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DESCRIPTION

[0001] This International Patent Cooperation Treaty Patent Application is a continuation-in-part of United States Patent Application No. 12/151,156, filed May 2, 2008, hereby incorporated by reference in the entirety herein.

I. TECHNICAL FIELD

[0002] Generally, a computer implemented remote monitoring system which generates a viewable reduced byte data representation for each one of a plurality of analyzed instrument signals. Specifically, a flow cytometer remote monitoring system which generates a viewable reduced byte data representation for each one of a plurality analyzed flow cytometer signals.

II. BACKGROUND

[0003] Flow cytometry systems can be utilized to analyze at least one of a plurality of particles. The plurality of particles is typically a population of biological particles such as sperm cells, stem cells, blood cells, bacteria, or the like. The analysis of a population of particles can occur at analysis rates of between about 10,000 particles per second and about 200,000 particles per second depending on the of the population of particles and the manner of analysis. The analysis of individual particles in a population can provide information relating to the presence or absence or the amount of one or more particle characteristics. The relative presence or absence or the amount of one or more particle characteristics may be used to as the basis on which to differentiate individual particles of an analyzed population into two or more discrete subpopulations of particles. The discreet subpopulations of particles can then be separated from the main population of particles and isolated as discrete subpopulations of particles as further described herein.

[0004] US 2003/0078703 A1 discloses an example of a flow cytometer. In the flow cytometer of US 2003/0078703 A1 measured signals are converted into list mode data. The list mode data are transferred to a database server. A client computer can access, view and analyse the data stored in the database.

[0005] The operator of a flow cytometer may rely on the use of a viewable data representation to make decisions about the operation of the flow cytometer device. Since the flow cytometer device can be analyzing many hundreds of millions of particles per hour and may be further sorting many millions of cells per hour, the viewable data representation may be designed to show the flow cytometer operator a continuously updated viewable data representation. The continuously updated viewable data representation may include analysis data of a fraction of the population of particles analyzed along with operating parameters for the flow cytometer

device updated in discrete analysis intervals. For example, the viewable data representation may be updated every 100 milliseconds, in the form of histograms of the most recent 10 seconds of particle analysis data.

[0006] The flow cytometer operator relies on the viewable data representation to both manage and control procedures and parameters for particle analysis and sorting but to also to control the hardware configuration of the flow cytometer device with regard to three dimensional positioning of components such as the fluidic nozzle, beam shaping optics and optical focus for detection, and the like. The flow cytometer operator may also control the rate of droplet formation, the amplitude of the energy used in droplet formation and the voltage applied to droplet streams without use of the viewable data representation; although these types of adjustments can change the scale and precision of measurements being made in analysis of the population of particles and result in changes to the viewable data representation displayed to the flow cytometer operator. Accordingly, the viewable data representation provides a source of real time information utilized by the flow cytometer operator to adjust particle analysis and flow cytometer device parameters.

[0007] The viewable data representation generated during operation of the flow cytometer device can be generated in an image format selected by the flow cytometer operator which can be configured by selection of particular data masks and data sets which populate such data masks which are typically provided as histograms. Although the analysis data generated by a flow cytometer device may be collected and stored in a memory element of the flow cytometer device as raw data files, the operation of a flow cytometer device to assess over a duration of time the operating condition, adjust analysis and hardware parameters to optimize the operating condition, or trouble shoot the operating condition for software or hardware problems may require access to a substantial portion or all of the history and detail of the viewable data representation. However, use of a substantial portion of the history and detail the viewable data representation during operation of a flow cytometer device, or other similar analysis device, can present certain problems.

[0008] One substantial problem with using a substantial portion of the history and detail of viewable data representation may be competition for computer processing capacity of the flow cytometer device (or other devices which provide similar viewable data representations) resulting in a delay updating display of the viewable data representation which in certain instances can appear as interrupted or non-continuous display of the viewable data representation. In certain instances depending on the flow cytometer device, attempts to access and utilize stored viewable data representations or attempts to control the functionalities of the flow cytometer device remotely can interfere with the normal operation and particle analysis of the flow cytometer device.

[0009] Another substantial problem may be that the viewable data representation is stored in files that subdivide some user defined amount of detail in screens per second and in seconds per file. These file management and data storage requirements can overload the computer processing unit ("CPU"), read only memory ("RAM"), and the local memory storage capacity of

individual flow cytometer devices, especially when such flow cytometer devices are equipped with versions of CPU, RAM, operating systems ("OS"), and other programs that were not designed to operate at speeds required for use of large file sizes associated with viewable data representations in form of images, video, histograms, or similar technology of electronically capturing, recording, processing, storing, transmitting, and reconstructing a sequence of still images.

[0010] Another substantial problem may be that viewable data representations in the form of image files and video files, or the like, may be too large to be utilized in the lesser bandwidths of local area networks ("LAN") or virtual private networks ("VPN") to effectively transmit to computers outside of the LAN.

[0011] Another substantial problem may be that the effort, time, and cost of providing sufficient storage space in memory for all historical viewable data representation at a level of resolution sufficient to be useful upon retrieval is too great when compared to the value of the viewable data representation stored.

[0012] Another substantial problem may be that conventional methods to capture, recording, processing, storing, transmitting, or reconstruct images to provide conventional viewable data representations are not scalable. For example, while conventional methods of providing a viewable data representation may be practical in regard to one flow cytometer device, it may not be possible or practical in the context of providing a viewable data representation for a plurality of flow cytometer devices such as 50 or 100 flow cytometer devices (or even a greater number of devices) at a single location or in a LAN environment, or in a distributed network of flow cytometer devices, or a plurality of distributed LAN environments connected in a wide area network ("WAN") environment.

III. DISCLOSURE OF INVENTION

[0013] The invention is defined by the independent claims.

IV. BRIEF DESCRIPTION OF THE DRAWINGS

[0014]

Figure 1 is a block diagram of hardware means and network means of a particular embodiment of the invention.

Figure 2 is a block diagram of hardware means and network means of another particular embodiment of the invention.

Figure 3 is a block diagram of a particular device for the analysis and sorting of a plurality of

particles.

Figure 4 is a block diagram of hardware means which can be utilized in a particular embodiment of the invention.

Figure 5 is block diagram of hardware means which can be utilized in a particular embodiment of the invention.

Figure 6 is block diagram of hardware means which can be utilized in a particular embodiment of the invention.

Figure 7 is a block diagram which shows a particular method of archiving reduced byte data representations generated by a particular embodiment of the invention.

Figure 8 is a diagram of a viewable data representation generated by a particular embodiment of the invention which includes a flow cytometer.

Figure 9 is a diagram of a plurality of viewable reduced byte data representations generated by a particular embodiment of the invention remotely monitoring a plurality of flow cytometers.

V. MODE(S) FOR CARRYING OUT THE INVENTION

[0015] Generally, computer implemented remote monitoring system which generates a viewable reduced byte data representation for each one of a plurality of analyzed instrument signals. Specifically, a flow cytometer remote monitoring system which generates a viewable reduced byte data representation for each one of a plurality analyzed flow cytometer signals.

[0016] Referring generally to Figures 1-9, the inventive instrument or flow cytometer remote monitoring system (1) may be described herein in terms of functional block components, screen shots, and various process steps. It should be appreciated that such functional blocks may be realized by any number of hardware or software components configured to perform the specified functions. For example, the inventive flow cytometer remote monitoring system (1) may employ various integrated circuit components which function without limitation as memory elements, processing elements, logic elements, look-up tables, or the like, which may carry out a variety of functions under the control of one or more microprocessors or other control devices.

[0017] Similarly, the software elements of the present invention may be implemented with any programming or scripting language such as C, C++, Java, COBOL, PERL, Labview, or any graphical user interface programming language, extensible markup language (XML), Microsoft's Visual Studio .NET, Visual Basic, or the like, with the various algorithms or Boolean Logic being implemented with any combination of data structures, objects, processes, routines or other programming elements. Further, it should be noted that the present invention might

employ any number of conventional wired or wireless techniques for data transmission, signaling, data processing, network control, and the like.

[0018] It should be appreciated that the particular implementations shown and described herein are illustrative of the invention and its best mode and are not intended to otherwise limit the scope of the present invention in any way. Indeed, for the sake of brevity, conventional data networking, application development and other functional aspects of the systems (and components of the individual operating components of the systems) may not be described in detail herein. Furthermore, the connecting lines shown in the various Figures contained herein are intended to represent exemplary functional relationships or physical couplings between the various elements. It should be noted that many alternative or additional functional relationships or physical connections may be present in various embodiments of the inventive flow cytometer remote monitoring system (1).

[0019] As will be appreciated by one of ordinary skill in the art, the present invention may be embodied as a method of data processing, a data processing system, a device for data processing, a computer program product, or the like. Accordingly, the present invention may take the form of an entirely software embodiment, an entirely hardware embodiment, or an embodiment combining aspects of both software and hardware. Furthermore, the present invention may take the form of a computer program product on a computer-readable storage medium having computer-readable program code means embodied in the storage medium. Any suitable computer-readable storage medium may be utilized, including hard disks, CD-ROM, optical storage devices, magnetic storage devices, ROM, flash RAM, or the like.

[0020] The present invention may be described herein with reference to screen shots, block diagrams and flowchart illustrations of the flow cytometer remote monitoring system (1) to describe computer programs, applications, or modules which can be utilized separately or in combination in accordance with various aspects or embodiments of the invention. It will be understood that each functional block of the block diagrams and the flowchart illustrations, and combinations of functional blocks in the block diagrams and flowchart illustrations, respectively, can be implemented by computer program instructions. These computer program instructions may be loaded onto a general purpose computer, special purpose computer or other programmable data processing apparatus to produce a machine, such that the instructions which execute on the computer or other programmable data processing apparatus implement the functions specified in the flowchart block or blocks.

[0021] These computer program instructions may also be stored in a computer-readable memory that can direct a computer or other programmable data processing apparatus to function in a particular manner, such that the instructions stored in the computer-readable memory produce an article of manufacture including instruction means which implement the function specified in the flowchart block or blocks. The computer program instructions may also be loaded onto a computer or other programmable data processing apparatus to cause a series of operational steps to be performed on the computer or other programmable apparatus to produce a computer-implemented process such that the instructions which execute on the

computer or other programmable apparatus provide steps for implementing the functions specified in the flowchart block or blocks.

[0022] Accordingly, functional blocks of the block diagrams and flowchart illustrations support combinations of means for performing the specified functions, combinations of steps for performing the specified functions, and program instruction means for performing the specified functions. It will also be understood that each functional block of the block diagrams and flowchart illustrations, and combinations of functional blocks in the block diagrams and flowchart illustrations, can be implemented by either special purpose hardware based computer systems which perform the specified functions or steps, or suitable combinations of special purpose hardware and computer instructions.

[0023] Furthermore, while embodiments of the inventive flow cytometer remote monitoring system (1) may be described in the context of monitoring a flow cytometer (2), the invention is not so limited, and the data management functionalities and image management functionalities of inventive flow cytometer remote monitoring system (1) can be utilized in the context of monitoring a numerous and wide variety of instruments including but not limited to chromatographs, spectrophotometers, computed topographs, computed tomographs, or the like. The term "flow cytometer" for the purposes of the embodiments of the invention described herein generally means any device configured to count, examine, or sort microscopic particles suspended in a stream of fluid such counting, examination or sorting can be based upon single or multiple parametric analysis of the physical or chemical characteristics of single cells flowing through an optical or electronic detection apparatus. A flow cytometer (2) can as non-limiting examples be configured to provide a single stream of fluid in which particles are entrained for analysis or a plurality of streams of fluid each stream of fluid entraining particles for analysis with particles in each such fluid stream interrogated by one or more laser beams each of which can be emitted by a corresponding one laser device or can be produced by splitting a single laser beam into a plurality of laser beams for interrogation of one fluid stream or a plurality of fluid streams. The signal generated by the optical or electronic detection apparatus for each stream can be processed as single channel of data or multiple channels of data at independent rates using one or a plurality of processors in parallel. A non-limiting example of a flow cytometer (2) device suitable for use in embodiments of the inventive flow cytometer remote monitoring system (1) can be a MOFLO[®] SX or MOFLO[®] SX XDP flow cytometer available from Dako Colorado, Inc. or can be flow cytometers available from Icyte, Becton Dickinson, Cytopenia, Partec, or the like; the invention is not so limited. Certain flow cytometer (2) devices can be utilized for FACS flow cytometry for sorting of heterogeneous mixtures of particles; however, the invention is not limited to the utilization of any particular type or kind of a flow cytometer (2) device.

[0024] For the purposes of the present invention, ranges may be expressed herein as from "about" one particular value to "about" another particular value. When such a range is expressed, another embodiment includes from the one particular value to the other particular value. Similarly, when values are expressed as approximations, by use of the antecedent "about," it will be understood that the particular value forms another embodiment. It will be

further understood that the endpoints of each of the ranges are significant both in relation to the other endpoint, and independently of the other endpoint.

[0025] Moreover, for the purposes of the present invention, the term "a" or "an" entity refers to one or more of that entity; for example, "a viewable data representation" refers to one or more of those viewable data representations. As such, the terms "a" or "an", "one or more" and "at least one" can be used interchangeably herein. Furthermore, the term "selected from the group consisting of" refers to one or more of elements in the list that follows, including combinations of two or more of the elements.

[0026] First referring primarily to Figure 1, a block diagram provides a general overview of elements (each element further described below) which can be used to implement a particular non-limiting embodiment of the inventive flow cytometer remote monitoring system (1). A first computer user (25) can through use of a command input device (23) such as keystroke of a computer keyboard or a mouse send commands to the processing unit (17) of a first computer (16) to deliver computer operating instructions to a flow cytometer (2)(or other type of computer controlled device or instrument). The flow cytometer (2) can generate a signal (3) which varies based upon the amount of at least one particle characteristic (4) of a plurality of particles (5). The signal (3) can be analyzed by the first computer (16) which can further function to generate a viewable data representation (28) of the analyzed signal which can be viewed by the first computer user (25) on a first computer monitor (26).

[0027] For the purposes of this invention the term "viewable data representation" means an intermittently updated graphical display of data generated by a device viewable by a computer user (25), including, as non-limiting examples, chromatograms, computed tomographies, histograms or the like. As a specific non-limiting example, see Figure 8 which shows a viewable data representation (28) generated having the form of a histogram (29) showing the separation of a plurality of sperm cells (9) into X chromosome bearing sperm cells (14) and Y chromosome bearing sperm cells (15).

[0028] The first computer user (25) can view the viewable data representation (28) to understand the functional condition of the flow cytometer (2)(or other device or instrument). The first computer (16) can further function to generate a plurality of time bound data representation files (35) of the viewable data representation (28) which can be temporarily stored in a memory element (18) of the first computer (16).

[0029] A second computer (37) can provide an image processor (38) which functions to convert the temporary copies of the plurality of time bound data representation files (35) into a first plurality of reduced byte data representation files (39) and a second plurality of reduced byte data representation files (40) retrievably stored in the second computer memory element (41). The second computer (37) as part of a LAN (50) can upon request serve all or part of the first plurality of reduced byte data representation files (39) or the second plurality of reduced byte data representation files (40) to a third computer (42) over a WAN (52).

[0030] Now referring specifically to Figure 2, which provides a particular example of the general hardware means and network means of the second computer (37) including a LAN file server (78) dedicated to provide the functions of converting the temporary copies of the plurality of time bound data representation files (35) into a first plurality of reduced byte data representation files (39) and a second plurality of reduced byte data representation files (40) retrievably stored in the second computer memory element (41) for a flow cytometer or a plurality of flow cytometers in a LAN environment. The embodiment can further provide a second local file server (79) which supports other processing functions in the LAN environment. The embodiment can further provide a WAN file server (80) which further supports the functions of the a LAN file server (78) of converting the temporary copies of the plurality of time bound data representation files (35) into a first plurality of reduced byte data representation files (39) and a second plurality of reduced byte data representation files (40) retrievably stored in the second computer memory element (41) for a flow cytometer or a plurality of flow cytometers in the LAN environment.

[0031] Again referring primarily to Figure 1, a third computer user (81) operating the third computer (42) can generate a request to receive all or part of the first plurality of reduced byte data representation files (39) or the second plurality of reduced byte data representation files (40) retrievably stored in the second computer memory element (41) of any one of a plurality of LANs (50). The files can be transferred from the second computer (37) utilizing a WAN communications device (56) over WAN logical connections (55). An image generator (45) provided by the third computer (42) can generate a viewable reduced byte data representation (46) of all or part of the first plurality of reduced byte data representation files (39) or the second plurality of reduced byte data representation files (40) transferred from the second computer (37) which can be displayed on a third computer monitor (49)(for example see Figure 9 which shows the viewable reduced byte data representation (46) generated from a corresponding two of the plurality of reduced byte data representation files (39) or the second plurality of reduced byte data representation files (40) retrievably stored in the second computer memory element (41) of any one or more of a plurality of LANs (50)) in the form of a histogram showing the separation of sperm cells into X chromosome bearing and Y chromosome bearing populations).

[0032] Now referring primarily to Figures 1 and 3, certain embodiments of the inventive flow cytometer remote monitoring system (1) can in part include a flow cytometer (2). The flow cytometer (2) can function to produce a signal (3)(whether analog, analog converted to digital, or digital) which varies whether in frequency, amplitude, or both frequency and amplitude) based upon change in at least one particle characteristic (4) among a plurality of particles (5). The plurality of particles (5) can be biological particles such as cells, sperm cells, organelles, chromosomes, deoxyribonucleic acids (DNA), ribonucleic acids (RNA), DNA fragments, RNA fragments, proteins, protein fragments, peptides, oligonucleotides, or the like, but can also include non-biological particles such as beads, styrene beads, or the like, or as mixtures of biological particles, mixtures of non-biological particles, or mixtures of biological and non-biological particles. The term "at least one particle characteristic" for the purposes of this invention means at least one part, component, or differentially modified part or component

common to at least a portion of the plurality of particles (5) entrained in the fluid stream (6) which varies in kind or amount between the plurality of particles (5).

[0033] As one non-limiting example, the plurality of particles (5) can be a plurality of sperm cells (9) and the at least one particle characteristic (4) can be the amount of deoxyribonucleic acid ("DNA") (10) contained in each of the plurality of sperm cells (9). The amount of DNA (10) can vary based upon whether the particular one of the plurality of sperm cells (9) contains an X chromosome or a Y chromosome. The X chromosome contains a greater amount of DNA (10) than the corresponding Y chromosome regardless of the male mammal from which the plurality of sperm cells (9) is obtained. Sperm cells (9) can be obtained from any male mammal including for example, a bovid, an ovine, an equine, a pig, a cervid, a canid, a felid, a rodent, a whale, a rabbit, an elephant, a rhinoceros, a primate, or the like, as well as from certain male non-mammal species.

[0034] Certain kinds of flow cytometer (2) devices operate to entrain each of the plurality of particles (5) in a fluid stream (6) which exits a nozzle (7) oscillated to produce droplets (8) in the fluid stream (6). Prior to the break off point for each of the droplets (8) each of the plurality of particles (5) in the fluid stream (6) passes through an interrogation means (11) to generate interrogation event rates of between about 10,000 per second and about 200,000 per second. Typically, the interrogation means (11) includes one or more laser beams (12) through which each of the plurality of particles (5) fall. Each interrogated one of the plurality of particles (5) can in response to interrogation by the laser beam(s) absorb or emit an amount of light (12A). For example, DNA can be quantitatively stained with a dye or fluorochrome such as Hoechst 33342. The stained DNA can emit an amount of light (12) in response to being interrogated with a laser beam. X chromosome bearing sperm cells (14) typically emit a greater amount of light (12A) than Y chromosome bearing sperm cells (15) because each X chromosome bearing sperm cell (14) contains a greater amount of stained DNA than a Y chromosome bearing sperm cell (15).

[0035] The amount of light (12A) emitted from the interrogated one of the plurality of particles (5) can be received by a photomultiplier element (13). The photomultiplier element (13) converts the received amount of emitted light (12A) into a signal (3) which correspondingly varies based upon change in the amount of emitted light (12A). In the analysis of a plurality of sperm cells (9) with a flow cytometer (2), the signal (3) generated can vary based upon the difference in the amount of light (12) generated by X chromosome bearing sperm cells (14) and Y chromosome bearing sperm cells (15) when passed through the interrogation means (11).

[0036] Now referring primarily to Figures 1, 2 and 4, the flow cytometer (2)(or other instrument) can be coupled to, integral with, or provide a first computer (16) having a processing unit (17), a memory element (18), and a bus (19) which operably couples components of the first computer (16), including, without limitation the memory element (18) to the processing unit (17). The first computer (16) may be a conventional computer, a distributed computer, or any other type of computer capable of delivering instructions to a flow cytometer

controller (or other instrument controller) which functions to operate the flow cytometer (or other instrument); the invention is not so limited. The processing unit (17) can comprise without limitation one central-processing unit (CPU), or a plurality of processing units which operate in parallel to process digital information, or a digital signal processor (DSP) plus a host processor, or the like. The bus (19) can be without limitation any of several types of bus configurations such as a memory bus or memory controller, a peripheral bus, and a local bus using any of a variety of bus architectures. The memory element (18) can without limitation be a read only memory (ROM), or a random access memory (RAM), or both. A basic input/output system (BIOS)(20), containing routines that assist transfer of data between the components of the first computer (16), for example during start-up, can be stored in ROM. The first computer (16) can further include a hard disk drive for reading from and writing to a hard disk a magnetic disk drive for reading from or writing to a removable magnetic disk, an optical disk drive for reading from or writing to a removable optical disk such as a CD ROM, or other optical media.

[0037] The hard disk drive, magnetic disk drive, and optical disk drive can be connected to the bus (19) by a hard disk drive interface, a magnetic disk drive interface, and an optical disk drive interface, respectively. The drives and their associated computer-readable media provide nonvolatile storage of computer-readable instructions, data structures, program modules and other data of the first computer (16). It can be appreciated by those skilled in the art that any type of computer-readable media that can store data that is accessible by the first computer (16), such as magnetic cassettes, flash memory cards, digital video disks, Bernoulli cartridges, random access memories (RAMs), read only memories (ROMs), and the like, may be provided by a first computer (16) used in embodiments of the inventive flow cytometer remote monitoring system (1).

[0038] The first computer (16) can further include an operating system (21) and a flow cytometer controller and particle analysis application (22) which may be stored on or in the hard disk, magnetic disk, optical disk, ROM, in RAM by a particular embodiment of a first computer (16) or alternately the functionalities of the a flow cytometer controller and particle analysis application (22) may be implemented as an application specific integrated chip (ASIC) or field programmable gate array (FPGA), or the like, or combinations or permutations thereof.

[0039] A first computer user (25) can enter commands and information into the first computer (16) through one or more command input device(s) (23) such as a keyboard and pointing device such as a mouse. Other command input devices (23) can include a microphone, joystick, game pad, scanner, or the like. These and other command input device(s) (23) are often connected to the processing unit (17) through a serial port interface (24) that can be coupled to the bus (19), but may be connected by other interfaces, such as a parallel port, game port, or a universal serial bus (USB). A monitor (26) or other type of display device can also be connected to the bus (19) via interfaces such as a video adapter (58), or the like. In addition to the monitor (24), the first computer (16) can further include other peripheral output devices (59), such as speakers and printers.

[0040] A click event occurs when the first computer user (25)(or other computer user)

activates or operates at least one function of the flow cytometer controller and particle analysis application (22), or other program or other application function, through the use of a command which for example can include pressing or releasing the left mouse button while a pointer is located over a control icon displayed on the monitor (26). However, it is not intended that a "click event" be limited to the press and release of the left button on a mouse while a pointer is located over a control icon. Rather, the term "click event" for the purposes of this invention broadly encompasses any manner of command by the first computer user (25) through which a function of the operating system (21) or the flow cytometer controller and particle analysis application (22) is activated or performed, whether through clickable selection of one or a plurality of control icon(s), voice command, keyboard stroke(s), mouse button, touch screen, touch pad, or otherwise.

[0041] Again referring primarily to Figures 1, 2 and 4, the first computer (16) and the flow cytometer controller and particle analysis application (22) can in part function to provide a signal analyzer (27) which intermittently or continuously converts the signal (3) produced by the flow cytometer (2) into a viewable data representation (28) of change in the at least one particle characteristic (4) of the plurality of particles (5) analyzed. The viewable data representation (28) can be continuously or intermittently displayed on the monitor (26) or updated upon elapse of a short interval of time such as 100 milliseconds. As a non-limiting example, the signal analyzer (27) coupled to a flow cytometer (2) which interrogates a plurality of sperm cells (9) in a corresponding plurality of droplets (8) can generate a viewable data representation (28) in the form of a histogram (29) which varies based on a frequency of Y chromosome bearing sperm cells (14) or a frequency of X chromosome bearing sperm cells (15) identified within the plurality of sperm cells (9). Certain embodiments of the signal analyzer (27) can further function to establish parameters and timed events by which the plurality of particles (5) can be separated, parsed or divided based upon the presence, absence, or amount of the at least one particle characteristic (4).

[0042] As a non-limiting example, a flow cytometer (2) such as a MOFLO[®] SX can be used to separate or sort the plurality of particles (5) into, discreet sub-populations based upon at least one particle characteristic (4). Subsequent to exiting the nozzle (7), the fluid stream (6) can break into droplets (8) each of which can contain a corresponding one each of the plurality of particles (5). Based on the above-described analysis of each of the plurality of particles (5) in the fluid stream (6), the droplets can be differentiated based on at least one particle characteristic (4) and separated by applying a charge (whether positive or negative) to each one of the droplets (8) analyzed and then deflecting the trajectory of each of the droplets (8) by passing the droplets (8) through a pair of charged plates (31). The trajectory of the positively charged droplets can be altered for delivery to a first container (32) and the trajectory of the negatively charged droplets can be altered for delivery to a second container (33). Uncharged droplets are not deflected and can be delivered to a third container (34) or to a waste stream. With respect to the separation of a plurality of sperm cells (10), a plurality of particle populations (30) can include X chromosome bearing sperm cells (14) isolated in the first container (32) and Y chromosome bearing sperm cells (15) isolated in the second container (33).

[0043] Again referring primarily to Figure 4, the first computer (16) can further provide an image representation generator (34) which can generate one or a plurality of time bound data representation files (35) of the viewable data representation (28) of change in at least one particle characteristic (4) of the plurality of analyzed particles (5) without generating an amount of lag in displaying the viewable data representation (28) or in the operation of functionalities of the flow cytometer (2). The term "lag" for the purposes of this invention means a reduction in performance or a delay in the generation of the viewable data representation (28) or a delay in a function of the flow cytometer (or other instrument) due to competition by various portions of the operating system (21) or the flow cytometer controller and particle analysis application (22) or other program or application for support by the a processing unit (17).

[0044] Typically, the plurality of time bound data representation files (35) of the viewable data representation (28) will include a plurality of bit map image representations (36) or screen shots of the viewable data representation (28) of change in at least one particle characteristic (4) of the plurality of analyzed particles (5) intermittently generated at a pre-determine rate. The pre-determined rate at which each of the plurality of time bound data representation files (35) can be generated will typically be variably adjustable between about 0.1 seconds and about 5 seconds, although other lesser or greater pre-determined rates can be selected so long as the rate does not generate a lag in analyzing the signal (3) generated by the flow cytometer (2) or the function of the flow cytometer (2). The time bound data representation files (35) can be stored temporarily in the memory element (18) of the first computer (16). Typically, each of the plurality of time bound data representation files (35) (generally as a plurality of bit map representations (36)) comprise a file of between about three megabytes and about six megabytes, although the invention is not so limited and each file can include a lesser or greater number of bytes.

[0045] Certain embodiments of the first computer (16) provided with a flow cytometer (2)(or other analysis device) may not provide the functionalities required to capture or generate the plurality of time bound data representation files (35) above-described. In that case a time bound data representation file generator (22A) in the form of a small software application can be stored in the memory element (18) and can function to periodically capture the viewable data representation (28) as above-described to provide the time bound data representation files (35) which can be transferred to the second computer (37).

[0046] Now referring primarily to Figure 5, the inventive flow cytometer remote monitoring system (1) can further include a second computer (37) linked to the first computer (16). The second computer (37) can provide the same or similar hardware means and software means as the first computer (16) although only a memory element (41) is shown along with sufficient hardware means and software means which function to provide an image processor (38). The image processor (38) can function to generate a first plurality of reduced byte data representation files (39) which correspond to each of the plurality of time bound data representation files (35) served from the first computer (16) upon request by the second computer (37). For the purposes of this invention the term "a reduced byte data representation

files" (39) comprises an image file of a lesser bytes than the corresponding time bound data representation file. Typically, each of the plurality of reduced byte data representation files will comprise between about one hundred kilobytes and about two hundred kilobytes (although the invention is not so limited and each file can include a lesser or greater number of bytes). As one example the reduced byte data representation file (39) can be in the form of a .jpeg, which includes substantially lesser bytes than for example a Windows Bitmap format which can be one form of the plurality of time bound image representation files (35) transferred from the first computer (16) to the second computer (37).

[0047] The image processor (38) can further function to generate a second plurality of reduced byte data representation files (40) of the plurality of time bound data representation files (35) served from the first computer (16) upon request by the second computer (37) each one having fewer bytes than the corresponding one of said first plurality of reduced byte data representation files (39). The second plurality of reduced byte data representation files (40) each one having fewer bytes than the corresponding one of the first plurality of reduced byte data representation files (39) can each provide as one non-limiting example an image file of between about two kilobytes and about four kilobytes (although the invention is not so limited and each file can include a lesser or greater number of bytes). Again, as one example the image file can be in the form of a .jpeg.

[0048] The first plurality of reduced byte data representation files (39) and the second plurality of reduced byte data representation files (40) can be generated in parallel from the corresponding plurality of time bound data representation files (35) or the second plurality of reduced byte data representation files (40) can be produced from the corresponding plurality of first plurality of reduced byte data representation files (39). Each of the first plurality of reduced byte data representation files (39) and the second plurality of reduced byte data representation files (40) can be retrievably stored in the second computer memory element (41) of the second computer (37).

[0049] Now referring primarily to Figure 8, the viewable data representation (28) generated by the flow cytometer particle analysis application (22) of the flow cytometer (2)(or other particle analysis application of another type of device) can further include viewable parametric data elements (82) of numeric data type. As a non-limiting example, in the context of a viewable data representation (28) generated by an embodiment of the flow cytometer particle analysis application (22) for a flow cytometer (2), the viewable parametric data elements (82) can include one or more of sample number, operator identification number, date, drop delay status, temperature status, humidity status, laser power status, network status, particle analysis start time, particle analysis stop time, time elapsed, event rate, total events, particle count, percent particles collected, percent particles aborted, percent particles coincident, or the like. In the context, in which the plurality of particles (5) analyzed are a plurality of sperm cells (9), the viewable parametric data elements (82) can further include current percent live, average percent live, current percent dead, average percent dead, current percent purity, average percent purity, percent compare sorter peer, percent, percent compare bull peer, or the like. The viewable parametric data elements (82) included in the viewable data representation (28)

can be converted by the image processor (38) as part of the viewable data representation (28) into a part of a corresponding one of the plurality of time bound data representation files (35) as above described and then converted to a corresponding one of the first plurality of reduced byte data representation files (39) and the second plurality of reduced byte data representation files (40), as above described.

[0050] Now referring to Figure 6 , the inventive flow cytometer remote monitoring system (1) can further include a third computer (42). The third computer (42) can provide the same or similar hardware means and software means as the first computer (16) or sufficient hardware means and software means to function to provide a reduced byte data representation file selection element (43) which can function to generate a request for a selected portion of the first plurality of reduced byte data representation files (39) or a selected portion of the second plurality of reduced byte data representation files (40) to the second computer (37)(the term selected portion can include one or all of the first or second plurality of reduced byte data representation files (39)(40)). The selected portion of the first plurality of reduced byte data representation files (39) or a selected portion of the second plurality of reduced byte data representation files (40) can be stored in a third computer memory element (44). The third computer (42) can further provide an image generator (45) which functions to display the selected portion of the first plurality of reduced by data representation files (39) or the second plurality of reduced byte data representation files (40), or both, in serial order to provide a viewable reduced byte data representation (46) of change in the at least one particle characteristic (4) of said plurality of analyzed particles (5). The a reduced byte data representation file selection element (43) can further include a time period selection element (47) which allows selection of a time bound portion of the first or the second plurality of reduced byte data representation files (39)(40) generated between a first time point and a second time point of the viewable data representation (28).

[0051] The image generator (45) of the third computer (42) can further include a viewing rate selector (48) which can function to allow variably adjusted selection of a viewing rate at which to view the viewable reduced byte data representation (46) of change in the at least one particle characteristic (4) of said plurality of analyzed particles (5). As to certain embodiments of the image generator (45) the viewing rate selector (48) allows variably adjustable selection of an accelerated rate at which the viewable reduced byte data representation (46) of change in said at least one particle characteristic of said plurality of particles (5) analyzed can be serially displayed on a third computer monitor (49). As to certain embodiments of the image generator (45) the viewing rate selector (48) allows variably adjustable selection of a decelerated rate at which the viewable reduced byte data representation (46) of change in said at least one particle characteristic of said plurality of analyzed particles can be serially displayed on the third computer monitor (49).

[0052] Now referring primarily to Figure 1, certain embodiments of the inventive flow cytometer remote monitoring system (1) can further include a local area network (50) ("LAN") at a first location (51) which includes local area network logical connections (53) between the flow cytometer (2) and the first computer (16)(or a plurality of flow cytometers (2) each coupled

to a corresponding plurality of first computers (16)) and the second computer (37). These logical connections (53) can be achieved by a local area network communication device (54) coupled to or a part of the first computer (16) or the second computer (37) or both. As to certain embodiments of the invention there can be a plurality of local area networks (50) each established at a plurality of discrete locations (51 A).

[0053] Again referring primarily to Figure 1, certain embodiments of the inventive flow cytometer remote monitoring system (1) can further include a wide area network (52) ("WAN") such as the Internet which includes wide area network logical connections (55) which allows communication between the third computer (42) established at a second location (56) discrete from the local area network (50) at the first location (51) or the plurality of first locations (51A) and the second computer (37) of any local area network (50). This configuration allows the third computer (42) to retrieve from any second computer (42) any one, a portion of, or all of the plurality of time bound data representation files (35), the a plurality of bit map image representations (36), the first plurality of reduced byte data representation files (39), or the second plurality of reduced byte data representation files (40) from the second computer memory element (41) for display and viewing as above described without generating an amount of lag in analyzing the signal (3) from the at least one flow cytometer (2) by the first computer (16).

[0054] When included in a WAN (52), the second computer (37) and the third computer (42) can further include a wide area network communications device (56) such as a modem for establishing communications over the WAN (52)(such as the Internet (57)). The wide area network communications device (56) can be internal or external to the second computer (37) and the third computer (42) and can be connected to the bus (19) via a serial port interface (24). In a WAN (52) environment, the second computer memory element (41) can comprise a plurality of second computer memory elements (41) coupled to the second computer (37) via the WAN which allows distributed retrievable storage of any one, a portion of, or all of the plurality of time bound data representation files (35), the a plurality of bit map image representations (36), the first plurality of reduced byte data representation files (39), or the second plurality of reduced byte data representation files (40). It can be appreciated that the LAN communication device (54), the WAN communications device (56), the LAN logical connections (53), and the WAN logical connections (55) and shown and described are exemplary and other hardware means or logical connection means, and communication means can be utilized for establishing a communications link between the second computer (16) and the third computer (42).

[0055] As but one non-limiting example of providing WAN connectivity between any second computer (37) and any third computer (42) for the purposes above-described a pc-engines WRAP1E circuit boards can run Valemount Networks StarOS router software on a 32meg Compact Flash (CF) memory card to provide a Virtual Distribution System (VDS) tunnel (similar to a Virtual Private Network or VPN) connected to a central router server and each second computer (37) local area network connects as a client to the main central station server to manage all network routing and class based queueing for the entire WAN.

[0056] While the computer means and the network means shown in Figures 1-6 can be utilized to practice the invention including the best mode, it is not intended that the description of the best mode of the invention or any preferred embodiment of the invention be limiting with respect to the utilization of a wide variety of similar, different, or equivalent computer means or network means to practice embodiments of the invention which include without limitation hand-held devices, such as personal digital assistants or camera/cell phone, multiprocessor systems, microprocessor-based or programmable consumer electronics, network PCs, minicomputers, mainframe computers, PLCs, or the like, in various permutations and combinations.

[0057] Now referring primarily to Figures 5 and 7, the second computer (37) can further include an image archive management module (60) which functions to delete, modify, or otherwise reduce file size of the plurality of time bound data representation files (35), the first plurality of reduced byte data representation files (39), or the second plurality of reduced byte data representation files (40) whether individually or collectively to minimize the memory space in which image files are retrievably stored in the second computer memory element (41). Each one of the plurality of time bound data representation files (35) can be converted into a corresponding one of the first plurality of reduced byte data representation files (39) and a corresponding one of the second plurality of reduced byte data representation files (40). Each one of the second plurality of reduced byte data representation files (40) can be stored in the second computer memory (41) in the LAN (51) until requested by the third computer (42) and each one of the first plurality of reduced byte data representation files (39) can also be stored in the second computer memory (41) until requested by the third computer (42) or until modified or deleted by function of a data storage minimization application (61) which governs image resolution priority (62) and further functions to govern image time value priority (63).

[0058] Now referring primarily to Figure 7, which provides a graph which plots image resolution priority (62A) against image time value priority (63A) with respect to each one of the first plurality of reduced byte data representation files (39) and each one of the second plurality of reduced byte data representation files (40). As can be understood from the Figure, each of the first plurality of reduced byte data representation files (39) can be correspondingly converted into a first plurality of reduced resolution reduced byte data representation files (64) and each of the second plurality of reduced byte data representation files (40) can be correspondingly converted into a second plurality of reduced resolution reduced byte data representation file (65). Additionally, any one of the first plurality of reduced byte data representation files (39), any one of the second plurality of reduced byte data representation files (40), any one of the first plurality of reduced resolution reduced byte data representation files (64), and any one of the second plurality of reduced resolution reduced byte data representation file (65) can reproduced at different times, and then progressively categorized as image time values (66, 67, and 68; 72, 73, and 74; 69, 70, and 71; and 75, 76, and 77 respectively) assigned lesser image time value priority (63A) with elapsed time. In this case, images with greater image time value priority (63A)(for example 66, 72, 69, and 75) can be categorically more important than images with lesser image time value priority (63A) (for

example 68, 74, 71, and 77). Similarly, any one of the of the first plurality of reduced byte data representation files (39), any one of the second plurality of reduced byte data representation files (40), and any one of the categories having lesser image value priority (63A) can be progressively rewritten as images of lesser image resolution value priority (62A) and then progressively categorized as image resolution values (72, 75, 73, 76, 74 and 77) assigned lesser image resolution value priority (62A) with decreased resolution.

[0059] The number of image value categories (82) could be unlimited but will generally be less than one dozen. Generally, categories with lesser image time value priority (63A) (for example 68, 74, 71, and 77) or lesser image resolution value priority (62A), or both lesser image time value priority (63A) and lesser image resolution value priority (63A)(for example 74 or 77) can be stored on low cost WAN file server (80)(see Figure 2 for example) while files of greater image time value (for example 66, 72, 69, and 75), can be stored in the second computer memory element (41) of a dedicated LAN file server (78) or general purpose LAN file server (79) (see Figure 2 for example). Simultaneous deletion and rewriting of image files can occur by function of the data storage minimization application (61) to produce the image value categories (82).

[0060] As can be easily understood from the foregoing, the basic concepts of the inventive flow cytometer remote monitoring system (1) may be embodied in a variety of ways. As such, the particular embodiments or elements of the invention disclosed by the description or shown in the figures or tables accompanying this application are not intended to be limiting, but rather exemplary of the numerous and varied embodiments generically encompassed by the invention or equivalents encompassed with respect to any particular element thereof. In addition, the specific description of a single embodiment or element of the invention may not explicitly describe all embodiments or elements possible; many alternatives are implicitly disclosed by the description and figures.

[0061] It should be understood that each element of an apparatus or each step of a method may be described by an apparatus term or method term. Such terms can be substituted where desired to make explicit the implicitly broad coverage to which this invention is entitled. As but one example, it should be understood that all steps of a method may be disclosed as an action, a means for taking that action, or as an element which causes that action. Similarly, each element of an apparatus may be disclosed as the physical element or the action which that physical element facilitates. As but one example, the disclosure of a "monitor" should be understood to encompass disclosure of the act of "monitoring" - - whether explicitly discussed or not -- and, conversely, were there effectively disclosure of the act of "monitoring", such a disclosure should be understood to encompass disclosure of a "monitor" and even a "means for monitoring." Such alternative terms for each element or step are to be understood to be explicitly included in the description.

REFERENCES CITED IN THE DESCRIPTION

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Patent documents cited in the description

- US12151156B [0001]
- US20030078703A1 [0004] [0004]

Patentkrav

1. Fremgangsmåde til at fjernovervåge et flowcytometer, hvilken fremgangsmåde omfatter trinnene:

- 5 a) at producere et signal (3) fra det nævnte flowcytometer (2), som varierer baseret på tilstedeværelsen, fraværet eller mængden af mindst en partikelkarakteristik (4) i et antal analyserede partikler (5);
- b) at analysere det nævnte signal (3) med en første computer
10 (16), der er koblet til det nævnte flowcytometer (2) for kontinuerligt at konvertere det nævnte signal (3) til en synlig datarepræsentation (28) af ændring i tilstedeværelsen, fraværet eller mængden af den nævnte mindst ene partikelkarakteristik (4) i det nævnte antal analyserede
15 partikler (5),
hvor den synlige datarepræsentation (28) betyder et med mellemrum opdateret grafisk display, der er genereret af en indretning, som kan ses af en første computerbruger (25) for at forstå den funktionelle tilstand af flowcytometeret (2);
- 20 c) at levere, med den nævnte første computer (16), i seriel rækkefølge midlertidige kopier af et antal tidsbestemte datarepræsentationsfiler (35) af den nævnte synlige datarepræsentation (28) af ændringen i tilstedeværelsen, fraværet eller mængden af den nævnte mindst ene
25 partikelkarakteristik (4) af det nævnte antal analyserede partikler (5) til en anden computer (37) uden at generere en forsinkelse i analyse af det nævnte signal (3) fra det nævnte mindst ene flowcytometer (2), hvor et antal tidsbestemte datarepræsentationsfiler (35) af den nævnte synlige
30 datarepræsentation (28) indbefatter et antal bitmap-repræsentationer (36) eller skærbilleder af den synlige datarepræsentation (28) af ændring i tilstedeværelsen, fraværet eller mængden af den nævnte mindst ene partikelkarakteristik (4) i det nævnte antal analyserede
35 partikler (5), der med mellemrum er genereret ved en forudbestemt hastighed af den nævnte første computer, og hvor forsinkelsen omfatter en forsinkelse i genereringen af den synlige datarepræsentation (28) eller i en funktion af

flowcytometeret;

- d) at behandle de nævnte leverede midlertidige kopier af et antal tidsbestemte datarepræsentationsfiler (35) med den nævnte anden computer (37) for tilsvarende at generere et
- 5 første antal byte-reducerede datarepræsentationsfiler (39), der er genfindeligt lagret i et andet computerlagerelement (41);
- e) at behandle de nævnte leverede midlertidige kopier af et antal tidsbestemte datarepræsentationsfiler (35) med den
- 10 nævnte anden computer (37) for at generere et andet antal byte-reducerede datarepræsentationsfiler (40), der er genfindeligt lagret i et andet computerlagerelement (41), hvor hver én har færre bytes end den tilsvarende én af det nævnte første antal byte-reducerede datarepræsentationsfiler;
- 15 f) at levere, med den nævnte anden computer (37), en anmodet del af det nævnte andet antal byte-reducerede datarepræsentationsfiler (40) til en tredje computer (42); og
- g) at vise hver én af den nævnte del af det nævnte andet antal af byte-reducerede datarepræsentationsfiler (40) i seriel
- 20 rækkefølge med den nævnte tredje computer (42) for at tilvejebringe en synlig byte-reduceret datarepræsentation (46) af ændring i tilstedeværelsen, fraværet eller mængden af den nævnte mindst ene partikelkarakteristik (4) af det nævnte antal analyserede partikler (5) til en tredje computerbruger
- 25 (81).

2. Fremgangsmåde til fjernovervågning af et flowcytometer som beskrevet i krav 1, hvor det nævnte trin at levere, med den nævnte første computer (16), i seriel rækkefølge
- 30 midlertidige kopier af et antal tidsbestemte datarepræsentationsfiler (35) af den nævnte synlige datarepræsentation (28) af ændring i tilstedeværelsen, fraværet eller mængden af den nævnte mindst ene partikelkarakteristik (4) i det nævnte antal analyserede
- 35 partikler (5) til en anden computer (37) uden at generere en forsinkelse i analyse af det nævnte signal (3) fra det nævnte mindst ene flowcytometer (2), omfatter trinnet med at levere, med den første computer (16), i seriel rækkefølge midlertidige

kopier af et antal tidsbestemte datarepræsentationsfiler (35) af den nævnte synlige datarepræsentation (28) af ændring i tilstedeværelse, fraværet eller mængden af den nævnte mindst ene partikelkarakteristik (4) af det nævnte antal analyserede partikler (5) til en anden computer (37) for at reducere eller undgå en forsinkelse i analyse af det nævnte signal (2) fra det nævnte mindst ene flowcytometer (37).

3. Fremgangsmåde til fjernovervågning af et flowcytometer (2) som beskrevet i krav 1, hvor det nævnte trin til at producere et signal (3) fra det nævnte flowcytometer (2), som varierer baseret på tilstedeværelsen, fraværet eller mængden af mindst en partikelkarakteristik (4) i et antal analyserede partikler (5) omfatter trinnet at producere et signal (3) fra det nævnte flowcytometer (2), som varierer baseret på en mængde deoxyribonukleinsyre (10), der er indeholdt i hver af et antal analyserede partikler (5).

4. Fremgangsmåde til fjernovervågning af et flowcytometer som beskrevet i krav 3, hvor trinnet at producere et signal (3) fra det nævnte flowcytometer (2), som varierer baseret på en mængde deoxyribonukleinsyre (10), der er indeholdt i hver af et antal analyserede partikler (5) omfatter trinnet at producere et signal (3), som varierer baseret på tilstedeværelsen eller fraværet af et X-kromosom eller et Y-kromosom i hver af det nævnte antal sædceller (9).

5. Fremgangsmåde til fjernovervågning af et flowcytometer som beskrevet i krav 4, hvor den nævnte synlige datarepræsentation (28) omfatter mindst et histogram (29), der varierer baseret på hyppigheden af det nævnte antal Y-kromosom-bærende sædceller (9) eller X-kromosom-bærende sædceller (9).

6. Fremgangsmåde til fjernovervågning af et flowcytometer som beskrevet i krav 1, hvor hver af det nævnte antal bitmap-repræsentationer omfatter en billedfil på mellem omkring tre megabyte og omkring seks megabyte.

7. Fremgangsmåde til fjernovervågning af et flowcytometer som beskrevet i krav 1, hvor det nævnte første antal byte-reducerede datarepræsentationsfiler (39) omfatter et første
5 antal billedfiler hver på mellem omkring hundrede kilobyte og omkring to hundrede kilobyte.

8. Fremgangsmåde til fjernovervågning af et flowcytometer som beskrevet i krav 7, hvor det nævnte andet antal byte-reducerede datarepræsentationsfiler (40), der hver én har
10 færre bytes end den tilsvarende én af det nævnte første antal byte-reducerede datarepræsentationsfiler (39), omfatter et andet antal billedfiler hver på mellem omkring to kilobyte og omkring fire kilobyte.

9. Fremgangsmåde til fjernovervågning af et flowcytometer som beskrevet i krav 8, hvor det nævnte trin at levere, med den nævnte anden computer (37), en anmodet del af det nævnte
15 andet antal byte-reduceret datarepræsentationsfiler (40) til en tredje computer (42) omfatter trinnet at levere det nævnte andet antal byte-reducerede datarepræsentationsfiler (40) via et fjerndatanetværk (52) til en tredje computer (42).

10. Fremgangsmåde til fjernovervågning af et flowcytometer som beskrevet i krav 11, hvor det nævnte trin at vise hver én
25 af den nævnte del af det nævnte andet antal byte-reducerede datarepræsentationsfiler (40) i seriel rækkefølge med den nævnte tredje computer (42) yderligere omfatter trinnet at vælge en hastighed, ved hvilken at se den nævnte synlige byte-reducerede datarepræsentation (46) af ændring i
30 tilstedeværelsen, fraværet eller mængden af den nævnte mindst ene partikelkarakteristik (4) af det nævnte antal analyserede partikler (5).

11. Fremgangsmåde til fjernovervågning af et flowcytometer som beskrevet i krav 10, hvilken fremgangsmåde yderligere omfatter trinnet at vise hver én af den nævnte del af det
35 nævnte andet antal byte-reducerede datarepræsentationsfiler

(40) i seriel rækkefølge med den nævnte tredje computer (42) for at tilvejebringe en accelereret eller decelereret synlig byte-reduceret datarepræsentation (46) af ændring i tilstedeværelsen, fraværet eller mængden af den nævnte mindst ene partikelkarakteristik (4) af det nævnte antal analyserede partikler (5).

12. Fremgangsmåde til fjernovervågning af et flowcytometer som beskrevet i krav 10, hvilken fremgangsmåde yderligere omfatter trinnet at vælge den nævnte del af det nævnte andet antal byte-reducerede datarepræsentationsfiler (40) for at vise med den nævnte tredje computer (42).

13. Fremgangsmåde til fjernovervågning af et flowcytometer som beskrevet i krav 12, hvilket trin at udvælge den nævnte del af det nævnte andet antal byte-reducerede datarepræsentationsfiler (40) til at vise med den nævnte tredje computer (42) yderligere omfatter trinnet at vælge en tidsbestemt del af det nævnte andet antal byte-reducerede datarepræsentationsfiler (40), der er genereret mellem et første tidspunkt og et andet tidspunkt af den nævnte synlige datarepræsentation (28).

14. Fremgangsmåde til fjernovervågning af et flowcytometer som beskrevet i krav 1, hvilken fremgangsmåde yderligere omfatter trinnet at etablere, ved et første sted (51), der har et første lokalt netværk (50), den nævnte anden computer (37), som er forbundet til et antal af den nævnte første computer (16), der hver tilsvarende er koblet til én af et antal af det nævnte flowcytometer (2), hvilken nævnte anden computer (37) yderligere tilvejebringer et hukommelseselement (41), som tillader genfindelig lagring af det nævnte første antal byte-reducerede datarepræsentationsfiler (39) og det nævnte andet antal byte-reducerede datarepræsentationsfiler (40), som svarer til det nævnte antal af tidsbestemte datarepræsentationsfiler (35), der er leveret af hver én af det nævnte antal af den nævnte første computer (16).

15. Fremgangsmåde til fjernovervågning af et flowcytometer som beskrevet i krav 14, hvilken fremgangsmåde yderligere omfatter trinnet at etablere, ved et antal steder (51A), et tilsvarende antal lokale netværkssøgninger, hvilke omfatter
5 den nævnte anden computer (42), der er forbundet til det nævnte antal af den nævnte første computer (16), som hver er tilsvarende koblet til én af det nævnte antal af det nævnte flowcytometer (2), hvilken nævnte anden computer (37) yderligere tilvejebringe et hukommelselement (41), som
10 tillader genfindelig lagring af det nævnte første antal byte-reducerede datarepræsentationsfiler (39) og det nævnte andet antal byte-reducerede datarepræsentationsfiler (40), der svarer til det nævnte antal tidsbestemte datarepræsentationsfiler (35), som er leveret af hver én af
15 det nævnte antal af den nævnte første computer (16).

16. Fremgangsmåde til fjernovervågning af et flowcytometer som beskrevet i krav 15, hvilken fremgangsmåde yderligere omfatter trinnet at tilvejebringe et fjernnetværk (52), der
20 indbefatter det nævnte antal lokalnetværk ved det nævnte antal steder (51A), hvilket nævnte første lokalnetværk (50) indbefatter den nævnte tredje computer.

17. Fremgangsmåde til fjernovervågning af et flowcytometer
25 som beskrevet i krav 16, hvilken fremgangsmåde yderligere omfatter trinnet selektivt at genfinde over det nævnte fjerndatanetværk (52) en del af det nævnte andet antal byte-reducerede datarepræsentationsfiler (40) svarende til det nævnte antal tidsbundne datarepræsentationsfiler (35), der er
30 leveret af én af det nævnte antal første computere (16) af det nævnte første lokalnetværk (50) eller af et andet af det nævnte antal lokale netværk.

18. Fremgangsmåde til fjernovervågning af et flowcytometer
35 som beskrevet i krav 17, hvor trinnet at vise hver én af den nævnte del af det nævnte andet antal byte-reducerede datarepræsentationsfiler (40) i seriel rækkefølge med den nævnte tredje computer (42) for at tilvejebringe en synlig

byte-reduceret datarepræsentation (46) af ændring i tilstedeværelsen, fraværet eller mængden af den nævnte mindst ene partikelkarakteristik (4) af det nævnte antal analyserede partikler (5) omfatter trinnet samtidigt at vise hver selektivt genfundet nævnte del af det nævnte andet antal byte-reducerede datarepræsentationsfiler (40), der svarer til det nævnte antal tidsbestemte datarepræsentationsfiler (35), der er leveret af én det nævnte antal første computere (16) af det nævnte andet antal af det nævnte antal lokalnetværk.

10

19. Fjernovervåget flowcytometer, hvilket flowcytometer omfatter:

a) et flowcytometer (2), som producerer et signal (3), der varierer baseret på tilstedeværelsen, fraværet eller mængden af mindst en partikelkarakteristik (4) i et antal analyserede partikler (5);

b) en første computer (16), der er koblet til det nævnte flowcytometer (2), hvor den nævnte første computer (16) tilvejebringer en signalanalysator (27), som kontinuerligt omdanner det nævnte signal (3), der er produceret af det nævnte flowcytometer (2) til en synlig datarepræsentation (28) af en ændring i tilstedeværelsen, fraværet eller mængden af den nævnte mindst ene partikelkarakteristik (4) i det nævnte antal analyserede partikler (5), og hvor den nævnte første computer (16) yderligere tilvejebringer en billedrepræsentationsgenerator (34), som genererer et antal tidsbestemte datarepræsentationsfiler (35) af den nævnte synlige datarepræsentation (28) af ændring i tilstedeværelsen, fraværet eller mængden af den nævnte mindst ene partikelkarakteristik (4) af det nævnte antal analyserede partikler (5) uden at generere en forsinkelse i analyse af det nævnte signal (3), der er produceret af det nævnte mindst ene flowcytometer (2), hvor den synlige datarepræsentation (28) betyder et med mellemrum opdateret grafisk display, der er genereret af en indretning, som kan ses af en første computerbruger (25) for at forstå den funktionelle tilstand af flowcytometeret (2), hvor et antal tidsbestemte datarepræsentationsfiler (35) af den nævnte synlige

datarepræsentation (28) indbefatter et antal bitmap-repræsentationer (36) eller skærbilleder af den synlige datarepræsentation (28) af ændring i tilstedeværelsen, fraværet eller mængden af den nævnte mindst ene partikelkarakteristik (4) i det nævnte antal analyserede partikler (5), der med mellemrum er genereret ved en forudbestemt hastighed, og hvor en forsinkelse omfatter en forsinkelse i genereringen af den synlige datarepræsentation (28) eller i en funktion af flowcytometeret;

10 c) en anden computer (37), der er forbundet til det første computer (16), som tilvejebringer en billedprocessor (38), der tilsvarende generer et første antal af byte-reducerede datarepræsentationsfiler (39) af midlertidige kopier af det nævnte antal tidsbestemte datarepræsentationsfiler (35), som
15 er leveret fra den nævnte første computer (16), hvilke første antal byte-reducerede datarepræsentationsfiler (39) er genfindeligt lagret i et andet computerlagerelement (41), og hvor billedprocessoren (38) yderligere genererer et andet antal byte-reducerede datarepræsentationsfiler (40) af
20 midlertidige kopier af det nævnte antal tidsbestemte datarepræsentationsfiler (35), der er leveret fra den nævnte første computer (16), hvor hver én har færre bytes end den tilsvarende ene af det nævnte antal byte-reduceret datarepræsentationsfiler, hvilket andet antal byte-reducerede
25 datarepræsentationsfiler (40) er genfindeligt lagret i et andet computerlagerelement; og

d) en tredje computer (42), som tilvejebringer et byte-reduceret datarepræsentationsudvælgelseselement (43), som genererer en anmodning om en udvalgt del af det nævnte andet
30 antal byte-reducerede datarepræsentationsfiler (40) til den nævnte anden computer (37), og hvor den tredje computer (42) yderligere tilvejebringer en billedgenerator (45), som viser den nævnte udvalgte del af det nævnte andet antal byte-reducerede datarepræsentationsfiler (40) i seriel rækkefølge
35 for at tilvejebringe en synlig byte-reduceret datarepræsentation (46) af ændring i tilstedeværelsen, fraværet eller mængden af den nævnte mindst ene partikelkarakteristik (4) af det nævnte antal analyserede

partikler (5).

20. Fjernovervåget flowcytometer som beskrevet i krav 19,
hvor det nævnte signal (3) fra det nævnte flowcytometer (2)
5 varierer baseret på en mængde deoxyribonukleinsyre (10), der
er indeholdt i hver af det nævnte antal analyserede partikler
(5).

21. Fjernovervåget flowcytometer som beskrevet i krav 20,
10 hvor det nævnte signal (3) fra det nævnte flowcytometer (2)
varierer baseret på tilstedeværelsen eller fraværet af et X-
kromosom eller et Y-kromosom i hver af et antal sædceller (9).

22. Fjernovervåget flowcytometer som beskrevet i krav 21,
15 hvor den nævnte synlige datarepræsentation (28) omfatter
mindst et histogram (29), der varierer baseret på en hyppighed
af det nævnte antal Y-kromosom-bærende sædceller (9) eller X-
kromosom-bærende sædceller (9).

20 23. Fjernovervåget flowcytometer som beskrevet i krav 19,
hvor det nævnte antal tidsbestemte datarepræsentationsfiler
(35) af den nævnte synlige datarepræsentation (28) af ændring
i tilstedeværelsen, fraværet eller mængden af den nævnte
mindst ene partikelkarakteristik (4) af det nævnte antal
25 analyserede partikler (5) omfatter et antal bitmap-
billedrepræsentationer af den nævnte synlige
datarepræsentation (28) af ændring i tilstedeværelsen,
fraværet eller mængden af den nævnte mindst ene
partikelkarakteristik (4) af det nævnte antal analyserede
30 partikler (5).

24. Fjernovervåget flowcytometer som beskrevet i krav 23,
hvor hver af det nævnte antal bitmap-repræsentationer omfatter
en billedfil på mellem omkring tre megabyte og omkring seks
35 megabyte.

25. Fjernovervåget flowcytometer som beskrevet i krav 24,
hvor det nævnte antal første byte-reducerede

datarepræsentationsfiler (39) omfatter et første antal billedfiler hver på mellem omkring hundrede kilobyte og omkring to hundrede kilobyte.

5 26. Fjernovervåget flowcytometer som beskrevet i krav 25, hvor det nævnte andet antal byte-reducerede datarepræsentationsfiler (40), der hver har færre bytes end den tilsvarende én af det nævnte første antal byte-reducerede datarepræsentationsfiler (39), omfatter et andet antal
10 billedfiler hver på mellem omkring to kilobyte og omkring fire kilobyte.

27. Fjernovervåget flowcytometer som beskrevet i krav 26, hvilket flowcytometer yderligere omfatter et fjernnetværk (52)
15 over hvilket en tredje computer (42) genererer en anmodning om en udvalgt del af det nævnte andet antal byte-reducerede datarepræsentationsfiler (40) til den nævnte anden computer (37).

20 28. Fjernovervåget flowcytometer som beskrevet i krav 27, hvor den nævnte tredje computer (42) yderligere omfatter en visningsrateudvælger (48), der er koblet til den nævnte billedgenerator (45), som tillader valg af en hastighed, ved hvilken at se den nævnte byte-reducerede datarepræsentation
25 (46) af ændring i tilstedeværelsen, fraværet eller mængden af den nævnte mindst ene partikelkarakteristik (4) af det nævnte antal analyserede partikler (5).

29. Fjernovervåget flowcytometer som beskrevet i krav 28,
30 hvor visningsrateudvælgeren (48) yderligere tillader valg af en accelereret eller decelereret synlig byte-reduceret datarepræsentation (46) af ændring i tilstedeværelsen, fraværet eller mængden af den nævnte mindst ene partikelkarakteristik (4) af det nævnte antal af analyserede
35 partikler (5).

30. Fjernovervåget flowcytometer som beskrevet i krav 29, hvor det byte-reducerede datarepræsentationsudvælgelseselement

(43) yderligere indbefatter et tidsperiodeudvælgelseselement (47), som tillader valg af en tidsbestemt del af det nævnte andet antal byte-reducerede datarepræsentationsfiler (40), der er genereret mellem et første tidspunkt og et andet tidspunkt af den nævnte synlige datarepræsentation (28).

31. Fjernovervåget flowcytometer som beskrevet i krav 19, hvor den nævnte første computer (16) forbinder til den nævnte anden computer (37) i et lokalt netværk (50).

10 32. Fjernovervåget flowcytometer som beskrevet i krav 31, hvor den nævnte første computer (16) omfatter et antal første computere (16), der hver er tilsvarende koblet til én af et antal flowcytometre (2).

15 33. Fjernovervåget flowcytometer som beskrevet i krav 32, hvor den nævnte billedgenerator (45) yderligere tilvejebringer et sammenfaldende billedvisningselement, som tillader sammenfaldende visning af et antal af de nævnte udvalgte dele af det nævnte andet antal byte-reducerede datarepræsentationsfiler (40) i seriel rækkefølge for at tilvejebringe en synlig byte-reduceret datarepræsentation (46) af ændring i tilstedeværelsen, fraværet eller mængden af den nævnte mindst ene partikelkarakteristik (4) af det nævnte antal analyserede partikler (5) for hver én af de nævnte antal flowcytometre (2).

20 34. Fjernovervåget flowcytometer som beskrevet i krav 33, hvor den nævnte tredje computer (42) forbinder til det nævnte lokale netværk (50) over et fjernnetværk (52).

30 35. Fjernovervåget flowcytometer som beskrevet i krav 34, hvor det nævnte fjernnetværk (52) omfatter Internettet (57).

DRAWINGS

1

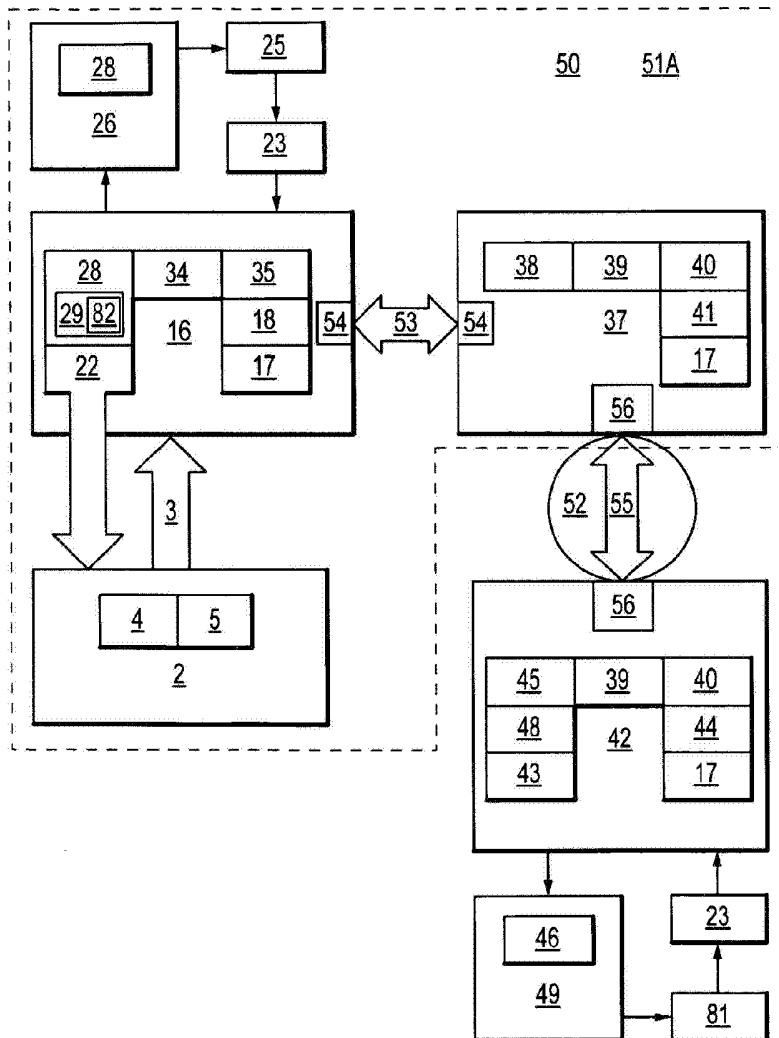


FIG.1

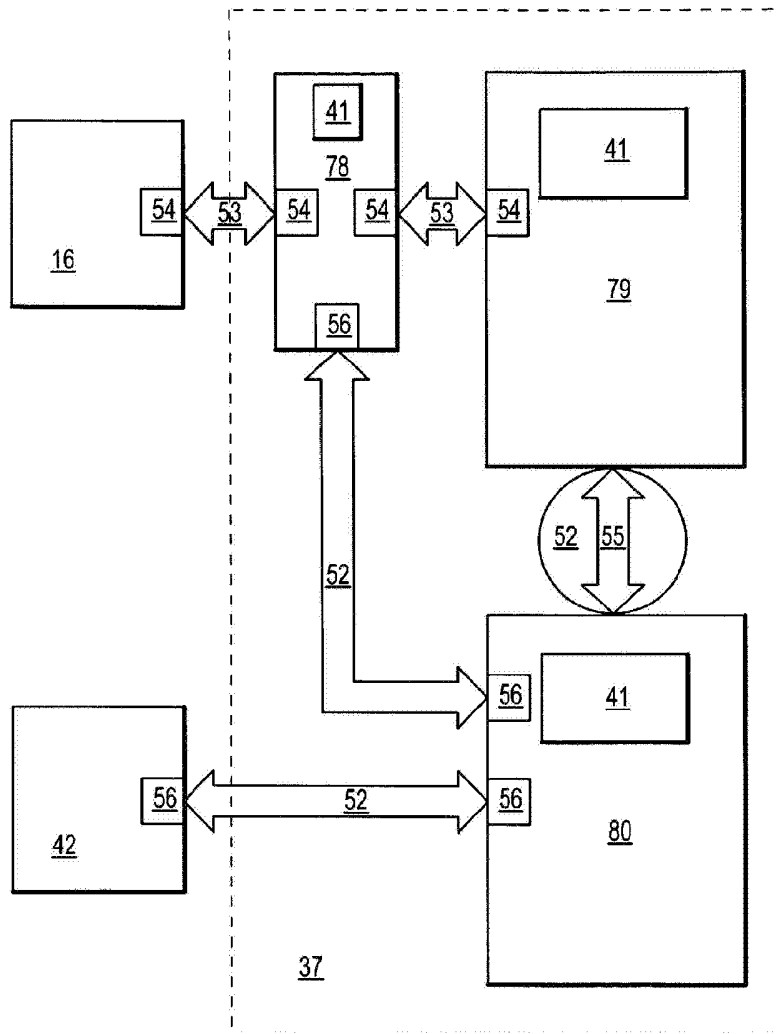


FIG.2

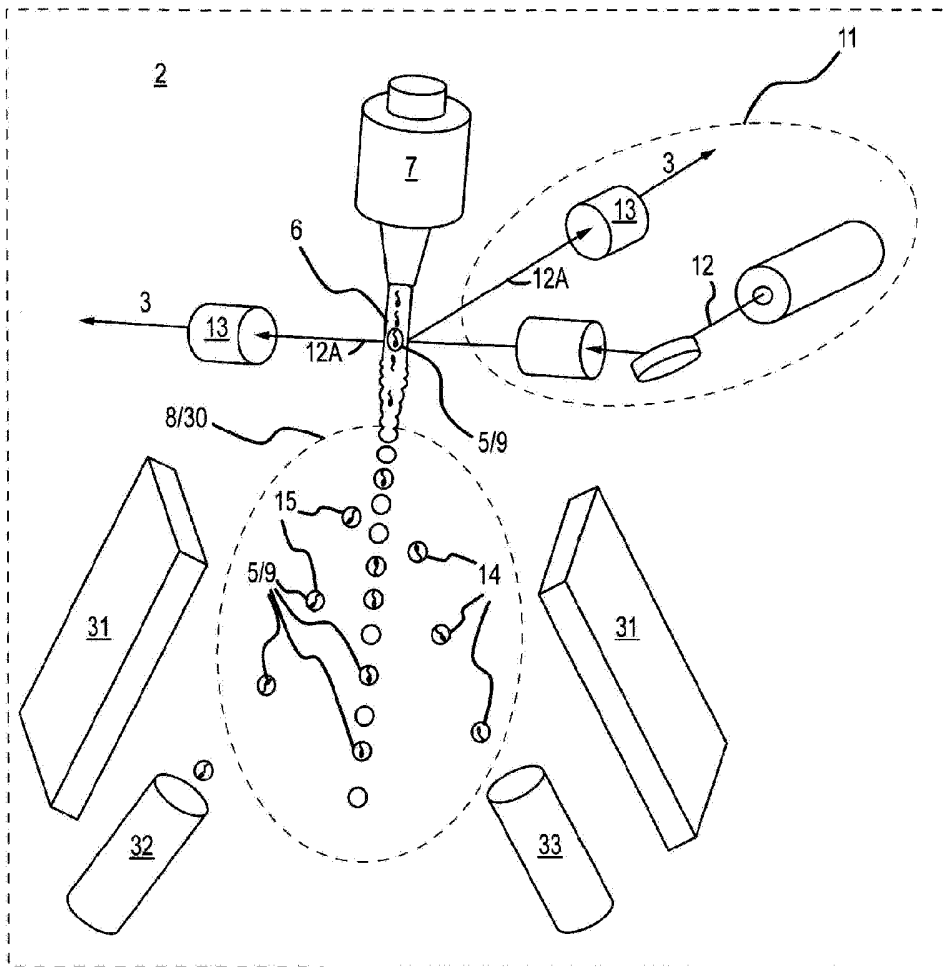


FIG. 3

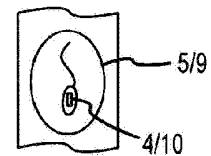


FIG. 3A

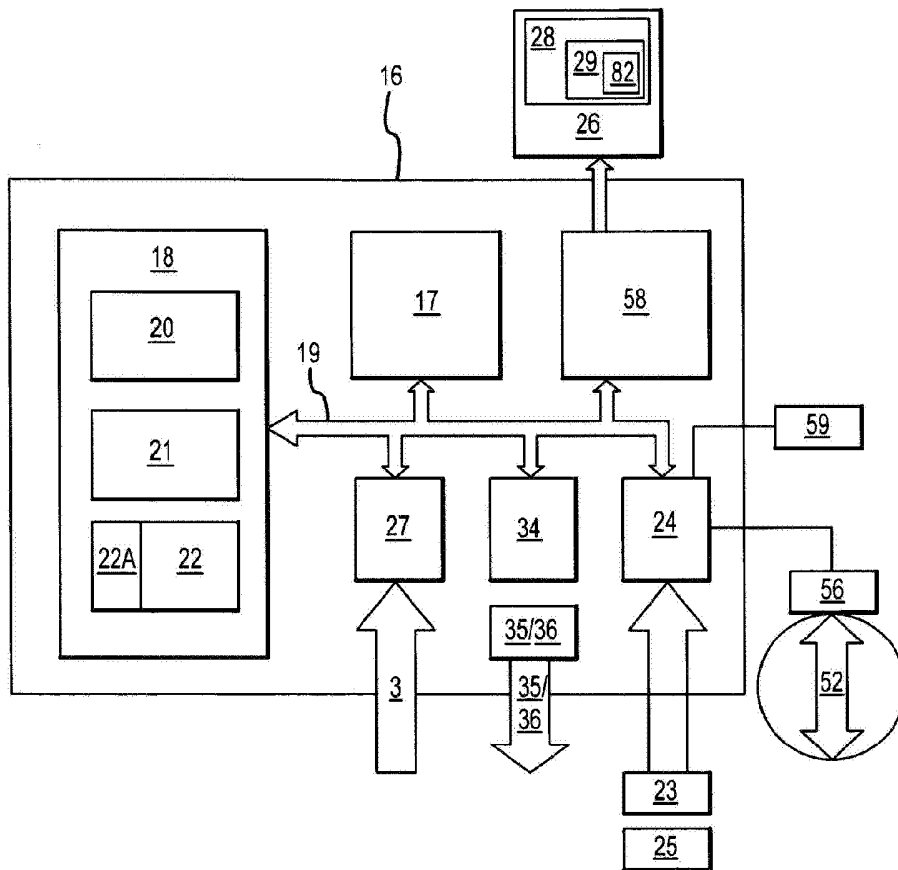


FIG.4

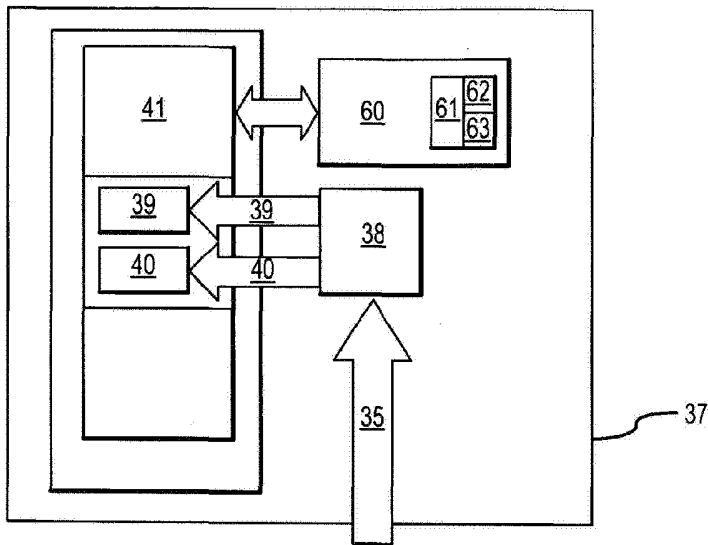


FIG.5

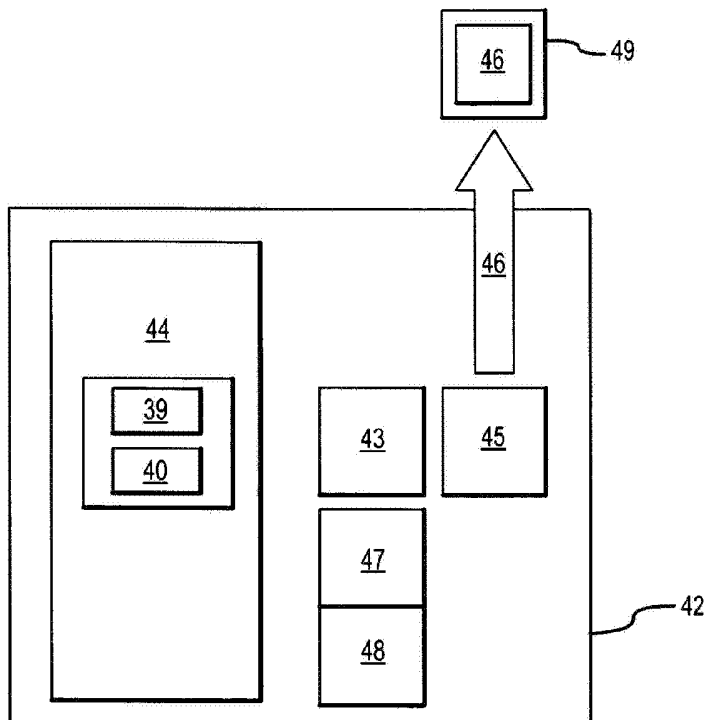


FIG.6

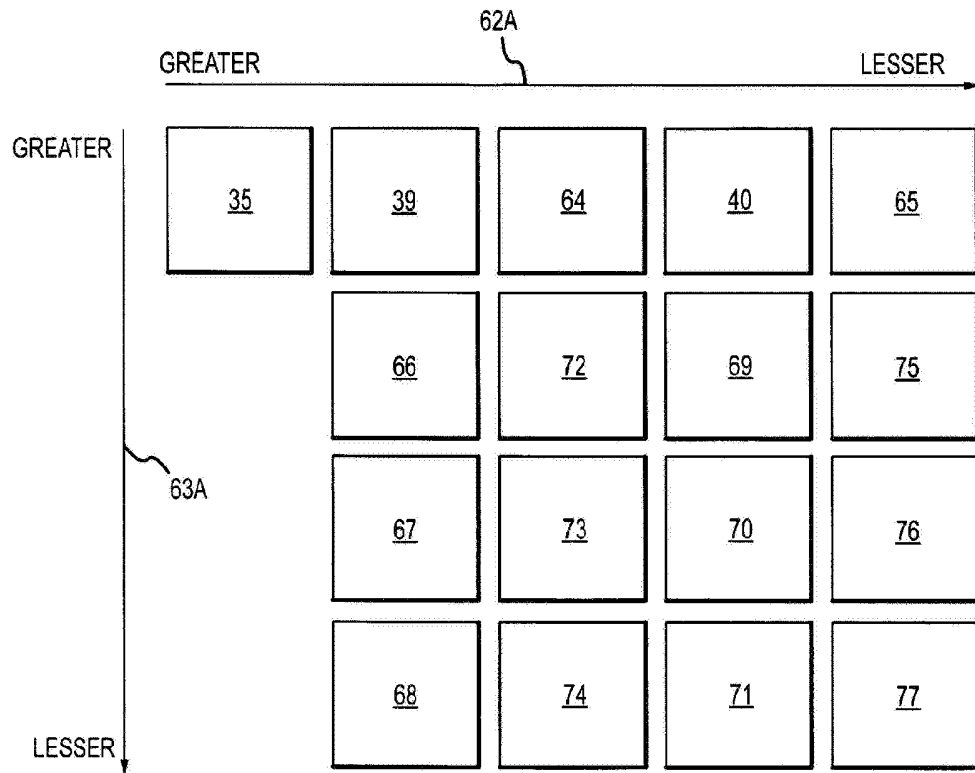


FIG.7

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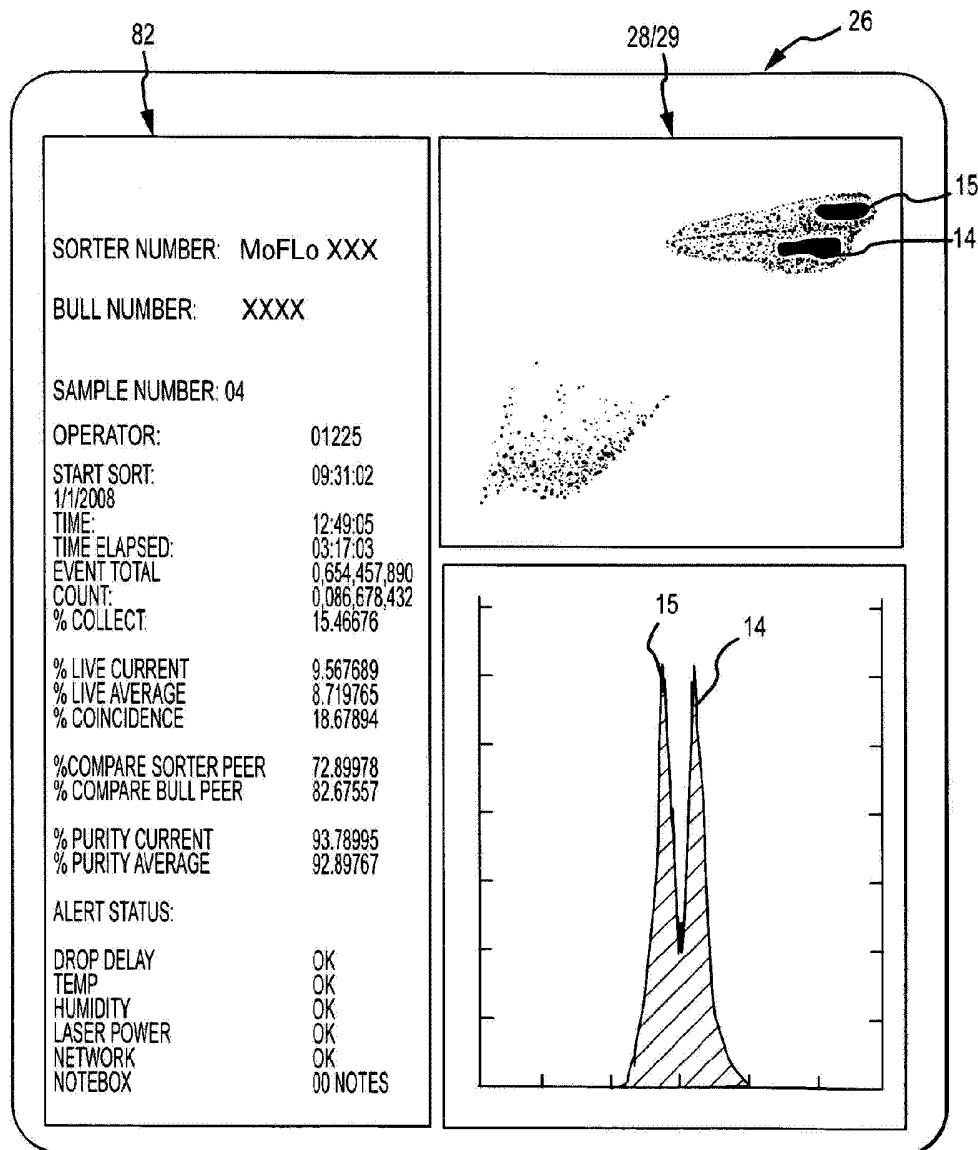


FIG.8

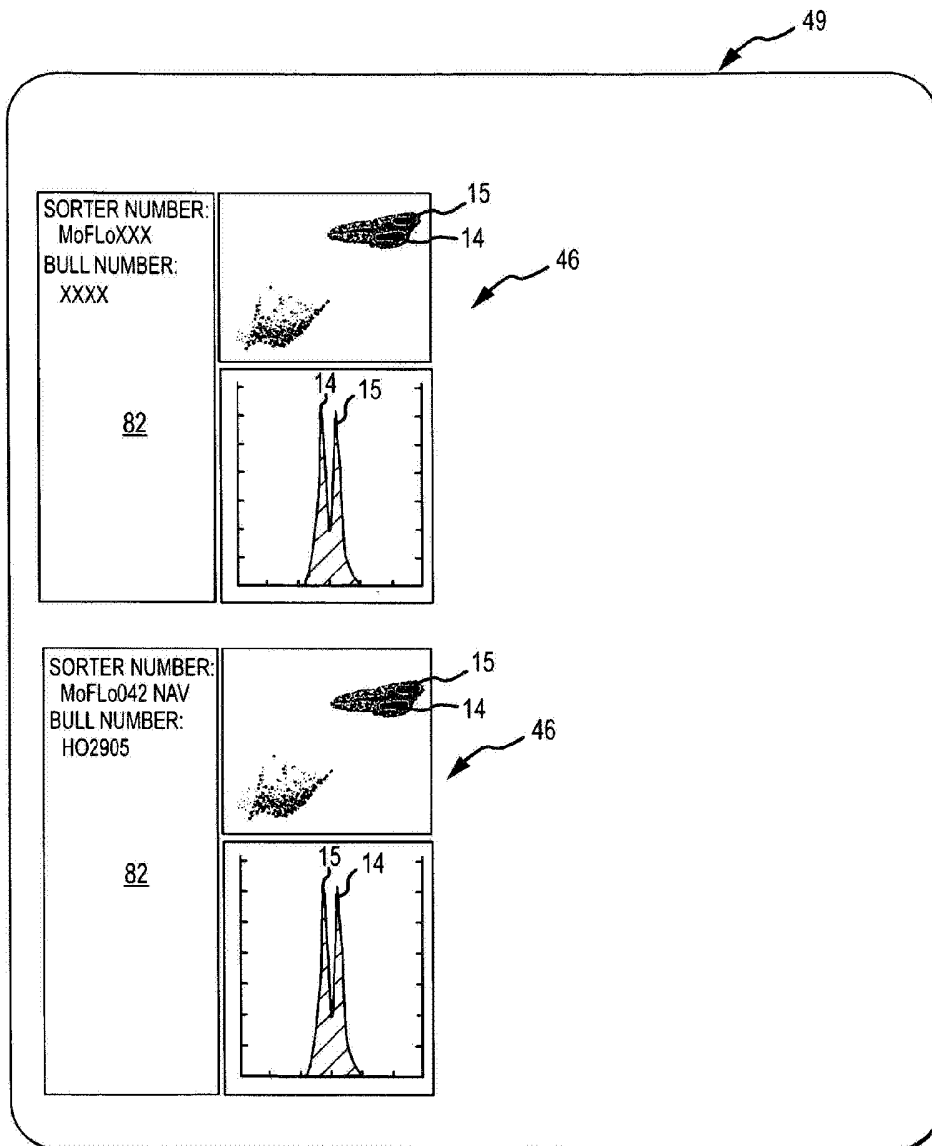


FIG.9