

# QUANTITATIVE STUDY OF THE CONTACT BETWEEN B CELLS AND ANTIBODIES-FUNCTIONALIZED OIL DROPLETS USING MICROFLUIDIC TRAPS

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In the immune system, B cells have the function to produce antibodies after the binding of B cell receptors with antigens presented at the surface of other cells, like dendritic cells for example [1]. This specific recognition leads to B cell spreading and antigen accumulation at the contact of the two cells [2]. In previous works, the kinetic of the antigen accumulation has not been fully studied [3]. Moreover, the B cell mechanical forces implied in the recognition processes are not known.

In our work, we used antibodies-functionalized oil droplets as antigen presenting cells. As liquid objects, they have the advantage of allowing antibody accumulation at their surface and potentially deformation, permitting force measurement. We put them into contact with B cells using a dedicated microdevice with adapted microfluidic traps to see the « time zero » of contact and also to increase the statistics of contact. Results from the kinetic studies and perspective on future developments will be presented.

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[3] C. Ketchum, H. Miller, W. Song, A. Upadhyaya, Ligand Mobility Regulates B Cell Receptor Clustering and Signaling Activation, **Biophys. J.** 106 (2014) 26–36.

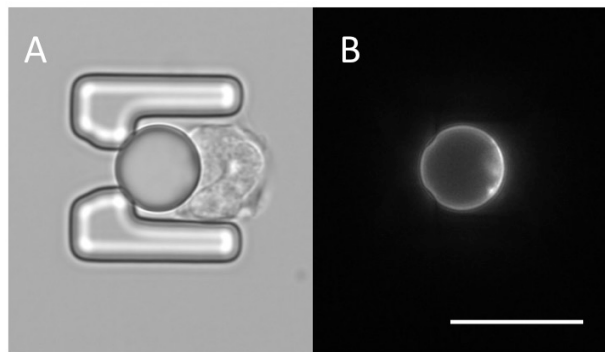


Image showing a B cell in contact with an antibody-functionalized oil droplet in a microfluidic trap. The trap has been first loaded with the functionalized oil droplet and the B cell has been loaded in second. The time of the image is 7 minutes after contact. (A) Microscope image in bright field. (B) Microscope image in fluorescence of the droplet showing an accumulation of streptavidin at the contact between the droplet and the cell. (Bar=20 $\mu$ m)