



JAPAN MADE QUALITY

Disposable Hemocytometer (4-Chambers)



Improved Neubauer Cell Counter

- Designed for rapid loading
- No washing, no labor, save time
- Same grid as standard one
- High calculation accuracy
- Made of durable and high grade plastic

More detail

https://www.funakoshi.co.jp/exports_contents/80019

Code	Product name	Unit
521-10	Disposable Hemocytometer (4-chambers)	20 slides / 1 case
		50 slides / 1 case
		200 slides / 4 cases



Improved Neubauer Cell Counter Counting Method for Culture Cells



[Sample Preparation]

- 1) Prepare cell suspension. When counting adherent cells, disperse cells by cell detachment solution such as Accutase or Accumax (Innovative Cell Technologies, Inc.) for accurate counting.
- Take 100 µL of cell suspension and transfer to another tube. Add 100 µL of prepared Trypan Blue (0.3 - 0.4%) solution to the tube. (This makes x 2 diluted solution).
 Note : Dilution rate should be optimized if cell numbers are high.
- 3) Pipet the solution gently.

[Counting]

1) Take 10 µL of trypan blue stained cell suspension. Inject the solution from sample inlets slowly.

Note : High speed injection may cause leak to other chambers.

- 2) Count all cells on compartment ① ④
- If the cells are on border lines, count those on 2 border lines only. (Red line, see Fig.2)
- · Count Live cells (Unstained) and / or Dead cells (Stained)
- 3) Calculate cell numbers
 - < Cell numbers in 1 mL >
- Live cells = (All live cells / counted compartments) x dilution rate x 10^4
- Dead cells = (All dead cells / counted compartments) x dilution rate x 10⁴
- Viability (%) = Live cells / All cells (=Live cells + Dead cells) x 100
- [Tips and Notes]
- For accurate counting, adjust cell numbers to 100 500 cells / 1 mm².
- Empty chamber may have small objects, but this does not affect to count.
- This is reference only. Detail counting method or for more information, please see experiment protocol book.