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# In-silico perturbation

## **Material**

NCBI dataset: https://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE90546

Paper Adamson B, Norman TM, Jost M, Cho MY et al. A Multiplexed Single-Cell CRISPR Screening Platform Enables Systematic Dissection of the Unfolded Protein Response. Cell 2016 Dec 15;167(7):1867-1882.e21. PMID: 27984733

### **Methods**

#### Gene chose

I chose 3 genes which show difference between *ATF6\_IRE1\_pMJ152* and *PERK\_IRE1\_pMJ154* to evaluate whether using tfs to reconstruct the genes is reliable.

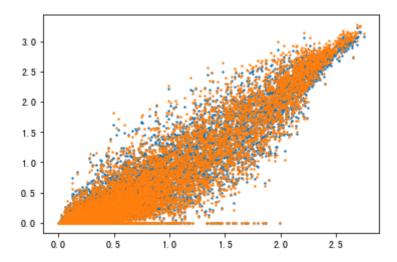
## **Result Visualization**

Test Dataset {'ATF6\_IRE1\_pMJ152', 'ATF6\_PERK\_IRE1\_pMJ158',

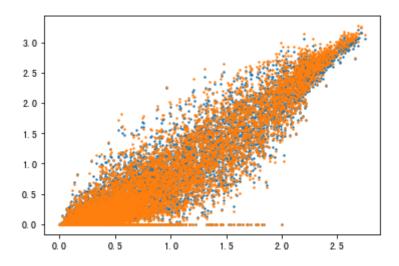
'ATF6\_PERK\_pMJ150',
'PERK\_IRE1\_pMJ154'}

We show the reconstruction of genes by transcription factors in an in-silico perturbation dataset. Two visualizations are included, one comparing predicted genes with real genes (blue), and the other comparing other processed genes with real genes (orange).

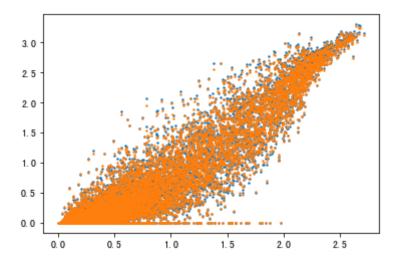
# PERK\_IRE1\_pMJ154



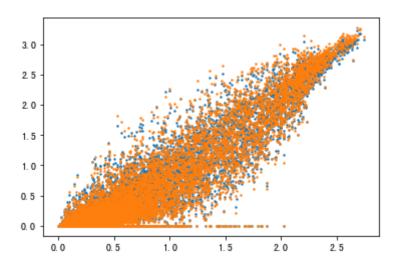
# ATF6\_PERK\_pMJ150



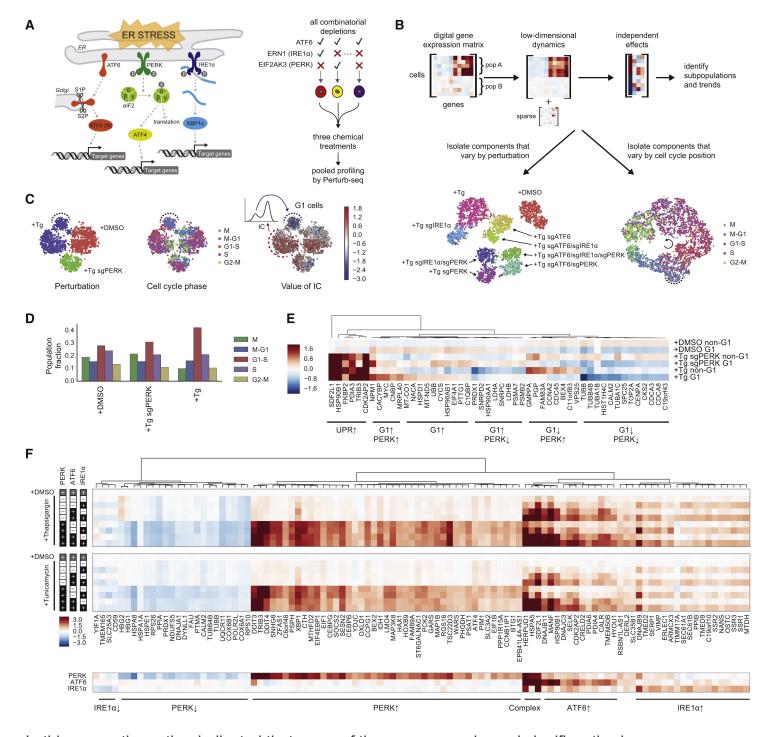
ATF6\_IRE1\_pMJ152



ATF6\_PERK\_IRE1\_pMJ158



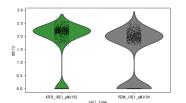
Gene

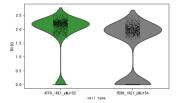


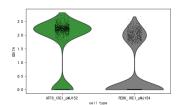
In this paper, the author indicated that some of the genes experienced significantly change.

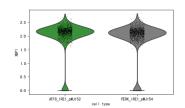
I chose ATF6\_IRE1\_pMJ152, PERK\_IRE1\_pMJ154 groups to find out whether the difference genes will be reconstructed.

#### Original

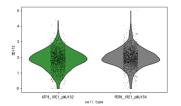


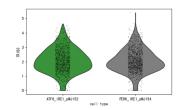


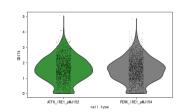


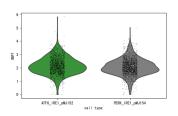


### **Predicted**

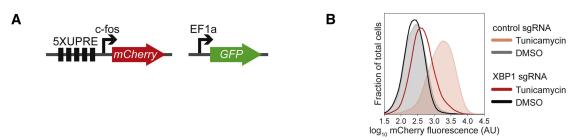


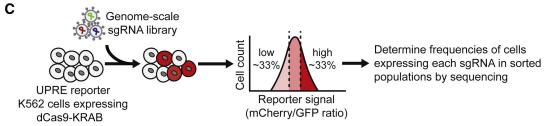


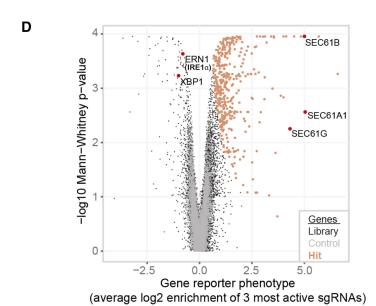


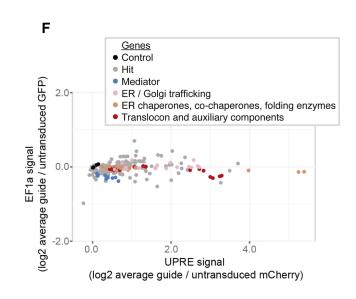


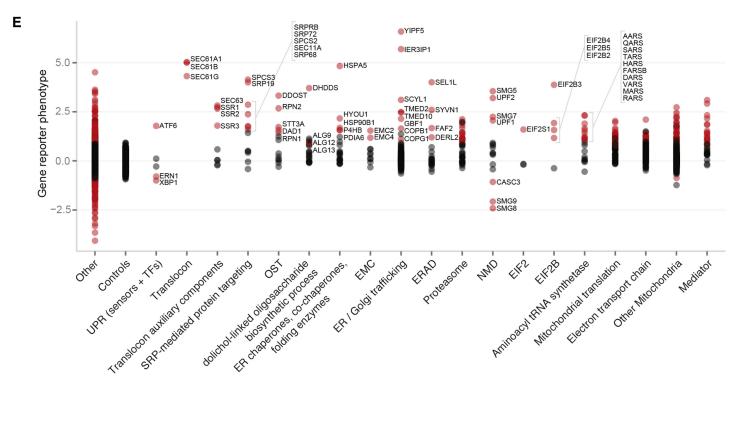
# sveral genes concerning IRE1











In the paper it said when the IRE1 was perturbed 3 main genes (**SEC61B SEC61A1 SEC61G**) will be suppressed, however it is not very obivious when I used the original data to construct the tsne map.

