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(54) Title: SOFTWARE INTEGRATED FLOW CYTOMETRIC ASSAY FOR QUANTIFICATION OF THE HUMAN POLY-
MORPHONUCLEAR LEUKOCYTE FcγRI RECEPTOR (CD64)

(57) Abstract: The invention relates a method of quantifying CD64 and CD 163 expression in leukocytes and, specifically to a kit
for use with a flow cytometer including a suspension of quantitative fluorescent microbead standards, fluorescent labeled antibodies
directed to CD64 and CD 163, and analytical software. The software is used to take information on the microbead suspension and
fluorescent labeled antibodies from a flow cytometer and analyse data, smooth curves, calculate new parameters, provide quality
control measures and notify of expiration of the assay system.



WO 2006/055816 A2

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**SOFTWARE INTEGRATED FLOW CYTOMETRIC ASSAY FOR
QUANTIFICATION OF THE HUMAN POLYMORPHONUCLEAR
LEUKOCYTE FcγRI RECEPTOR (CD64)**

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Field of the Invention:

This invention relates to the field of cell surface antigens and quantitation of expression, in particular CD64 and CD163, which can be quantified using flow cytometric instrumentation, fluorescent labeled antibodies, microbead standards and integrated software.

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Background:

Sepsis is a significant health problem in the United States with an estimated nearly 750,000 new cases annually [1]. Sepsis is the 11th leading cause of death and the leading cause of mortality in the non-coronary intensive care unit with a mortality rate of 30-50% [1-3]. The frequency of sepsis has increased over 135% in a ten-year interval and is predicted to continue to rise with the increase in antibiotic resistance [1, 4]. In addition to the fully developed cases of sepsis, there are millions of patients on an annual basis in the U.S. alone that suffer severe infection or the clinical manifestations similar to sepsis that require diagnostic evaluation. Ironically, although treatment of sepsis has evolved in the last decades with newer therapeutic options of improved antibiotics and more novel approaches, such as inhibitors of the inflammatory response, little has changed to improve diagnosis and therapeutic monitoring [1-8]. Yet the medical detection and confirmation of patients suffering from infections or sepsis requires laboratory blood tests.

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The laboratory diagnosis of infection and sepsis still relies on the same basic modalities that have been available since the 1970s or before. Clinical decisions regarding the likelihood of infection or sepsis are typically made on leukocyte counts, the presence of myeloid immature forms in the peripheral blood (bands or “left-shift”), laboratory tests for “acute phase reactants”, such as C reactive protein, and

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clinical findings, such as fever. The cellular diagnostic parameters are typically problematic and lack specificity, particularly in infants [9, 10] and the elderly [11]. Furthermore, recent evidence indicates that leukocyte counts may have no significant influence on clinical decisions in patients with abdominal pain regarding surgery [12] and band counts provide little or no diagnostic information as to the probability of infection in a given patient [13, 14]. C reactive protein (CRP) levels, although not a new diagnostic test, have increasingly been advocated as a more objective diagnostic assay and therapeutic monitor, particularly in the neonatal and elderly population [15-19]. The advantages of CRP values, such as lack of interpretative subjectivity and improved sensitivity and specificity, are most evident in patient populations where studies have demonstrated a poor diagnostic utility of leukocyte counts and immaturity indices, such as in neonates, the elderly, and immuno-compromised patients. However, reliance on CRP values for sepsis or infection detection is also problematic in patients with co-morbid disease process. Being one of many acute phase reactants, CRP becomes elevated in any disease process with tissue injury, not just infection [16, 18, 20, 21]. Another limitation of CRP is a lack of specificity in distinguishing viral from bacterial infection. This lack of specificity as to the etiology of an infectious process has lead to the recent advocacy by some for the diagnostic utility of procalcitonin (PCT) measurements as a more specific indicator of bacterial infection [22-24]. Similarly there has been enthusiasm in the last ten years for potential diagnostic utility of cytokine levels, such as IL-6, and tumor necrosis factor (TNF) levels in septic patients [25-31]. But such studies generally have failed to demonstrate significant diagnostic utility for such tests and they additionally proved problematic due to the short half-life and sample stability of such inflammatory mediators. Molecular diagnostics have also made promises of newer approaches for early detection or prognosis, but still have fallen short due to technical limitation in turn around time, sensitivity, or cost [32, 33]. PCT, cytokines, TNF, and other molecular diagnostics currently remain experimental and none has been FDA cleared for *in vitro* diagnostic use to date.

The optimal diagnostic for sepsis and severe infection should be directed to early and specific changes related to the systemic acute inflammatory response. One of the major problems with the current laboratory tests for detecting the acute

inflammatory response related to infection or tissue damage is the lack of specificity [14].

Summary of the Invention:

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The invention relates a method of quantifying CD64 and CD163 expression in leukocytes and, specifically to a kit for use with a flow cytometer including a suspension of quantitative fluorescent microbead standards, fluorescent labeled antibodies directed to CD64 and CD163, and analytical software. The software is used to take information on the microbead suspension and fluorescent labeled antibodies from a flow cytometer and analyse data, smooth curves, calculate new parameters, provide quality control measures and indicate the expiration of the assay system.

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The integrated software-fluorescent microbeads standards system is a powerful tool for monitoring, optimizing fluorescence intensity measurements and increasing sensitivity. The integration of software, quantitative standards, and fluorescent-labeled antibodies allows for improved reproducibility in measurements of CD64 and CD163 on leukocytes between different production lots of the assay system. An entire calibration and analysis can be completed in less than five minutes, and a summary report generated on a printer. All results are automatically recorded in a cumulative file history to provide comprehensive documentation of an instrument's performance.

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The software is matched to specific lots of microbead standards and fluorescent-labelled mixtures of monoclonal antibodies, since all fluorescent molecules are subject to degradation and variability in conjugation efficiencies between production lots. Thus, for each batch of microbeads and fluorescent reagents, the associated software program includes information on the fluorescence intensity of each population of microbeads within the batch. When the kit approaches its expiration date, the software alerts the operator and further interrogates the users to verify the assay lot in use and the type of flow cytometric instrument employed as part of the specimen analysis. One of the novel features of this invention is that standardization between production lots is monitored and calibrated through the use

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of biologic specimens, specifically human blood samples with leukocytes of varying levels of CD64 and CD163 expression. The bias or difference in leukocyte CD64 and CD163 expression between different production lots is normalized by a factor determined by regression analysis between lots and changes in the fluorescent value assigned to the micorbeads within the software. These monitoring comparisons of different production lots of the assay kit done on different biologic samples will determine whether the microbeads are within the required range of fluorescence intensity to insure levels of imprecision between assay lots of $\leq 5\%$. Daily calibration of flow cytometers using the microbeads and software of the invention allows objective comprehensive monitoring of fluorescence intensity without complications due to reagents, cell preparation techniques, operating conditions, or stained cell samples. Once the calibration is done, the actual amount of fluorescent antibody bound by the specimen cells may be determined through the use of external standards, such as SRM 8640 produced by the National Institute of Standards and Technology (NIST). This measurement may be used to calculate the number of CD64 molecules (receptors) using fluorescent labeled antibodies or determine an index of antigen expression that is comparable to absolute molecular expression, which in turn, is useful in early diagnosis of infection .

In particular, the invention includes a kit for automated performance analysis to quantify CD64 and CD163 expression in a leukocyte and calibration for a flow cytometer, said kit used with a flow cytometer or a blood cell counter with fluorescence detection capabilities, blood specimens, a programmable computer and a display screen, said kit comprising:

- (a) at least one or more monoclonal antibodies directed to CD64 conjugated to fluorescein isothiocyanate (FITC) or similar fluorochrome;
- (b) at least one or more monoclonal antibodies directed to CD163 conjugated to phycoerythrin or similar fluorochrome;
- (c) a suspension of microbeads having reliable fluorescence stability over a finite period of time;
- (d) at least one or more fluorochromes, said fluorochrome having a useful life, said useful life being designated in said kit; and

(e) software operating on said computer comprising 1) information on the fluorescence intensity of each fluorescent reagent and microbeads, said software operating with said computer to control calculation of calibration and fluorescence information tailored to said microbeads, and display on said screen an expiration date correlated to the useful life of said microbeads to an operator calibrating said flow cytometer with said microbeads, wherein said software ceases operation on said computer if the operator does not accurately acknowledge the instrument model and correct assay kit with an expiration time, whereupon said flow cytometer cannot be calibrated with said software and said microbeads and the operator must utilize another suspension of microbead populations and software tailored for use on said computer with said another suspension to calibrate the flow cytometer; 2) the ability to acquire data on blood specimens using fluorescence, forward angle light scatter ("FS") and side or wide angle light scatter ("SS") and thus automatically gate or identify specific cell subpopulations, smooth the data, locate peaks in fluorescence intensity corresponding to each microbead and cell specific populations, construct linear regression plots, determine the fluorescence threshold intensity of the flow cytometer, and create histograms or graphic displays; 3) the ability to find cell specific clusters using iterative cluster finding algorithms; 4) the ability to perform automated data normalization; and 5) the ability to calculate the CD64 and CD163 indices.

Thus, the CD64 assay of the present invention represents a significant improvement in the diagnosis, therapeutic monitoring, and management of individuals of all ages with serious infections. Another equally significant societal benefit of the CD64 assay is that by providing a more sensitive and specific assay of the human sepsis and infection, there should also be a secondary effect of reducing the spread of antibiotic resistance of bacteria, a growing worldwide health concern.

Other and further aspects, features and advantages of the present teachings will be apparent from the following description of the various embodiments of the present teachings given for the purpose of disclosure.

Brief Description of the Figures:

Figure 1 is a diagrammatic representation of PMN CD64 expression in patients with culture proven infection (diamonds) are elevated in over 97.5% of individuals compared to the normal low value seen with healthy controls (circles); PMN CD64 values do not correlate with the neutrophil leukocyte count and are increased in those patients with infection, but normal neutrophil count.

Figure 2 represents flow cytometric analysis of human blood sample with normal (top panel) and elevated (bottom panel) neutrophil CD64 quantified using bead calibrators to derive PMN CD64 Index units; the other blood leukocyte populations of lymphocytes, monocytes, as well as neutrophils (shown in gated region in figure) are identified through software data analysis using iterative cluster finding algorithms and both anti-CD64 FITC and anti-CD163 PE staining of the samples.

Figure 3 demonstrates the software analysis of samples where the defined internal cellular controls to not “pass” or demonstrate the expected result of lymphocytes lacking CD64 expression (top panel), which is typically due to improper instrument set-up or addition of incorrect reagents, and monocytes lacking significant expression of CD64 (bottom panel), which is usually the result of insufficient anti-CD64 reagent added to the blood sample. The automated flags or popup boxes alert the user to problems with the technical performance of the assay.

Figure 4 illustrates the excellent correlation of the Trillium PMN CD64 index to the bead calibrated method of measuring PMN CD64 expression using MESF units is seen in two separate clinical studies.

Figure 5 illustrates that the level of PMN CD64 is correlated with the clinical impression in patients seen in a hospital emergency room (lower left), suggesting a role in the CD64 assay in screening for a systemic inflammatory reaction and triaging patients to the appropriate medical management setting and therapy; the same degree of correlation between clinical score (defined in lower right) and neutrophil count or CRP levels was not observed in the same study (upper left and right).

Figure 6 illustrates the that interlot correlation is high, but that there is a difference in the PMN CD64 index due to differences in the microbead fluorescence and antibody fluorochrome conjugation efficiency (upper left panel). However the invention of using the Leuko64 software to assign microbead fluorescence values based upon interlot comparison on multiple biologic blood samples allows for good interlot correlation as well as a low interlot imprecision (upper right panel). The invention also allows for the same improvement of interlot imprecision for the monocyte CD163 index with the software (lower panels).

Detailed Description of the Invention:

The inventors of the present invention and others have reported clinical studies indicating that PMN CD64 expression represents a specific diagnostic indicator of an active acute inflammatory response [34-43]. Unlike many other activation related antigenic changes on neutrophils (PMN) and other leukocytes, such as CD11b, CD18, CD16, CD45RA, and CD62L, neutrophil CD64 has negligible expression in the health individual and shows no evidence of sensitivity to blood sample manipulation [35, 37]. In contrast to many other myeloid antigens, such as CD16 or CD11b, PMN CD64 expression in both healthy individuals and those with upregulated CD64 expression is not altered by cell activation by chemotactic factors, PMA, or *in vitro* exposure to bacteria [35, 37, 38, 44, 45]. Additionally the expression of PMN CD64 is stable in anti-coagulated blood samples for up to 48 hours [35, 45]. Additionally previous studies by Trillium have shown normal levels of CD64 during otherwise healthy pregnant women, in contrast to other PMN and monocyte antigens, such as CD11b, CD14, CD32, and CD62L, which have been reported to show a significant change in women during pregnancy [46, 47]. Thus indicating the PMN CD64 measurements are practical as a diagnostic test, allowing performance in any laboratory setting for both outpatients and inpatients. PMN CD64 up-regulation by inflammatory related cytokines, such as G-CSF and interferon gamma (IFN- γ) occurs in a the short time scale of 4 to 6 hours for cell surface expression and 1 to 3 hours for detectable mRNA increases by Northern blot analysis [35, 40, 48-50]. This upregulation of PMN CD64 appears to directly correlate with enhanced bactericidal [40] and anti-fungal [51] activity. Thus, given that normal neutrophil expression of

CD64 is less than 1000 sites per cell and that detectable up-regulation can be found within 4 to 6 hours after an acute tissue injury, dissemination of infection, or administration of recombinant interferon gamma or G-CSF, early diagnosis is possible. Therefore, PMN CD64 quantitation represents an ideal candidate as an improved diagnostic compared to leukocyte counts, myeloid immaturity assessment, and acute phase reactants for the reasons listed in Table 1 and data in Figure 1. In summary, quantitative PMN CD64 measurements offer the potential to be the first, clinically useful diagnostic cell-based parameter of a systemic acute inflammatory response.

TABLE 1. Rationale for clinical utility of PMN CD64 assay.

- PMN CD64 expression is negligible in the healthy state and increases under the influence of inflammatory related cytokines (IFN- γ , IL-12, G-CSF)
- Increased PMN CD64 expression results in enhanced antibody-mediated functional responses (phagocytosis, oxidative burst, bactericidal activity) in PMNs – pathophysiologically significant change
- Measurements stable in routinely collected and handled anti-coagulated blood samples for 30 hours at room temperature and 48 hours refrigerated
- Assay performed within one hour, requires small volume, batch analysis possible
- PMN CD64 increase highly correlated with sepsis, infection or systemic acute inflammation
- Level of PMN CD64 barometer of systemic inflammatory response – potential for therapeutic monitor role in antibiotic use
- CD64 up-regulation not surrogate for absolute neutrophil count, band count, CBC instrument flagging, Sedimentation Rate, C reactive protein
- Patients of all ages and immuno-suppressed conditions capable of CD64 up-regulation
- High specificity - PMN CD64 expression is not elevated in:
 - Malignancy of myeloid cells (CML, MPD, MDS)
 - Any drug therapy (other than cytokines)
 - Clinical conditions with localized tissue damage (myocardial ischemia, uncomplicated surgery, and exercise injury)
 - Pregnancy

Flow cytometric quantitative techniques have been the methodology employed in the majority of studies looking at CD64 expression in myeloid cells. This instrumentation, although considered a high complexity form of clinical testing, is currently available in university medical centers, large tertiary care community hospitals, and reference diagnostic laboratories in the U.S. and most industrialized

countries. The methodology most refined with regard to clinical applications has been developed by the inventors of the present invention, and have validated a whole blood lysis technique using directly conjugated anti-CD64 and standardized fluorescence bead calibrators [35, 37, 40, 52, 53]. The current methodology requires 5 50 μ l or less of whole blood, thus being amenable to testing of patients of all ages. The assay precision has been validated on replicates as having a coefficient of variation of less than 5% [35], thus being quite competitive in precision with other clinical laboratory assays (Figures 2, 3).

10 Of equal practical importance for a clinical diagnostic test, the quantitative PMN CD64 results are stable at least 30 hours at room temperature and up to 72 hours for anticoagulated blood held at refrigerated temperatures. The results can quantified in terms of mean equivalent soluble fluorescence units (MESF) derived from the difference between binding of CD64 monoclonal antibody and isotype control
15 antibody run under parallel conditions. The day to day and assay to assay results are calibrated through the use of Quantum 26 FITC beads (formerly Flow Cytometry Standards, San Juan, PR, now Bang Labs, Fishers, IN) and computer data analysis with Leuk64 software (Verity Software House, Topsham, Maine). Alternatively with accurate fluorochrome to protein (F/P) ratio determinations of the monoclonal
20 antibody reagent, the MESF units can be converted to actual average antibody binding or antigen (if the monoclonal reagent is monomeric) sites per cell. The problem with these approaches is that, although the flow cytometric assays for CD64 using beads for quantitation are sensitive and relatively precise, the cost and technical time and complexity required for a PMN CD64 clinical assay is relatively high (>\$50 per test)
25 and are subject to measurement imprecision between lots of beads and antibodies. The present invention is a novel method of quantifying PMN CD64 expression, which appears to be equivalent to previous flow cytometric bead assays with less cost to the assay and marked reduction in lot to lot imprecision. The novel method for CD64 quantitation, termed the PMN CD64 Index, is based upon a ratio of the anti-CD64
30 antibody binding on neutrophils to lymphocytes, a leukocyte cell population lacking CD64 expression, or to a microbead suspension. Progress to date indicates the PMN CD64 Index is substantially equivalent to the more complex CD64 quantitation using bead calibrators (Figure 4).

Studies with quantitative PMN CD64 measurements to date have indicated a high degree of specificity of greater than >98% for detection of systemic infection [35, 38, 41, 42, 54]. Interestingly along with the sensitivity for detecting the acute inflammatory response in conjunction with infection there is only a weak correlation with neutrophilia, sedimentation rate, and/or the presence of bands [35, 37], suggesting additional diagnostic information is afforded by CD64 measurements compared to standard laboratory testing modalities. Given there are multiple clinical explanations for elevated neutrophil counts, the potential for PMN CD64 expression to provide increased diagnostic sensitivity and specificity of infection, along with its ability to be performed in a clinical setting in a matter of hours, it should find clinical utility. Although clinical blood samples show a weak to moderate tendency for increased PMN CD64 expression in parallel with increasing neutrophil counts or band counts [35], clinical samples are consistently identified where PMN CD64 expression and leukocyte enumeration are discordant. Clinical samples with elevated absolute neutrophil counts, yet exhibiting normal CD64 expression, are commonly seen in three clinical situations. The situations being glucocorticoid or steroid therapy, the early post-surgical period, and chronic myeloproliferative disorders, in which treating physicians have learned to expect elevated PMN counts without the necessity for further therapeutic action or concern for infection. Observations to date of over 100 cases of chronic myeloid leukemia, chronic myeloproliferative disorders, and myelodysplastic syndromes have shown no increased CD64 expression on mature myeloid cells, except in such patients with documented evidence of infection or other recognizable or acute inflammatory process [45, 55, 56]. Thus, one additional potential clinical application of PMN CD64 expression could also be used to distinguish acute inflammatory responses from hematological malignancies resulting in neutrophilia. The corollary situation, where elevated PMN CD64 is observed in parallel with normal or decreased neutrophil counts is observed most commonly in clinical situations where patients have received myelosuppressive drugs, such as cancer chemotherapy, and yet have clinical symptoms or laboratory evidence of infection. Additional studies are required to validate the diagnostic utility of quantitative PMN CD64 measurements in specific clinical situations and to obtain FDA clearance as an *in vitro* diagnostic test. However, previous published work does indicate the level of PMN CD64 expression to correlate with both clinical probability

of infection [35, 37-39, 41-43] and the type of infection (gram negative bacteria inducing higher PMN CD64 expression compared to gram positive bacteria) [42].

Additionally, as shown in Figure 5 the magnitude of PMN CD64 expression appears lower in localized compared to systemic infections [37, 42]. In summary, the present invention relating to PMN CD64 diagnostic tests: 1) function in a triage role for patients having clinical symptoms suggestive of infection; 2) provide quantitative assessment of a systemic inflammatory response; 3) distinguish between inflammatory and leukemia-related causes for elevated leukocyte counts; 4) serve as a therapeutic monitor in patients with sepsis or infection.

Components Used in the Present invention Assay:

The following is a list of components and reagents that are used in the assay for inflammation and detection of tissue injury by quantifying CD64 expression in polymorphonuclear leukocytes:

(A) at least one or more monoclonal antibodies directed to CD64 conjugated to fluorescein isothiocyanate (FITC) or similar fluorochromes;

(B) at least one or more monoclonal antibodies directed to CD163 conjugated to phycoerythrin or similar fluorochromes;

(C) a suspension of polystyrene microbeads with at least one or more fluorochromes, one of which is identical to that conjugated to the CD64 monoclonal antibodies, environmentally sensitive to the same factors affecting the fluorescence of antibodies bound to cells, and traceable to an external standard, such as the NIST SRM 8640.

(D) the Leuko64 software, which allows for fluorescence value assignment to the microbeads and calibration of the quantitation of measurements of CD64 and CD163 expression and cellular cluster gating independent of user interaction.

Description of the Leuko64 Software:

Although the current assay for PMN CD64 can be performed using third party software capable of analyzing flow cytometric data, this approach leaves some degree of subjectivity in data analysis to end users, who typically have a range of skills.

Thus, there remains the potential for errors or subjectivity in the assay result. Hence a preferable approach to minimize the user's subjectivity in data analysis and to streamline the assay is to include software for flow cytometric data with the CD64 assay. The Leuko64 CD64 assay kit includes a software that will indicate the outdate of the kit in parallel that of the assay reagent. The software has the ability to find cell clusters (lymphocytes, monocyte, and neutrophils) and perform automated data normalization and calculation of the PMN CD64 index. Additionally, the program can perform some specified analysis to serve as quality control functions, for example to validate that the monocyte population has the expected CD64 expression detected by the fluorescent antibody reagents, thus serving as an internal positive control for each specimen, and to validate that the lymphocyte population has no significant CD64 expression, thereby serving as an internal negative control for each specimen. Most importantly, the program allows for rapid data analysis from a variety of commercial clinical flow cytometry instruments, provide quality control related "flags" to the operator, and improve the standardization of data analysis to further minimize assay imprecision. A preferred embodiment of the invention is the integration of the software and bead fluorescence value assignment for CD64 and CD163 index values, which can be traceable to reference standards of fluorescence for specific fluorochromes. This value assignment of beads using a regression correlation between assay lots on biologic cell samples, such as blood leukocytes, provides for a low degree of imprecision between assay lots, thereby improving diagnostic utility in clinical practice.

Exemplification:

The examples herein are given for the purpose of illustrating various embodiments of the invention and are not meant to limit the present invention in any fashion. All documents mentioned herein are incorporated by reference in their entirety.

One skilled in the art will readily appreciate that the present invention is well adapted to carry out the objects and obtain the ends and advantages mentioned, as well as those inherent therein. It will be apparent to those skilled in the art that various modifications and variations can be made in practicing the present invention without

departing from the spirit or scope of the invention. Changes therein and other uses will occur to those skilled in the art which are encompassed within the spirit of the invention as defined by the scope of the claims.

5 Example 1: Leukocyte CD64 Assay:

The method for of flow cytometric analysis of blood samples for PMN CD64 is outlined briefly, and provided in detail by Davis et al [34-37, 40, 52, 53, 58, 59]. The current procedure for flow cytometric CD64 staining and analysis is as follows:

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Trillium Diagnostics Leukocyte CD64 Assay Staining Procedure:

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1 Dilute the 10X Trillium Lyse (Reagent B) 1:10 by mixing 1 part of the concentrated Reagent B with 9 parts filtered distilled water. Make a volume sufficient for anticipated number of tests (1.0mL is required for each sample). The final pH of the diluted Reagent B should be 7.40 ± 0.02 ; adjusted with NaOH or HCl if necessary. Diluted or 1X Trillium Lyse is stable for 1 week at room temperature (20-26°C) or 30 days at 2-8°C.

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2 Diluted lyse solution must be between 20°C and 37°C when used. Cold solution may result in poor lysis.

3 Aliquot 50µL of Leuko64 Reagent A into a 12 x 75 mm polystyrene tube.

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4 Aliquot 50µL of mixed anticoagulated blood sample with white blood count $<25 \times 10^9$ cells/L (dilute as required) to the tube containing Reagent A, gently mix or vortex, and incubate 10 minutes in the dark at room temperature.

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5 Add 1.0mL of 1X Trillium Lyse (diluted Reagent B) and thoroughly vortex. Let stand for 15 minutes in the dark at room temperature. Intermittent vortexing or mixing enhances lysis.

6 Add 5µL Leuko64 beads (Reagent C) to each tube, vortex, and analyze on flow cytometer using instrument set-up and analysis protocol below. Prepared samples should be held at 2-8°C, shielded from light until analyzed. Analysis should be performed within 24 hours of staining.

CD64 Instrument Calibration Bead Specimen:

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FLOW CYTOMETER SET-UP

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1 Add 5µL Leuko64 beads (Reagent C) to a tube containing 0.5mL 1X Trillium Lyse (Reagent B diluted 1:10 with filtered distilled water, pH 7.40 ± 0.02). This will be used to establish PMT voltage and scatter settings on the instrument.

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2 Set up 4 two-parameter histograms:

FS (lin) vs SS (log)

CD64 FITC vs SS (log)

CD163 vs SS (log)

CD163 PE vs CD64 FITC

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And 3 one-parameter histograms:

FL1 - CD64 FITC

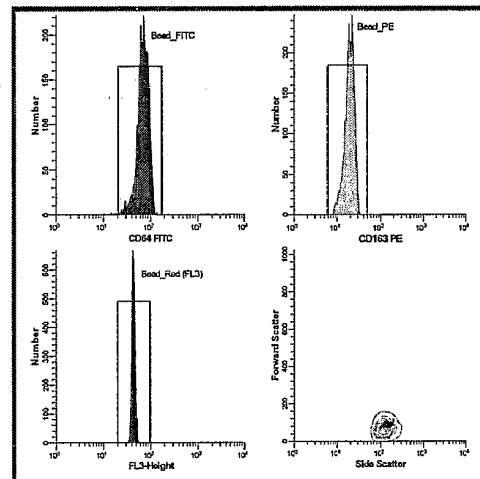
FL2 - CD163 PE

FL3 (the PMT with filter setup able to detect 685nm)

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3 Turn OFF all compensation settings.

- 4 Run the bead suspension and make the following adjustments on the one-parameter histograms at 1024 channel resolution (see diagram):
- The center of the peak on the FL1 (FITC) axis should be at the end of the second decade of fluorescence intensity (MFI = ~90).
 - The center of the peak on the FL2 (PE) axis should be at a MFI = ~20 (the first tic in the second decade of fluorescence intensity).
 - The center of the peak on the FL3 axis should be midscale in second decade (MFI = ~30).
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- 10 5 On the FS vs SS histogram, the bead population should be positioned at the start of the third decade on log side scatter signal, and at channel 85-120 on forward or low angle scatter signal (on 256 scale use channel 20-25).
- 15 6 The threshold to exclude platelets and red cell debris should be set on SS (log) using lymphocytes. The discriminator to exclude debris and small particles should be set just to the left of the lymphocytes.
- 20 7 Acquisition and Storage settings should be adjusted to allow for collection of 50,000 ungated events. Setting resolution at 1024 is generally recommended, but is required when using Beckman Coulter FC-500™ or Becton Dickinson FACSCanto™ cytometers.
- 8 Save the settings and acquisition template.



Flow Cytometric Acquisition:

- 30 1. Perform the flow cytometer's daily instrument QC program and determine that the cytometer is ready for specimen acquisition.
2. Call up the CD64 protocol and run the stained specimens, including the beads.
- 35 3. Acquire at least 50,000 events and save list mode file for analysis with the Leuko64 software.

Analysis of Listmode Files:

1. Leuko64 software which is the software program used for the calculation of results.
2. Call up the bead mix listmode file and following the directions in the Leuko64 manual to initiate the Leuko64 Program
3. Call up the first listmode file to be analyzed. Confirm that the coarse gates around the leukocyte populations of lymphocytes, monocytes, and granulocytes are positioned correctly. If needed, the gate region can be adjusted to surround the appropriate cell population.
4. Analysis of specimen list mode files can be analyzed in a step by step fashion or by using the automated batch analysis function. Analysis results can be stored in the database and printing of the results is suggested to allow for subsequent documentation and review of the appropriate gating of the leukocyte subpopulations.
5. If the analysis is stopped due to either the lymphocyte or monocyte flag or warning dialog box appearing, the user should investigate possible causes for these adverse findings and consider reanalysis and/or restaining of the sample if necessary.

20 Limitations of Procedure:

1. It is essential to acquire adequate numbers of bead events (500 or more). Inadequate numbers of beads may cause invalidate CD96 and CD163 index results, which in turn will may result in false positive or negative results.
2. Some specimens do not lyse completely. However we are interested in the Neutrophil and bead populations, so the specimens can still be run. If the discriminator setting is not optimized, the listmode file may have insufficient bead or cell events to be statistically robust.
3. The assay requires sufficient anti-CD64 reagent added to the tube, this could be verified by examination of the monocyte population, which can serve as an internal positive control as this populations should always stain "positive". If that is not the case, the software analysis monocyte flag will be activated.

35 **Components of Invention**

I. Reagent A

Leuko64 Reagent A:	Lot #L64-03-07	Bottled conc (mg/mL)	Dilution Fac
Anti-CD64 clone 32.2 FITC (ATCC: HB9469)	0206113.1T1	0.68	diluted 1:272
Anti-CD64 clone 22 FITC	0104031.1T4	1.47	diluted 1:980
Anti-CD163 PE clone 48 or 158 (ATCC: HB 10714)	lot 0103203.2E5	0.93	diluted 1:115
PBS w/ 0.1% BSA + 0.01% NaN ₃ (Sigma)			

40 Reagent A is used at a volume of 0.050 mL per test. Thus for a 250 test kit the total volume needed is 12.50 mL plus 1.25 mL for wastage. Bottle in two containers with a file volume of 7.5 -10 mL. Bottles should be flat bottom amber glass composition with leak proof screw cap.

II. Leuko64 Reagent B: 10X Trillium Lyse Solution

- NH₄Cl (potassium chloride) 80.0 g Sigma # P-9541
 - EDTA Trisodium Salt 10.0 g Sigma # ED3SS
 - KH₂PO₄ (potassium phosphate) 1.0 g Sigma # P-5379
- 5 • QS to 1000 ml with reagent grade H₂O
- Filter using 0.2µm-pore vacuum filter Nalgene # 295 4520
- Adjust pH to 7.4 when solution reaches room temperature (use aliquots and aseptic tech
- To avoid contaminating the filtered solution)
- Store in refrigerator at 2-8°C
- 10 • Expiration 12 months

Reagent B is used at a volume of 1.0 mL per test after dilution (diluted form to have one week outdate). Thus for a 250 test kit the total volume needed is 25 mL plus 15 mL for wastage. Bottle in a container with a fill volume of 50 mL. Bottle can be of

15 plastic composition (nalgene) with flat bottom and screw cap.

- III. Leuko64 Reagent C: Calibrator Beads (NIST traceable reference material)
- Polystyrene 5.2 µm beads tagged with a proprietary embedded red dye (starfire red)
- 20 and surface coated with FITC (NTRM to NIST SRM 1932)
- Suspended in an isotonic buffered solution containing BSA, sodium azide, Tween 20, pH 7.4.
- Final bead concentration at $4-6 \times 10^6$ per mL.

Bead Diluent:

Tween 20	10uL
NaN ₃	0.013g
PBS	100mL
pH = 7.4	

Make a 1:20 dilution of each into PBS with 0.01% Tween and 2mM NaN₃

FITC Beads:

Beads	250uL
Bead Diluent(above)	4750uL

- 25 Reagent C is used at a volume of 0.005 mL per test. Thus for a 250 test kit the total volume needed is 1.25 mL plus 0.25 mL for wastage and instrument calibration.
- Bottle in one container with a file volume of 2.0 mL. Bottles should be flat bottom amber glass composition with leak proof screw cap.
- 30 Reagent C will be provided for each lot following QC by Trillium Diagnostics, LLC and OEM manufacturing by Bangs, Inc.

IV. Leuko64 Software

- 35 Lot specific Leuko64 software will be provided in the kit in a standard CD disc. The CDs will be provided for packaging after QC validation by Trillium Diagnostics, LLC

and Verity Software House. The software provides a lot-specific value assignment for the calibration beads and subsequently the final PMN and Monocyte CD64 Index values; this corrects for any lot-to-lot variations in bead production or monoclonal antibody fluorochrome labeling. Software validation procedure is in software folder.

5

All publications and patent applications disclosed herein, including US Patent Nos. 4,918,004 and 5,380,663 are incorporated into this application by reference in their entirety, which can be used in the instant invention.

10

One skilled in the art will readily appreciate that the present invention is well adapted to carry out the objects and obtain the ends and advantages mentioned, as well as those inherent therein. It will be apparent to those skilled in the art that various modifications and variations can be made in practicing the present invention without departing from the spirit or scope of the invention. Changes therein and other uses will occur to those skilled in the art which are encompassed and equivalent within the spirit of the invention as defined by the scope of the claims.

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We claim:

1. A kit for automated performance analysis to quantify CD64 and CD 163 expression in a leukocyte and calibration for a flow cytometer, said kit used with a flow cytometer, blood specimens, a programmable computer and a display screen, said kit comprising:
 - (a) at least one or more monoclonal antibodies directed to CD64 conjugated to fluorescein isothiocyanate (FITC) or similar fluorochromes;
 - (b) at least one or more monoclonal antibodies directed to CD163 conjugated to phycoerythrin or similar fluorochromes;
 - (c) a suspension of polystyrene microbeads with at least one or more fluorochromes, one of which is identical to that conjugated to the CD64 monoclonal antibodies, environmentally sensitive to the same factors affecting the fluorescence of antibodies bound to cells, and traceable to an external standard, such as the NIST SRM 8640;
 - (d) at least one or more fluorochromes, said fluorochrome having a useful life, said useful life being designated in said kit; and
 - (e) software operating on said computer comprising
 - 1) information on the fluorescence intensity of each fluorescent reagent and microbeads, said software operating with said computer to control calculation of calibration and fluorescence information tailored to said microbeads, and display on said screen an expiration date correlated to the useful life of said microbeads to an operator calibrating said flow cytometer with said microbeads, wherein said software ceases operation on said computer at said expiration date, whereupon said flow cytometer cannot be calibrated with said software and said microbeads and the operator must utilize another suspension of microbead populations and software tailored for use on said computer with said another suspension to calibrate the flow cytometer;
 - 2) the ability to acquire data on blood specimens using fluorescence signals, FS and SS and thus, automatically gate and smooth the data, locate peaks in fluorescence intensity corresponding to each microbead and cell specific populations, construct linear regression plots, determine the fluorescence threshold intensity of the flow cytometer, create histograms;
 - 3) the ability to find cell specific clusters;
 - 4) the ability to perform automated data normalization; and
 - 5) the ability to calculate the CD64 and CD163 index on specific cell populations.

2. A kit according to claim 1, wherein the monoclonal antibody directed to CD64 is Clone 22.
- 5 3. A kit according to claim 1, wherein the monoclonal antibody directed to CD64 is Clone 32.2 (ATCC Number HB-9469).
4. A kit according to claim 1, wherein the monoclonal antibody directed to CD163 is Clone Mac2 (ATCC Number HB-10714).
- 10 5. A kit according to claim 1, wherein the fluorochrome is fluorescein isothiocyanate (FITC).
6. A kit according to claim 1, wherein the microbeads are comprised of polystyrene.
- 15 7. A kit according to claim 1, wherein the cell specific cluster comprised of lymphocytes.
8. A kit according to claim 1, wherein the cell specific cluster comprised of
- 20 monocytes.
9. A kit according to claim 1, wherein the cell specific cluster comprised of neutrophils.
- 25 10. A kit according to claim 1, wherein the software further comprises the ability to perform quality control functions.
11. A kit according to claim 10, wherein the quality control function is to validate that the monocyte population has the expected CD64 expression detected by the
- 30 fluorescent monoclonal antibodies and, consequently serve as an internal positive control for each specimen.
12. A method for quantifying human leukocyte FcγRI receptor (CD64) expression comprising;

(a) providing a flow cytometer, blood specimens, a programmable computer and a display screen;

(b) providing a combination of reagents which allow for the detection of the FcγRI
5 receptor; and

(c) providing a software operating on said computer comprising the ability to acquire data on blood specimens using fluorescence, FS and SS and thus create histograms, automatically gate and smooth the data, locate peaks in fluorescence intensity
10 corresponding to each microbead and cell specific populations, construct linear regression plots' determine the fluorescence threshold intensity of the flow cytometer; to find cell specific clusters the ability to perform automated data normalization and the ability to calculate the CD64 index.

15 13. A method of claim 12 wherein the leukocyte is a polymorphonuclear leukocyte.

14. A method for quantifying human leukocyte, in particular monocyte, CD163 expression comprising;

(a) providing a flow cytometer, blood specimens, a programmable computer and a
20 display screen;

(b) providing a combination of reagents which allow for the detection of the CD163 or the hemoglobin-haptoglobin complex receptor; and

(c) providing a software operating on said computer comprising the ability to acquire data on blood specimens using fluorescence, FS and SS and thus create histograms, automatically gate and smooth the data, locate peaks in fluorescence intensity
25 corresponding to each microbead and cell specific populations, construct linear regression plots' determine the fluorescence threshold intensity of the flow cytometer; to find cell specific clusters the ability to perform automated data normalization and the ability to calculate the CD163 index.

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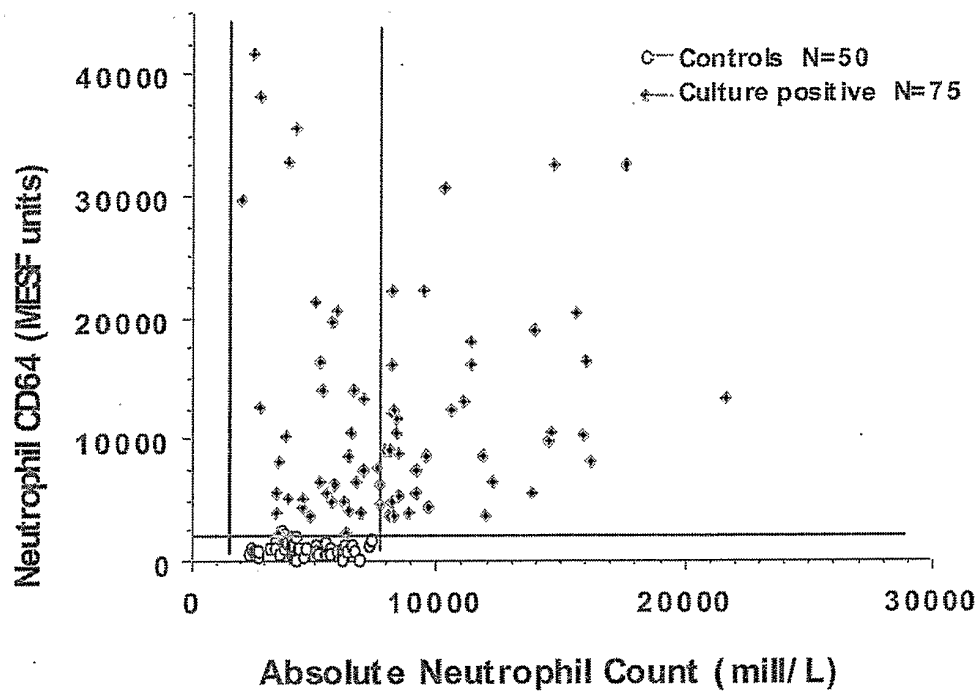


Figure 1.

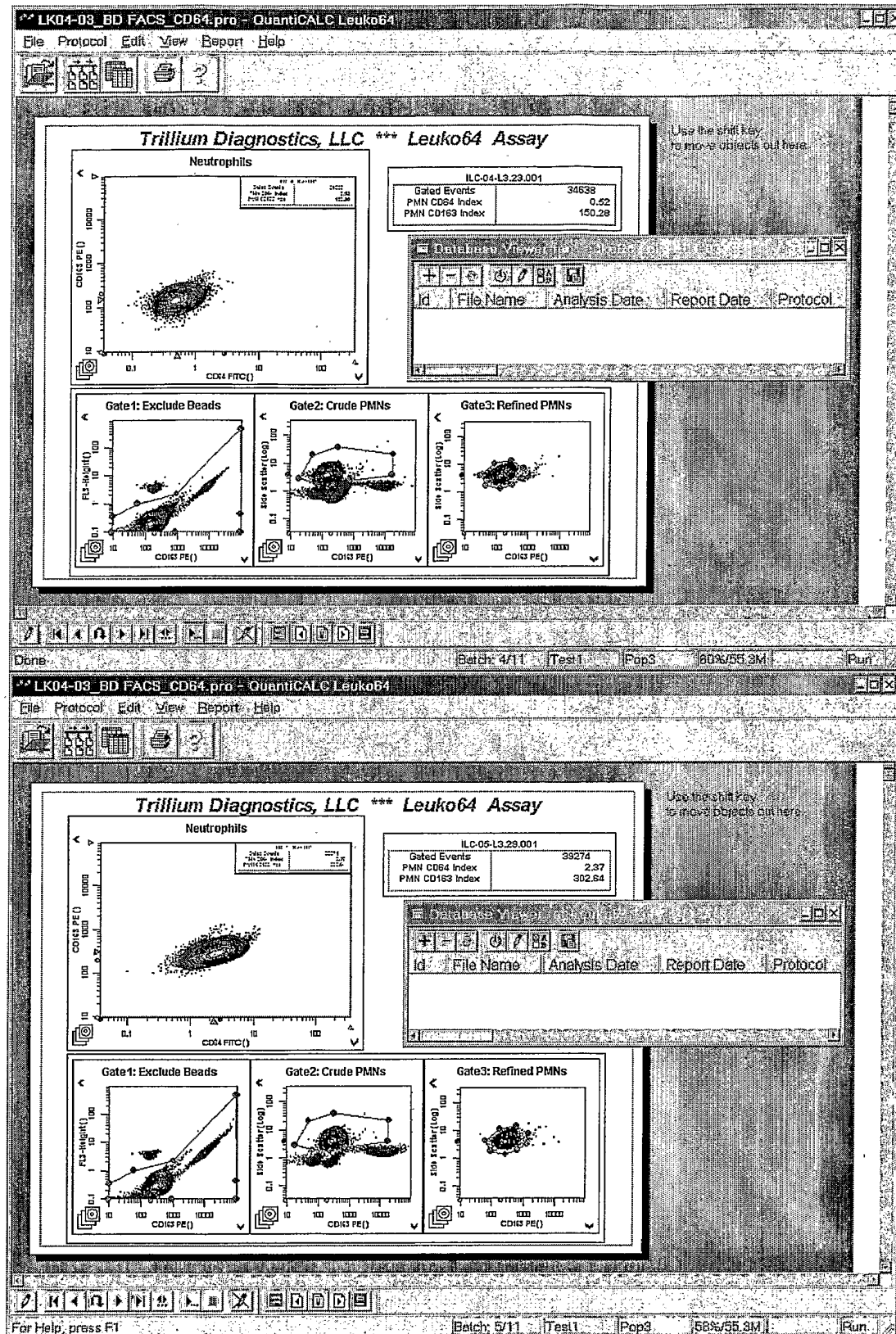


Figure 2

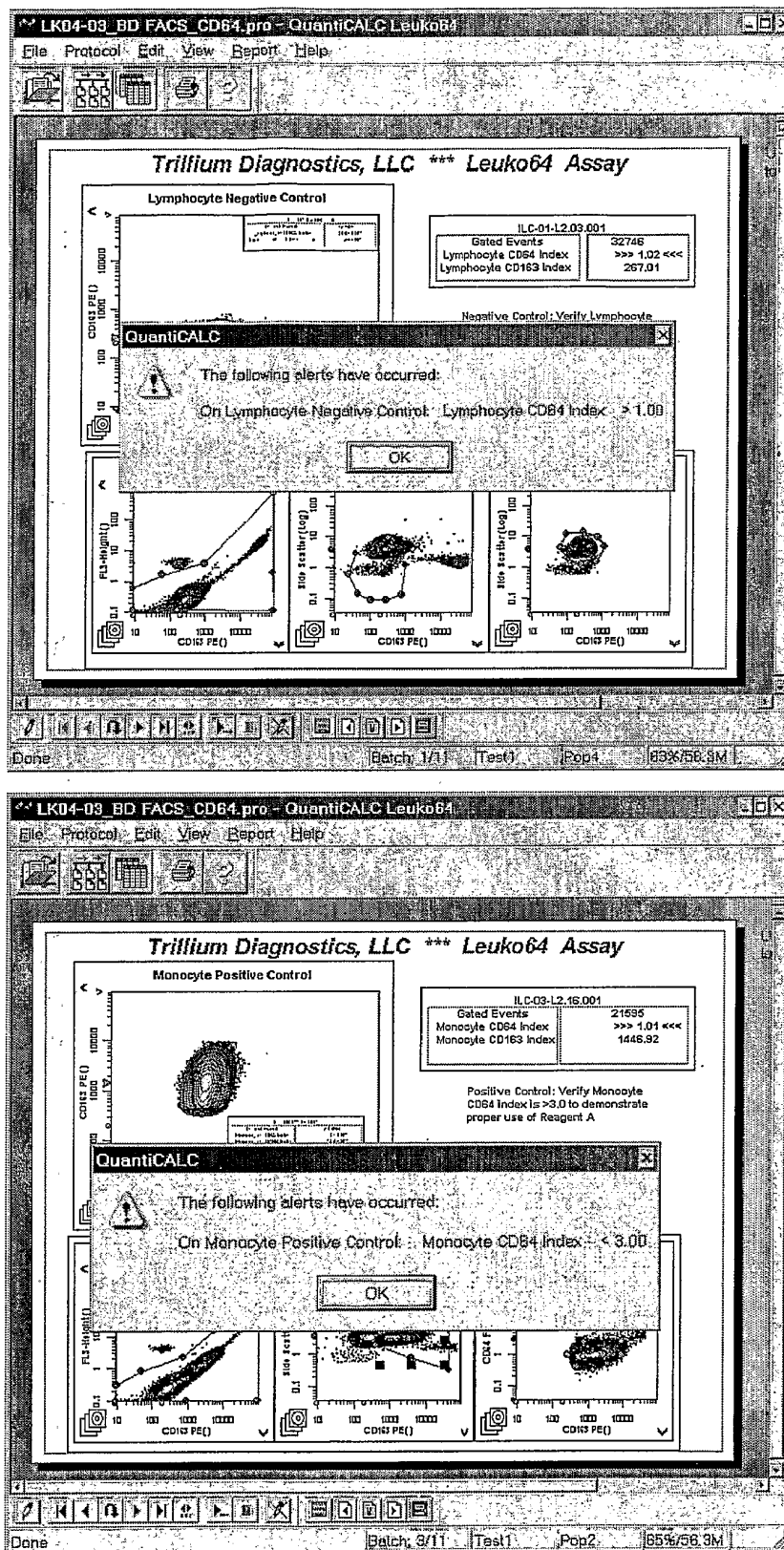


Figure 3.

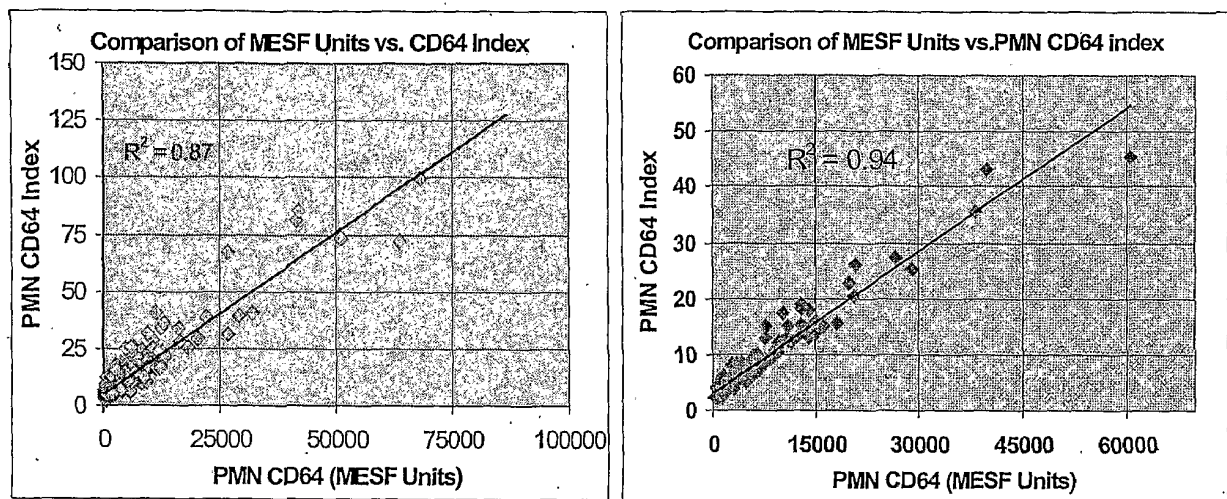


Figure 4

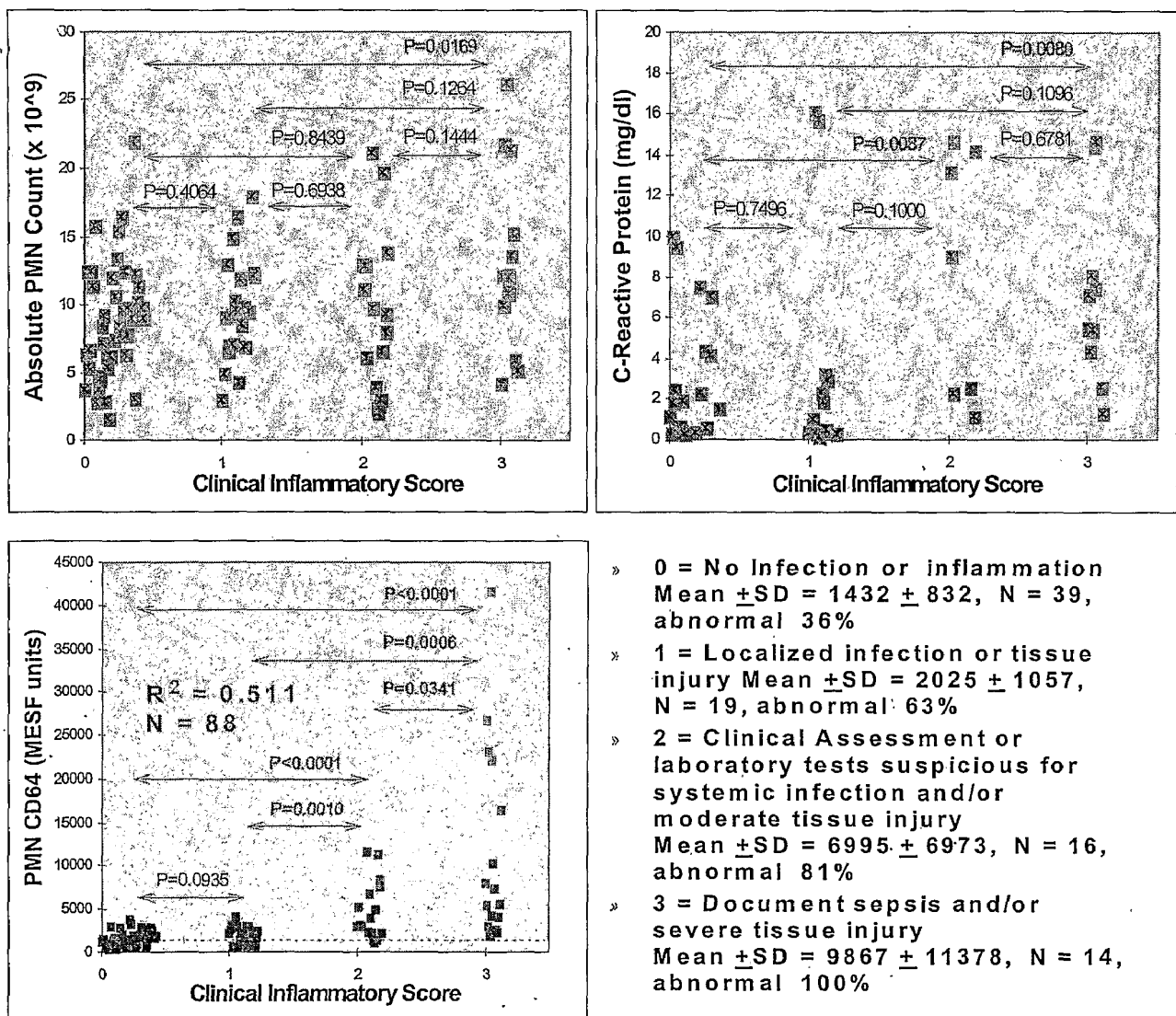
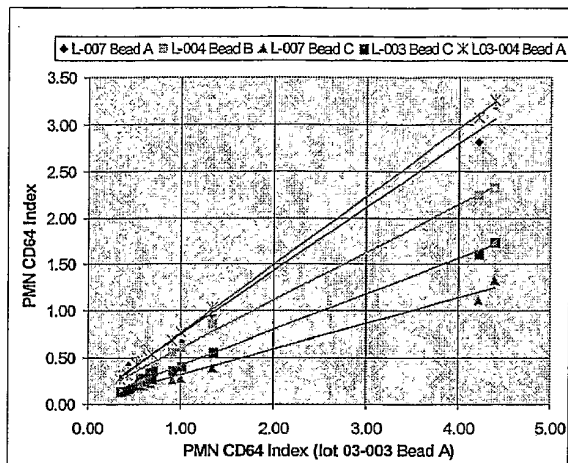


Figure 5

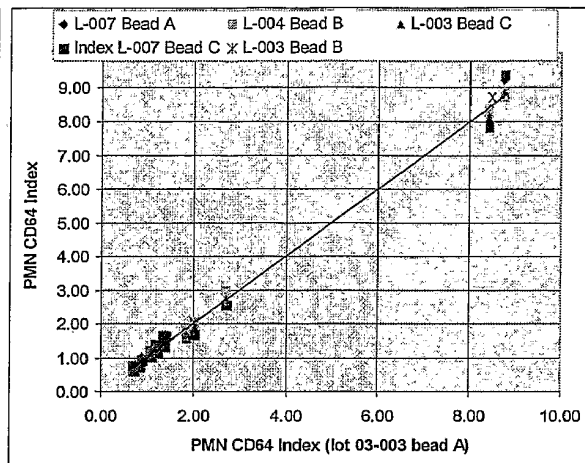
Leuko64 Software: Value assignment of beads within software allows lot to lot standardization of CD64 Index.

Lot comparisons without software



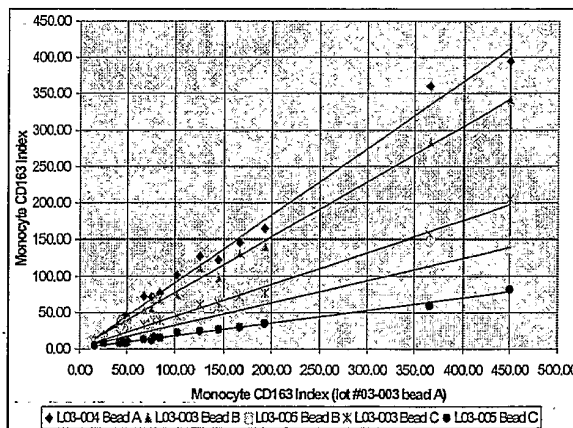
Lot Comparison	Regression	r ²
03-003 A vs 03-003 B	$y = 0.7355x + 0.0212$	0.9996
03-003 A vs 03-007 A	$y = 0.6867x + 0.0568$	0.9956
03-003 A vs 03-004 B	$y = 0.5159x + 0.0841$	0.9984
03-003 A vs 03-003 C	$y = 0.3837x + 0.0325$	0.9958
03-003 A vs 03-007 C	$y = 0.2754x + 0.0438$	0.9884

Lot comparisons using software

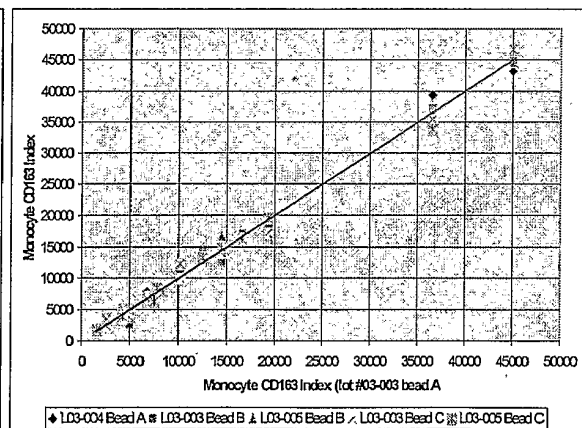


Lot Comparison	Regression	r ²
03-003 A vs 03-007 A	$y = x + 6E-05$	0.9956
03-003 A vs 03-003 B	$y = x - 4E-06$	0.9996
03-003 A vs 03-004 B	$y = x - 3E-05$	0.9984
03-003 A vs 03-003 C	$y = x + 0.0001$	0.9958
03-003 A vs 03-007 C	$y = x - 0.0002$	0.9884

Leuko64 Assay: Software value assignment of bead fluorescence values also allows lot to lot CD163 Index standardization



Lot Comparison	Regression	r ²
03-03 A vs 03-04 A	$y = 0.9154x - 0.3291$	0.9916
03-03 A vs 03-03 B	$y = 0.7582x + 1.6349$	0.9926
03-03 A vs 03-05 B	$y = 0.3053x + 2.4133$	0.9948
03-03 A vs 03-03 C	$y = 0.4376x + 1.427$	0.9952
03-03 A vs 03-05 C	$y = 0.1718x + 1.7214$	0.9911



Lot Comparison	Regression	r ²
03-03 A vs 03-04 A	$y = 1.0017x - 84.496$	0.9918
03-03 A vs 03-03 B	$y = 1.0016x - 11.74$	0.9927
03-03 A vs 03-05 B	$y = 0.9998x + 0.0144$	0.9948
03-03 A vs 03-03 C	$y = 0.9999x - 0.007$	0.9952
03-03 A vs 03-05 C	$y = 1.0000x - 0.0091$	0.9911

Figure 6

专利名称(译)	软件集成流式细胞分析法用于人体多形核白细胞FcRI受体 (CD64) 的定量分析		
公开(公告)号	EP1836557A4	公开(公告)日	2009-01-21
申请号	EP2005851842	申请日	2005-11-18
[标]申请(专利权)人(译)	延龄诊断		
申请(专利权)人(译)	延龄诊断, LLC		
当前申请(专利权)人(译)	延龄诊断, LLC		
[标]发明人	DAVIS BRUCE H BAGWELL C BRUCE HUNSBERGER BENJAMIN		
发明人	DAVIS, BRUCE, H. BAGWELL, C., BRUCE HUNSBERGER, BENJAMIN		
IPC分类号	G06F7/00 G01N33/48 G01N33/537		
CPC分类号	G01N33/6872 G01N33/537 G01N33/56972 G01N33/582 G01N2333/70535 G01N2333/70596 G01N2496/05 G01N2800/26		
优先权	60/629654 2004-11-19 US		
其他公开文献	EP1836557A2		
外部链接	Espacenet		

摘要(译)

本发明涉及定量白细胞中CD64和CD163表达的方法，尤其涉及一种与流式细胞仪一起使用的试剂盒，该试剂盒包括定量荧光微珠标准物的悬浮液，针对CD64和CD163的荧光标记抗体以及分析软件。该软件用于从流式细胞仪上获取微珠悬浮液和荧光标记抗体的信息，并分析数据，平滑曲线，计算新参数，提供质量控制措施并通知测定系统到期。

