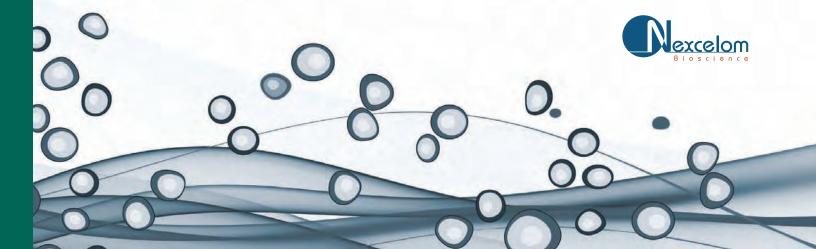
# Cellometer® K2

# Image Cytometer for Cell Counting & Analysis





# Cellometer K2 Image Cytometer

# **Optimized Analysis of Primary Cells**



#### Features of the Cellometer K2

**Dual Fluorescence and Bright Field Imaging:** staining of both live and dead cells in heterogeneous samples

**User-Friendly Software and Assay Selection:** Enhanced inter-operator reproducibility, minimal training, auto-save option

Fast Results: Obtain cell images, counts, size measurements, and viability calculations in 60 seconds

Small Sample Size: Only 20 µl of sample

**Broad Dynamic Range:** Measurable concentration range of 1 x  $10^5$  to 1 x  $10^7$  cells/mL using Nexcelom's proprietary de-clustering function

Many Compatible Dyes: Trypan blue, AO, PI, EB, 7AAD, AO/PI, AO/EB, Calcein AM, CFDA, Calcein AM/PI, CFDA/PI



### **Advantages of Cellometer Image Cytometer**

#### Cell Imaging

- Verify cell morphology and counted live/dead cells
- Export cell images for presentations and publications

#### Pattern Recognition Software

- Accurately count cells in clumps
- Count irregular-shaped cells
- Eliminate debris from cell counts
- Differentiate cells based on size

#### Automated Data Management

- Pre-set assays and automated reports
- Archive sample images and auto-save results

#### Maintenance-free System

- Disposable counting chambers no wash steps
- No required instrument maintenance

Learn why thousands of users, including the top ten pharmaceutical companies, trust Cellometer.

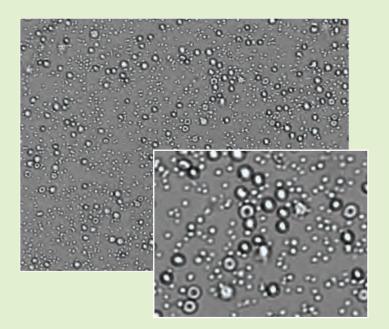
On-Line Demonstrations are completed in just 20 to 30 minutes and provide an overview of how Cellometer works using existing images of cells that interest you.

On-Site Demonstrations are a convenient way to test a Cellometer system for a specific application. An experienced Applications Specialist will arrive at your lab for a hands-on session to test your cells and show how Cellometer can enhance your workflow.

Technical Seminars are an excellent way to introduce Cellometer systems to a lab group or collaborators in different laboratories within an organization. A trained biologist will discuss and demonstrate the capabilities and advantages of Cellometer image cytometry.

Call 978-327-5340 or E-mail info@nexcelom.com today to schedule a free demonstration or technical seminar.





#### PBMC Analysis in the Presence of Red Blood Cells

Measure PBMCs from whole blood without lysing. Obtain baseline PBMC concentration and viability prior to biomarker studies

#### **Nucleated Cell Concentration & Viability**

Evaluate cord blood and bone marrow samples

#### **GFP Transfection Efficiency & Viability**

Quickly and easily monitor DNA, RNA, and siRNA transfection

#### Analysis of Clumpy & Irregular-Shaped Cells

Nexcelom's proprietary pattern-recognition software enables accurate analysis of >98% of mammalian cell types

#### **Cell Line Analysis**

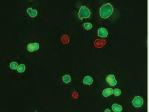
Automatically capture fluorescent cell images, concentration, Trypan blue or PI viability, and mean diameter in 60 seconds!

#### **Analysis of Cells from Heterogeneous Samples**

- Whole Blood
- Peripheral Blood
- Cord Blood
- Bone Marrow
- Bronchoalveolar Lavage (BAL)

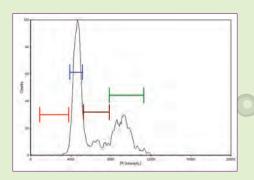
#### Primary Hepatocytes: Cell Count and Viability





#### **Cell Based Assays**

- Cell Cycle
- Apoptosis
- GFP



## Proven Performance in Many Research Areas



- Clinical Immunology: PBMCs
- DMPK: Primary Hepatocytes
- Regenerative Medicine: Stem Cells
- Transplantation: Nucleated Cells
- Vaccine Development: Splenocytes
- Oncology: Cell Lines, Cell Cycle, Apoptosis
- Basic Research: Primary Cells / Cell Lines / GFP

Optimized for Primary Cell Analysis

PBMCs

Splenocytes

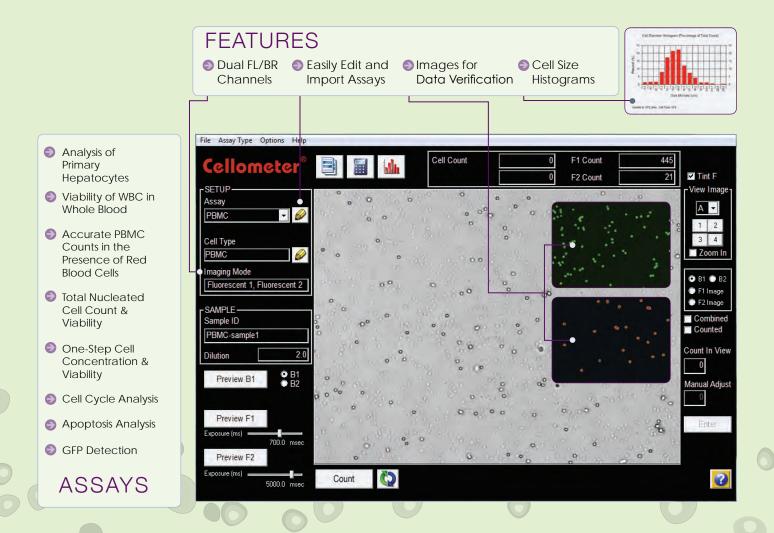
Cell Cycle

Cells

Apoptosis Conrega

Contact Nexcelom regarding your cell type

# Cellometer K2 Image Cytometer for Cell Counting & Analysis from Nexcelom Bioscience



#### **How It Works**





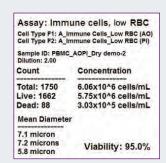
Pipette 20 µl of Cell Sample



Insert Counting Chamber



Select Assay & Click Count



Get Results

# Dual-Fluorescence for Primary Cell Viability in Heterogeneous Samples

Live / Dead Cell Concentration using AO / PI

**Dual-Fluorescence Viability**, using acridine orange (AO) and propidium iodide (PI), is the recommended method for accurate viability analysis of primary cells, such as PBMCs, splenocytes, and stem cells, in samples containing debris and unwanted non-nucleated cell types including red blood cells.

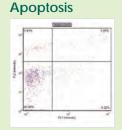
Acridine orange (AO) and propidium iodide (PI) are nuclear staining (nucleic acid binding) dyes. AO is permeable to both live and dead cells and stains all nucleated cells to generate green fluorescence. PI enters dead cells with compromised membranes and stains all dead nucleated cells to generate red fluorescence.

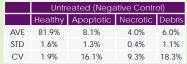
Because mature mammalian red blood cells do not contain nuclei, only live and dead mononuclear cells produce a fluorescent signal. There is no need to lyse red blood cells, saving time and eliminating an extra sample preparation step.

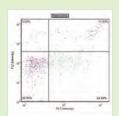


#### Export to FCS Express\* for Flow-Like Data Output









	Treated (Positive Control)						
	Healthy	Apoptotic	Necrotic	Debris			
AVE	58.8%	24.7%	13.8%	2.7%			
STD	1.9%	1.1%	1.2%	0.2%			
CV	3.2%	4.3%	9.0%	9.2%			

FCS Express 4 Flow Cytometry software is a product of De Nova Software

### Performance of the Cellometer K2 Image Cytometer

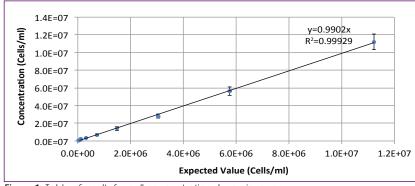


Figure 1. Table of results for cell concentration dynamic range

#### Concentration Dynamic Range

Figure 1 depicts the dynamic range for cell concentration measurements on Cellometer K2. This data set was taken on a concentration series of cultured Jurkat cell line.

Samples from 1 x 10<sup>5</sup> – 1 x 10<sup>7</sup> cells/ml can be counted without further dilution.

The %CV at each concentration was below 10%.

**Viability Dynamic Range** The viability dynamic range is 0 - 100% for Cellometer K2 Image Cytometer using dual fluorescence AO/PI stain.

Sample	N Value	Average Live Cell Concentration	% Viability	CV of Concentration	CV of Viability
Jurkat	24	3.61E+06	92.2%	8.9%	1.0%
Human PBMC	10	5.94E+06	96.0%	4.7%	0.5%
Mouse Splenocyte	10	1.86E+07	88.6%	5.6%	0.7%

Figure 2. Table of results for cell concentration and viability using AOPI

Consistency and Repeability The results indicate the accuracy of the Cellometer K2 instrument in assessing the viability of Jurkat cells using AOPI for cell viability. Jurkat, human PBMC, mouse splenocytes were tested at 24, 10, and 10 sample replications, respectively. The viability average was calculated and plotted. The results show the reliability and accuracy of the Cellometer K2 in measuring cell concentration and viability of mammalian cells.

Cellometer Counting Chamber									
Catalog #	Description	Size	Unit						
CHT4-PD100-003	Standard chamber thickness. Packed in microscope slide boxes. Ready to use.	Case of 500 slides for 1,000 counts (10 individual boxes)	1 Case						
CHT4-SD100-014	Standard chamber thickness. Packed with protective film on both sides. Remove protective film before use.	Case of 900 slides for 1,800 counts	1 Case						
CHT4-PD300-003	3x standard chamber thickness. Packed in microscope slide boxes. Ready to use.	Case of 500 slides for 1,000 counts (10 individual boxes)	1 Case						
See www.nexcelom.com/products for updated product selections									

#### Cellometer Cell Counters, Cell Analysis Systems & Image Cytometry

Nexcelom offers a wide range of Cellometer systems developed and optimized for specific applications and cell types.









For more information, visit
www.nexcelom.com

#### Contact us at: Nexcelom Bioscience 360 Merrimack Street, Building 9 Lawrence, MA 01843, USA

Email: info@nexcelom.com							
Phone: 978.327.5340							
Fax: 978.327.5341							

Features	T #	Automat	ed Cell	Counte	rs	Image Cytometers				
	Mini	Auto T4	Auto 1000	Auto 2000	X4 (10x)	X1	X2	K2	Vision CBA	Visio CB/ (10)
Cell / Sample Type										
Objective Magnification	4X	4X	4X	4X	10X	10X	10X	4X	5X	10)
Cell Line	Х	Х	Х	X				Х	Х	
Cultured Primary Cells	Х	X	Х	Х				Х	Х	
Algae					X					Х
Platelets					Х		Х			Х
Low Cocnentration Cell Lines				Х				Х	Х	
Yeast (Clean Sample)					X	X	X			
Primary cells (Messy Sample*)				X				Х	Х	
PBMCs, Splenocytes, Stem Cells				Χ				Х	Х	
Yeast (Messy Sample)							X			Х
Hepatocytes								Х	Х	
Adipocytes***				Χ				Х	Х	
Cell-Based Assay **						X	X	Х	Х	Х
Apoptosis (Annexin V-FITC/PI)								Х	Х	Х
Apoptosis (Caspase Activity)								Х	Х	Х
Autophagy (CytolD-green)									Х	Х
Cell Proliferation (CFSE)									Х	Х
Cell Cycle (PI)						Х	Χ	Х	Х	Х
GFP Transfection				Χ			Χ	Х	Х	Х
YFP Transfection									Х	Х
RFP Transfection									Х	Х
Mitochondrial Potential (JC-1)									Х	Х
Multi-drug Resistance (ABC Transporter)									х	х
Surface Marker Analysis									Х	Х
Vitality (Calcein-AM/PI)							X	Х	Х	Х

A messy sample is a heterogeneous sample containing unwanted cell types, such as red blood cells, in addition to the cells of interest. FCS Express 4 license must be purchased in order to perform Cell Based Assay or Image Cytometry analysis

\*Cellometer CHT4-PD300 slides are required for cells greater than 80µm in diameter

