



# Center for Biologics Evaluation and Research ADVANCING REGULATORY SCIENCE



## Novel Machine Learning Technique Identifies Populations of Stimulated Mesenchymal Stromal Cells with Immunosuppressive Activity

A technique developed at FDA for computer-assisted analysis of thousands of images of stimulated human mesenchymal stromal cells (MSCs) identifies morphological characteristics that could serve as quality control markers during large-scale manufacturing of immunosuppressive cell therapies.

**“Morphological profiling using machine learning reveals emergent subpopulations of interferon- $\gamma$ -stimulated mesenchymal stromal cells that predict immunosuppression”**

*Cytotherapy, 2019;21:17-31*

Ross A. Marklein<sup>1,2</sup>, Matthew W. Klinker<sup>1</sup>, Katherine A. Drake<sup>3</sup>, Hannah G. Polikowsky<sup>3</sup>, Elizabeth C. Lessey-Morillon<sup>1</sup>, & Steven R. Bauer<sup>1</sup>

<sup>1</sup>Division of Cellular and Gene Therapies, Center for Biologics Evaluation and Research (CBER), US Food and Drug Administration, Silver Spring, Maryland, USA,

<sup>2</sup>School of Chemical, Materials, and Biomedical Engineering, University of Georgia, Athens, Georgia, USA, and <sup>3</sup>Cytobank, Inc., Mountain View, California, USA

### The potential medical benefits of immunosuppressive MSCs

Treatment of immune-system-based disorders

- **Therapies:** Immune system disorders, such as graft-versus-host disease, type 1 diabetes, and multiple sclerosis.
- **Tissue and organ transplants:** Preventing immune rejection of tissue grafts and organ transplants.

### The problem this work addresses

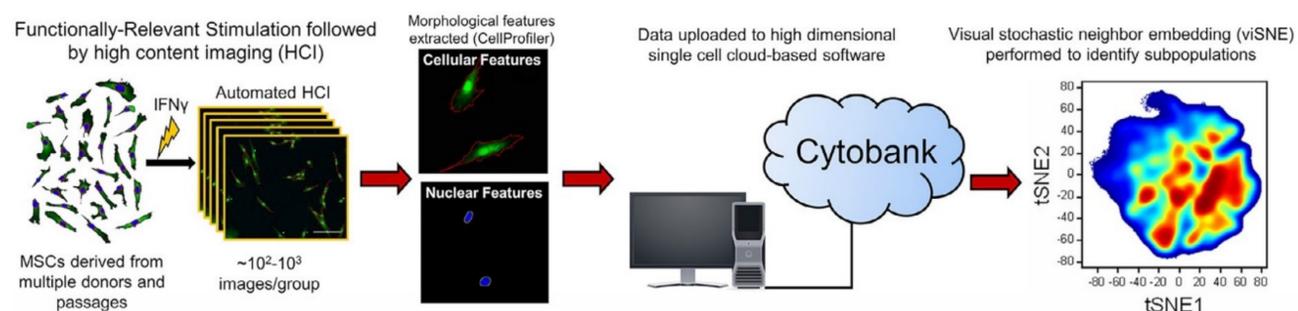
Functional heterogeneity is a barrier to many promising cell therapies

- **Functional heterogeneity:** Donor-to-donor and intra-population variation in the ability of MSCs to suppress immune system activity after being stimulated in vitro with Interferon-gamma (IFN- $\gamma$ ).
- **Wide variations in clinical trial outcome:** Outcomes from clinical trials using stimulated MSCs obtained from different human donors and grown in different bioreactors by different manufacturers vary widely, which complicates studies of potential MSC-based immunosuppressive therapies.

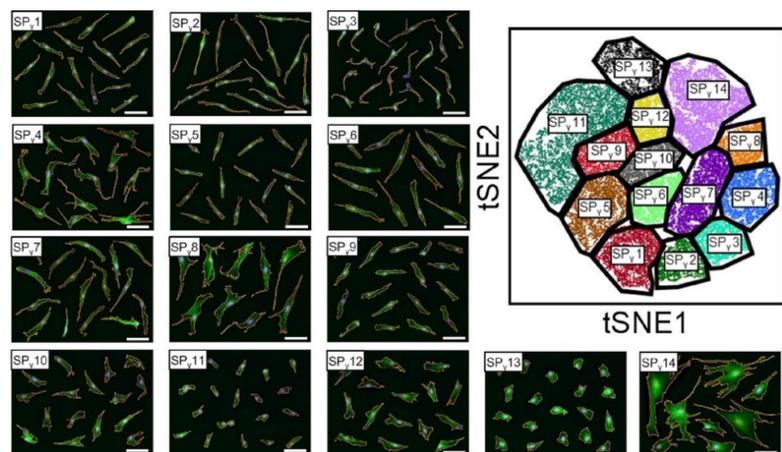
### One potential solution to the problem of functional heterogeneity

Functionally-relevant morphological profiling (FRMP) and Visual stochastic neighbor embedding (viSNE)

- **FRMP:** 1) Quantifies subcellular, single cell, or multicell morphological attributes of MSCs (e.g., cell perimeter, cell radius, nuclear perimeter; 2) identifies correlations between the number of cells in distinct morphological subpopulations and the degree of immunosuppression displayed by different preparations of MSCs.
- **viSNE:** Machine learning technique that identifies specific subpopulations of stimulated MSCs that share certain morphological attributes.
- **Quality control markers:** FRMP/viSNE enables identification of specific morphological attributes that predict whether any one of several populations of MSCs responded to IFN- $\gamma$  by developing the ability to suppress two key immune T cells, CD4+ and CD8+, before that ability is assessed in vitro.



Organizing cells based on single-cell morphological data. 1) Derive MSC lines from multiple donors and at different passages, stimulate the MSCs with 0, 10 and 50 ng/mL IFN- $\gamma$  for 24 h, and stain MSCs for cellular and nuclear features; 2) acquire hundreds of images for each cell line/passage/stimulation; 3) acquire high-dimensional single-cell morphological features data using custom Cell Profiler pipeline; 4) send data to cloud-based software (Cytobank); 5) analyze data using viSNE.



### How viSNE is used to find MSCs with immunosuppressive potential

Grouping MSCs according to morphological traits linked to immunosuppressive activity

- **Grouping MSCs according to shared morphological traits:** viSNE was used to group stimulated MSCs into 14 different clusters, based on which traits each cell shares with other MSCs in that cluster.
- **Linking morphological traits with immunosuppressive activity:** viSNE links each cluster of MSCs having shared morphological traits to the relative ability of those clusters to suppress CD4+ and CD8+ T cells. This identifies the sets of morphological characteristics that reflect immunosuppressive activity

‘Morphological catalog’ of IFN- $\gamma$ -stimulated MSCs highlights distinct morphological subpopulations.

### The potential role of FRMP/viSNE in stem-cell-based therapies

Supporting large-scale manufacturing by understanding functional heterogeneity

- **Facilitating large-scale production in bioreactors:** FRMP/viSNE could facilitate identification of immunosuppressive MSC preparations and optimize manufacturing methods.
- **Broader therapeutic uses of FRMP:** Identifying subpopulation and frequency of stem cells stimulated by other molecules to acquire therapeutic potential for treating a variety of disorders.

FRMP/viSNE could lead to better understanding of different MSCs obtained from different human donors and grown in different bioreactors by different manufacturers. Such understanding could support optimized production of safe and effective MSC-based immunosuppressive therapies. This strategy might also be successfully used

to assess how other types of stem cells respond to different growth factors, thus providing critical insights into how best to develop stem-cell-based therapies for a variety of diseases.