows point to the injection site in the front footpad and the location of the auxiliary lymph node. FIG. 10E depicts an image obtained from lock-in detection of emission from FNDs from the same images used to generate FIG. 10D (bottom inset, red in overlay) overlaid on the unprocessed image obtained with the magnetic field off (top inset, white in overlay). In the partially dissected mouse, localization of the FNDs to the lymph node can be clearly seen. FIG. 10F depicts the pixel values as a function of time corresponding to the selected points in FIG. 10B and FIG. 10D. The pixels selected were over (1) the axillary lymph node and (2) a negative control on the skin in FIG. 10B and (3) the axillary lymph node and (4) a negative control on a rib in FIG. 10D. Signal modulation as a result of the applied magnetic field is clearly visible in the lymph node through the skin, as well as when the chest cavity was opened. Meanwhile, the skin and rib showed random signal as would be expected for the negative control.

[0119] Female athymic (nu/nu) mice were purchased from Charles River Laboratories at 4-6 weeks of age and housed in a specific pathogen-free American Association for Laboratory Animal Care approved facility. All experiments were approved by the National Cancer Institute's Animal Care and Use Committee. Mice were anesthetized using gas mixtures of 1.5-2.5% isoflurane in O₂ to maintain a respiration rate of ~30 bpm during the injection procedure. A volume of 10 µL of an ~80 mg/mL silica-coated nanodiamond (~100nm) solution in PBS (pH7.4) was intradermally injected into the front foot pad of each mouse. Previous studies have shown that the primary draining LNs from this injection site are the axillary and lateral thoracic LNs.

[0120] At 24 hours post-injection, the mice were sacrificed and optical imaging was performed using a MAESTROTM CRi spectroscopic optical camera (excitation filter 523 nm, emission filter 675 nm long pass). Images were also taken with magnetic modulation in a UVP® BIOSPECTRUM® Imaging System equipped with a BIOLITE® MultiSpectral Light Source (excitation filter 525 nm, emission filter 650 nm long pass) with additional illumination provided by a green laser (Z-Bolt® #DPSS-100) achieving ~1 mW/cm² intensity. Each mouse was imaged in the instrument on a non-magnetic stage under which the permanent magnet (Catalog# DZ08-N52, K&J Magnetics) could be slid in and out. Images were captured in series with magnetic fields off and on and movies stitched together in ImageJ software available at <http://rsb.info.nih.gov/ij>.

[0121] FIG. 11A is an image of a lymph node (LN) injected with silica-coated FNDs without applied magnetic field. FIG. 11B is an image of the same field of view during application of a magnetic field of ~100 Gauss. FIG. 11B was then subtracted from FIG. 11A. FIG. 11C depicts the sum of 20 such subtracted images. The FNDs are observed as bright spots in the otherwise dark field. FIGS. 11D and 11E depict the sums of 20 such subtractions when magnetic field was always OFF and ON, respectively.

[0122] Female athymic (nu/nu) mice were purchased from Charles River Laboratories of Wilmington, Massachusetts at 4-6 weeks of age and housed in a specific pathogen-free American Association for Laboratory Animal Care approved facility. Animals were used between 6 and 8 weeks of age, and experiments were approved by the National Cancer Institute's Animal Care and User Committee. Lymphadenectomy of the axillary and lateral thoracic lymph nodes (LN) of the mice was performed. The excised LN was injected with silica-coated FNDs. The LN was

then fixed in 10% formalin for 3 days, washed with PBS, and mounted on a SUPERFROST PLUS® slide (obtained from VWR) using PERMOUNT® medium (obtained from Fisher Scientific) to adhere the coverslip.

[0123] Background-free images were obtained by using a Neodymium permanent magnet (obtained from K&J Magnetics of Jamison, Pennsylvania). A CARL ZEISS® LSM5 LIVE microscope was used to obtain the movies. Each frame of the movie was a scanning confocal image with 1 s time resolution. A 10X ZEISS® objective with 0.3 NA (EC Plan-Neofluar) was used to excite the FNDs and collect the FND emission. Samples were excited at 532 nm. Emission was filtered using a long pass filter (bandpass of 560-675 nm) and detected using a PMT.

Working Example# 5 – Background-Free Imaging Using Fluorescence Lifetime Imaging (FLIM)

[0124] FIG. 12A is a two-photon FLIM lineage of a different region of the same LN used in FIGS. 11A-11E. Longer lifetimes (indicated by red color) are believed to be due to FNDs, for which the reported lifetimes range from ~10-20 ns. FIG. 12B is a background-free image of the same field of view obtained by subtracting images without and with a magnetic field, and adding 10 such subtractions.

[0125] A LEICA® TCS SP5 scanning confocal system was customized to obtain both the fluorescent lifetime imaging microscopy (FLIM) image (FIG. 12A) and the background-free confocal image in which a Neodymium permanent magnet was used to apply the magnetic field (~100 Gauss) (FIG. 12B). For FIG. 12B, 10 subtractions of images with and without magnetic fields were added to obtain the final image. An oil immersion objective (LEICA® HCX PL APO CS 40.0X 1.25 NA Oil UV) was used both to excite and to collect the emission. For the FLIM image, the sample was excited at 930 nm using a SPECTRA-PHYSICS® MAI TAI® laser. For FLIM imaging, emission from the FNDs in the LN was filtered using a band pass filter (BP 607-683 nm) and was detected using a PICOQUANT™ PICOHARP™ 300 Time-Correlated Single Photon Counting (TCSPC) system fitted with single photon avalanche diode (SPAD). For the scanning confocal image (400 Hz scanning speed) of the same field of view (FOV), the sample was excited at 561 nm and the emission was filtered using band pass filter (600-789 nm) and was detected using a PMT.

[0126] The present subject matter provides unique, elegant methods that can be easily incorporated with existing microscope or animal imaging systems without coming in contact

with the sample. No complicated and error-prone spectral unmixing is necessary. Fluorescence of commonly-used stains and labels cannot be modulated selectively using a magnetic field. Chemical or optical modifications of commonly used labels such as GFP and dyes are possible, but these modifications are difficult to implement, are invasive, and can possibly induce unwanted changes.

[0127] While the present subject matter has been described with reference to the above embodiments, it will be understood by those skilled in the art that various changes can be made and equivalents can be substituted for elements thereof without departing from the scope of the subject matter. In addition, many modifications can be made to adapt a particular situation or material to the teachings of the subject matter without departing from the essential scope thereof. Therefore, it is intended that the subject matter not be limited to the particular embodiment disclosed as the best mode contemplated for carrying out this subject matter, but that the subject matter will include all embodiments falling within the scope of the appended claims.

[0128] The functions of several elements may, in alternative embodiments, be carried out by fewer elements, or a single element. Similarly, in some embodiments, any functional element may perform fewer, or different, operations than those described with respect to the illustrated embodiment. Also, functional elements (*e.g.*, modules, databases, computers, and the like) shown as distinct for purposes of illustration may be incorporated within other functional elements, separated in different hardware or distributed in a particular implementation.

[0129] FIG. 13 shows the elements of a modulated magnetic field fluorescence imaging system. A biological sample of interest is mounted on the AFM stage 11. An apparatus for applying a modulated magnetic field is mounted over the AFM stage. The magnet field applicator is controlled by AFM controller 10 controlled by AFM computer 13. The position of the magnetic field is monitored using XYZ closed-loop scanner 9. A condenser and trans-illuminator are mounted above the AFM stage and illuminate the biological sample.

[0130] A total internal reflection fluorescence (TIRF) imaging apparatus is mounted below the AFM stage 11. TIRF illumination can be provided by an Ar/Kr laser 23 optically coupled to filter 16, telescope 15, optical fiber 18, and TIRF illuminator 19. Radiation from TIRF illuminator 19 is incident on a beam splitter, which directs the Ar/Kr laser radiation through a lens to focus on the biological sample. Ar/Kr laser 23 is cooled with water cooling system 24.

[0131] Alternatively, the sample can be irradiated with a xenon-arc lamp such as a Lambda LS illuminator (Sutter Instrument Co.). Radiation from the xenon arc 25 can be collimated using Epi-IRM illuminator 12, directed onto a beam splitter and focused onto the biological sample.

[0132] Fluorescence is collected by the confocal microscope or other suitable imaging system and the fluorescence detected using CCD cameras 14 and video camera 22. Imaging optics are controlled and imaging data acquired using microscope computer 8.

[0133] The modulated magnetic field fluorescence imaging system shown in FIG. 13 illustrates the general elements of a suitable imaging system. In general, the system is configured such that a time-modulated magnetic field is applied to the biological sample. The magnetic field may be applied to the entire biological sample or a portion of the biological sample. The system is configured to irradiate the entire biological sample or a portion of the biological sample to cause the biological sample to fluoresce. The system is further configured to collect the fluorescence.

[0134] In certain embodiments, fluorescent nanodiamonds are used to image a biological sample. A biological sample can be labeled with functionalized fluorescent nanodiamonds. The time-varying magnetic field is applied at a magnetic strength sufficient to mix the ± 1 and 0 triplet states at a high field strength.

[0135] FIG. 14 is a schematic summarizing the elements of a modulated magnetic field fluorescence imaging system and includes a modulated magnetic field apparatus, an imaging instrument, a specimen such as a biological sample, and a software analysis module. The modulated magnetic field can be applied using a permanent magnet or an electromagnet. In certain embodiments, a permanent magnet can be moved with respect to the specimen to provide a time-varying magnetic field. In certain embodiments, an electromagnet is used to apply a time-varying magnetic field with different waveforms.

[0136] An imaging instrument referred to in FIG. 14 can include suitable microscope imaging optics such as used in confocal microscopy, light microscopy, TIRF microscopy, and FLIM microscopy, or animal imaging systems such as transillumination scanners or reflectance scanners. As appropriate, the imaging instrument can include, for example, optical filters, lenses, collimators, and beam splitters. An imaging instrument can also include devices for detecting fluorescence and optically imaging the sample. Optics for irradiating the sample to

induce fluorescence and translation stages for moving the magnet, optics, radiation source, and/or imaging components are also included.

[0137] The various components can be controlled by software. Image processing can involve comparison of images obtained within and without a magnetic field applied to the sample or involve comparing the fluorescence obtained at low and high magnetic field strength.

[0138] FIG. 15 shows various configurations of a magnet with respect to a sample stage. For example, in certain embodiments an electromagnet can be mounted above, below and/or to the sides of the sample stage. An electromagnet can be used to apply a time-varying magnetic field. In another configuration, a permanent magnet can be mounted above, below, and/or to the sides of the sample stage and physically moved toward and away from the sample stage to change the strength of the magnetic field applied to the sample. As shown in the third scheme, a permanent magnet can be mounted on an apparatus configured to rotate the magnet about the sample to provide a time-varying magnetic field. In certain embodiments, a permanent magnet can be stationary and a magnetic shield can be disposed between the magnet and the sample to provide a time-varying magnetic field.

[0139] FIG. 16 is a schematic showing the general elements of a small animal optical imaging instruments. The imaging instrument includes a light source, a specimen such as a biological sample, a detector, software for controlling the light source and the detector(s) and a user interface. The light source can include multiple excitation filters for controlling the wavelength range of the excitation irradiation and the detector can include multiple filters for selecting the wavelength range of the fluorescence.

[0140] FIG. 17 is a schematic showing the general elements of a small animal imaging instrument for obtaining time-modulated magnetic field fluorescent images. The instrument shown in FIG. 17 includes a light source, specimen such as a biological sample, a detector(s), software for controlling the light source and detector(s), a user interface, and a software interface for controlling a magnetic field applicator for applying a magnetic field to the specimen. The light source can include multiple excitation filters for controlling the wavelength range of the excitation irradiation and the detector can include multiple filters for selecting the wavelength range of the fluorescence. In certain embodiments, the excitation wavelength range and the detection wavelength range are selected to optimize the contrast of magnetic field modified NV-center nanodiamond fluorescence. The magnetic field applicator can include a permanent magnet

or an electromagnet configured to apply a time-varying magnetic field to the specimen. The software addition can be used to correlate magnetic field-dependent images of the specimen with optical images of the specimen, and to perform image processing to provide images of the specimen. In the case of imaging based on magnetic field modified NV-center nanodiamond fluorescence images of the specimen or a portion of the specimen obtained with a magnetic field and without a magnetic field or with a lower intensity magnetic field can be compared reduce or eliminate background fluorescence that does not originate from NV-center fluorescent nanodiamonds.

[0141] FIG. 18 shows another embodiment of a small animal imaging instrument for magnetic field modulation including a light source, a specimen, a detector, a magnetic field, applicator, software for controlling the light source, the detector, and the magnetic field applicator, and a user interface. In this embodiment the light source includes a source and/or optical system configured to optimize excitation of the NV-center fluorescent nanodiamond triplet state and a detector and optics including, for example, a filter, selected to optimize detection of NV-center fluorescent nanodiamond fluorescence.

[0142] FIGS. 19-21 show various configurations of a light source, specimen, detector, and magnetic field applicator. FIG. 19 shows a light source, specimen, and detector in a linear configuration with the magnetic field applied from below a rotating stage. FIG. 20 shows the light source, specimen and detector in a confocal configuration in which the excitation source and the detector(s) are on the same side of the specimen and with the magnetic field applicator on the opposite side of the specimen. FIG. 21 shows an example of imaging instrumentation in which detectors are located around the specimen and with a single light source an magnetic field applicator. In certain embodiments the detectors, light source, and magnetic field applicator can be rotated around the sample. In certain of the embodiments illustrated in FIG. 21, there can be multiple light sources and magnetic field applicators.

[0143] Modulated magnetic field fluorescence imaging apparatus can be stationary systems in which the specimen or biological sample is mounted on a stage and the imaging instrumentation is a standalone system. In certain embodiments, an imaging apparatus provided by the present disclosure can be a portable device such as a hand-held scanner. As shown, for example, in FIG. 22, a clinical hand-held scanner can include a light source, a magnetic field applicator, a camera or detector, software configured to obtain modulated magnetic field

fluorescence images of a patient, and an interface between the clinician and the software. In certain embodiments, the elements of the scanner are optimized to obtain images of magnetic field modified NV-center nanodiamond fluorescence.

[0144] In certain embodiments, the fluorescent nanodiamonds are silica coated. Silica-coated nanodiamonds and methods of preparing silica-coated nanodiamonds are disclosed, for example, in WO 2014/014970 A1, which is incorporated by reference in its entirety. Similar methods can be used to silica coat fluorescent nanodiamonds.

[0145] Silica-coated fluorescent nanodiamonds can be functionalized with probes, affinity molecules, or other entities that can bind to specific targets of a biological sample. These functionalized silica-coated fluorescent nanodiamonds can be applied to a biological specimen such as a tissue sample or cells to stain the specimen with the functionalized nanodiamonds. In other embodiments, functionalized nanodiamonds can be administered in vivo and the magnetically modified fluorescence of the functionalized silica-coating nanodiamonds measured through the skin of a patient. In certain embodiments, following in vivo administration of functionalized nanodiamonds to a subject, tissue or biological fluid can be removed from the subject and evaluated for the presence of functionalized nanodiamonds. Functionalized silica-coated nanodiamonds can be used to probe targets similar to ways in which fluorescent probes are used.

[0146] While certain embodiments according to the present subject matter have been described, the present subject matter is not limited to just the described embodiments. Various changes and/or modifications can be made to any of the described embodiments without departing from the spirit or scope of the present subject matter. Also, various combinations of elements, steps, features, and/or aspects of the described embodiments are possible and contemplated even if such combinations are not expressly identified herein.

CLAIMS

What is claimed is:

1. An imaging system comprising:
an imaging stage for mounting a specimen;
a radiation source configured to excite the triplet excited state of fluorescent nanodiamonds and directed on specimen;
a fluorescence detector configured to detect triplet-triplet fluorescent nanodiamond fluorescence and configured to detect fluorescence from the specimen; and
an apparatus configured to apply a time-varying magnetic field to the specimen.
2. The imaging system of claim 1, comprising a specimen mounted on the imaging stage.
3. The imaging system of claim 1, wherein the radiation source is selected from a laser and a xenon arc lamp.
4. The imaging system of claim 1, wherein the fluorescent nanodiamonds comprise silica-coated fluorescent nanodiamonds.
5. The imaging system of claim 1, wherein the fluorescent nanodiamonds comprise functionalized silica-coated fluorescent nanodiamonds.
6. The imaging system of claim 1, wherein the apparatus configured to apply a time-varying magnetic field comprises a permanent magnet.
7. The imaging system of claim 1, wherein applying the time-varying magnetic field comprising moving the permanent magnet with respect to the specimen.
8. The imaging system of claim 1, wherein the apparatus configured to apply a time-varying magnetic field comprises an electromagnet.

9. The imaging system of claim 1, wherein the specimen is selected from a patient, an animal, a tissue, and a cell.
10. The imaging system of claim 1, comprising an imaging processing unit.
11. The imaging system of claim 1, wherein the image processing unit is configured to compare images obtained with a first applied magnetic field with fluorescence images obtained with a second applied magnetic field.
12. The imaging system of claim 1, wherein the first magnetic field and the second magnetic field have a different intensity.
13. The imaging system of claim 1, wherein the imaging system is portable.
14. The imaging system of claim 1, wherein the imaging system is configured to image a portion of the specimen having a dimension from 1 cm to 10 cm.
15. The imaging system of claim 1, wherein the imaging system is configured to image a portion of the specimen having a dimension of at least 1 cm.
16. The imaging system of claim 1, wherein the imaging system is configured to image a portion of the specimen having a dimension of less than 0.1 cm.
17. The imaging system of claim 1, wherein the imaging system is configured to image a portion of the specimen having a dimension from 0.01 cm to 0.1 cm.
18. The imaging system of claim 1, wherein the radiation source and the detector are configured to move with respect to the specimen.

19. The imaging system of claim 1, wherein the radiation source, the detector, and the apparatus configured to apply a time-varying magnetic field are configured to move with respect to the specimen.

20. The imaging system of claim 1, wherein the detector comprises a plurality of detectors.

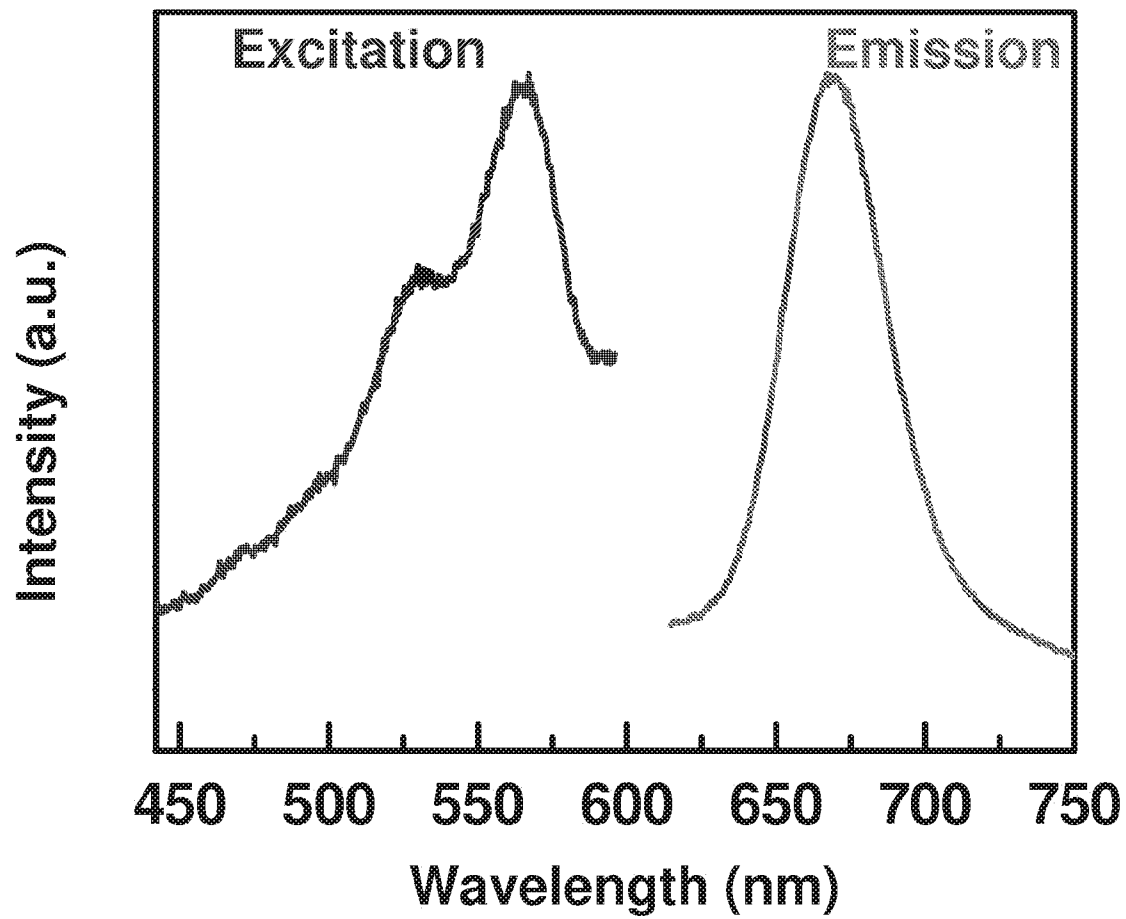


FIG. 1

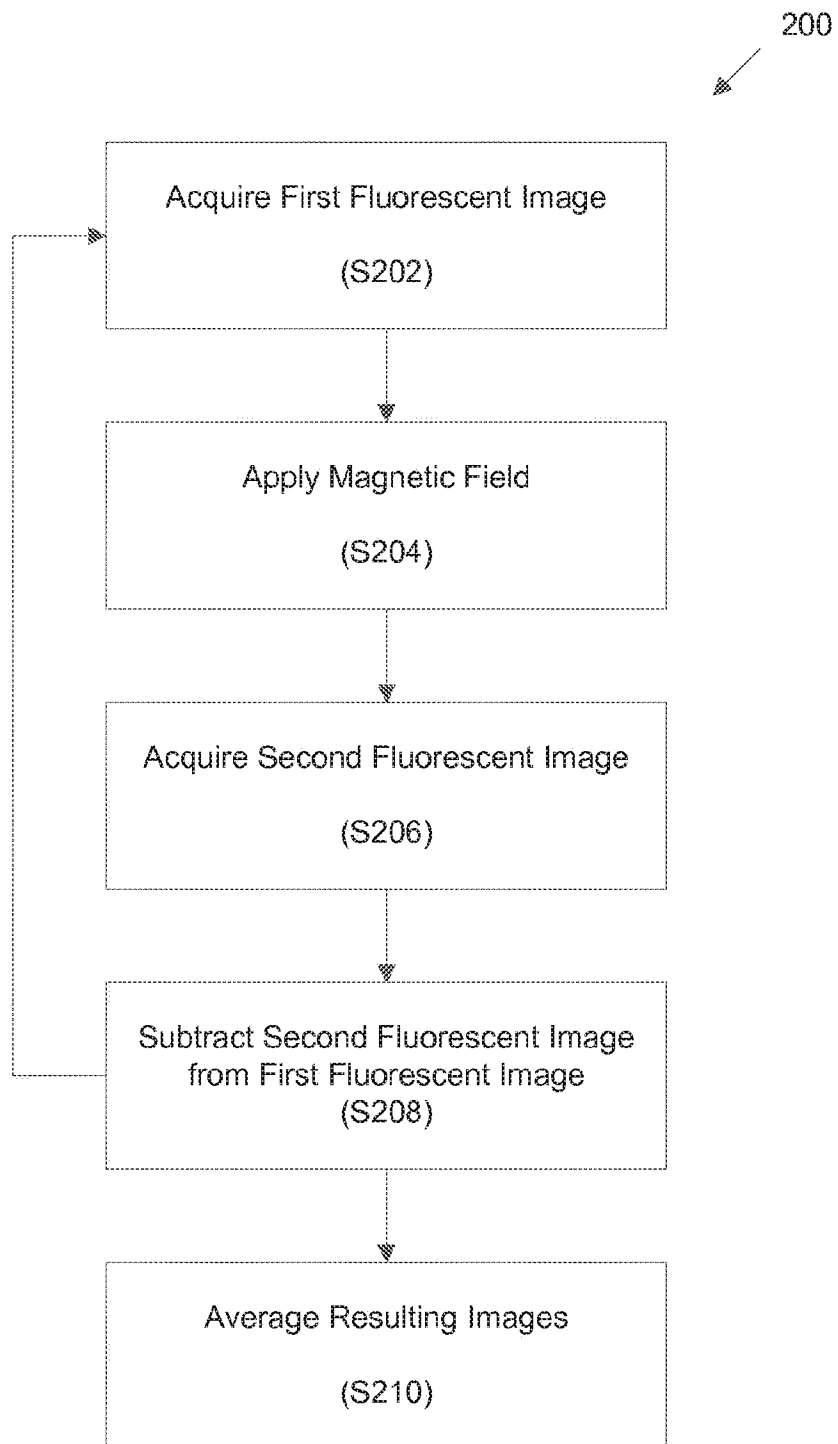


FIG. 2

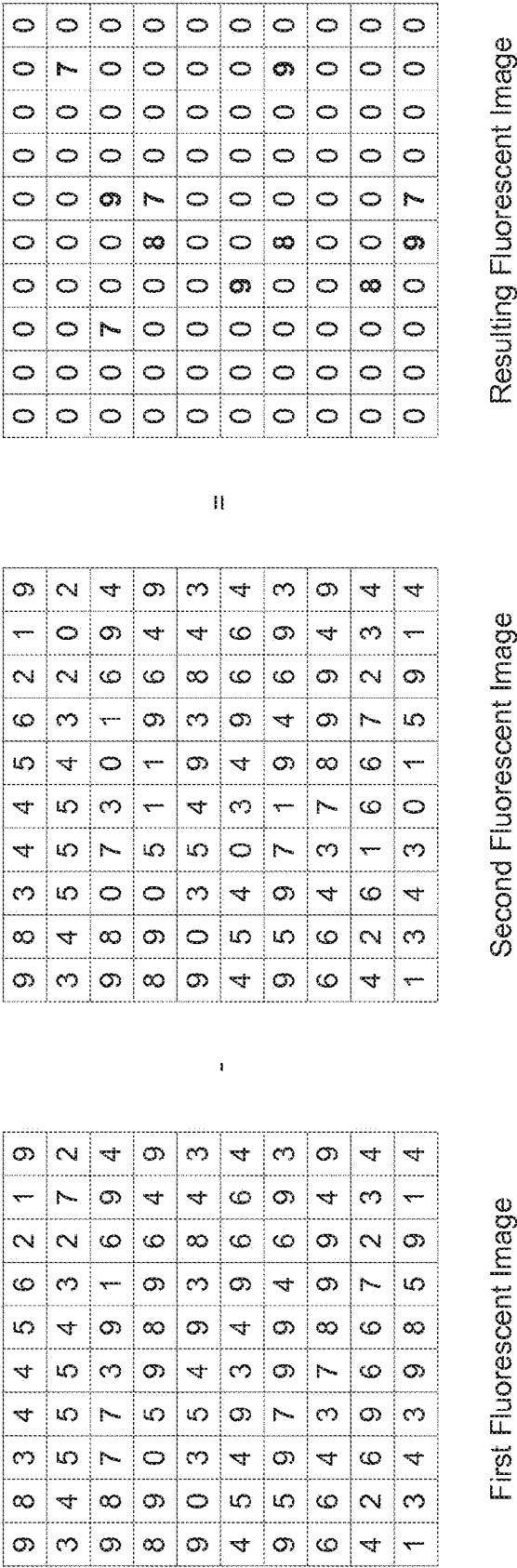


FIG. 3

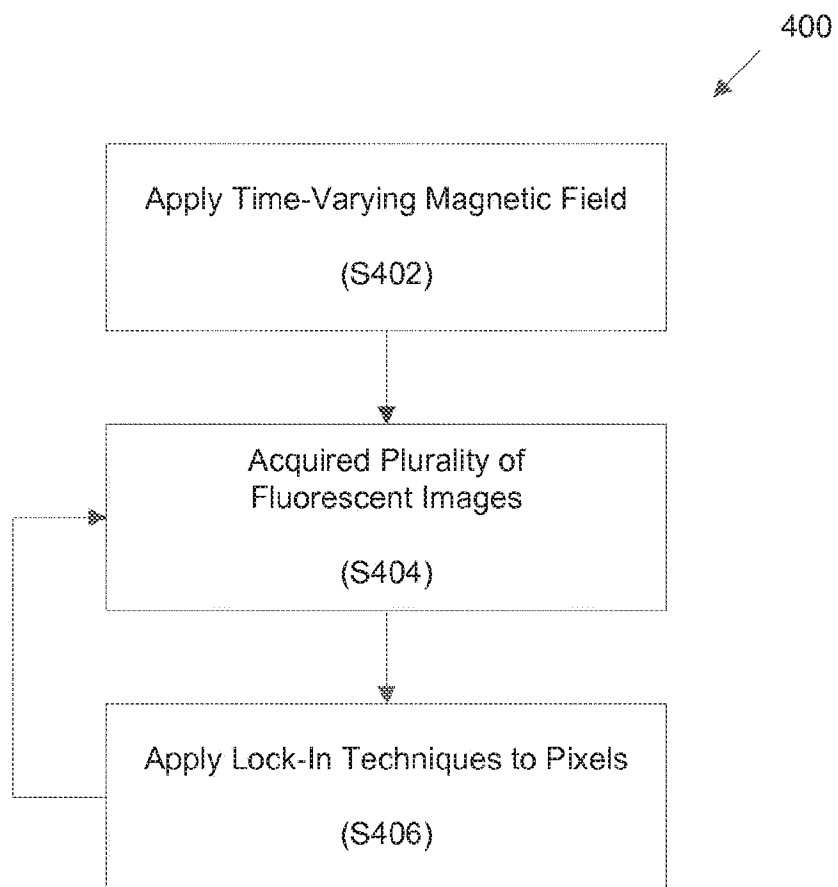


FIG. 4

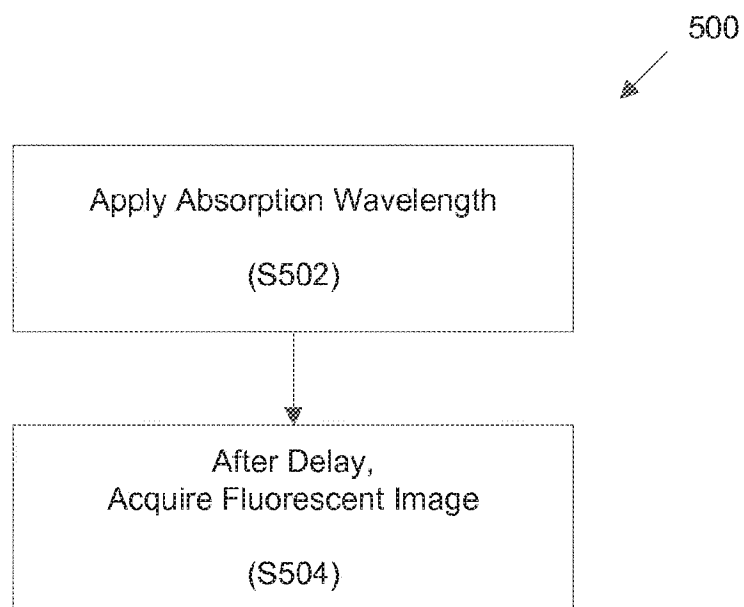


FIG. 5

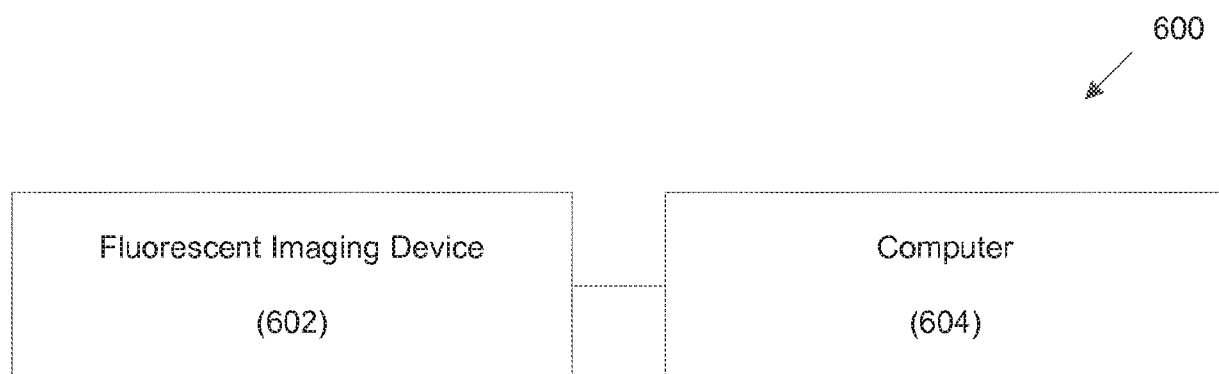


FIG. 6

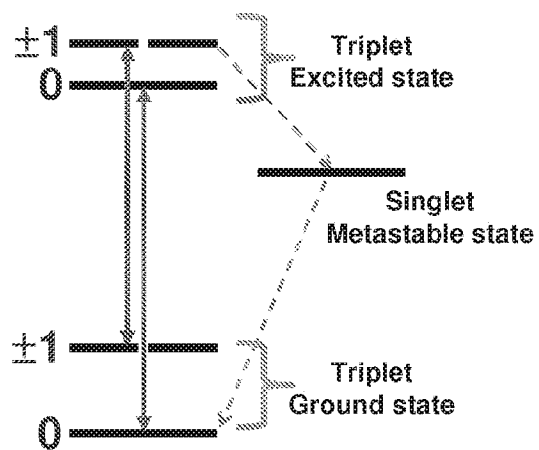


FIG. 7A

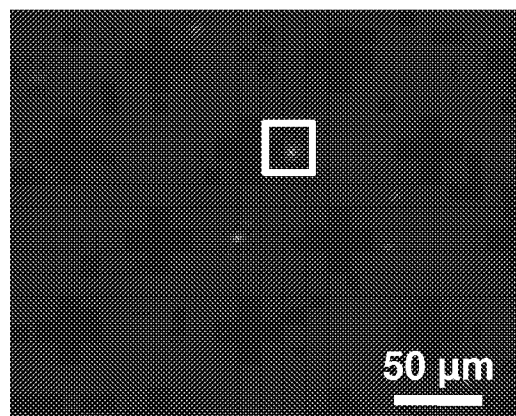


FIG. 7B

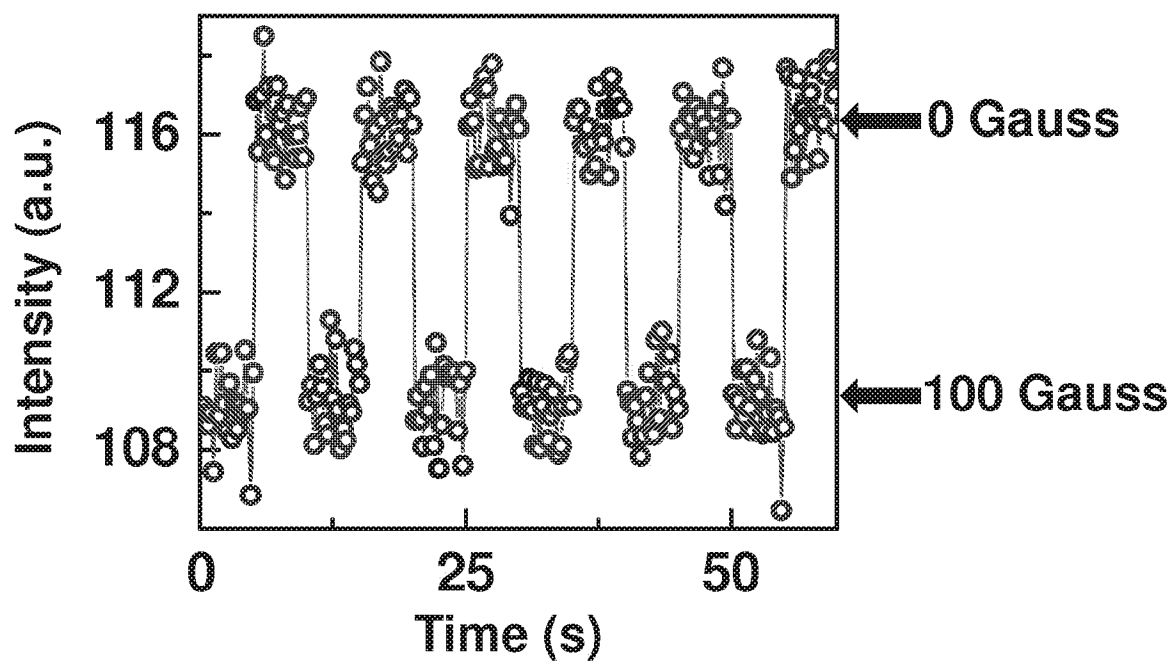


FIG. 7C

FIG. 8A

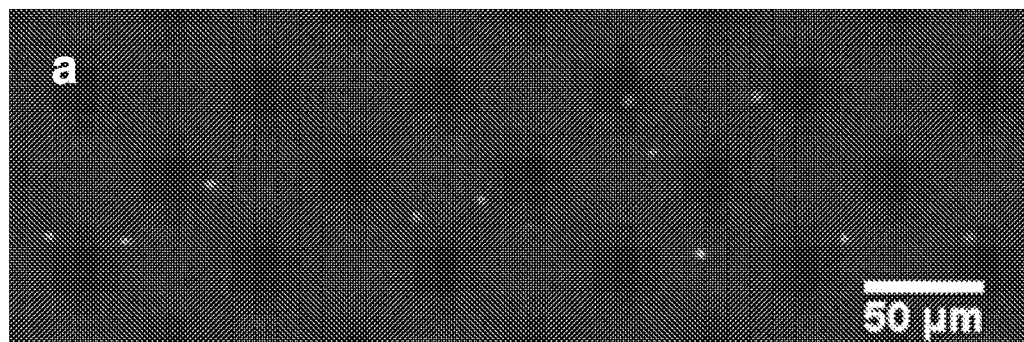


FIG. 8B

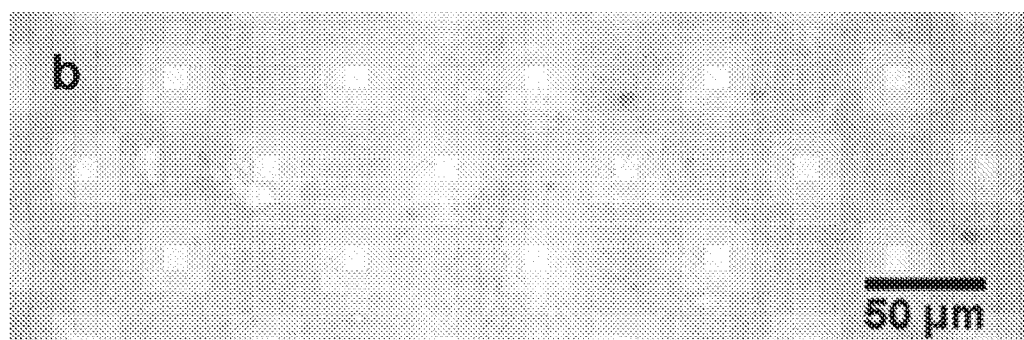
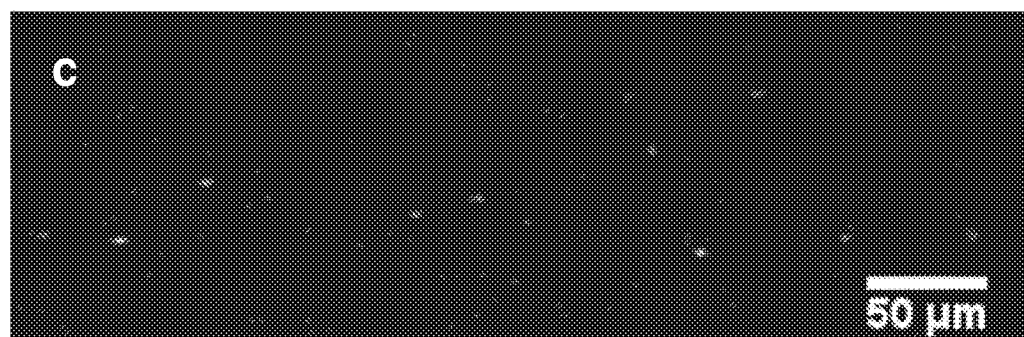


FIG. 8C



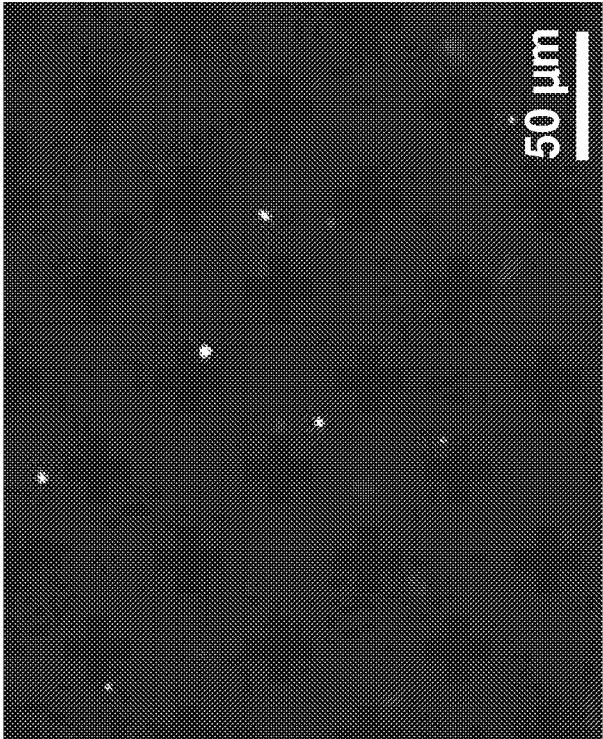


FIG. 9B

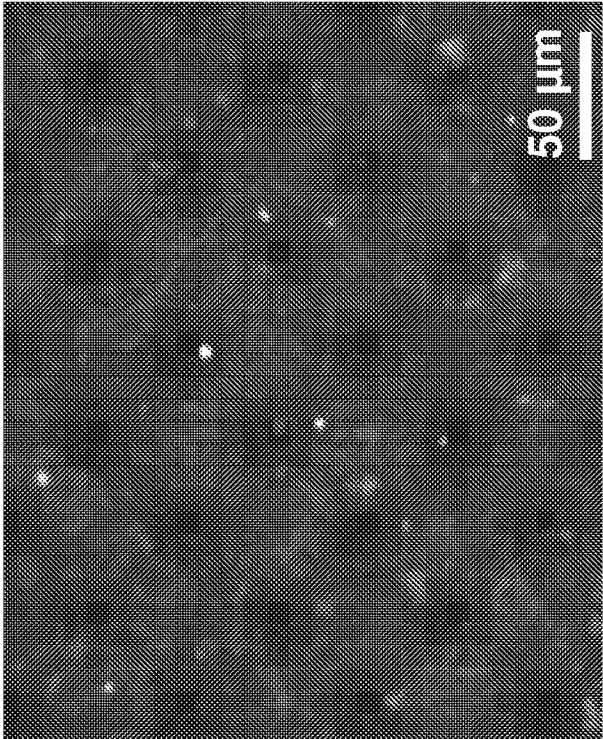
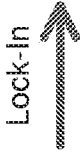


FIG. 9A

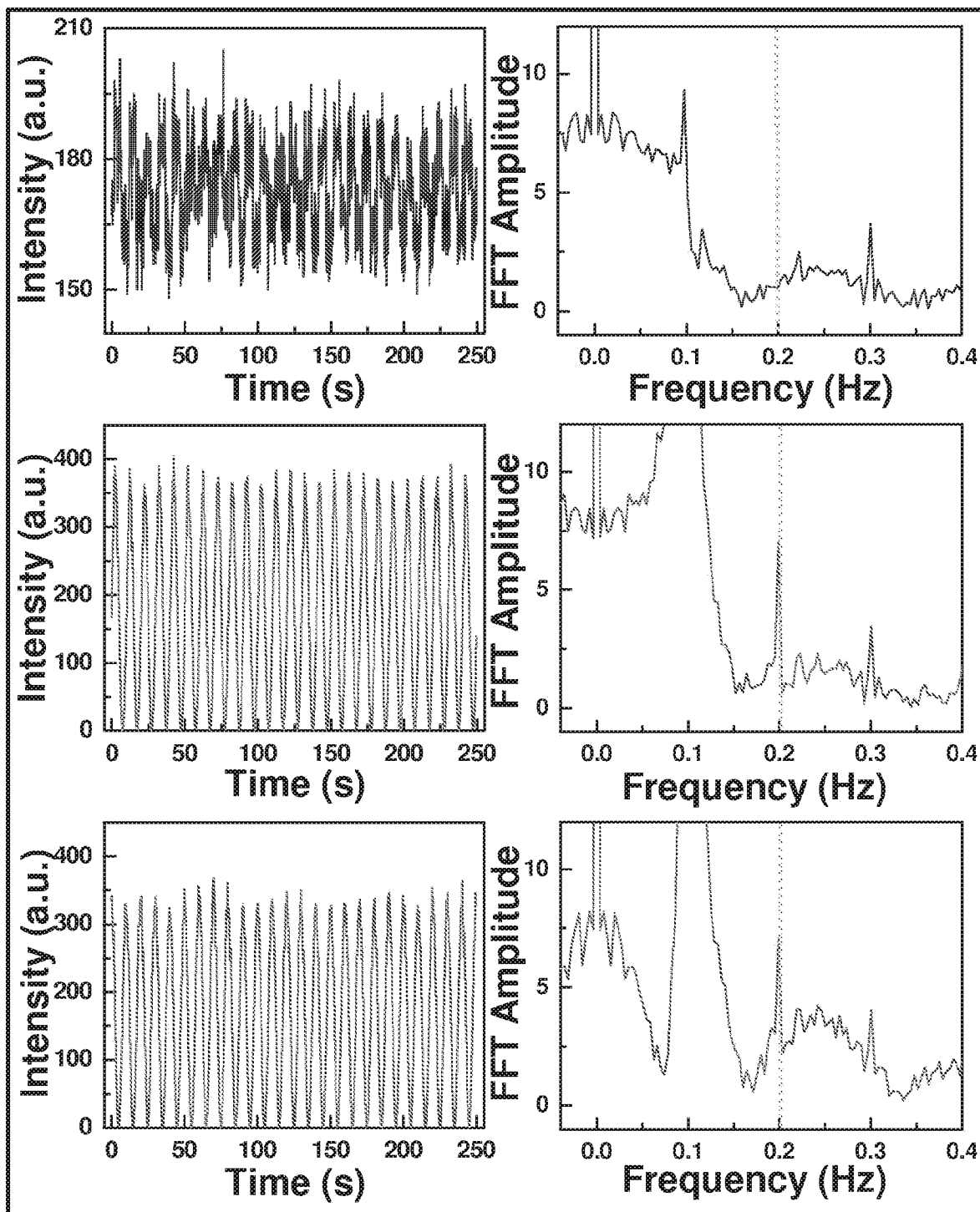


FIG. 9C

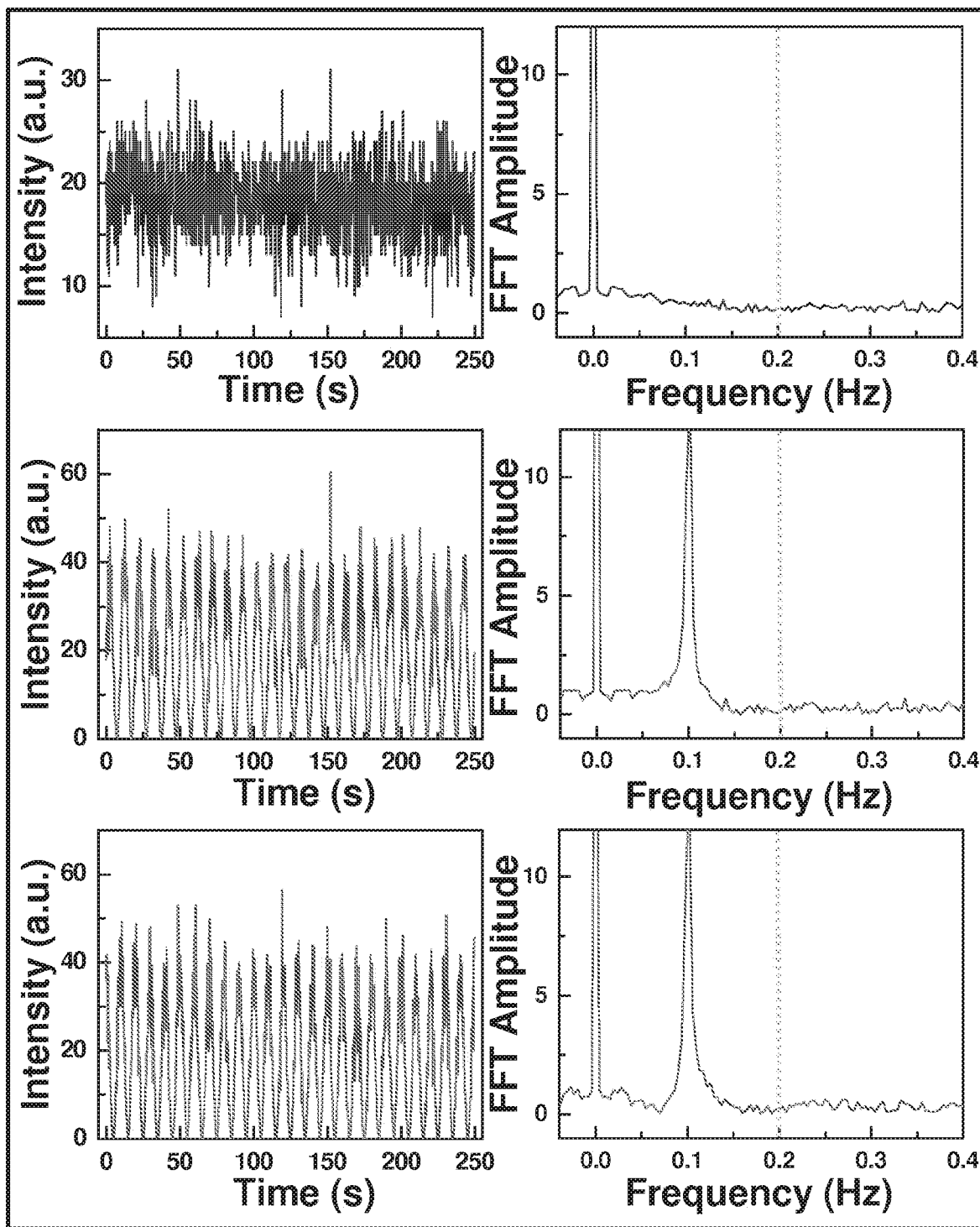


FIG. 9D

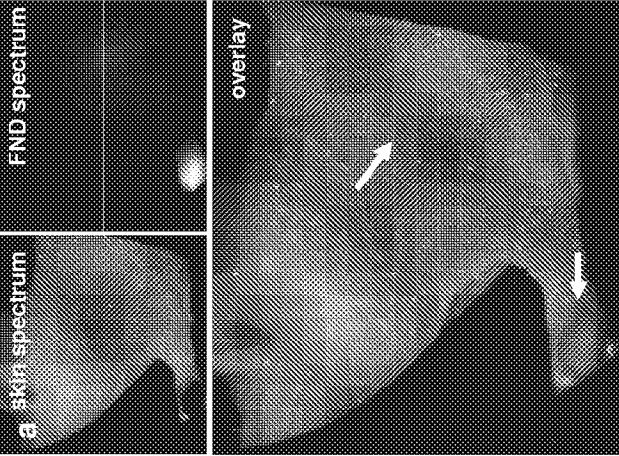


FIG. 10A

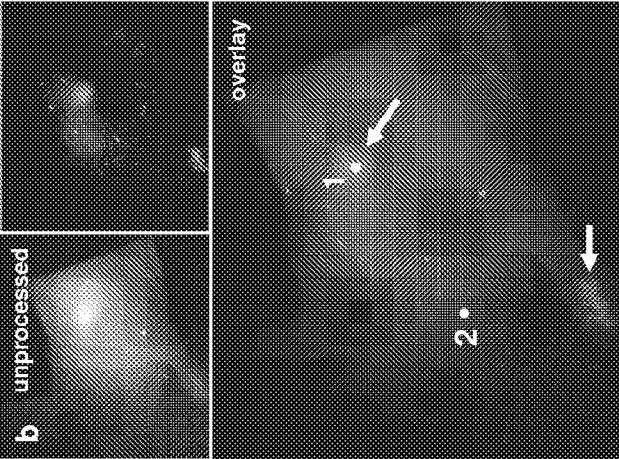


FIG. 10B

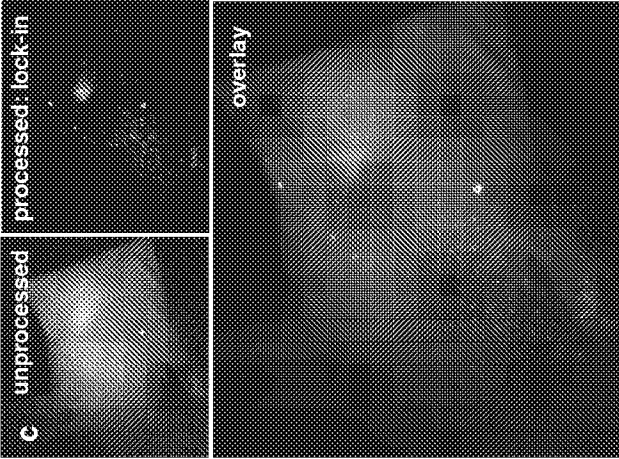


FIG. 10C

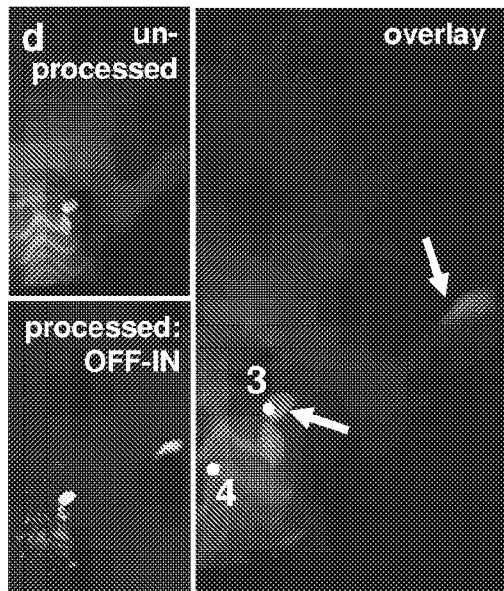


FIG. 10D

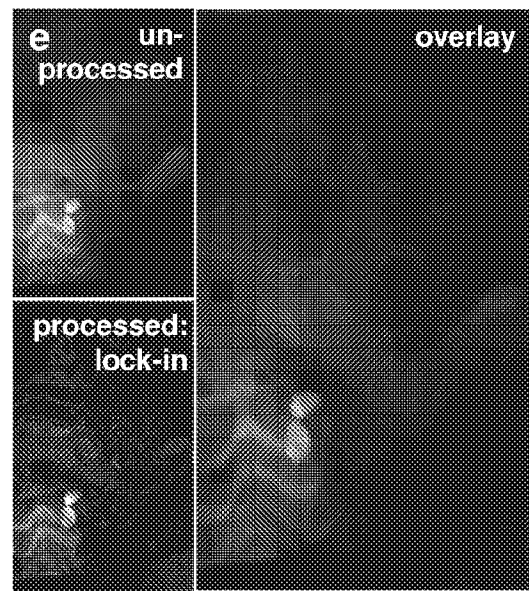


FIG. 10E

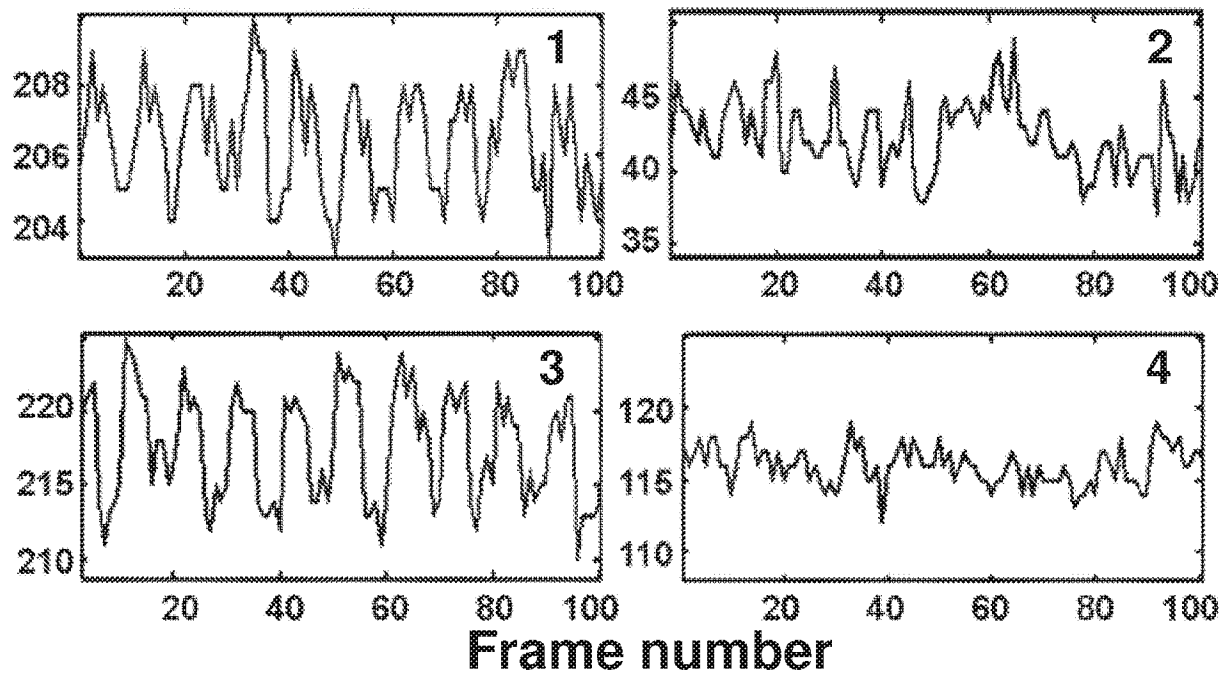


FIG. 10F

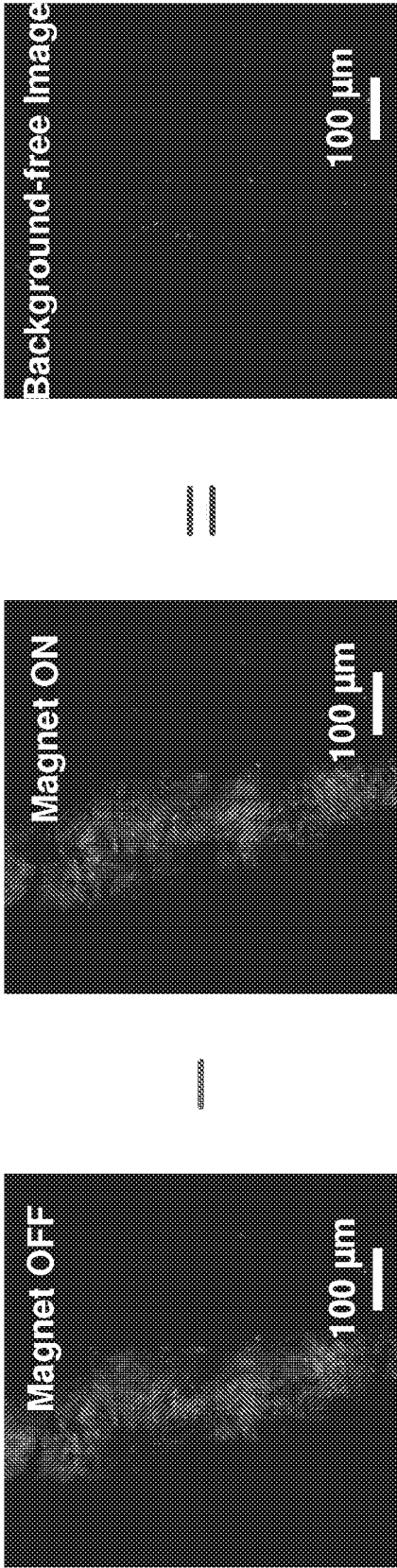


FIG. 11C

FIG. 11B

FIG. 11A

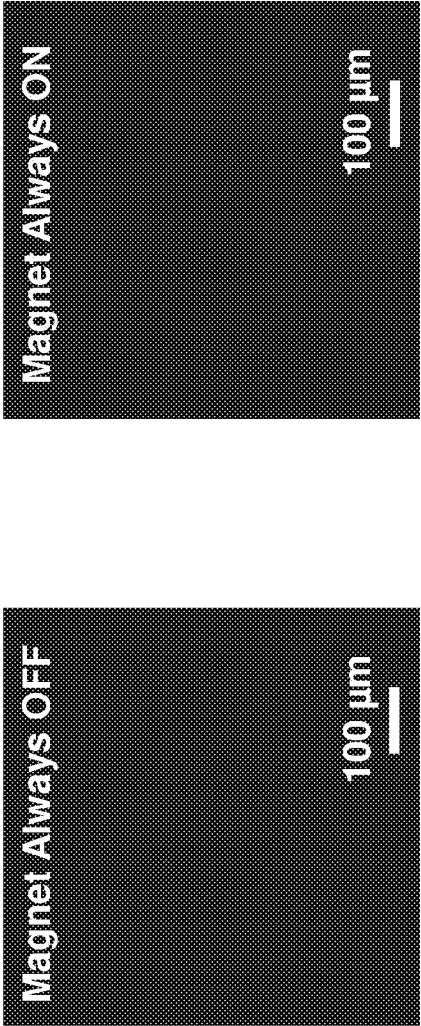


FIG. 11D

FIG. 11E

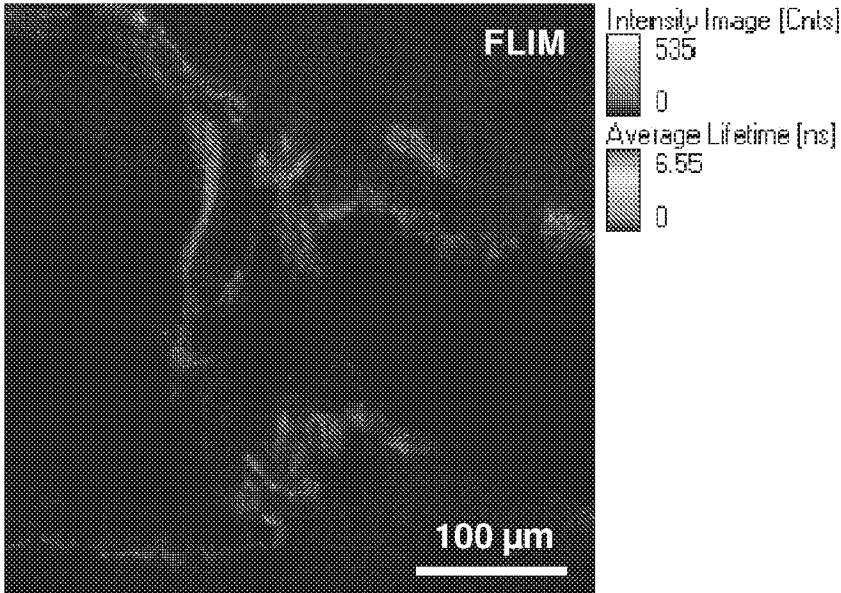


FIG. 12A

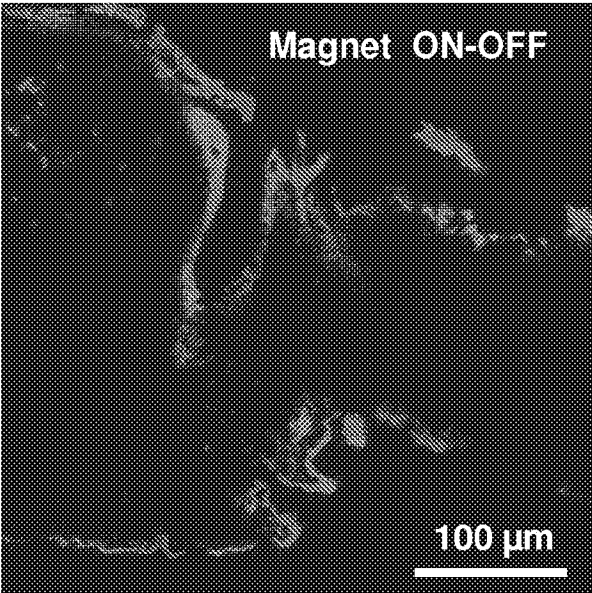


FIG. 12B

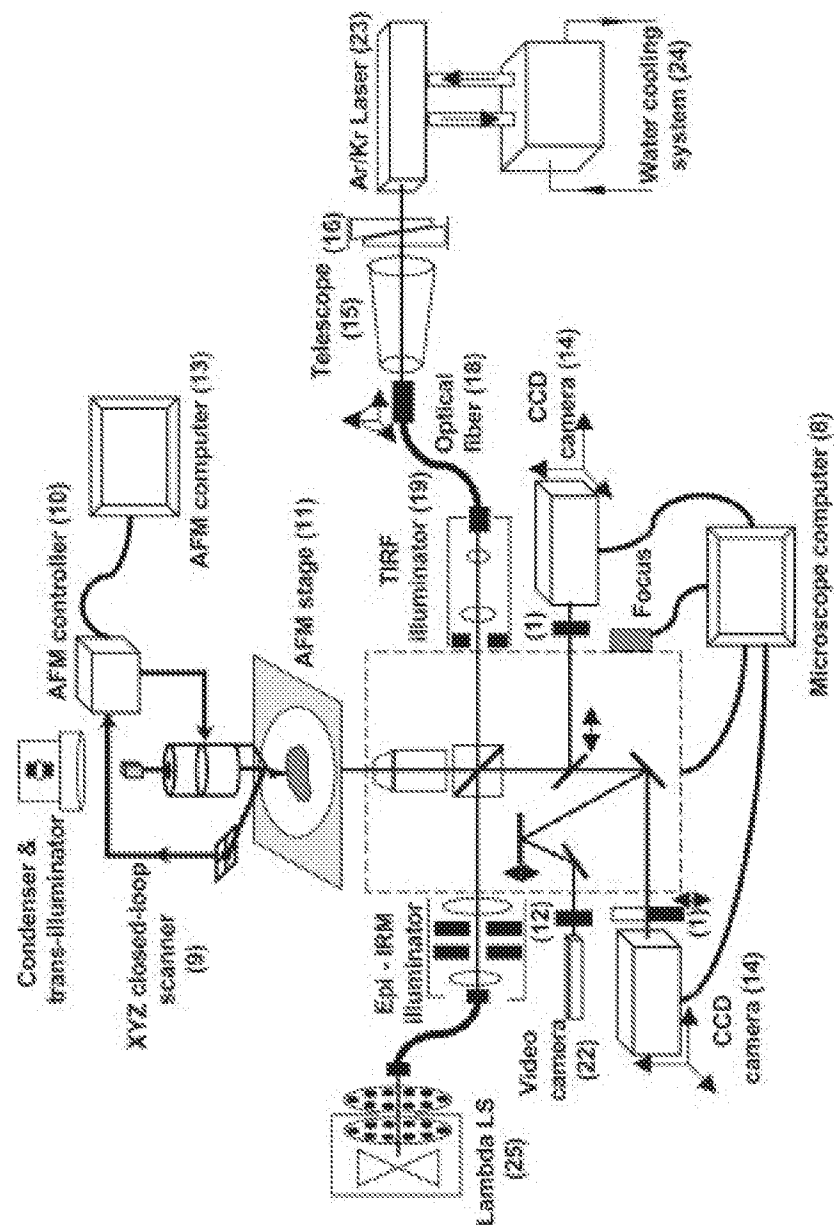
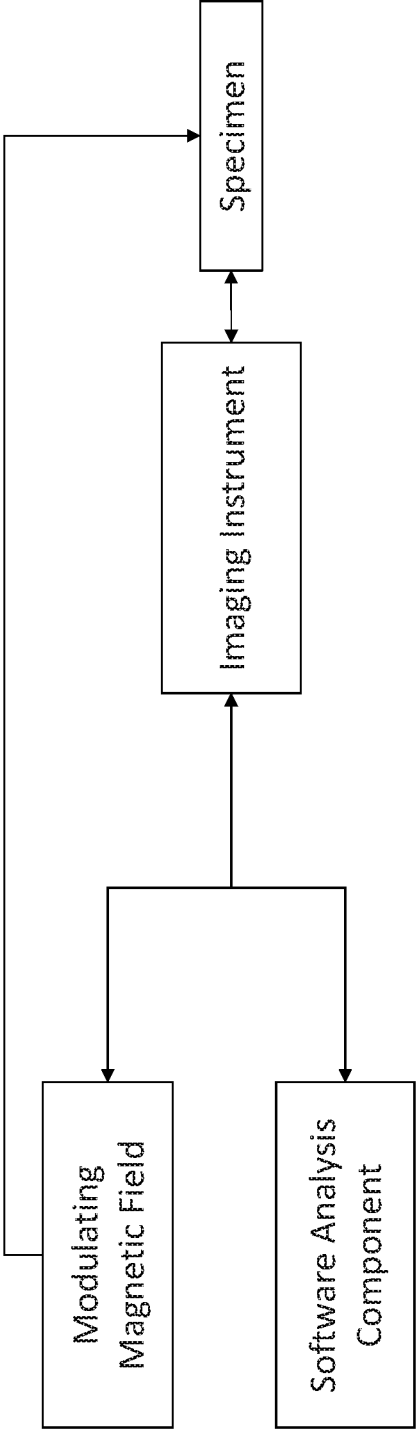


FIG. 13

FIG. 14



Modulating Magnetic Field

- Applied by permanent magnetic that slides or rotates in and out or by an electromagnet that can be modulated to have fields with different wave functions

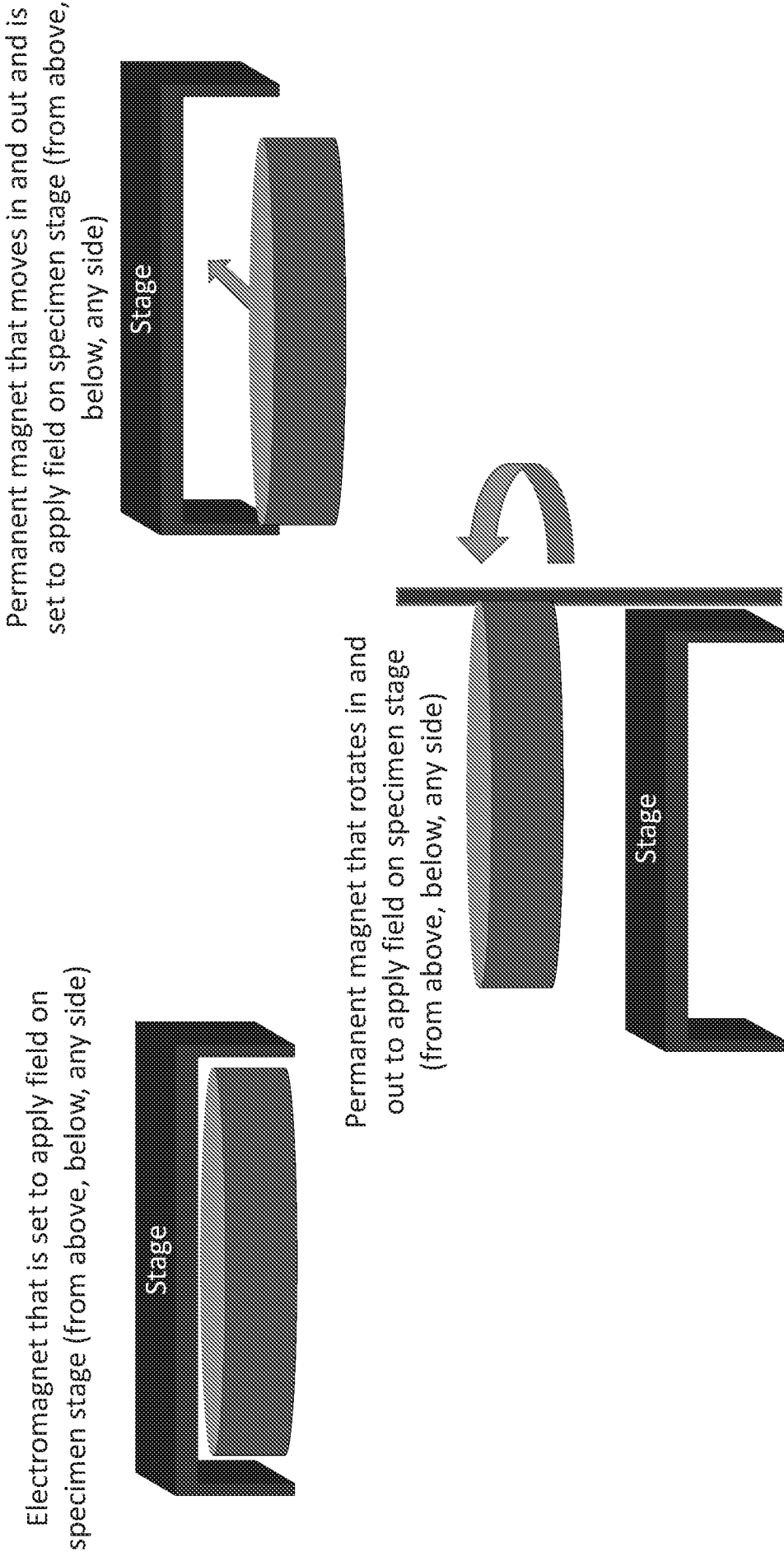
Imaging Instrument

- Includes various types of microscopes (confocal, light, TIRF, FLIM, etc), animal scanners (transillumination, reflectance, etc)
- Includes the image software components of optical imaging (i.e. contrast, filtering, unmixing, etc), as well as the physical components (i.e. cameras, filters, stage, lasers, etc)

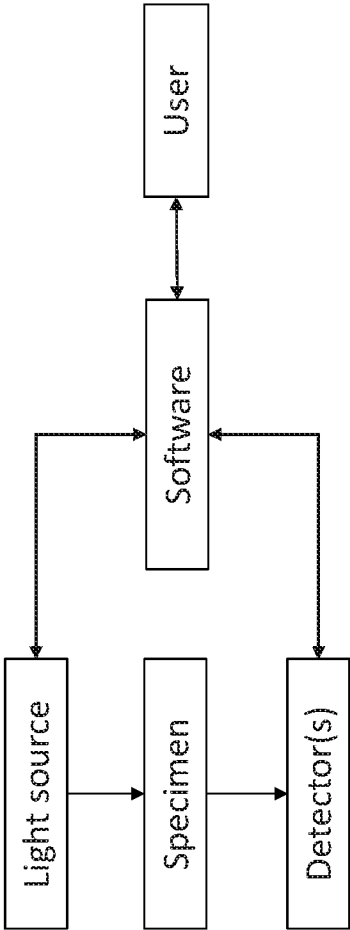
Software Analysis Component

- Image processing with subtraction or lock-in

FIG. 15



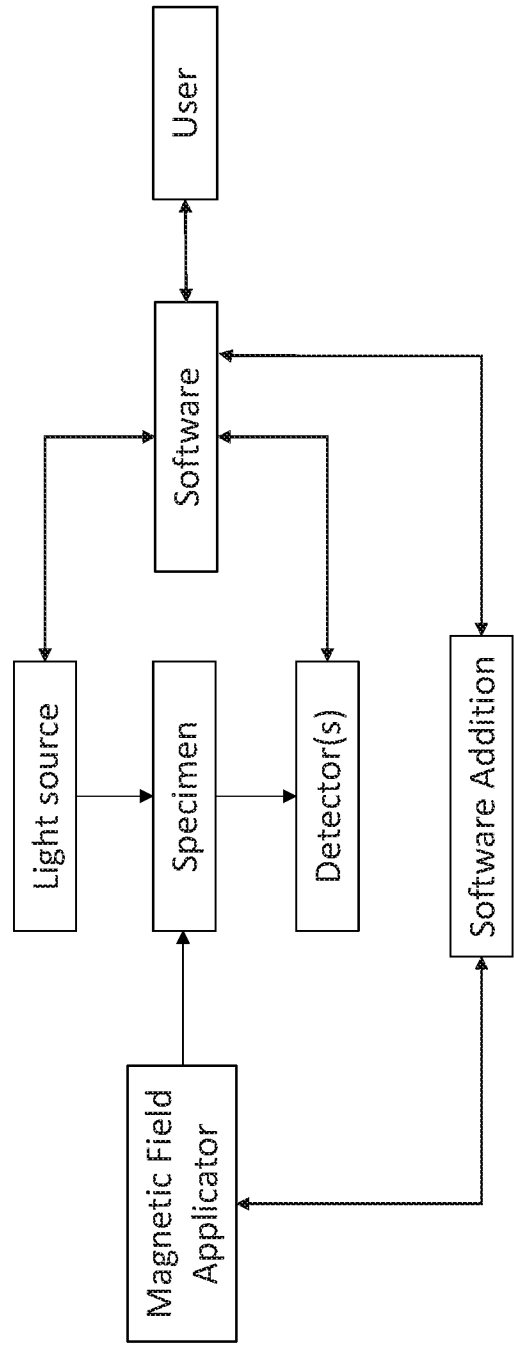
Standard Optical Microscopy or Small Animal Imaging Instrument



Light Source: multiple excitation filters
Detector: multiple emissions filters

FIG. 16

FIG. 17



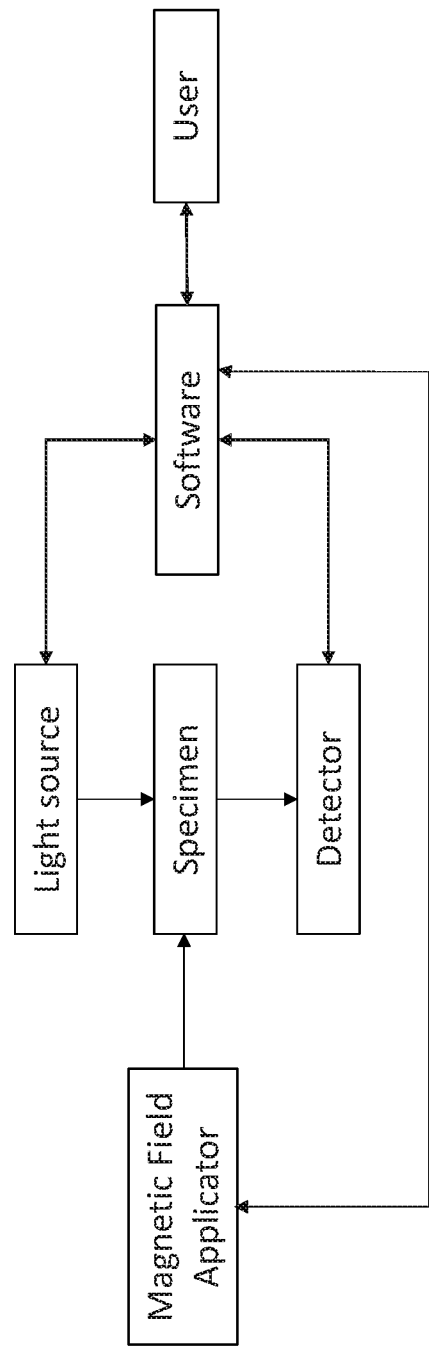
Light Source: multiple excitation filters

Detector: multiple emissions filters

Magnetic Field Applicator: permanent magnetic physical modulated (rotating or sliding mechanism) or electromagnet

Software Addition: correlation of image capture with magnetic field application, background subtraction (previous patent), image correction if necessary

FIG. 18



Light Source: single optimized excitation filter
Detector: single optimized emissions filter
Magnetic Field Applicator: permanent magnetic physical modulated (rotating or sliding mechanism) or electromagnet

FIG. 19

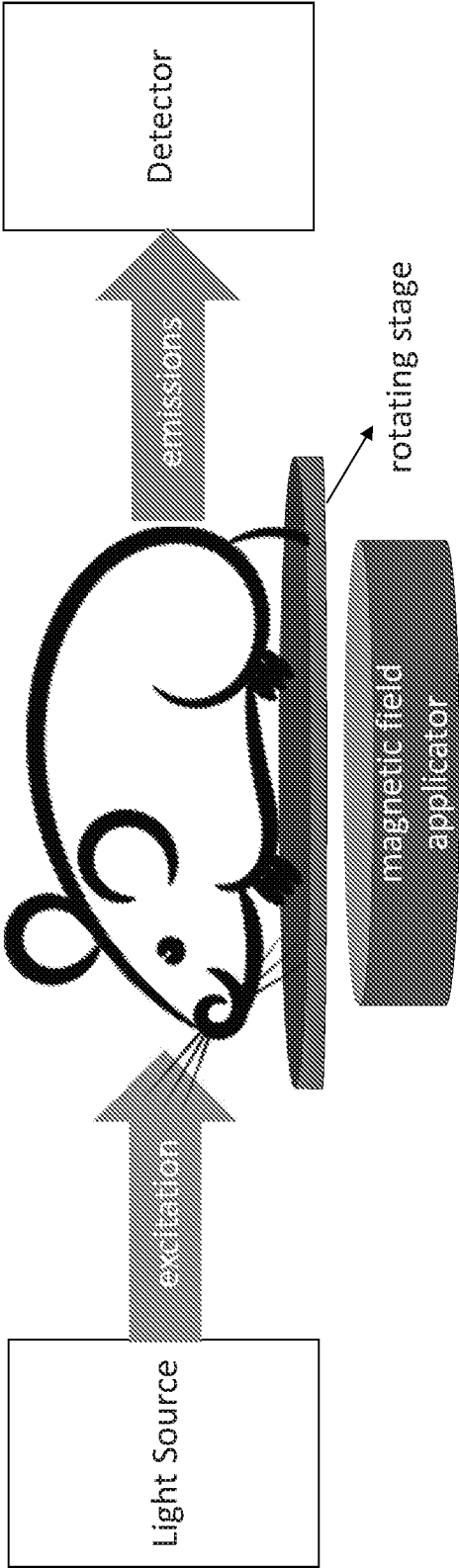
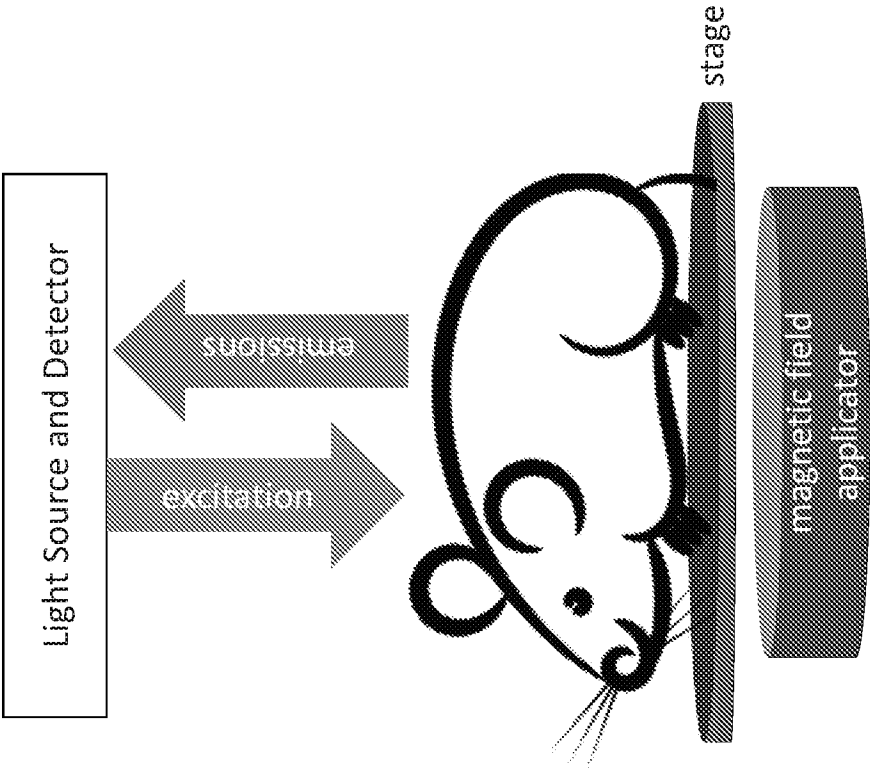


FIG. 20



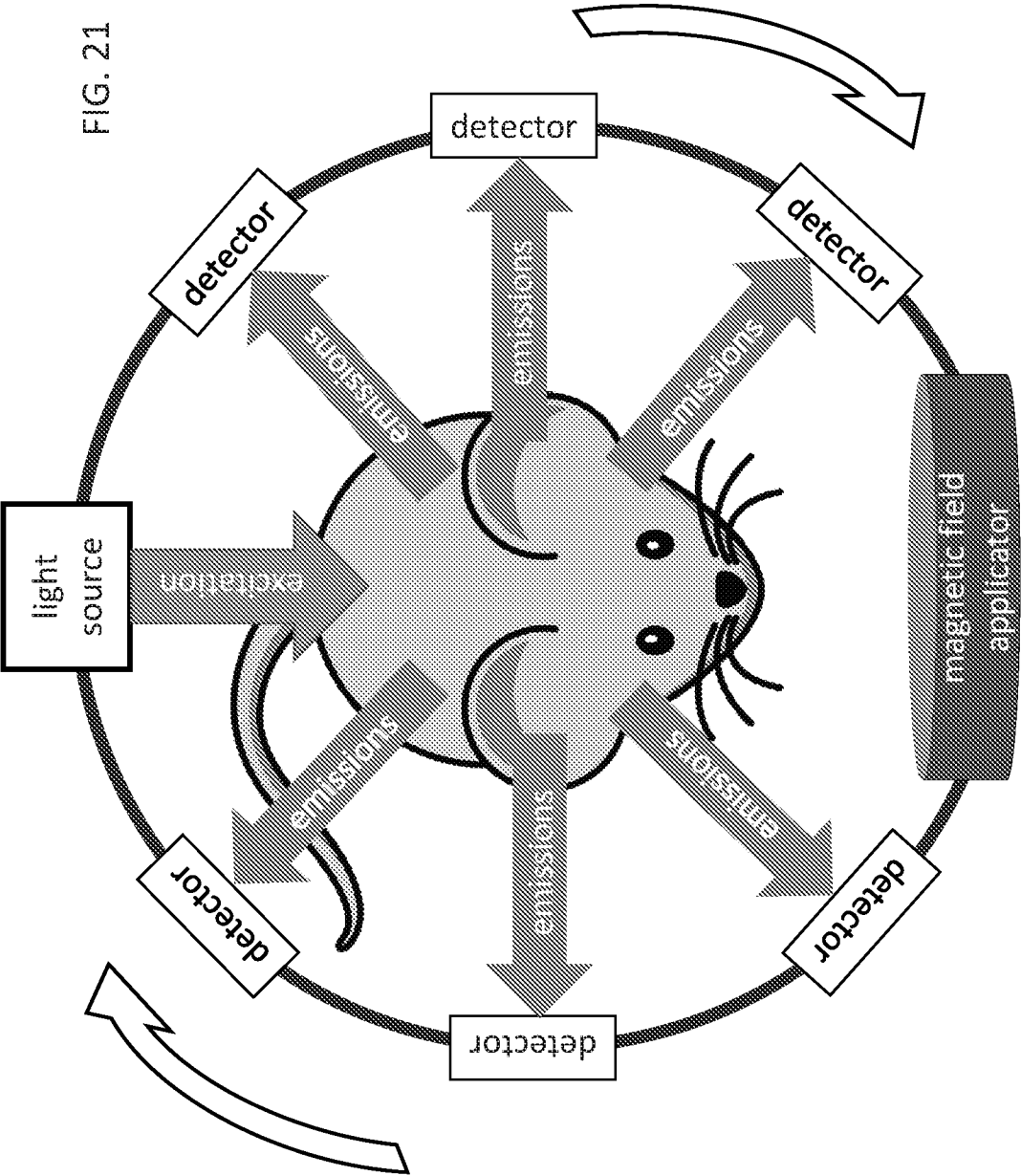
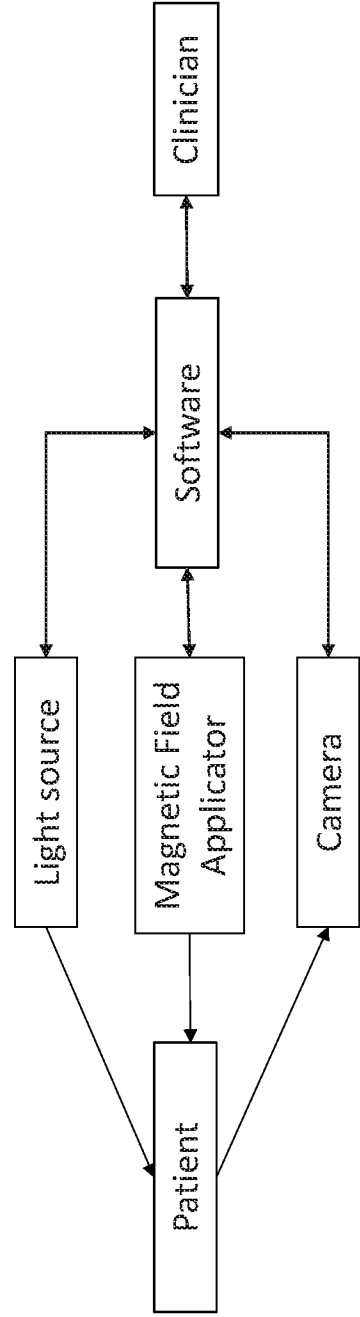


FIG. 21

Clinical Hand-Held Scanner



Light Source: single optimized excitation filter
Camera: with single optimized emissions filter

FIG. 22

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US16/26791

A. CLASSIFICATION OF SUBJECT MATTER

IPC(8) - G01N 21/64; G01Q 60/54, 70/14 (2016.01)

CPC - G01N 21/6458; G01Q 70/14; G01R 33/022

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)

IPC(8) Classifications: G01N 21/64; G01Q 60/54, 70/14 (2016.01)

CPC Classifications: G01N 21/6458; G01Q 70/14; G01R 33/022

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)

PatSeer (US, EP, WO, JP, DE, GB, CN, FR, KR, ES, AU, IN, CA, INPADOC Data); Google Scholar; IEEE; EBSCO

Keywords used: imaging; fluorescence detector; triplet-triplet fluorescent nanodiamond; time-varying magnetic field

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	US 2014/0099007 A1 (THE UNITED STATES OF AMERICA, AS REPRESENTED BY THE SECRETARY, DEPARTMENT OF HEALTH AND HUMAN SERVICES) April 10, 2014; abstract; figure 6, 7A; paragraphs [0005]; [0075]; [0110]	1-20
Y	US 3,843,939 A (GROFF, R) October 22, 1974; abstract; figure 2; column 6, lines 1-19	1-20
Y	US 2014/0180073 A1 (ANTICANCER INC.) June 26, 2014; abstract; paragraphs [0010]; [0030-0031]; [0039]	13
Y	US 2014/0350395 A1 (SHACHAF, C et al.) November 27, 2014; abstract; paragraphs [0054]; [0091]; [0139]	14-17
Y	US 2010/0090127 A1 (YEKTA, A et al.) April 15, 2010; figure 1A; paragraphs [0030]; [0076]	18, 20
Y	US 2008/0230715 A1 (NIELSEN, T et al.) September 25, 2008; abstract; paragraphs [0005]; [0018]; [0024]	19
P,Y	WO 2015/199940 A1 (GOOGLE INC.) December 30, 2015; whole document	1-20

☐ 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

08 June 2016 (08.06.2016)

Date of mailing of the international search report

11 JUL 2016

Name and mailing address of the ISA/

Mail Stop PCT, Attn: ISA/US, Commissioner for Patents

P.O. Box 1450, Alexandria, Virginia 22313-1450

Facsimile No. 571-273-8300

Authorized officer

Shane Thomas

PCT Helpdesk: 571-272-4300

PCT OSP: 571-272-7774