

**All-India Institute of Medical Sciences**  
**Ansari Nagar, New Delhi-29**  
(RESEARCH SECTION)

Ref. No. 29/Prop/Path./AKD/18-19/RS

Dated: 11.01.2019

**Subject: Purchase of Dynamic Light Scattering (DLS) for the Deptt. of Pathology, AIIMS, New Delhi-29 on proprietary basis-  
Inviting comments thereon.**

\*\*\*\*\*

The request has been received from **Dr.A.K.Dinda, Professor, Pathology, AIIMS** to purchase the subjected item from **M/s.Aimil Ltd. (Mgf. M/s.Malvern Pnalytical** on proprietary basis. The proposal submitted by **M/s.Aimil Ltd.** and Performa Invoice and Departmental PAC certifications are attached.

The above documents are being uploaded for open information to submit objections, comments, if any, from any manufacturer regarding proprietary nature of the equipment/item within issue of 15 days giving reference **No. 29/Prop/Path./AKD/18-19/RS**. The comments should be received by office of Stores Officer (RS), Research Section at AIIMS on or before **25/01/2019 upto 12:00 noon**, failing which it will be presumed that any other vendor is having no comment to offer and case will be decided on merits.

**STORES OFFICER (RS)**

**Encl: Related documents enclosed.**


**1. PAC Certificate enclosed.**

**2. Performa Invoice**

ALL INDIA INSTITUTE OF MEDICAL SCIENCE  
ANSARI NAGAR, NEW DELHI-110029

PROPRIETARY/SPECIFIC BRANT GOODS CERTIFICATE

- |  |  |
|--|--|
| 1. Item/Type/Model No. required  | Dynamic Light Scattering<br>Detector   |
| 2. Is the item a spare part attachment or accessory for existing equipment.  | NO   |
| 3. Name of the manufacturers/ supplier of the item proposed by the indenter.   | M/s Malvern Panalytical  |
| 4. Are they sole manufacturers/sole Distributors of the item   | YES  |
| 5. Is there any other item with similar/equivalent specification available in the market to meet the job requirement envisaged. If the answer is yes, why the same can't be procured. Demanding Officer should bring out Comparative functional advantages/cost effectiveness of the recommended item from those offered by other. | NO   |
| 6. What were the efforts made to locate alternative source of supply or use other substitutes.   | THE ITEM IS PROPRIETARY OF THE FIRM  |
| 7. Why open/limited tender can't be resorted to, for locating alternative source.  |  |
| 8. Are the proprietary items certifying that the rates are reasonable or not.  | YES, RATES ARE REASONABLE  |
| 9. Any other justification for procuring item from single source.  | THE FUNCTIONING OF THE EQUIPMENT FULFILLS THE CRITERIA OF PI'S WORK. NO OTHER MANUFACTURERS HAVE COMPARITIVE FULFILLMENTS. |

  
Signature of Indenter  
(Demanding Officer)  
Dr. A.R. DIXIT  
MD, PhD

I certify that the item 'at' Sr. No. 1 above is required to be procured on single tender basis as the source of supply is definitely known/the specified brand proposed was advantage in meeting our functional requirements and limited tender system could be dispensed with as they would serve no useful purpose in this particular case.

  
u/1

डा० ललित कुमार/Dr. LALIT KUMAR  
आचार्य एवं अध्यक्ष/Professor & Head  
विशेष अणुचिकित्सा विभाग/Dept. of Medical Oncology  
आ० वि०आ०एच०आई०एम०एच० (आ०आ०एच०)

  
Anand W. Barwad  
Professor  
Department of Chemistry  
All India Institute of Medical Sciences

**TECHNICAL SPECIFICATIONS FOR DYNAMIC LIGHT SCATTERING SYSTEM WITH ZETA POTENTIAL**

1. PC controlled system for measurement of Particle size and Zeta Potential of dispersed particles/molecules in solution.

Correlator: High sensitive correlator with minimum 1024 physical size channels or above.

The optics must be fully pre-aligned with no user adjustment required.

Measurement Type: Particle size and Zeta potential

**Particle Size:**

- a. Measurement Principle: Dynamic Light Scattering
- b. Size Range: 0.3nm to 10.0 microns (hydrodynamic dia.) or better
- c. Sample Conc. Range: 0.00001% to 40% (% volume) or better
- d. Accuracy: Better than +/- 2% of NIST traceable latex standards
- e. Precision/repeatability: Better than +/- 2% of NIST traceable latex standards
- f. Particle Volume: 12  $\mu$ L (minimum) or lower
- g. **Scattering Angle (Back):** Fixed 170° or better. Preferably system should take measure at variable positions within the cell, to minimize multiple scattering effect.
- h. System should have Adaptive Correlation : It should have the ability to automatically and dynamically classify correlation sub runs on the fly into steady state or transient events.
- i. System should have Optical Filter Wheel with Integrated Narrow Band Filter and DDLS polarisers - the narrow band filter should not be permanently installed, so it can be user selected to remove fluorescence when present on a sample by sample basis without suppressing signal when the sample does not fluoresce.

**Zeta Potential**

- a. Measured Principle: Electrophoretic light scattering and should be capable to measure Aqueous/Polar Solvents and Oils, High Salts/ Non-Polar solvents based systems. It should use the method of Phase Analysis Light Scattering (PALS) to improve the repeatability of measurements of low mobility samples
- b. Measured range: +/- 500mV
- c. Size Range for Zeta Potential: 5 nm to 100 microns or above.
- d. Minimum sample volume : 40  $\mu$ L (minimum) or lower
- e. Maximum sample conductivity : 260 mS/cm
- f. Disposable cells should be quoted to remove any cross contamination

**Light Source:** He-Ne / Diode laser 633nm with 4 mW power or lower

**Detector:** Avalanche Photo Diode for high sensitivity measurements.

**Temperature control range:** 2-100°C or better

**Sampling Cell:** Cuvette cell

**Measurement time:** 1-2 minutes or less

*[Signature]*  
Medical Superintendent  
A.I.I.M.S. Hospital  
New Delhi-110029

*[Signature]*  
Dr. A.K. DINDA  
MD, PhD  
Professor  
Dept. of Pathology

*[Signature]*  
Dr. Parthiv  
Professor  
Department of Biochemistry  
All India Institute of Medical Sciences  
New Delhi-110029

*[Signature]*  
Dr. Adarsh W. Barwad  
Assistant Professor  
Department of Pathology  
All India Institute of Medical Sciences  
New Delhi

*[Signature]*  
Dr. Lalit Kumar  
Assistant Professor & Head  
Department of Medical Oncology  
All India Institute of Medical Sciences  
New Delhi-110029

**Output:**

- a. For particle size: differential/cumulative distributions; values for sizes at given percentages, fits to distribution models
- b. For zeta potential: plot of zeta potential distribution, mean zeta value

**Software:**

- a. The software should be Windows XP/Window 7 or 10 based. It should provide result such as particle size and zeta potential measurement determination.
- b. The software should have in built diagnostic features like Size Quality Report and Zeta Quality Report with the ability to provide expert advice based on the raw data that has been accumulated. The raw data should be available later for analysis & use with other software modules.

**Light Weight & Portable Instrument**

**User friendly Software based on Microsoft Windows 7 or 10**

2. **Computer:** Windows @ 7 or 10,64bit OS, 4th generation i7 Processor, 8GB physical memory and 1TB hard drive & DVD DriveWide Screen Monitor & Software supports for the equipment
3. **12mm square disposable polystyrene cuvettes with stoppers: 100 numbers**
4. **Disposable Zeta Potential cells should be quoted to remove any cross contamination: 20 numbers**
5. **General:**
  - Power requirements: 240V, 50 HZ
  - Operating environment temperature: 20-35 degree C
  - Operating environment humidity – 40 to 90%
  - Should be free from any vibration
  - Vendor to quote for One year Warranty and additional AMC cost for the system
  - Prices need to be quoted in FOR AIIMS.

**6. ANY OBJECTIONS AGAINST THIS PROPRIETARY PURCHASE MAY KINDLY MEET PROF. A. K. DINDA ALONG WITH THE DEMONSTRATIONS AT ROOM NO. 3006, DEPT. OF PATHOLOGY, CONVERGENCE BLOCK, AIIMS WITHIN 15 DAYS OF PUBLICATIONS OF THE SPECIFICATIONS ON AIIMS WEBSITE**

Medical Superintendent  
AIIMS Hospital  
New Delhi  
Dr. A.K. DINDA  
The officer in-charge, Dept. of Medical Sciences  
AIIMS, New Delhi  
Dr. A.K. DINDA, MD, PhD  
Professor  
Dept. of Pathology  
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Dr. Adarsh W. Barwad  
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Dr. Parag Mehta  
Professor  
Department of Biochemistry  
AIIMS, New Delhi



Prof. Amit Kumar Dinda, MD, PhD  
Department of Pathology, All India Institute of Medical Sciences, New Delhi  
Secretary, Indian Society of Nanomedicine (ISNM)  
Ex-President, Indian Society of Renal & Transplant Pathology (ISRTP)  
Ex-Vice-President, Society for Tissue Engineering and Regenerative Medicine (India) (SABOI)  
Fellow, Electron Microscopy Society of India (EMSI)

26.12.2018

Store Officer,  
Research Section,  
AIIMS, New Delhi -110029

Sub: Justification for Purchase of DLS on Proprietary basis

Dear Sir,

In reference to the project no. I hereby request you to purchase the DLS on proprietary basis due to the following justifications as mentioned:

1. Non-Invasive Back Scatter (NIBS) - Patent No. EP884580, US6016195, JP 11051843
2. High and Low Frequency Electrophoresis (M3) - Patent No. EP 1154266, US7217350, JP04727064
3. Light Scattering Measurement using simultaneous detection - Patent No. EP 2235501, CN102066901, US20090251696

**JUSTIFICATIONS:**

**1. Non-Invasive Back Scatter (NIBS)**

Non-invasive back scatter is sensitive as the particle being measured is directly on the surface of the fiber and all the light scattered back in detection solid angle is taken up by the multimodal fiber placed in the suspension (invasive).

All Dynamic Light Scattering equipments use this standard Back Scatter Technique where use of a focusing lens allows both the laser and detector fiber to be placed outside the suspension and light scattered only at one particular angle and dimension is picked up by a single mode optical fiber and quantified.

Malvern Zetasizer PRO uses a patented technique of using multimode optical fiber and gradient index lens to pick up all the scattered photons at a particular back scatter conical angle (surface of a cone with backscatter angle vertex).

Malvern Zetasizer PRO can detect all the rays scattered at the designated conical angle (175o) while other competitor instruments can detect only one ray in the designated angle; hence Malvern Zetasizer PRO is more sensitive in photon detection and correlation estimation.

**2. High and Low Frequency Electrophoresis (M3)**

Different instruments use different techniques to overcome the effect of electrode fouling and polarization during zeta potential measurement in high ionic strength suspensions. Malvern's Zetasizer PRO uses their own patented M3PALS technique. Malvern Zetasizer PRO can discriminate upto three zeta peaks and also give the percentage of nanoparticles conforming to those zeta values (as required in our specifications) while other competitor instruments can only give the mean zeta potential and its standard deviation. It fails to identify subpopulations of nanoparticles with different zeta potentials within the same sample. Also by using the patented diffusion barrier technique, Malvern Zetasizer PRO can handle high ionic suspension without electrode fouling.

Room No. 1084, 1st Floor, Teaching Block, Department of Pathology  
All India Institute of Medical Sciences, New Delhi - 110 029



Prof. Amit Kumar Dinda, MD, PhD  
Department of Pathology, All India Institute of Medical Sciences, New Delhi  
Secretary, Indian Society of Nanomedicine (ISNM)  
Ex-President, Indian Society of Renal & Transplant Pathology (ISRTP)  
Ex-Vice-President, Society for Tissue Engineering and Regenerative Medicine (India) (SABOI)  
Fellow, Electron Microscopy Society of India (EMSI)

### 3. Light Scattering Measurement using simultaneous detection

Other competitors instruments can detect scattering at 3 angles (back, side & forward), it is NOT SIMULTANEOUS; i.e. detection does not occur at both the channels at the same time. While large particles (agglomerates) scatter more in the forward direction and smaller particles in backward direction, Malvern Zetasizer Pro can detect scattering in both directions simultaneously and whenever the forward scattering is more as in agglomerates, the back scatter reading at that instant is disregarded (real-time or post-processing gating) for calculation of photon correlation. By this method, Malvern Zetasizer PRO is superior to existing similar instruments by preventing the final size calculation being biased towards the higher side by the presence of contaminating agglomerates and also gives a numerical aggregation index to judge the quality of the preparation.

### 4. Malvern Zetasizer PRO has Optical Filter Wheel with Integrated Narrow Band Filter and DDLS polarisers:

This means the narrow band filter is not permanently installed, so it can be user selected to remove fluorescence when present on a sample by sample basis without suppressing signal when the sample does not fluoresce. The horizontal and vertical polarisers allow collection of particle size data in Backscatter and therefore perform DDLS measurements that may highlight such physical phenomena as rotational diffusion or asymmetry in samples. This is particularly useful in samples such as Gold/Quantum Dots without compromising the sensitivity for other samples.

Also, the said equipment is considered as standard for initial characterization of nanoparticles across the globe and has been cited in more than 1000s of publications.

Keeping above points in mind, I request you to kindly permit us to purchase this equipment under proprietary head.

Kindly do the needful at the earliest.

Thanking you,  
Sincerely,

Prof. A. K. Dinda

Dr. A.K. DINDA  
MD, PhD  
Professor  
Deptt. of Pathology  
All India Institute of Medical Sciences  
Ansari Nagar, New Delhi-110 029

Room No. 1084, 1st Floor, Teaching Block, Department of Pathology  
All India Institute of Medical Sciences, New Delhi - 110 029

To Whom it may concern,

Date of Issue: 20/11/2018

Re. Letter of Unicity/Proprietary Certificate

This letter certifies that the Zetasizer Ultra (ZSU5700)\* and Zetasizer Pro (ZSU5800) of Malvern Panalytical Limited UK, which uses the techniques of dynamic light scattering and electrophoretic light scattering to measure the size, mobility and zeta potential of particles in dispersions is, at the date of issue, UNIQUE for the following characteristics and/or performances:

- \*MADLS/B - Multi Angle Dynamic Light Scattering ability to perform multiple DLS measurements at multiple angles and solve the complex correlation function to produce a single DLS result independent of angle or concentration (assuming within measurable range). This gives high resolution i.e. the ability to resolve distributions up to 2:1 (standard DLS is not consistently able to resolve at 3:1). Results are reported by Intensity (at 173 degrees) and as angular-independent Volume distributions.
- \*MADLS Particle Concentration - ability to measure particle concentrations using a calibration free, ensemble approach, based on the light scattering intensity of individual size modes to provide a measure of particle concentration of the sample modes in particles/mL.
- High and Low Frequency Electrophoresis (M3-PALS) an apparatus able to determine the zeta potential distribution of a particle dispersion contained in a cell/cuvette free of the effects of electro-osmosis.
- Adaptive Correlation - the ability to automatically and dynamically classify correlation sub runs on the fly into steady state or transient events. This has multiple advantages - speeds measurement up two to three fold, gives more stable and repeatable results typically with less sample preparation, increases measurement precision five fold and increases the sensitivity to small amounts of large material by tracking transient events separately to those more representative of the sample. Adaptive Correlation requires no assumptions to be made by the user, works across all size ranges measurable and at all angles measurable with no data rejection.
- Option to utilize Laser Doppler Electrophoresis Using a Diffusion Barrier to minimise sample degradation, decrease volume requirements to 20µl and increase robustness of measurements in high conductivity media.
- \*Low volume Capillary Sizing Cell - reduces sample volume to 3 µl, but in a high quality disposable glass capillary cell, and increases upper measurable size limit to 10 µm (latex) without the need for density matching of the sample dispersant to particle density.
- Optical Filter-Wheel (Patent Pending) with Integrated Narrow Band Filter and DOLS polarisers - this means the narrow band filter is not permanently installed, so it can be user selected to remove fluorescence when present on a sample by sample basis without suppressing signal when the sample does not fluoresce. The horizontal and vertical polarisers allow collection of particle size data in

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Medical Superintendent

*A.K.D.*  
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Dr. Adarsh W. Borwad  
Assistant Professor  
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AllMS, New Delhi

Backscatter and therefore perform DOLS measurements that may highlight such physical phenomena as rotational diffusion or asymmetry in samples.

- Deep Learning AI - Gives real time size data quality analysis and advice on how to use the data or corrective action for the sample. Supports novice users with expert data analysis and advice.

Patents Granted:

High and Low Frequency Electrophoresis (M3-PALS)

- EP1154264
- US7217330
- JP04723044

Light Scattering Measurements using Simultaneous Detection

- EP2235501
- CN102544901
- JP2011523451
- US20090251696

Diffusion Barrier Method

- WO2012060272A1
- JP2013544003

Patents Pending:

MADLS  
Adaptive Correlation  
Low Volume Capillary Cell  
DOLS Automatic Filter Wheel

\*Ultra only

On behalf of Malvern Panalytical Ltd.

Darrell Bancars - Product Manager - Nanomaterials

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Medical Superintendent

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*Dr. Adarsh W. Borwad*  
Dr. Adarsh W. Borwad  
Assistant Professor  
Department of Pathology  
AllMS, New Delhi



US006016195A

United States Patent [19] Peters

[11] Patent Number: 6,016,195  
[45] Date of Patent: Jan. 18, 2000

[54] FIBER OPTIC DEVICE FOR DETECTING THE SCATTERED LIGHT OR FLORESCENT LIGHT FROM A SUSPENSION

[73] Inventor: Rainer Peters, Langen, Germany  
[73] Assignee: AIV-Laser-Vertriebsgesellschaft mbH, Langen, Germany

[21] Appl. No.: 09/094,777

[22] Filed: Jun. 15, 1998

[30] Foreign Application Priority Data  
Jun. 15, 1997 [DE] Germany 197 25 211

[51] Int. Cl.<sup>7</sup> G01N 21/00

[52] U.S. Cl. 356/342; 356/337; 356/73.1

[58] Field of Search 356/342; 337; 356/73.1

[56] References Cited

U.S. PATENT DOCUMENTS  
4,099,875 7/1978 McMahon 356/342  
5,141,512 8/1992 Thompson 356/218

OTHER PUBLICATIONS  
Wise, H. and Hoon, D., "Single-mode fibers in fiber-optic quasielastic light scattering: A study of the dynamics of concentrated latex dispersions", May 15, 1991, J. Chem. Phys. 94(10), American Institute of Physics, pp. 6429-6443.

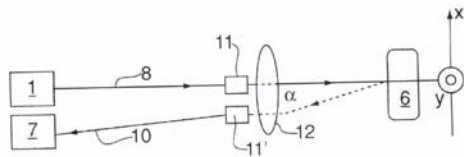
Amari, Rafat R. et al., "Microsolvation characterization by the use of a nonresonant backscatter fiber optic probe", J. Appl. Optics, vol. 32, No. 21, pp. 3822-3827, 1993.  
Rogers, Richard B. et al., "A Compact Laser Light Scattering Instrument for Microgravity Research", 1996, Photon Correlation & Scattering, 1996 Technical Digest Series, vol. 14, pp. 40-42.  
Ricks, Jaroslav, "Dynamic light scattering with single-mode and multimode receivers", May 20, 1993, Applied Optics, vol. 32, No. 15, pp. 2860-2875.

Primary Examiner—Robert Kim  
Assistant Examiner—Reginald Ratliff  
Attorney, Agent, or Firm—Kenyon & Kenyon

ABSTRACT

[57] The present invention relates to a fiber optic detector for detection of scattered light or fluorescent light from a liquid suspension, comprising: a lighting optical fiber for transmitting light to a lighting optical fiber outlet; a first gradient index lens disposed at the lighting optical fiber outlet to parallelize light; a detecting optical fiber for transmitting back-scattered light from a detecting optical fiber inlet; a second gradient index lens disposed at the detecting optical fiber inlet to parallelize back-scattered light; and at least one lighting optical fiber on a point and for focusing light back-scattered from the point to the second gradient index lens of the detecting optical fiber for transmission by the detecting optical fiber.

20 Claims, 3 Drawing Sheets



Dr. *[Signature]* NDA  
MD, PhD  
Professor  
Dept. of Pathology  
All India Institute of Medical Sciences  
Ansar Nagar, New Delhi-110 029

*[Signature]*  
1/18/98





(12) **United States Patent**  
**McNeil-Watson et al.**

(16) **Patent No.:** US 7,217,350 B2  
(45) **Date of Patent:** May 15, 2007

(54) **MOBILITY AND EFFECTS ARISING FROM SURFACE CHARGE**

4,101,220 A \* 7-1978 Bean et al. 204-549  
4,351,709 A \* 9-1982 Gostz 204-549  
5,245,290 A \* 9-1993 Cannon et al. 324-457

(75) **Inventors:** Fraser Keith McNeil-Watson, Worcesterhire (GB); Malcolm Trevor Connah, Worcesterhire (GB)

(73) **Assignee:** Malvern Instruments Limited, Worcesterhire (GB)

(\* ) **Notice:** Subject to any disclaimer, the term of this patent is extended or adjusted under 35 U.S.C. 154(b) by 1163 days.

(21) **Appl. No.:** 09/843,339

(22) **Filed:** Apr. 26, 2001

(65) **Prior Publication Data**  
US 2002/0040851 A1 Apr. 11, 2002

(30) **Foreign Application Priority Data**  
Apr. 29, 2000 (GB) 0010377.0

(51) **Int. Cl.**  
**C02F 1/469** (2006.01)  
**C07K 1/26** (2006.01)  
**B01D 57/02** (2006.01)  
**B01D 59/02** (2006.01)

(52) **U.S. Cl.** 204/549; 204/601; 204/603; 204/452; 204/456; 204/451  
(58) **Field of Classification Search** 204/549; 204/601; 603; 452; 455; 451  
See application file for complete search history.

(56) **References Cited**  
U.S. PATENT DOCUMENTS  
4,097,153 A \* 6-1978 DeKoenig 204-549

FOREIGN PATENT DOCUMENTS

WO WO8802482 A 4-1988  
WO WO 93/04363 3-1993  
WO WO93/04363 A 3-1993

\* cited by examiner

*Primary Examiner*—Ling-Sui Choi  
(74) *Attorney, Agent, or Firm*—Baker Botts LLP

(57) **ABSTRACT**

A method of, and apparatus for, automatically determining an electric charge-related characteristic or derived parameter of particles in a dispersion or of a cell wall, comprises having a particle-containing dispersion provided in a cell. The dispersion is illuminated with light from a light source and detecting from a detection volume light scattered from the particles. An electric field is applied to the dispersion at a first frequency of change of direction of the electric field and an electric field is subsequently applied to the dispersion at a second frequency of change of direction of the electric field. First signals detected from the scattered light when the first frequency electric field is applied are used to provide first values related to the velocity distribution of the particles. A second velocity related signal or value derived from scattered light during the time that the second frequency electric field is applied is used to modify the velocity distribution related values to produce a modified particle velocity-related distribution, and the first frequency is low enough to obtain an acceptable resolution of the distribution of particle velocity-related values.

9 Claims, 11 Drawing Sheets

*Handwritten signature*  
Dr. A.K. DINDA  
MD, PhD  
Professor  
Dept. of Pathology  
All India Institute of Medical Sciences  
Ansari Nagar, New Delhi-110 029



(12) **United States Patent**  
**McNeil-Watson et al.**

(39) Patent No.: **US 7,217,350 B2**  
(45) Date of Patent: **May 15, 2007**

(54) **MOBILITY AND EFFECTS ARISING FROM SURFACE CHARGE**

(75) Inventors: **Fraser Keith McNeil-Watson**, Worcesterstate (OH); **Malcolm Trevor Connors**, Worcesterstate (OH)

(73) Assignee: **Malvern Instruments Limited**, Worcesterstate (OH)

(\* ) Notice: Subject to any disclaimer, the term of this patent is extended or adjusted under 35 U.S.C. 154(b) by 1165 days.

(21) Appl. No.: **09/843,330**

(22) Filed: **Apr. 26, 2001**

(65) **Prior Publication Data**

US 2002/0040851 A1 Apr. 11, 2002

(30) **Foreign Application Priority Data**

Apr. 29, 2000 (GB) 0010377.0

(51) Int. Cl. (2006.01)  
**C82F 1/269** (2006.01)  
**C87K 1/26** (2006.01)  
**B10B 5/002** (2006.01)  
**B10D 36/02** (2006.01)

(52) U.S. Cl. 2004/49, 204/601, 204/603, 204/452, 204/456, 204/451

(58) **Field of Classification Search** 204/541, 204/601, 603, 452, 456, 451  
See application file for complete search history.

(56) **References Cited**  
U.S. PATENT DOCUMENTS  
4,097,353 A \* 6/1978 DeLuggis 204,540

4,081,220 A \* 7/1978 Bost et al. 204,540  
4,311,709 A \* 9/1982 Guder 204,540  
5,247,299 A \* 9/1993 Cannon et al. 224,457

**FOREIGN PATENT DOCUMENTS**  
963 WO/98/2402 A 4/1998  
963 WO/98/3851 3/1993  
963 WO/98/4381 A 3/1993

\* cited by examiner  
**Primary Examiner**—Langshui Chai  
(74) Attorney, Agent, or Firm—Baker Botts LLP

(57) **ABSTRACT**

A method of, and apparatus for, automatically determining an electric charge-related characteristic or derived parameter of particles in a dispersion in of a cell wall, comprising having a particle-containing dispersion provided in a cell. The dispersion is illuminated with light from a light source and detecting from a detection volume light scattered from the particles. An electric field is applied to the dispersion at a first frequency of charge of direction of the electric field and an electric field is subsequently applied to the dispersion at a second frequency of charge of direction of the electric field. First signals detected from the scattered light when the first frequency electric field is applied are used to provide first values related to the velocity distribution of the particles. A second velocity-related signal or value derived from scattered light during the time that the second frequency electric field is applied is used to modify the velocity distribution related values to produce a modified particle velocity-related distribution; and the first frequency is low enough to obtain an acceptable resolution of the distribution of particle velocity-related values.

**9 Claims, 11 Drawing Sheets**

*[Signature]*  
**Dr. A.K. DINDA**  
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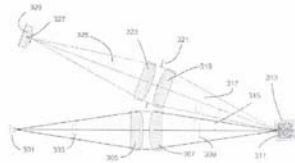


(12) United States Patent (10) Patent No.: US 9,435,726 B2  
Smart et al. (45) Date of Patent: Sep. 6, 2016

(54) DYNAMIC AND DEPOLARIZED DYNAMIC LIGHT SCATTERING COLLOID ANALYZER  
(71) Applicant: Scattering Solutions, Inc., Irvine, CA (US)  
(72) Inventors: Anthony E. Smart, Costa Mesa, CA (US); William V. Meyer, Lakewood, OH (US); Craig J. Saffell, Irvine, CA (US)  
(73) Assignee: Scattering Solutions, Inc., Irvine, CA (US)  
(\*) Notice: Subject to any disclaimer, the term of this patent is extended or adjusted under 35 U.S.C. 154(b) by 0 days.  
(21) Appl. No.: 14/797,146  
(22) Filed: May 8, 2015  
(65) Prior Publication Data  
US 2015/055688 A1 Dec. 10, 2015  
Related U.S. Application Data  
(63) Continuation of application No. 14/222,774, filed on Mar. 24, 2014, now Pat. No. 8,052,261, which is a continuation of application No. 12,961,079, filed on Aug. 23, 2009, now Pat. No. 8,717,562.  
(51) Int. Cl. G01N 15/02 (2006.01) G01N 15/10 (2006.01)  
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(54) Field of Classification Search CPC: G01N 15/0205; G01N 15/0211; G01N 15/1459; G01N 2013/0716; G01N 15/1434  
20 Claims, 14 Drawing Sheets

USPC Class: 356/336  
See application file for complete search history.  
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ABSTRACT  
Apparatus are described for measuring the characteristics of colloidal particles suspended in transparent media by Dynamic Light Scattering (DLS) and Depolarized Dynamic Light Scattering (DDLS) in regions where conventional measurements are difficult or impractical. Matching the diameter of an illuminating beam and an intersecting diameter of a field stop usage enables measurements in regions that include concentrated turbid suspensions that frequently appear non-linearly to those that multiple scattering typically gives a falsely low estimate of particle size. At the opposite extreme, where insufficient signal is available to determine either or both of the translational and/or rotational relaxation times of the particles, typically where they are too small, too few, or of insufficient refractive index difference from the medium to scatter enough light, measurements can be improved by: a) using a sufficiently large aperture such that many coherence areas fall upon the detector; and b) optical homodyne amplification of the scattered signal.



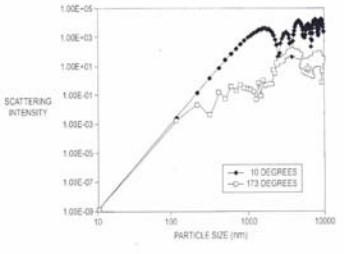
Optical Configuration for Small Matched FWHM  
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(54) LIGHT SCATTERING MEASUREMENTS USING SHELL ENSEMBLES OF PARTICLES  
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ABSTRACT  
Methods and apparatus for measuring particle characteristics are disclosed. In one aspect, an amount of light arising from interaction between light and a suspended sample is detected simultaneously with the acquisition of phase contrast from a different direction. At least one measure of particle characteristics can then be derived based at least in part on having both information from the steps of acquiring and detecting.



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