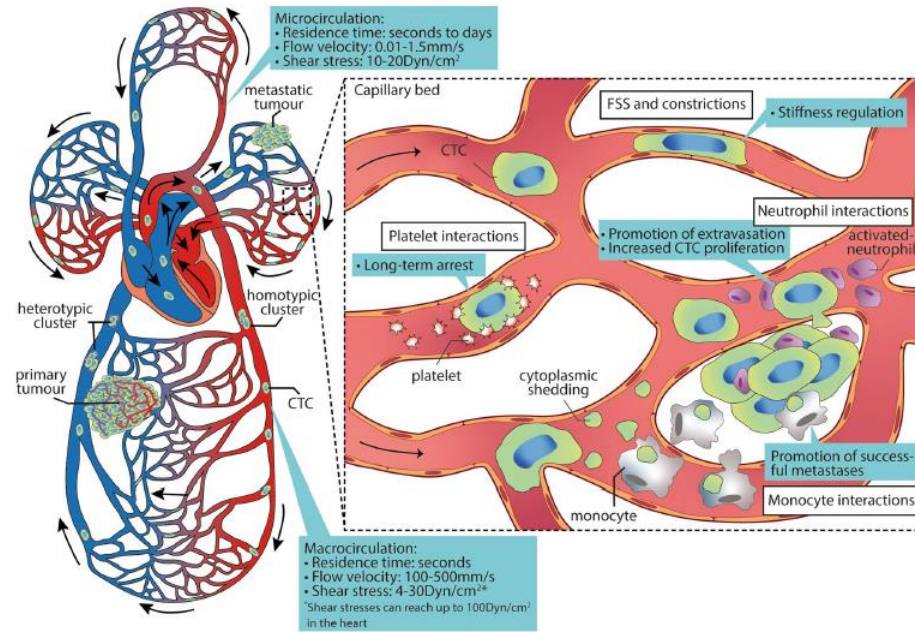
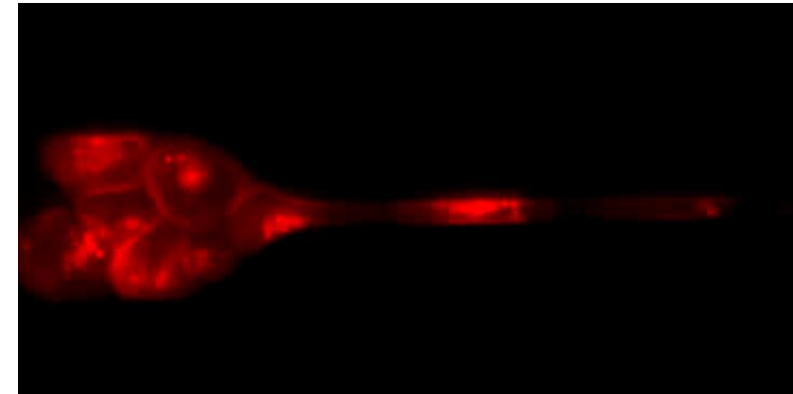
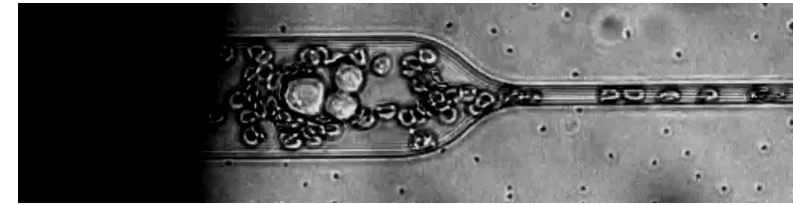


Metastasis

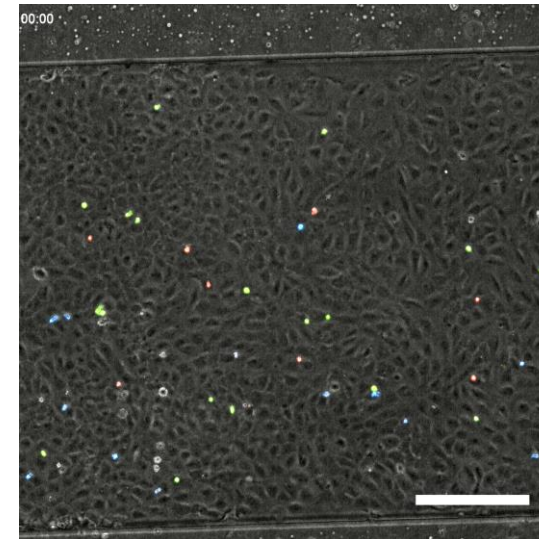
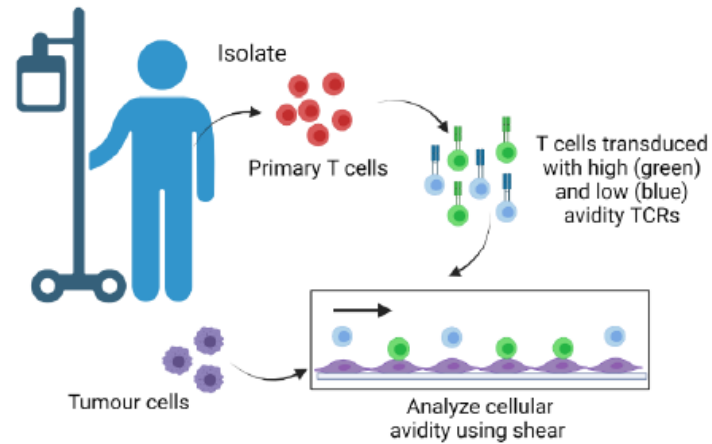


Vrynas, Paizal, Bakal & Au *Clin. & Exp. Met.* 2021



Au et al. *PNAS* 2016

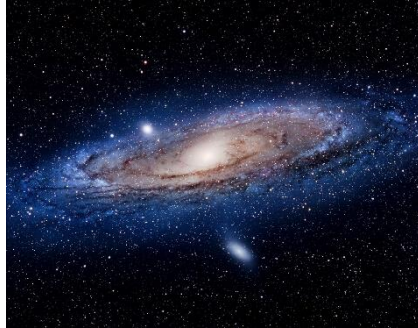
Drug Screening & Immunotherapy



Ashby et al. *Advanced Healthcare Materials* In Press

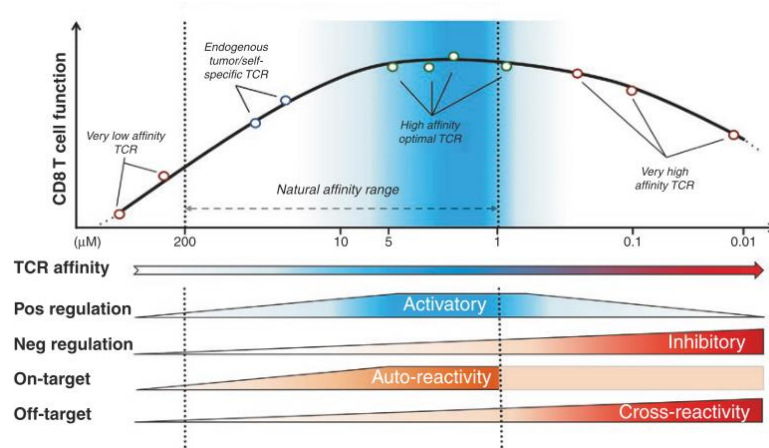
Microfluidic T Cell Selection by Cellular Avidity

TCR selection is challenging



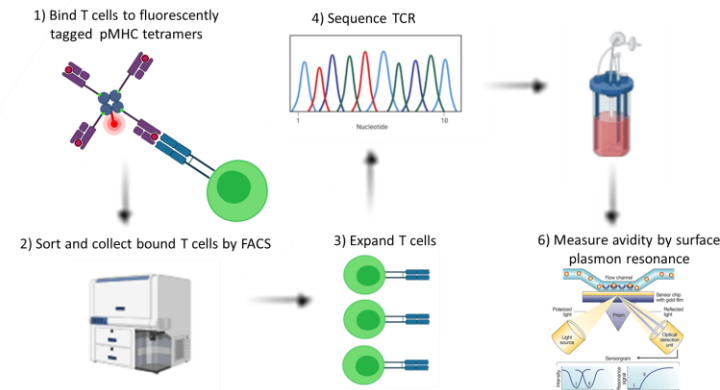
Nearly 1 trillion distinct TCRs in humans

Binding strength “Avidity” may aid selection of TCRs that provoke optimal immune responses



Schmid et al. *J. Immunology* 2017

Current methods are inefficient & not cell-cell



Julian Ashby
(Imperial)



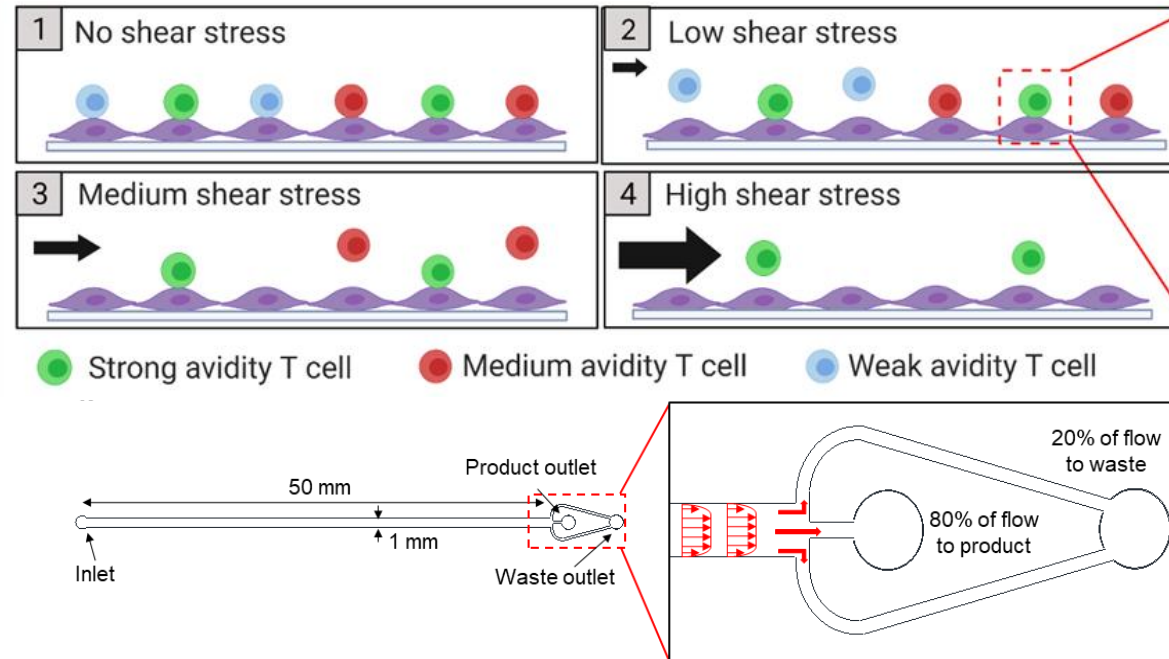
Li Tang
(EPFL)



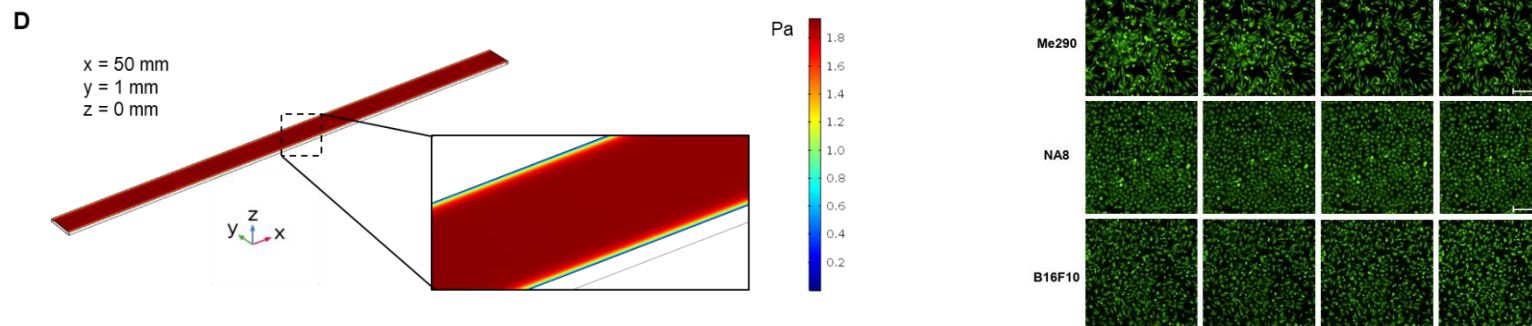
Julien Schmidt/Alexandre
Harari (CHUV – Lausanne)

Can Microfluidics Accelerate T Cell Selection in Solid Tumours?

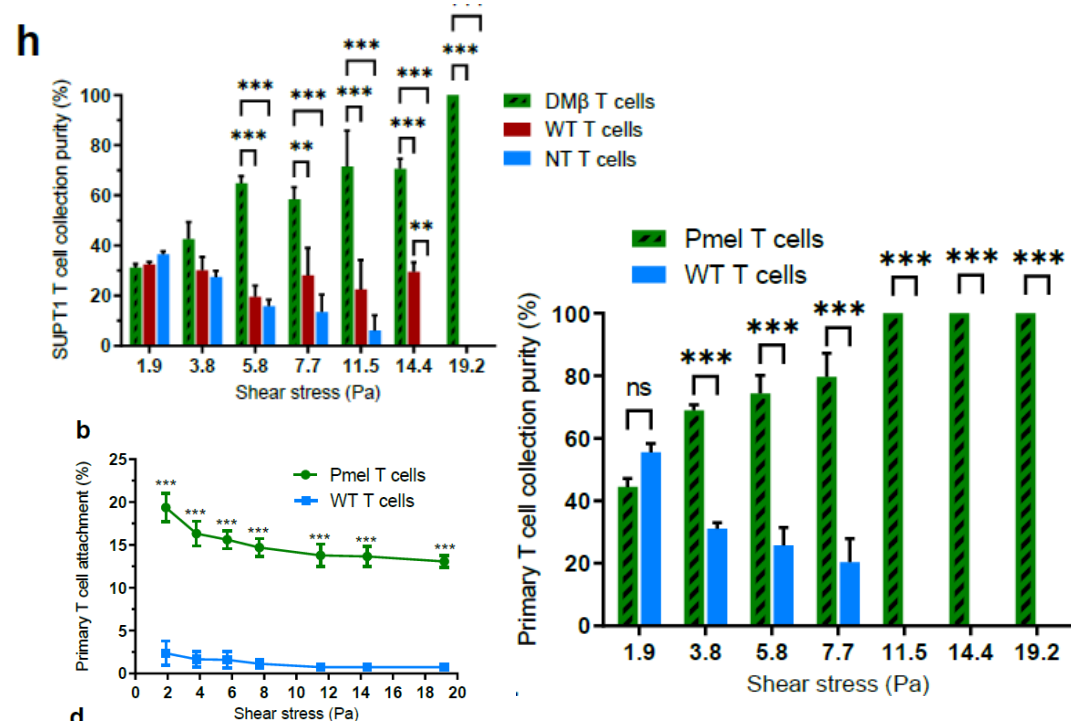
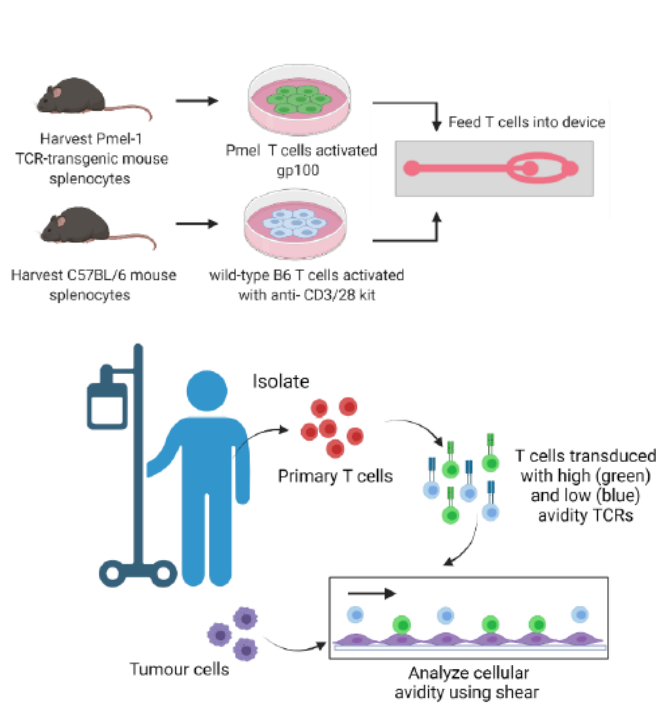
Fluid Shear Stress Based Selection



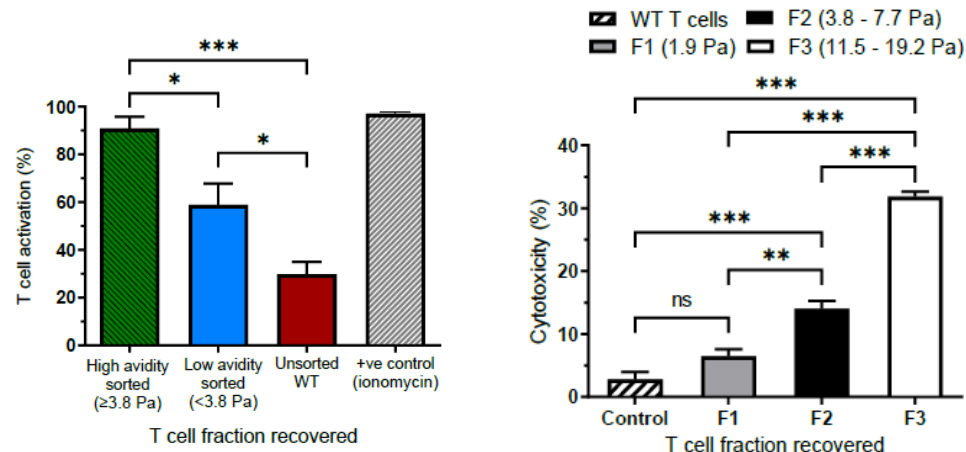
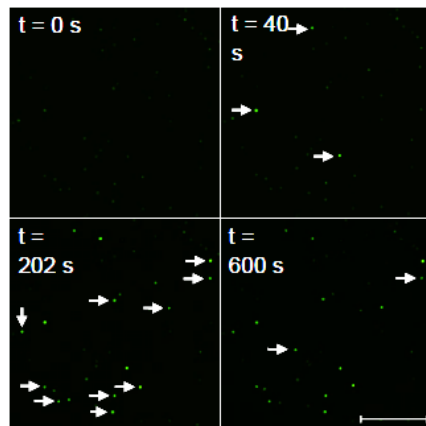
Requires: ~Uniform Shear Stress Distribution & Strong Tumour Cell Adhesion



Collects high avidity T cells w/ up to 100% Purity

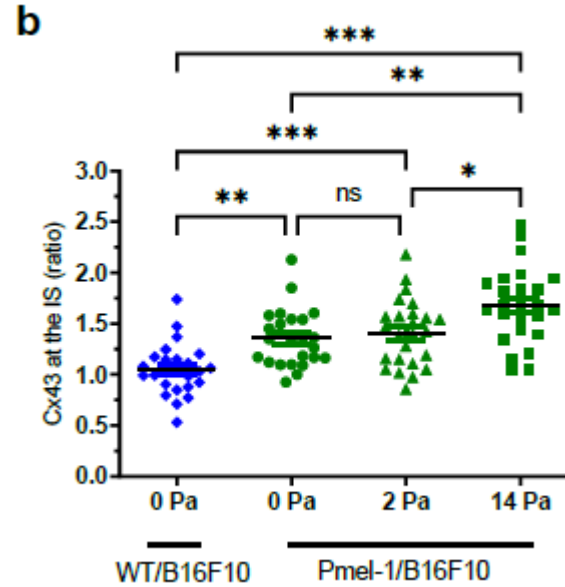
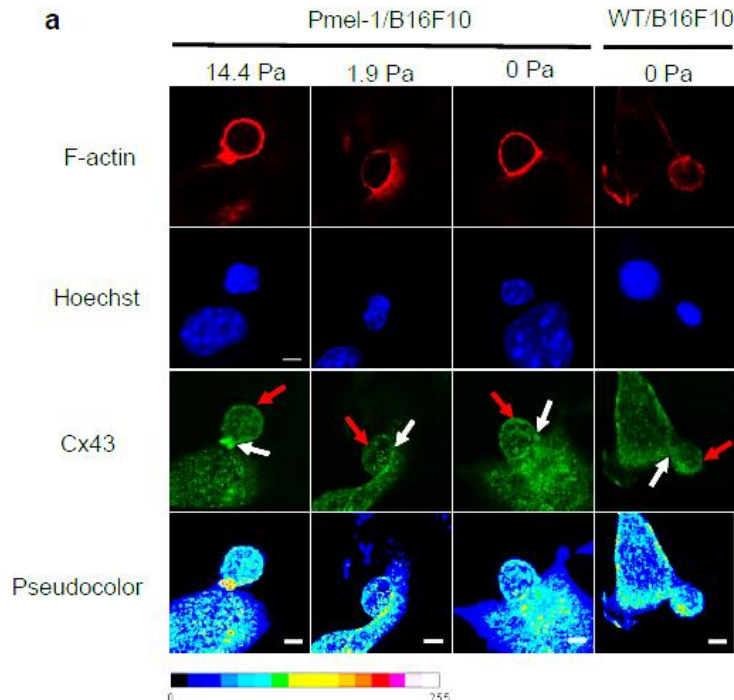


High avidity fractions provoke greater T cell activation & cytotoxicity



Holistic “Cellular Avidity” selection

Unlike multimer methods, probe all cell-cell interactions not just TCR-pMHC



High shear fractions have greater immunological synapse protein expression

- Selects high functional T cells by physiological cell-cell interactions
- Can probe up to 10,000 interacting cell pairs per device in <1 hour
- Recovers live-cell fractions based on avidity w/ up to 100% purity
- Compatible with supplemental fluorescent, luminescent & functional on-chip assays

Microfluidic opportunities in Pancreatic Cancer:

- Immunotherapy
- Drug screening & delivery
- Cancer cell migration
- Stromal interactions
- Tumour evolution & heterogeneity
- Dormancy & metastasis
- Tumour-on-chip & organ-on-chip models



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