# Precision Immunology

Co-Chairs: Holden T. Maecker, PhD – Stanford University Medical Center Drew M. Pardoll, MD, PhD – Johns Hopkins University School of Medicine

4:05 p.m. – 4:10 p.m.	Introduction Holden T. Maecker, PhD – <i>Stanford University</i>	
4:10 p.m. – 4:35 p.m.	Pre-existing Immunity and Treatment Outcome with Anti-PD1 Therapy in Melanoma Paul Tumeh, MD – <i>University of California Los Angeles</i>	
4:35 p.m. – 4:55 p.m.	Epitope Signatures and Melanoma Response to CTLA4 Therapy Timothy Chan, MD, PhD – <i>Memorial Sloan Kettering Cancer Center</i>	
4:55 p.m. – 5:15 p.m.	PD-1 Blockade in Mismatch Repair Deficient Tumors Luis A. Diaz, MD – Johns Hopkins Sidney Kimmel Cancer Center	
5:15 p.m. – 5:30 p.m.	Dissecting the Tumor Micro-Environment in Triple Negative Breast Cancer Identifies a Mutually Exclusive Expression Pattern of the Immune Co-Inhibitory Molecules B7-H4 and PD-L1 Kurt Schalper, MD, PhD – <i>Yale University</i>	

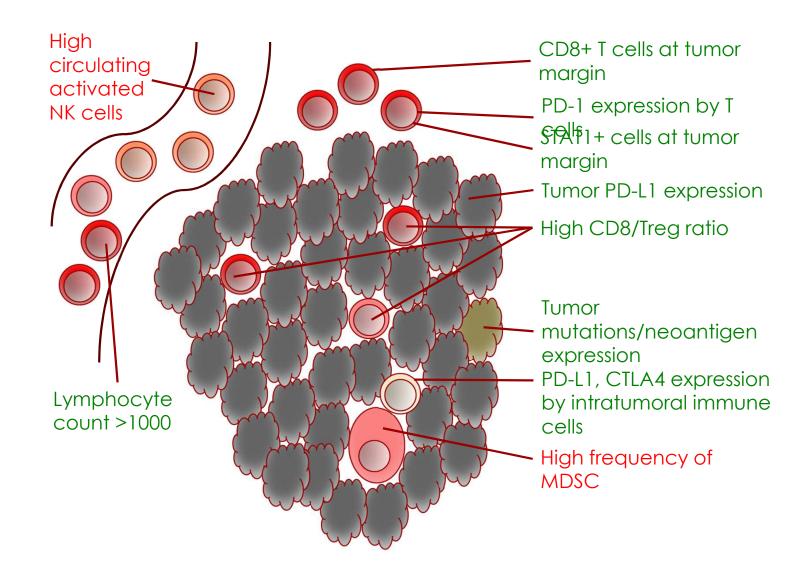
### What is Precision Immunology?

- Targeting the right therapy to the right group of patients, based upon immune metrics
  - Not one-size-fits-all
  - Not necessarily individualized medicine, but small enough groups to be precise

Same therapy for all patients	Precision therapy	Individual therapy for each patient
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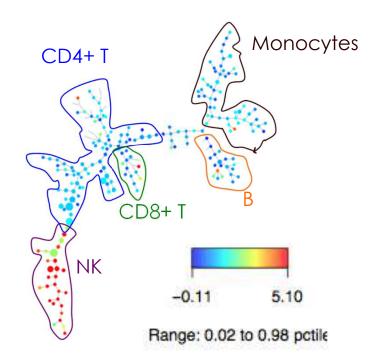
## Why measure the immune system?

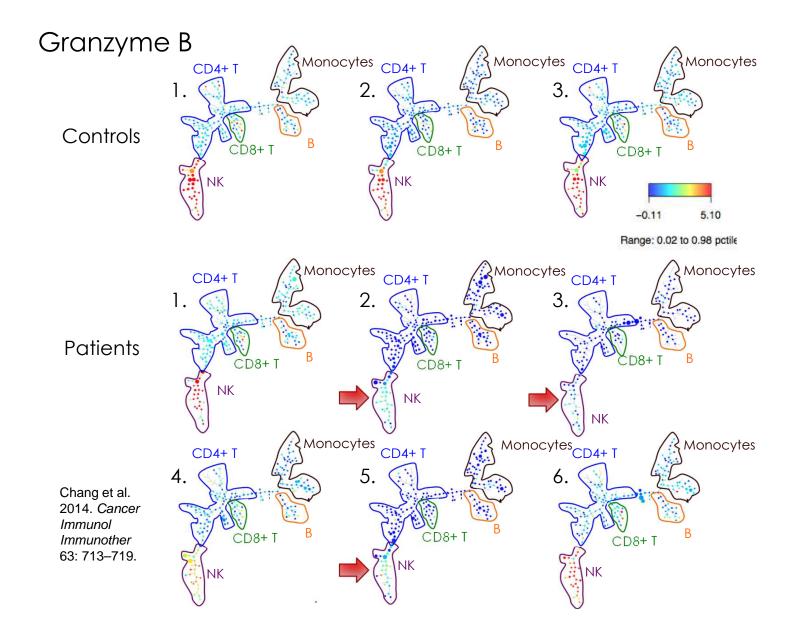
- Cancer patients are variably immunosuppressed (by their tumor, by prior therapy)
- Immunotherapies rely on the immune system to produce a response, yet we don't measure the relevant features of each patient's immune system
  - Global immunocompetence: PBMC subsets and functions
  - Tumor-specific: Antigen-specific T cells (tumor neoepitopes), tumor microenvironment features



### Looking for baseline predictors of response in PBMC

- Anti-EGFR (Cetuximab) as a targeted therapy in Kras<sup>wt</sup> head and neck cancer (Holbrook Kohrt)
- PBMC at baseline stimulated with PMA+ ionomycin
- CyTOF mass cytometry with 38 marker panel
- Clustering and display of data using SPADE





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