



# Quantitative characterization of cells using automated image analysis

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### Cellometer Automated Cell Counters: Simplest Solution to Automate Hemacytometer

### 30 seconds to get more accurate data



Cellometer Auto T4 Plus Automated Cell Counter

**Cellometer Auto M10** Automated Cell Counter with high magnification

#### **Cellometer Vision**

Automated Cell Counter with multimode detection (bright field and fluorescence)







### The importance of cell counting

- Application :
- Cell invasion/Migration transwell seeding
- Drug selection
- Condition testing- time course/dose
- Flowcytometry







### The importance of cell counting

• Hemacytometer













### Why need automated cell counter?



ED







17	>	Cellometer <sup>®</sup> AutoT4			Hemacytometer		
13		Concentration	n	CV	Concentration	CV	n
	Jurkat	1.17 × 10⁰	4	2.27%	1.11×10°	8.04%	3
	YAC-1	1.38 × 10 <sup>6</sup>	4	3.17%	1.41 × 10°	11.89%	3
	K562	6.36 x 10⁵	4	8.29%	5.58 x 10⁵	9.66%	3
	H460	1.32 × 10 <sup>6</sup>	4	5.57%	1.16 x 10⁴	20.65%	3
	NCI H128	2.46 x 10⁵	4	11.23%	1.50 x 10⁵	49.33%	3
	SW620	3.81 × 10⁵	4	2.79%	3.70 x 10⁵	19.49%	3
	Hut 102	9.18×10⁵	4	0.39%	7.68 x 10⁵	10.60%	3





### The importance of cell counting

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### The evolution of cell counting

• From Hemacytometer to Cellometer











Count and Analyze Cells in 3 Steps

**Step 1.** Pipette 20uL sample into a disposable Cellometer chamber

1.99 x 10<sup>6</sup>

**Step 2.** Place chamber into Cellometer

**Step 3.** Run Cellometer software to obtain cell concentration and viability automatically

*Typical cell lines take less than 30 seconds per sample.* 

•Cell concentration •Viability •Cell size •On screen image











### Principle of Cellometer is the image-based analysis

N Cellometer Auto I4	Сеll Туре			
File Cell Type Options <u>Analysis</u> Help	Cell Type Name Jurkat 🗖 Save as New Cell Type			
Display Imate Count	Detailed Description			
Cell Type Test Viability	Cell Diameter Min. 5 micron Cell Size			
Jurkat  Detroy	Cell Roundness 0.10 default 0.10 Cell Roundness			
nd HEK293	Contrast Enhancement 0.40 default: 0.40; range: 0 - 0.5; high value for light cells			
Hela HT1080 HT-29	✓ Modify Dead Cell Parameters         Sensitivity         1.0         hig         Dead cell analysis			
Jurkat	Uniformity 150 default: 150; range 100 - 255 higher value for non-uniform dead cells			
PBMC_human_fresh PBMC_human_fresh_high con	Very Dim Dead Cells Contrast Enhancement 0.60			
SF9 Vero	Modify Decluster Details     Decluster Edge Factor     0.5     hig     Docustor			
(micron)	Decluster Th Factor 1.0 def			
	Select Special Cell Type			
Cell Concentration 8.62 x10 <sup>5</sup>	Background Adjustment 1.0 default: 1.0; range 0 - 1.0 lower value to pick up dim cells			
	No Decluster     Print			
	Save Cancel			









Modify decluster details : •Counting declustering cell clumps



### Decluster off

Decluster on

#### Cellometer® 0 30 -SETUP-0 Test Viability Cell Type **Cell Size Analysis** -Jurkat Diameter File Name: C:\Documents and Settings\All Users\Application Data\Nexcelom\diameters.txt Sample ID **Cell Diameters** Histogram Jurkat

![](_page_11_Figure_1.jpeg)

![](_page_12_Picture_0.jpeg)

**PBMC** in Platelets

![](_page_12_Picture_1.jpeg)

### **Revolution of Cellometer** Advanced imaging technology :

- 1. Counting specific cells in a heterogeneous sample (ex: PBMC)
- 2. Counting declustering cell clumps
- 3. Counting irregular shaped cells (activated T cell)
- 4. Differentiation of cell population by size (SF9 virus infection )
- 5. New cell types are easily added via a New Cell

Type Wizard or manual configuration

![](_page_12_Picture_9.jpeg)

Irregular shaped cells

SF 9 insect cell - virus infection

Size Microns (um)

Cell Diameter Histogram

20 15

![](_page_13_Picture_0.jpeg)

![](_page_13_Picture_1.jpeg)

### **Correlation between T4 and hemacytometer**

 Breast cancer cell line concentration measured by T4 and hemacytometer

![](_page_13_Figure_4.jpeg)

![](_page_14_Picture_0.jpeg)

![](_page_14_Picture_1.jpeg)

Citation

![](_page_14_Figure_3.jpeg)

The percentage of viable cells was counted using the Nexcelom Cellometer Auto T4

The Journal of Immunology, 2009, 182; 6460-6469

![](_page_15_Picture_1.jpeg)

### Appendix 3 - Citations

20 articles found		Too many results? View tools to help narrow your search				
1-10 of	All Articles 20 articles	Articles Indexed by Subject View: <u>Next 10 Results</u> *				
Citation: 🧿	standard 🔾 condensed   View: 10 💌	per page   Update display				
For checked items below: I view abstracts I download to citation manager						
PROMOLECTION SCREENING	Product Focus: Automation and I J Biomol Screen, Dec 2006; 11: 1044 - 10 For more information, visit www.ve launched its CellometerTMinformatio Nexcelom Bioscience at www.nexcelom announces	Robotics 52. locity11.com. Nexcelom Bioscience, LLC has n and application notes are available from .com. The Automation Partnership (TAP)	<ul> <li>PDF (\$30.00)</li> <li>Find more like this</li> </ul>			
CONTRACTOR	HEAD AND NECK CANCER: Intratumoral Epidermal Growth F in Head and Neck Cancer: First H Antitumor Mechanisms Stephen Y. Lai, Priya Koppikar, Sufi M. The Seethala, Barton F. Branstetter, William E. Vivian W.Y. Lui, Dong M. Shin, Sanjiv S. Ag Myers, Jonas T. Johnson, Gordon Mills, Att J. Clin. Oncol., Mar 2009; 27: 1235 - 1242 in Lipofectamine 2000 (Invitrogen counted on the Nexcelom automated The cell proliferation experiments were	Factor Receptor Antisense DNA Therapy Juman Application and Potential omas, Erin E. Childs, Ann Marie Egloff, Raja R. Gooding, Ashok Muthukrishnan, James M. Mountz, garwala, Rita Johnson, Larry A. Couture, Eugene N. hanassios Argiris, and Jennifer R. Grandis . Carlsbad, CA). After 24 hours, viable cells were ell counter (Nexcelom Bioscience, Lawrence, MA). independently performed four times	<ul> <li>Abstract</li> <li>Full text</li> <li>PDF</li> <li>View Citation Map</li> <li>Find more like this</li> </ul>			
(\$40 for 14 days)	INFLAMMATION: Neisseria gonorrhoeae Activates the Signaling Activities of the NL Joseph A. Duncan, Xi Gao, Max Tze-Han H Stephen B. Willingham, Daniel T. Bergstra PY. Ting J. Immunol., May 2009; 182: 6460 - 6469 sample) and counted using the au Bioscience) according to the manufactur viable cells was counted using the Next from infected cells	the Proteinase Cathepsin B to Mediate RP3 and ASC-Containing Inflammasome Muang, Brian P. O'Connor, Christopher E. Thomas, Ih, Gary A. Jarvis, P. Frederick Sparling, and Jenny D. Mutomated Cellometer Auto T4 (Nexcelom rers protocols. Cellblue and the percentage of celom Cellometer Auto T4. B, Culture supernatants	<ul> <li>Abstract</li> <li>Full text (\$10)</li> <li>PDF (\$10)</li> <li>extra: Data Supplement</li> <li>View Citation Map</li> <li>Find more like this</li> </ul>			

![](_page_16_Picture_0.jpeg)

![](_page_16_Picture_1.jpeg)

#### **Current Customers:**

- Cellometer is used by hundreds of customers all over the world
- Biotech, Pharmaceutical, University, Academic and Government Research labs

![](_page_16_Figure_5.jpeg)

Discovering new boundaries

![](_page_17_Picture_0.jpeg)

![](_page_17_Picture_1.jpeg)

### **Product Comparison**

![](_page_17_Picture_3.jpeg)

	Auto T4/M10	Auto X4	Vision	
Counting Cell Lines			duo	trio
Cell line concentration				
Trypan blue viability				
Cell size measurements				
Using Flourescence to Count Primary Cells				
Primary Cells-Live/Total cell concentration	lu -			
Primary Cells- Viability				
Fluorescence Population Assays				2
Viability with fluorescence stains- Cell Lines				
GFP transfection rates				
Annexin-V Apoptosis				
Specialty Cells (Adipocytes, Hepatocytes)				

![](_page_18_Picture_0.jpeg)

![](_page_18_Picture_1.jpeg)

### Thanks for your attention !

![](_page_18_Picture_3.jpeg)