

Supporting Information

Antibiograms in five pipetting steps: precise dilution assays in sub microliter volumes with a conventional pipette

Ladislav Derzsi,^{‡a,b} Tomasz S. Kaminski^{‡a} and Piotr Garstecki^{*a}

a - Institute of Physical Chemistry, Polish Academy of Sciences, Kasprzaka 44/52, 01-224 Warsaw, Poland, E-mail: garst@ichf.edu.pl

b - University of Padova, Department of Physics and Astronomy, Via Marzolo 8, 35131 Padova, Italy

[‡] equal contribution.

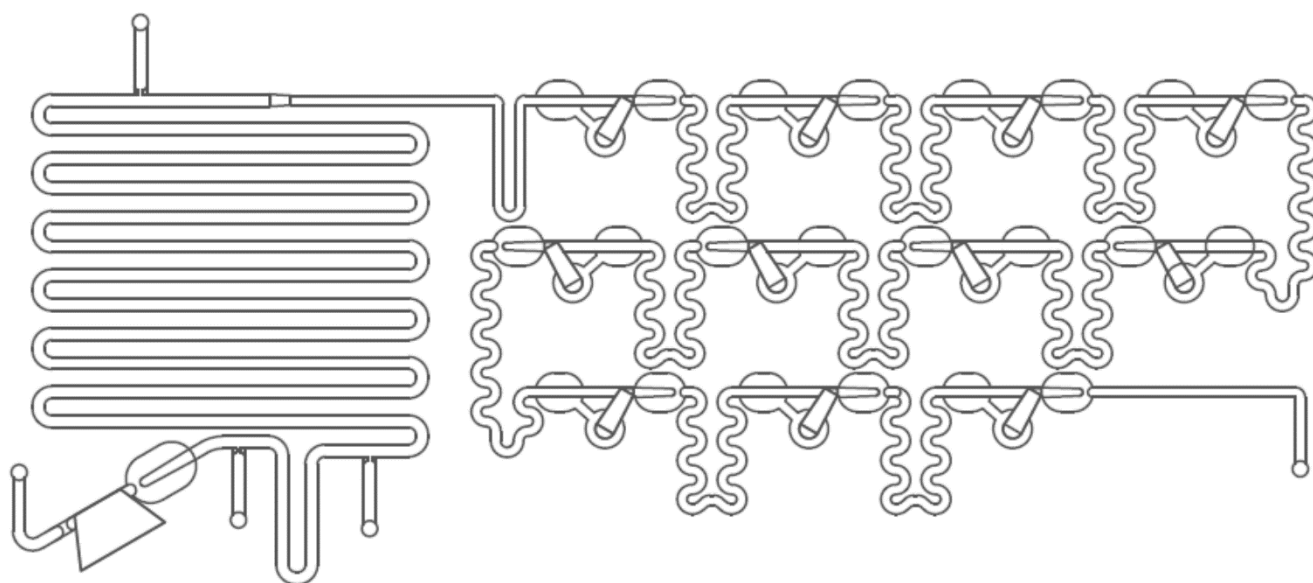


Fig. S1. Schematics of the microfluidic chip that we used for preparation of the gradient and determination of MIC.

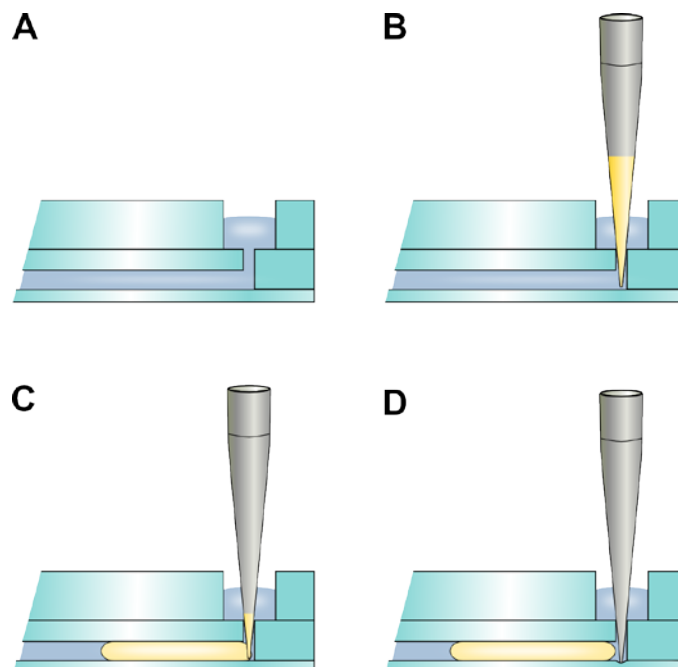


Fig.S2. A scheme depicting process of deposition of a sample into microfluidic channel with a automatic pipette.

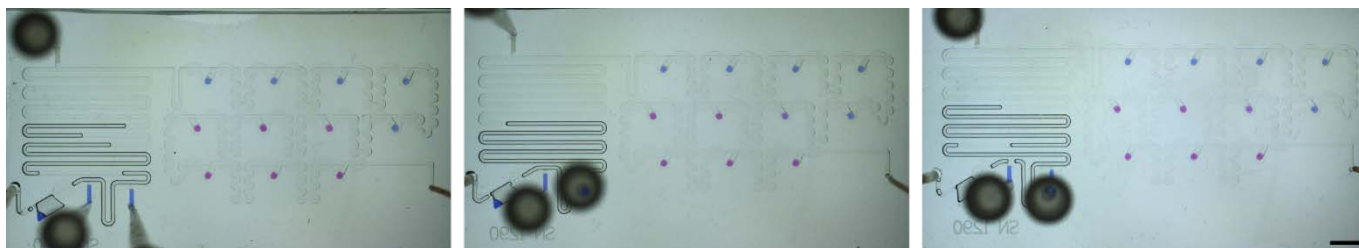
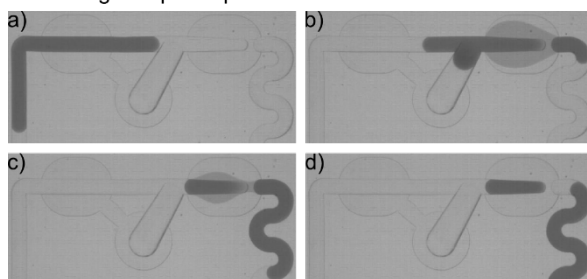


Fig.S3. Snapshots of 3 different chips depicting results of bubble formation during incubation. Scale bar is 5 mm.

I. Metering of 2 μ L droplet



II. Metering of 7 μ L droplet

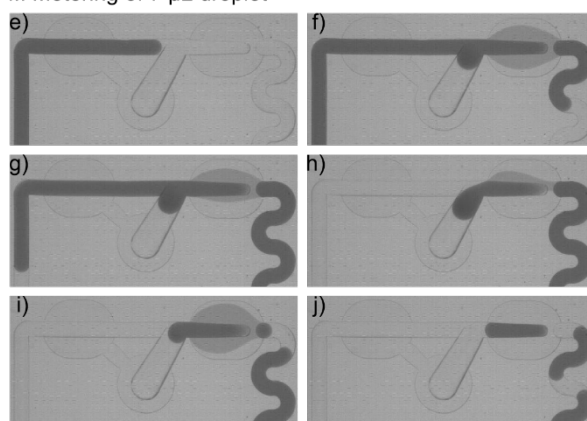


Fig. S4. Snapshots depicting the metering of small (a-d) and large (e-j) plugs . The large 7 μ L plug breaks before it flows entirely through the metering trap.