(54) PARTICLE FACILITATED TESTING

(71) Applicant: UNIVERSITY OF THE WEST OF ENGLAND, BRISTOL, Bristol (GB)

(72) Inventors: Richard William Luxton, Bristol (GB);
Janice Helen Kiely, Bristol (GB);
Patrick Wraith, Bristol (GB)

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(57) ABSTRACT

Magnetic particles are distributed across a fluid flow by applied magnetic field to interact with a test substance in fluid. Alternatively or additionally, particles, which may be magnetic, are combined with cells and energy, e.g. ultrasonic energy, is applied to cause the particles to create a lysate. Alternatively or additionally, the size of a quantity of magnetic particles is assessed by its impact on the tuning mechanism of a controlled oscillator that is affected by the particles.
FIG. 6

PARAMAGNETIC PARTICLES ADDED

FIG. 7

DIFFERENCE IN DETECTION SIGNAL VALUE

CONCENTRATION ng/ml
FIG. 8
FIG. 10
FIG. 12
PARTICLE FACILITATED TESTING FIELD

[0001] The invention relates to apparatus for, and methods of, testing fluid, and to apparatus for and methods of lysing cells using particles.

BACKGROUND

[0002] It is known to utilise magnetic particles to capture an analyte in a solution under test. Conventionally, the magnetic particles are coated with a substance to which the analyte will attach. A sensor surface in contact with the solution is provided with a similar coating and a magnetic field is applied to urge the magnetic particles onto the surface. Analyte bound to the magnetic particles then becomes attached to the sensor surface also. Thus, magnetic particles that have picked up analyte become immobilised on the sensor surface. An inductor located near to the sensor surface is used to quantify the number of magnetic particles that are so immobilised. The inductor forms part of a resonant electrical circuit. The resonant frequency of this circuit is determined in part by the inductance of this inductor and the inducance of the inductor is determined in part by the quantity of immobilised magnetic particles.

[0003] In the investigation of cell organelles and measurement of intracellular proteins, cells need to be disrupted or lysed, releasing the intracellular components for study. Freeze-thaw methods are commonly used to lyse both bacterial and mammalian cells. These methods involve freezing a cell suspension using a dry ice/ethanol bath or freezer and then thawing the material at room temperature or 37°C. This method of lysis causes cells to swell and ultimately break as ice crystals form during the freezing process and then contract during thawing. Multiple cycles are necessary for effective lysis, and the process can be time consuming. However, the freeze/thaw methods have been shown to release proteins located in the cytoplasm of bacteria effectively, and are recommended for the lysis of mammalian cells in some protocols.

[0004] Another approach commonly used to disrupt cells is to solubilise the cell membrane using a detergent. This has the added advantage of releasing membrane bound proteins but may dissociate protein complexes. Classically, physical methods have been used to disrupt cells, such as grinding tissue in a pestle and mortar or using a blade either as a scalpel or a liquidiser. There are some inherent disadvantages to mechanical lysis methods such as localized heating within a sample leading to protein denaturation and aggregation.

[0005] Ultrasound has also been used as a method of physical cell disruption which is based on the generation of high frequency pulses of pressure. Sonication (i.e. the process of disrupting the cell using sound waves) generates heat which may denature proteins, so the process should be performed in an ice bath. Some studies have shown that lysis using detergents to solubilise the cell membranes is more efficient at releasing intracellular protein than ultrasound.

BRIEF SUMMARY

[0006] According to an aspect of the invention, there is provided apparatus for lysing a cell, the apparatus comprising a chamber for holding the cell to be lysed and means for introducing energy into the chamber, wherein the chamber contains a plurality of particles which may be excited by the energy to enhance lysing of the cell.

[0007] The means for introducing energy into the chamber may comprise means for introducing sound waves into the chamber.

[0008] The means for introducing sound waves into the chamber may comprise a sonicator probe.

[0009] Additionally or alternatively, the means for introducing sound waves into the chamber may comprise an ultrasonic transducer.

[0010] The means for introducing energy into the chamber may be operable to introduce energy into the chamber in a pulsed manner.

[0011] The particles may be of a plastics material.

[0012] Alternatively, the particles may be of metal.

[0013] Alternatively, the particles may be of a combination of metal and a plastics material.

[0014] The plurality of particles may be provided with a binding agent to which components of a lysed cell may bind.

[0015] The chamber may comprise a sensor surface provided with a binding agent to which components of a lysed cell may bind.

[0016] A label may be provided to identify a complex formed when a component binds to the binding agent.

[0017] The label may comprise an enzyme.

[0018] The plurality of particles may be preferably in the range from approximately 0.1 μm to approximately 100 μm in diameter.

[0019] The plurality of particles are more preferably in the range from approximately 1 μm to approximately 20 μm in diameter.

[0020] In certain embodiments, the plurality of particles may be magnetic.

[0021] For example, the plurality of particles may be of a paramagnetic, ferromagnetic, diamagnetic or super-paramagnetic material.

[0022] The apparatus may further comprise sensing means for sensing the magnetic particles.

[0023] The apparatus may further comprise means for generating a magnetic field to draw the magnetic particles towards the sensing surface of the chamber.

[0024] The means for generating a magnetic field may comprise a permanent magnet.

[0025] Additionally or alternatively, the means for generating a magnetic field may comprise an electromagnet.

[0026] According to a further aspect of the invention, there is provided a method of lysing a cell, the method comprising introducing the cell into a chamber containing a plurality of particles and introducing energy into the chamber to excite the plurality of particles.

[0027] The energy introduced into the chamber may comprise sound waves.

[0028] The sound waves may be introduced into the chamber using a sonicator probe.

[0029] Additionally or alternatively the sound waves may be introduced into the chamber using an ultrasonic transducer.

[0030] The energy may be introduced into the chamber in a pulsed manner.

[0031] The particles may be of a plastics material.

[0032] Alternatively the particles may be of metal.

[0033] Alternatively, the particles may be of a combination of metal and a plastics material.
The plurality of particles may be provided with a binding agent to which components of a lysed cell may bind.

A label may be provided to identify a complex formed when a component binds to the binding agent.

The plurality of particles are preferably in the range from approximately 0.1 μm to approximately 100 μm in diameter.

The plurality of particles are more preferably in the range from approximately 1 μm to approximately 20 μm in diameter.

In certain embodiments, the plurality of particles may be magnetic.

For example, the plurality of particles may be of a paramagnetic, ferromagnetic, diamagnetic or super-paramagnetic material. Sensing means may be used to sense the magnetic particles.

A magnetic field may be generated to draw the magnetic particles towards the sensor surface of the chamber.

The magnetic field may be generated using means comprising a permanent magnet.

Additionally or alternatively, the magnetic field may be generated using means comprising an electromagnet.

**BRIEF DESCRIPTION OF THE DRAWINGS**

By way of example only, certain embodiments of the invention will now be described with reference to the accompanying drawings, in which:

FIG. 1 provides an overview of a fluid analysis system;

FIG. 2 shows in more detail the test unit of the system shown in FIG. 1;

FIG. 3 illustrates the distribution of magnetic particles within the measurement chamber of the test unit of FIG. 2 under certain magnetic field conditions;

FIG. 4 illustrates schematically the underside of the plate that is disposed at the bottom of the measurement chamber that is shown in FIG. 2;

FIG. 5 illustrates schematically the main elements of the measurement unit of the test unit that is shown in FIG. 2;

FIG. 6 illustrates the type of response that can be obtained from the measurement unit of FIG. 5 during an assay;

FIG. 7 illustrates a plot, for various test substance concentrations, of a metric that can be derived from an assay of the type shown in FIG. 6;

FIG. 8 illustrates a modified form of the test unit shown in FIG. 2;

FIG. 9 illustrates a modified form of fluid analysis system containing several test stations;

FIG. 10 illustrates a modified form of the test unit of FIG. 8 in which the measurement chamber is formed as a removable unit;

FIG. 11 illustrates schematically an alternative embodiment of a test unit;

FIG. 12 illustrates schematically the main elements of an alternative measurement unit which is used in the test unit that is shown in FIG. 11.

**DETAILED DESCRIPTION**

**Analyte Detection**

FIG. 1 shows an overview of a fluid analysis system 10. A pump 12 is provided with an inlet 14 for acquiring a sample of liquid that is to be examined by a test unit 16. After a sample of liquid has been acquired through inlet 14, the pump then operates to repeatedly circulate the liquid sample through the test unit 16 via tubes 18 and 20. The pump 12 pumps the liquid to the test unit 16 through tube 18 and the liquid returns from the test unit 16 to the pump 12 through tube 20. The test unit 16 is configured to detect the presence of a particular antigen in the liquid that is being pumped through the test unit 16. Henceforth, this antigen shall be referred to as the target antigen.

The test unit 16 is shown in more detail in FIG. 2. The main structure of the test unit 16 is provided by a block 22 of plastics material. A spherical measurement chamber 24 is formed in the centre of the block 22. Two bores 26 and 28 are formed in the block 22 to connect the measurement chamber 24 with the exterior of the block. The mouth that bore 28 presents to the exterior of the block 22 is connected to tube 18 and the mouth that bore 26 presents to the exterior of the block 22 is connected to tube 20. In this way, the pump 12 can pass the test liquid through the measurement chamber 24. The measurement chamber 24 is populated with particles of paramagnetic material, which are denoted in FIG. 2 by the black dots lying within the measurement chamber 24. The paramagnetic particles are treated with a coating of a particular antibody to which the target antigen will bind.

The measurement chamber 24 is preferably shaped so as to reduce the speed of the test liquid as it flows through the measurement chamber 24, to reduce disturbance to the paramagnetic particles, which will usually be manipulated to form specific configurations, as will be described below.

A square plate 31 is mounted in the bottom of the measurement chamber 24. The plate 31 has upper and lower major surfaces facing towards and away from the centre of.
the measurement chamber, respectively. The upper major surface of the plate 31 is covered with a coating 30 of the same antibody that has been applied to the paramagnetic particles. The lower major surface of the plate 31 is provided with an electrical coil which is connected to a measurement unit 32 by means of electrical connection 34.

Two cavities 36 and 38 are provided in the upper and lower surfaces of the block 22. A permanent magnet 40 is slidably mounted within cavity 36. A shaft 42 connects magnet 40 to a drive unit 44. The drive unit 44 is configured to act on the shaft 42 to vary the position of magnet 40 within cavity 36. That is to say, the drive unit can raise and lower the magnet 40 in the cavity 36 so as to vary the distance of the magnet 40 from the measurement chamber 24. Analogously, a permanent magnet 46 is slidably mounted in cavity 38 and can be moved by drive unit 48 by means of rod 50. The positions of the magnets 40 and 46 within the cavities 36 and 38 are governed by a control unit 52 that applies control signals to the drive units 44 and 48 through connections 54 and 56. Surface 58 constitutes the north pole of magnet 40 and surface 60 constitutes the south pole of magnet 46. The magnets 40 and 46 are closely fitted to their corresponding cavities 36 and 38 so that the pole faces 58 and 60 and the major surfaces of the plate 31 remain parallel with one another as the magnets are moved.

The positions of the magnets 40 and 46 relative to the centre of the measurement chamber 24 dictate the magnetic field that is experienced by the paramagnetic particles that are located within the measurement chamber. In order to promote the capture of any target antigen that is present within the test liquid that is flowing through the measurement chamber 24, the magnets 40 and 46 are positioned so as to generate within the measurement chamber 24 a magnetic field that causes the paramagnetic particles to distribute themselves across the measurement chamber in the manner of a sieve acting on the test liquid that is flowing through the measurement chamber 24. In this configuration, the paramagnetic particles form strands that extend across the flow within the measurement chamber 24 and generally attempt to extend between the pole faces 58 and 60, following the lines of magnetic force extending between the pole faces 58 and 60. These strands are illustrated schematically in FIG. 3 which shows the central portion of the block 22, focussing on the measurement chamber 24.

In FIG. 3, the strands are denoted by irregular vertical lines within the measurement chamber 24. Some strands, e.g. 62, may form extending from the upper surface of the measurement chamber. Other strands, e.g. 64, form extending from the base of the measurement chamber, which is effectively provided by the plate 31. Yet other strands, e.g. 66, may extend entirely between the base and upper surface of the measurement chamber 24. In this condition, the paramagnetic particles are distributed across the flow of the test liquid through the measurement chamber 24 which facilitates the paramagnetic particles’ capture, via their antibody coating, of target antigen in the test liquid. Accordingly, the state of the magnetic field required to place the paramagnetic particles in this condition shall be referred to as the “capture state”. The precise positions of the magnets 40 and 46 that are required to transform the magnetic field in the measurement chamber 24 into the capture state will depend upon various parameters of the precise design of the equipment and can be determined through experimentation. Examples of such parameters include the material and size of the paramagnetic particles, the material and size of the magnets 40 and 46 and the diameter of the measurement chamber 24.

The magnetic field within the measurement chamber 24 can also be adjusted to a so-called “collection state” in which the paramagnetic particles are drawn down to collect over the upper major surface of the plate 31. The collection state of the magnetic field can be achieved by moving the magnets 40 and 46 to their maximum and minimum distances, respectively, from the centre of the measurement chamber 24. When the magnetic field in the measurement chamber 24 is in the collection state, the paramagnetic particles are urged onto the antibody coating 30 on the plate 31. Some of the paramagnetic particles in contact with the coating 30 will have target antigen bound onto them. These particles can then become linked to the coating 30 by the target antigen that they carry and therefore become immobilised on the plate 31.

In order to examine the test liquid for the presence of the target antigen, the magnets 40 and 46 are moved to cycle the magnetic field in the measurement chamber 24 between the capture and collection states. When desired, the quantity of paramagnetic particles that have become attached to the antibody coating 30 on the plate 31 can be assessed electronically, as will now be explained.

**Sensing Arrangement**

FIG. 4 shows the lower major surface of the plate 31. An electrical coil 68 is provided on the lower major surface of the plate 31. A pair of supply conductors 70 extend from the coil and provide the connection 34 to the measurement unit 32. The coil 68 forms part of a voltage controlled oscillator (VCO), the remainder of which is housed within the measurement unit 32.

In FIG. 5, the circuitry that supplements coil 68 to form the VCO is indicated 72. From another perspective, coil 68 is simply an inductor that forms part of a VCO design and which has been located remote from the other components of the design. It should be noted, however, that the coil 68 and the measurement unit are, preferably, physically close to the measurement chamber 24. VCO designs that are suitable for adaptation in this manner are known to the skilled person. The VCO incorporates a variable capacitor 90, which is, for example, a variable capacitance diode. Together, the coil 68 and the capacitor 90 determine the frequency of the VCO’s output signal.

The measurement unit 32 also includes a crystal oscillator 74. The crystal oscillator 74 produces a 70 MHz output signal on line 76. The voltage controlled oscillator produces a signal on line 78 whose frequency the measurement unit 32 endeavours to maintain locked to a frequency of 70 KHz away from the output signal of the crystal oscillator 74. To achieve this end, output signals of the VCO and the crystal oscillator 74 are mixed together in a mixer 80 and resulting signal is provided on line 82 as one input to a phase detector 84. The other input to the phase detector 84 is provided over line 79 and is the output signal of a variable oscillator 81. The variable oscillator 81 and the phase detector 84 are integrated into a single package 85, which may also contain the other elements of the system of FIG. 5, with the exception of the crystal oscillator 74. The variable oscillator 81 is tuned so that the output signal that it provides on line 79 has a frequency of 70 KHz. The phase detector 84...
produces a DC voltage on line 86 that is proportional to the phase difference between its two input signals. This DC voltage is sensed on line 88 (and is referred to henceforth as a detection signal) and is also applied via line 92 to the variable capacitor 90 within the VCO. This voltage controls the capacitance of the variable capacitor 90, thereby tuning the frequency of the VCO’s output. It will be apparent to the skilled person that the elements shown in FIG. 5 are formed into a phase locked loop (PLL), or more accurately a frequency locked loop (FLL) for the purpose of locking the frequency of the output signal of the mixer 80 to 70 KHz, i.e. to the frequency of the output of the variable oscillator 81. This means that the PLL acts to maintain the output signal of the VCO at 69.93 MHz.

As mentioned earlier, coil 68 forms part of the voltage controlled oscillator that is the object of the PLL. The frequency of the output of the VCO that is supplied over line 78 is governed in part by the inductance of coil 68. In turn, the inductance of coil 68 is governed by the distribution of the paramagnetic particles within the measurement chamber 24 and in particular by the immobilisation of target antigen carrying paramagnetic particles on the coating 30. Accordingly, the voltage of the output of the phase detector 84 that is sensed on line 88 contains information about the behaviour of the paramagnetic particles and, in turn, about test antigen in the measurement chamber 24. In order to make deductions about test antigen in the measurement chamber 24, the voltage of the output of the phase detector 84 is recorded over time as the magnetic field within the measurement chamber 24 is varied. A typical assay will now be described.

Results

FIG. 6 shows a plot of a sandwich assay performed using apparatus according to the present embodiment of the invention. FIG. 6 plots the detection signal versus time. At the beginning of the measurement process, the measurement chamber 24 contains just a buffer solution and the detection signal value is A. Then, the magnets 40 and 46 are positioned so as to bring the magnetic field in the measurement chamber 24 into the collection state, which causes the detection signal value to change to B. Next, a quantity of liquid containing antibody coated paramagnetic particles mixed with target antigen is added to the buffer solution in the measurement chamber 24. This causes a marked drop in the detection signal value to C. At this point, the paramagnetic particles are clumped on the upper surface of the plate 31. The magnets 40 and 46 are then repositioned to change the magnetic field within the measurement chamber 24 to the capture state. This reduces the detection signal value to D. After 30 seconds, the magnets 40 and 46 are repositioned to change the magnetic field to the collection state, whereupon the detection signal value changes to E. After 30 seconds, the magnets 40 and 46 are repositioned to change the magnetic field back to the capture state such that the detection signal value changes to F. After 30 seconds, the magnets 40 and 46 are repositioned to change the magnetic field to the collection state, whereupon the detection signal value changes to G. After a further 30 seconds, the magnets 40 and 46 are drawn back as far as possible from the measurement chamber 24 and the detection signal value changes to H and the paramagnetic particles are allowed to relax on the upper surface of the plate 31 for 30 seconds. Then magnet 40 is driven to its point of closest approach to the chamber 24 whilst magnetic 46 is kept remote from the chamber 24. This causes the detection signal value to change to I. During this time, the proximity of magnet 40 causes any paramagnetic particles that are not bound to the coating on plate 31 to move away from the plate. Then, after 30 seconds, magnet 40 is retracted as far as possible from the sample chamber 24 such that the detection signal value changes to J.

In an alternative method, the paramagnetic particles are added to the buffer solution in the measurement chamber 24 before the magnets 40 and 46 are positioned so as to bring the magnetic field in the measurement chamber 24. In this method, the magnets 40 and 46 are positioned so as to cause the magnetic field to be in the capture state, causing the detection signal value to change to D immediately.

Various metrics can be derived from the time varying detection signal value shown in FIG. 6. For example, the following metrics could be used:

- J-H
- J-A
- (J-H)/A
- (J-H)/B
- (J-H)/(B-C)
- J-H for different known concentrations of target antigen. A curve 96, which has been fitted to the results, is shown. Such a curve can thereafter be employed to estimate the target antigen concentration in live test situations.

Further Analyte Detection Arrangements

Another embodiment of the invention is shown in FIG. 8. In this embodiment, the permanent magnets 40 and 46 have been replaced with electromagnets 98 and 100, the energisation of which is controlled by the control unit 52 in order to vary the magnetic field within the measurement chamber 24, e.g. to change the magnetic field from the collection state to the capture state.

FIG. 9 shows a further embodiment of the invention in which several sample chambers are used. Elements 104 to 110 each denote a test unit similar to test unit 16 of FIG. 1. However, in the embodiment of FIG. 9, the control and measurement functions associated with the plurality of sampling locations are collected into a single unit 112. This means, for example, that a single crystal oscillator can be used to provide a reference frequency for the VCOs associated with each of the sample chambers. Pump 102 takes in a volume of test liquid and circulates it through elements 104 to 110. It will be appreciated that elements 104 and 106 are placed in parallel whilst 108 and 110 are placed in series. The elements 104 to 110 need not all test for the same antigen. It is possible to utilise a temperature control system to keep multiple measurement chambers at the same temperature, should this be necessary given the types of test performed in those chambers (e.g. the temperature could be held the same within a group of chambers testing for the same antigen using the same antibody). A temperature control system can also be used to stabilize the temperature of those electronic components whose electrical properties or performance are temperature dependent (for example, components such as coil 68).
In the foregoing embodiments, a charge of the fluid under test is recirculated through the measurement chamber (or, as the case may be, chambers), and this is useful when attempting to detect a very low concentration of the target antigen. In other embodiments, however, it is possible to arrange that a given charge of test fluid is passed through a given measurement chamber just once. Additionally or alternatively, it is possible to hold a charge of test fluid with a given measurement chamber for a protracted period before perhaps processing another charge.

In the foregoing embodiments, antibody coated particles with attached antigen adhere to an antibody coating on a plate. Over time, it is possible that all of the magnetic particles will become adhered, or that no more particles can become adhered, resulting in the exhaustion of the measurement chamber. The measurement system can be configured to detect this condition (by monitoring the behaviour of the VCO that incorporates the coil that is associated with plate in question) and issue an appropriate indication to a user, who can take action to replenish the system. An embodiment in which replenishment is facilitated shall now be described.

FIG. 10 illustrates a variant 116 of the test unit of FIG. 8 in which the measurement chamber is formed as a removable unit 114. In FIG. 10, elements carried over from FIG. 8 retain the same reference numeral and shall not be described in detail again. Of course, the concept of rendering the measurement chamber replaceable is not limited to the particular type of test unit shown in FIG. 8 and could be applied to any type of test unit, within reason.

In test unit 116, there is a removable cell 114 in block 22. This cell contains the measurement chamber 24, and the paramagnetic particles and plate 31 within it, and also parts of connection 34, bore 26 and bore 28. The cell 114 and the block 22 are provided with appropriate electrical connectors at the interface between the cell and the block in order to complete connection 34 when the cell is installed in the block. Likewise, fluid-tight connectors are provided at that interface to complete bores 26 and 28 when the cell 112 is installed in the block 22. Thus, an incumbent cell 114 can be replaced at will, e.g. with a fresh cell of the same type (when it is desired to refresh an exhausted measurement chamber) or with a cell of a different type in which the paramagnetic particles and the plate 31 are coated differently (in order to switch to testing for a different antigen). During fabrication of such a cell, the paramagnetic particles and the plate are given coatings appropriate for the antigen that the cell is to detect. The paramagnetic particles can be doped into the measurement chamber of the cell with suitable stabilising agents to allow rapid dispersal of individual particles when they are rehydrated by test fluid entering the chamber. Examples of suitable stabilising agents include sucrose, trehalose, and other poly-ionic compounds.

FIG. 11 shows an alternative embodiment of a test unit of the system shown in FIG. 1. In this embodiment, the measurement chamber 120 is generally cylindrical, which helps to cause a controlled flow of fluid through the measurement chamber 120 and to reduce turbulence. The measurement chamber 120 is formed in the centre of a block 122 of a plastics material. Two bores 124, 126 are formed in the block 122 to connect the interior of the measurement chamber 120 with the exterior of the block 122, and the bores 124, 126 are tapered to assist in causing a controlled flow of fluid through the measurement chamber 126 and to reduce turbulence in the fluid. The mouth that bore 124 presents to the exterior of the block 122 is connected to tube 18 and the mouth that bore 126 presents to the exterior of the block is connected to tube 20. In this way, pump 12 can pass the test liquid through the measurement chamber 120. The measurement chamber 120 is populated with particles of paramagnetic material, which are denoted in FIG. 11 by the small circles lying within the measurement chamber 120. The paramagnetic particles are treated with a coating of a particular antibody to which the target antigen will bind.

A plate 128 is mounted in the bottom of the measurement chamber 120. The plate 128 has upper and lower major surfaces facing towards and away from the centre of the measurement chamber 120 respectively. The upper major surface of the plate 128 is covered with a coating 130 of the same antibody that has been applied to the paramagnetic particles. Disposed beneath the plate 128, externally of the measurement chamber 120, is an electrical coil 131 which is connected to a measurement unit 132 by means of an electrical connection 134.

Two cavities 136, 138 are provided in the upper and lower surfaces of the block 122. A permanent magnet 140 is slidably mounted within cavity 136. A shaft 138 connects permanent magnet 140 to a servo 144. The servo 144 is configured to act on the shaft 142 to vary the position of the magnet 140 within the cavity 136. That is to say, the servo 144 can raise and lower the magnet 140 in the cavity 136 so as to vary the distance of the magnet 140 from the measurement chamber 120. Analogously, a permanent magnet 146 is slidably mounted in cavity 138 and can be moved by a servo 148 by means of a shaft 150. The positions of the magnets 140, 146 within the cavities 136, 138 are governed by a control unit 152 that applies control signals to the servos 144, 148 through connections 154, 156. Surface 158 constitutes the north pole of the magnet 140 and surface 160 constitutes the south pole of the magnet 146. The magnets 140, 146 are closely fitted to their corresponding cavities 136, 138 so that the pole faces 158 and 160 and the major surfaces of the plate 128 remain parallel with one another as the magnets 140, 146 are moved.

As is the case for the test unit shown in FIG. 1, in the embodiment shown in FIG. 11, the positions of the magnets 140, 146 relative to the centre of the measurement chamber 120 dictate the magnetic field that is experienced by the paramagnetic particles within the measurement chamber 120. The magnets 140, 146 may be positioned so as to cause the paramagnetic particles to adopt a sieve-like configuration by forming into strands that extend across the flow within the measurement chamber 120.

The electrical coil 131 in this embodiment is positioned outside of the measurement chamber 120, but performs the same role as the electrical coil of the embodiment of FIG. 2 in detecting the number of paramagnetic particles that are bound to antibody coating 130 of the plate 128.

The system of FIG. 11 uses a detection unit, which is shown schematically in FIG. 12. The detection unit 160 is similar to the detection unit shown in FIG. 5, and thus like elements have the same reference numerals in FIG. 12. However, in the detection unit 160, the variable oscillator 81 is not present. Instead, the phase detector 162 and a phase shift unit 164 form a quadrature phase detector. In this arrangement the signal 82 is split into two components. One passes directly into one port of the phase detector 162 and the second is phase shifted by 90 degrees (at 70 KHz) and
tuned by an appropriate capacitor-inductance-resistor band-pass filter in phase shift unit 164, before passing into the second port of the phase detector 162.

[0103] In this arrangement, the phase shift of the phase shift unit 164 is frequency dependent. Therefore, if the signal deviates from 70 KHz then the phase shift will deviate from the basic value (i.e. the value of the phase shift at 70 KHz). For example, frequencies greater than 70 KHz could result in a phase shift greater than 90 degrees and frequencies less than 70 KHz could result in a phase shift of less than 90 degrees. The output signal 86 from the phase detector 162 is proportional to the phase difference between the two signal components and hence the level of deviation of the signal 82 from 70 KHz. The phase detector output signal 86 adjusts the variable capacitor to bring the frequency of the VCO back to a frequency of 70 KHz away from the output signal of the crystal oscillator 74.

[0104] It will be noted, from FIG. 6 for example, that when measurements are taken as the paramagnetic particles are drawn to the plate 31/128 by the action of the magnetic field generated by the magnets 40/46 and 140/146 that there is a step change in output voltage or frequency. This effect is caused by the proximity of the magnet 46/146, which is typically of Neodymium, to the electrical coil 68/131, which causes the inductance of the coil 68/131 to drop as the coil 68/131 approaches magnetic saturation. This has the effect of reducing the inductance of the coil 68/131, which tries to skew the frequency of signal 78.

[0105] The paramagnetic particles, which are typically of Magnetite or Ferrite, have the effect of increasing the permeability of the electrical coil 68/131, when in close proximity to the coil. This effectively increases the inductance of the electrical coil 68/131 and tends to try to lower the frequency of signal 78. Ferrite ceramics have the same effect on signal 78.

[0106] Thus, if the magnets 40/140 and 46/146 are given a tip made from Ferrite, or are coated with Ferrite, then the shift of the resonant frequency of the PLL/FLL circuit can be balanced out to a large extent. This results in good sensitivity to paramagnetic particles regardless of whether they are close to the electrical coil 68/131 or not.

[0107] Although the pole faces 58/160 and 601/60 are shown in FIGS. 2 and 11 as being flat, they may have a rounded profile to give an evenly distributed magnetic field at the flat surface of the plate 31/128, thus allowing an even layer of paramagnetic particles to form.

[0108] Although the example given above describes the use of the apparatus of the invention in performing sandwich assays, it will be appreciated by those skilled in the art that it can be used in performing other types of assays. For example, the apparatus could be used to perform a “displacement assay”, in which antigen coated paramagnetic particles are initially bound to the upper surface of the plate 31 and are displaced, on the introduction of a sample containing the target antigen into the measurement chamber 24, from the plate 31 due to competitive interaction between the target antigen and the antigen of the paramagnetic particles, resulting in a change in the detection signal.

[0109] Alternatively, the apparatus of the invention can be used to perform a “competitive assay”, in which a binding agent is attached to the upper surface of the plate 31. A first complementary binding agent, the target antigen, is introduced into the measurement chamber 24 with the sample, whilst a second complementary binding agent is attached to paramagnetic particles, and the first and second complementary binding agents compete to bind to the binding agent of the plate 31. The greater the concentration of the target antigen, the fewer paramagnetic particles will bind to the binding agent of the plate 31, and the detection signal will change accordingly.

Lysing Arrangement

[0110] Referring now to FIG. 13, an apparatus for cell lysis is shown generally at 200, and comprises a lysing chamber 202 for holding a liquid 204 containing cells 206 to be lysed. The lysing chamber 202 also contains a plurality of particles 208 to enhance lysis. One or more sonicator probes 210 are provided to introduce energy in the form of sound waves at ultrasound frequency into the chamber 202. Alternatively, one or more ultrasonic transducers may be integrated into the chamber 202 or positioned adjacent the chamber 122 to introduce the ultrasound energy into the chamber 202. Using particles of a suitable size and at a suitable density, ultrasound energy introduced into the chamber 202 by the sonicator probe 210 enables the particles 208 to acquire sufficient kinetic energy to lyse cells 206 mixed with the particles 208 in the chamber 202. The sonicator probe 210 may be activated in a continuous or pulsed fashion for a sufficient time to cause lysis of the cells 206 to occur.

[0111] Varying degrees of cell lysis can be achieved by adjusting one or more of the following parameters: the amount of ultrasound energy imparted, the type of particle 208 used, the concentration of the particles 208 or the size of the particles 208 used. The particles 208 should be of a size suitable to cause effective lysis. Preferably the particles 208 that are used to enhance the cell lysis are in the range of 0.1 μm-100 μm, or more preferably between 1μm-20 μm. The particles 208 should be used in a concentration range suitable to cause effective lysis of the amount of cells 206 in the chamber 202.

[0112] The particles 208 should be appropriately constructed and/or formed from material of appropriate density to cause cell lysis. For example, the particles 208 may be made from metal or a plastics material, or a combination of metal and a plastics material, or may be of any other suitable material.

[0113] By controlling the degree of cell lysis, various cell components, for example proteins and organelles, can be released from the cells 206. Alternatively, the cells 206 can be greatly disrupted to release enhanced levels of intracellular protein above and beyond that released using sonication alone.

[0114] Any type of cell, including mammalian cells, non-mammalian cells, plant cells, bacteria, yeasts and spores or a mixture thereof, may be disrupted using the apparatus and method described above with reference to FIG. 13.

[0115] The apparatus shown in FIG. 13 may be used to identify, quantify or separate a component of interest from lysed cells. In this application, the particles 208 are coated with a binding agent to which intracellular components may bind, so as to capture such intracellular components. The binding agent may be, for example, an antibody, a lectin, DNA, RNA, a receptor protein or any other binding agent or moiety. The intracellular component of interest may be, for example, a protein or a cell-organelle that binds specifically to the binding agent. The intracellular component that binds to the binding agent may be identified, quantified or sepa-
rated by using a label or reporter molecule which is associated with the intracellular component/binding agent complex formed during binding. For example, the label may be an enzyme which reacts with a suitable substrate to produce a coloured or fluorescent product. This reaction product may be used to identify, quantify or separate the intracellular component, as will be apparent to those skilled in the art.

[0116] FIG. 14 shows a modified version of the cell lysis apparatus of FIG. 12 in which means for magnetically detecting a target component of a cell, such as a protein or cell organelle, is provided. Elements common to this embodiment and the embodiment of FIG. 12 are denoted by like reference numerals. The lysing chamber 202 of this embodiment may form the measurement chamber of a fluid analysis system as described above with reference to FIGS. 1 to 11.

[0117] In this modified apparatus, the chamber 202 contains liquid 204 comprising a sample of cells 206 to be lysed, and a plurality of magnetic particles 220. The magnetic particles may be, for example, ferromagnetic, diamagnetic, paramagnetic or super-paramagnetic. The magnetic particles 220 are coated with a binding agent to which a target component, such as a protein or cell organelle, may bind. A sensor surface 222 is coated with a similar binding agent 224 to that used to coat the magnetic particles 220, such that the target component may bind to the binding agent on the sensor surface 222. A magnetic sensing means 226 is provided beneath the sensor surface. The magnetic sensing means 226 may be integrated into the lysis chamber 202, or may be positioned adjacent the lysis chamber 202. The magnetic sensing means 226 may be a magnetic coil or may be a resonant coil magnetometer, a magneto-resistive sensor, a micro-machined cantilever device or a superconducting quantum interference device, for example.

[0118] In use of the apparatus of FIG. 14, liquid containing the cells 206 to be lysed is placed in the chamber 202 with the magnetic particles 220 and a sonicator probe 210 is activated either continuously or in a pulsed manner for a time sufficient for lysis of the cells 206 to occur.

[0119] FIG. 15 shows the apparatus of FIG. 14 after lysis has occurred. Elements common to this Figure and FIGS. 13 and 14 are denoted by like reference numerals.

[0120] Lysing of the cells 206 by continuous or pulsed activation of the sonicator probe 210 produces lysed cells 230 and causes the target components such as proteins(s) and/or cell organelle(s) to bind to the binding agent that is used to coat the magnetic particles 220, to form a bound complex comprising the component (e.g. protein(s) or cell organelle(s)) of interest and magnetic particles, hereinafter referred to as “bound particles” 232.

[0121] FIG. 16 shows the apparatus of FIGS. 14 and 15 when an external magnetic force is used to manipulate the magnetic particles 220 after lysis has taken place. Again, common elements are denoted by like reference numerals. The external magnetic force may be provided by one or more permanent magnets, or by adjusting one or more electromagnets, for example. The magnets may be mounted externally of the chamber 202 or may be integrated into the chamber 202.

[0122] The externally applied magnetic force acts in the direction of the arrow 240 and is used to pull the bound particles 232 and the magnetic particles 220 towards the sensor surface 222, where the bound particles 232 bind to the binding agent 224 on the sensor surface 222. The bound particles 232 become cross-linked to the sensor surface 222, causing them to be immobilised on the sensor surface 222.

[0123] FIG. 17 shows the apparatus of FIG. 16 with the external magnetic force applied in the direction of the arrow 250 (i.e. the direction of the external magnetic field is reversed), and used to pull unbound magnetic particles 220 away from the sensor surface 222, thus leaving just the bound particles 232 attached to the binding agent 224 on the sensor surface 222, which allows the sensing means 226 to quantify the amount of bound particles 232 present. The amount of bound particles 232 detected by the sensing means 226 can then be used to determine the amount of the target components such as protein(s) and/or cell organelle(s) present.

[0124] Using the method and apparatus described above with reference to FIGS. 14 to 17, lysis of cells and identification, quantification or separation of intracellular components such as proteins and cell organelles to be performed in the same vessel.

[0125] Experiments carried out in relation to lysis of cells using particles will now be described.

Further Results

[0126] FIG. 18 plots the amount of total protein released after sonication of Jurkat cells in the lysis chamber when no particles are present, when particles of 2.8 μm diameter are present and when particles of 1 μm diameter are present.

[0127] FIG. 19 shows a plot of the dose response of a magneto-immunoassay to prostatic specific antigen (PSA) released from LNCAP cells by particle enhanced sonication.

[0128] FIG. 20 shows a Scanning Electron Microscope (SEM) image of Jurkat cells sonicated with no paramagnetic particles present.

[0129] FIG. 21 shows a SEM photo of Jurkat cells sonicated in the presence of 2.8 μm particles.

[0130] FIG. 22 shows a SEM photo of Jurkat cells sonicated in the presence of 1 μm particles.

[0131] In a specific example using the apparatus shown in FIG. 13, Jurkat cells were lysed using different sized particles in conjunction with ultrasound.

[0132] Jurkat cells were cultured in 75 mm² tissue culture flasks in sterile penicillin/streptomycin supplemented RPMI-1640 containing 10% newborn calf serum and L-glutamine and incubated in a humidified atmosphere at 37.5°C with CO₂. The cells were routinely passaged 1:4 (1 part cells: 4 parts growth medium) every 2 to 3 days. At 3 days post-passage, the cells were centrifuged for 5 minutes at 21°C at 1500 rpm. The cells were then re-suspended in 1 ml penicillin/streptomycin-supplemented RPMI (50 ml of PBS +5 ml L-Glutamine+5 ml Penicillin & Streptomycin to 500 ml of RPMI 1640). The cells were counted by the Trypan Blue exclusion method, in which a 20 μl sample of the cell suspension was diluted with 20 μl Trypan blue stain (0.2% w/v Trypan blue dissolved in PBS and stored at 4°C). The suspension was gently vortexed and 10 μl of the stained cells were counted using a haemocyтомeter.

[0133] To demonstrate the effect of particle size on the efficiency of cell lysis Jurkat cells were centrifuged at 200 g for 10 minutes and the supernatant was discarded. The pellets obtained were re-suspended in 1 ml phosphate buffer saline in the chamber, and 5 μl of 2.8 μm or 1 μm paramagnetic particles (Dynabeads) were added. The mixture was then treated with a sonicator probe for 1 minute. To prevent excessive heat generated by the probe, the sample was
immersed in an ice bath and the ultrasound was applied in multiple short bursts. The effect of the sonication with and without particles was quantified by measuring total protein released into the supernatant and the physical effect on the cells was studied using SEM (Scanning Electron Microscopy).

The addition of paramagnetic particles to the cells prior to sonication enhanced the amount of protein released from the cells in a given time. Moreover, paramagnetic particles of different sizes enhanced the protein released from the cells to different extents. Without paramagnetic particles the sonication process released 4 µg protein/10^6 cells, with the addition of 2.8 µm particles (Dynabeads), twice as much protein was released from the cells (8 µg/10^6) and 1.0 µm particles (Dynabeads) released approximately three and half times as much protein (14 µg/10^6), as is shown in FIG. 16. A significant difference was observed in the concentration of protein released by the sonication probe alone and combined with paramagnetic particles as shown by the total protein measurement (P<0.001).

Scanning Electron Microscopy (SEM) was used to evaluate the effect of sonication on cell morphology with and without the paramagnetic particles (see FIGS. 18, 19 and 20). Surprisingly, the different size of particles had a different effect on the cells. 2.8 µm particles appeared to cause coagulation of the intracellular proteins whereas 1.0 µm particles induced the formation of membranous like structures.

In a second example, the apparatus shown in FIGS. 14-17 was used to demonstrate the magnetic detection of intracellular prostatic specific antigen. In this case LNCAP cells were lysed in the presence of 1 µm paramagnetic particles which were previously coated with anti-PSA. The floor of the lysis chamber incorporated the magnetic sensor which had a second anti-PSA antibody immobilised on its surface. Following sonication in the presence of the paramagnetic particles, an external magnetic field was applied to all the paramagnetic particles down to the sensor surface. Particles which had PSA bound to the surface by the antibody interaction were cross-linked to the sensor surface by the binding of the second antibody immobilised on the sensor surface to the captured PSA molecule on the paramagnetic particle. The captured PSA acted as a biological bridge holding the particle on the surface through the immunological linkage. A second external magnetic field was applied to remove unbound paramagnetic particles prior to measurement. A resonant coil magnetometer, lying underneath the sensor surface, was used to detect the presence of paramagnetic particles attached to the sensor surface. FIG. 18 shows the dose response of a magento-immunoassay for PSA released from LNCAP cells by particle enhance lysis.

These examples demonstrate that paramagnetic particles used in magneto-biosensors can be used to enhance the release of intracellular proteins from the cells, as part of an integrated measuring system for the rapid measurement of intracellular proteins.

1. Apparatus for lyzing a cell, the apparatus comprising a chamber for holding the cell to be lysed and means for introducing energy into the chamber, wherein the chamber contains a plurality of particles which may be excited by the energy to enhance lysing of the cell.

2. Apparatus according to claim 1 wherein the means for introducing energy into the chamber comprises means for introducing sound waves into the chamber.

3. Apparatus according to claim 2 wherein the means for introducing sound waves into the chamber comprises a sonicator probe.

4. Apparatus according to claim 2 wherein the means for introducing sound waves into the chamber comprises an ultrasonic transducer.

5. Apparatus according to claim 1 wherein the means for introducing energy into the chamber is operable to introduce energy into the chamber in a pulsed manner.

6. Apparatus according to claim 1 wherein the particle are of a plastics material.

7. Apparatus according to claim 1 wherein the particles are of metal.

8. Apparatus according to claim 1 wherein the particles are of a combination of metal and a plastics material.

9. Apparatus according to claim 1 wherein the plurality of particles are provided with a binding agent to which components of a lysed cell may bind.

10. Apparatus according to claim 1 wherein the chamber comprises a sensor surface provided with a binding agent to which components of a lysed cell may bind.

11. Apparatus according to claim 9 wherein a label is provided to identify a complex formed when a component binds to the binding agent.

12. Apparatus according to claim 11 wherein the label comprises an enzyme.

13. Apparatus according to claim 1 wherein the plurality of particles are in the range from approximately 0.1 µm to approximately 100 µm in diameter.

14. Apparatus according to claim 13 wherein the plurality of particles are in the range from approximately 1 µm to approximately 20 µm in diameter.

15. Apparatus according to claim 1 wherein the plurality of particles are magnetic.

16. Apparatus according to claim 15 wherein the plurality of particles are of a paramagnetic material.

17. Apparatus according to claim 15 wherein the plurality of particles are of a ferromagnetic material.

18. Apparatus according to claim 15 wherein the plurality of particles are of a diamagnetic material.

19. Apparatus according to claim 15 wherein the plurality of particles are of a super-paramagnetic material.

20. Apparatus according to claim 15 further comprising sensing means for sensing the magnetic particles.

21. Apparatus according to claim 20 further comprising means for generating a magnetic field to draw the magnetic particles towards the sensing surface of the chamber.

22. Apparatus according to claim 21 wherein the means for generating a magnetic field comprises a permanent magnet.

23. Apparatus according to claim 21 wherein the means for generating a magnetic field comprises an electromagnet.

24. A method of lysing a cell, the method comprising introducing the cell into a chamber containing a plurality of particles and introducing energy into the chamber to excite the plurality of particles.

25. A method according to claim 24 wherein the energy introduced into the chamber comprises sound waves.

26. A method according to claim 25 wherein the sound waves are introduced into the chamber using a sonicator probe.

27. A method according to claim 25 wherein the sound waves are introduced into the chamber using an ultrasonic transducer.
28. A method according to claim 25 wherein the energy is introduced into the chamber in a pulsed manner.

29. A method according to claim 24 wherein the particles are of a plastics material.

30. A method according to claim 24 wherein the particles are of metal.

31. A method according to claim 24 wherein the particles are of a combination of metal and a plastics material.

32. A method according to claim 24 wherein the plurality of particles are provided with a binding agent to which components of a lysed cell may bind.

33. A method according to claim 24 wherein the chamber comprises a sensor surface provided with a binding agent to which components of a lysed cell may bind.

34. A method according to claim 33 wherein a label is provided to identify a complex formed when a component binds to the binding agent.

35. A method according to claim 34 wherein the label comprises an enzyme.

36. A method according to claim 24 wherein the plurality of particles are in the range from approximately 0.1 μm to approximately 100 μm in diameter.

37. A method according to claim 36 wherein the plurality of particles are in the range from approximately 1 μm to approximately 20 μm in diameter.

38. A method according to claim 24 wherein the plurality of particles are magnetic.

39. A method according to claim 38 wherein the plurality of particles are of a paramagnetic material.

40. A method according to claim 38 wherein the plurality of particles are of a ferromagnetic material.

41. A method according to claim 38 wherein the plurality of particles are of a diamagnetic material.

42. A method according to claim 38 wherein the plurality of particles are of a super-paramagnetic material.

43. A method according to claim 38 wherein sensing means are used to sense the magnetic particles.

44. A method according to claim 43 wherein a magnetic field is generated to draw the magnetic particles towards the sensor surface of the chamber.

45. A method according to claim 44 wherein the magnetic field is generated using means comprising a permanent magnet.

46. A method according to claim 44 wherein the magnetic field is generated using means comprising an electromagnet.

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