## INTERNSHIP PROPOSAL

## Mechanics of T lymphocytes

Laboratory name: Laboratoire d'Hydrodynamique (LadHyX), Ecole polytechnique	
CNRS identification code: UMR 7646	
Internship director'surname: Julien HUSSON	
e-mail: julien.husson@polytechnique.edu	Phone number: +33 1 69 33 52 82
Web page: http://cellmechanics.jimdofree.com	
Internship location: Laboratoire d'Hydrodynamique, Bâtiment Pôle Mécanique, Ecole	
polytechnique, Palaiseau, France	
Thesis possibility after internship: YES	
Funding: YES	If YES, which type of funding: ANR

## Title

Quantifying cell mechanical properties is important for better understanding various cellular processes. Focusing on the immune reaction at the single-cell level, we have shown that when a white blood cell (or leukocyte) contacts another cell to attack it or exchange information, the former becomes much stiffer and more viscous within minutes. Studying these mechanical changes in immune cells can help scientists better understand the mechanism by which immune cells identify threats such as cancer cells. Many aspects of the mechanical properties of immune cells remain unclear and require further characterization. This internship will contribute to this characterization.

We develop tools to probe the mechanical properties of cells via microindentation, i.e. by pressing them with micrometric beads or needles. Among other parameters, we can measure the tension of the actomyosin cortex underlying the cell membrane, which can fluctuate over time. One question to be addressed is whether these fluctuations are used by leukocytes to sense their environment and whether they can help them to react faster to stimuli brought by other cells.

Another aspect of the internship is motivated by some of our preliminary data showing that pressing a T lymphocyte (or T cell, a leukocyte at the core of immune response) with an adhesive microsphere leads to an apparent higher stiffness of the leukocyte than when the microsphere is not adherent. This effect might be either a purely mechanical artifact due to adhesive vs. non-adhesive boundary conditions, or might be a signature of a rapid mechanical response of the cell to adhesion. We will address this question by producing sets of microbeads with controlled adhesive strength and by inhibiting specific molecular components of the cell cytoskeleton.

During this internship with a major experimental component, we will use our micropipette-based single-cell rheometer to perform cell microindentation to quantify the viscoelastic properties of white blood cells. Cell and molecular biology know-how will be provided by biologists through established collaborations.

 Please, indicate which speciality(ies) seem(s) to be more adapted to the subject:

 Condensed Matter Physics: NO
 Soft Matter and Biological Physics: YES

 Quantum Physics: NO
 Theoretical Physics: NO