## ■ 1 INIT

# 2 build\_data\_sets

## 2.1 read protein data

LFQ: rel. quantification – for same protein across different samples

iBAQ: orthogonal comparison – for different proteins in the same sample

Intensity: supposed to be both

C:\ProgramData\Anaconda3\lib\site-packages\ipykernel\_launcher.py:2: DtypeWarning: Columns (5,6,2764,2765,2773,2774,2775,2776) have mixed types. Specify dtype option on import or se t low\_memory=False.

### 2.2 read mrna data

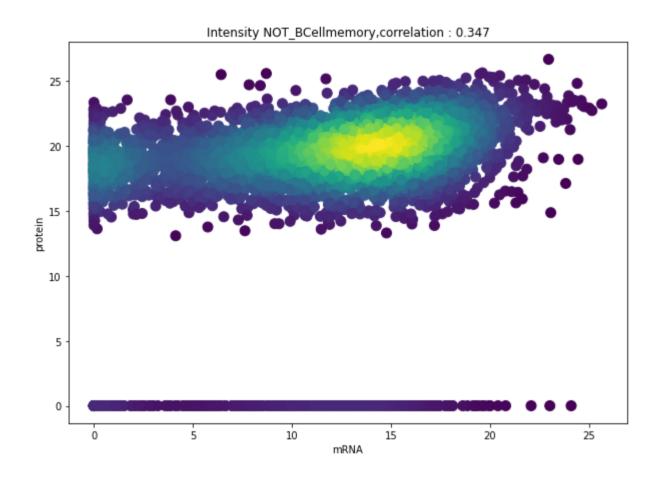
we start with mapping the cells annotations from the MassSpec to mRNA :

## 3 compare sets

## 3.1 compare all proteins to all genes

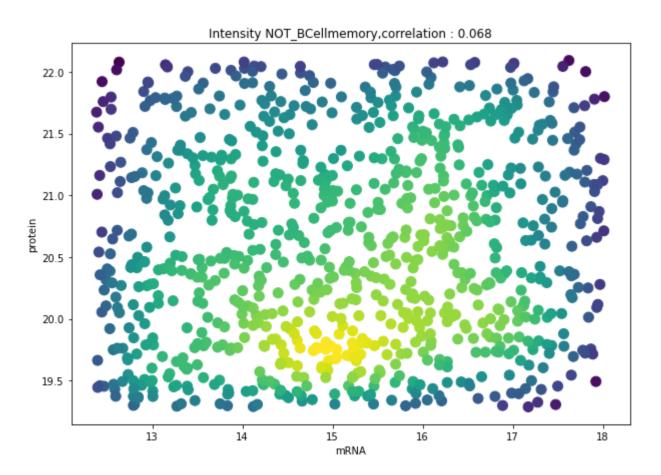
#### 3.1.1 med value of genes

Here we took the median value of each protein (between Jan versions) and compare them to the adjusted mRNA values :



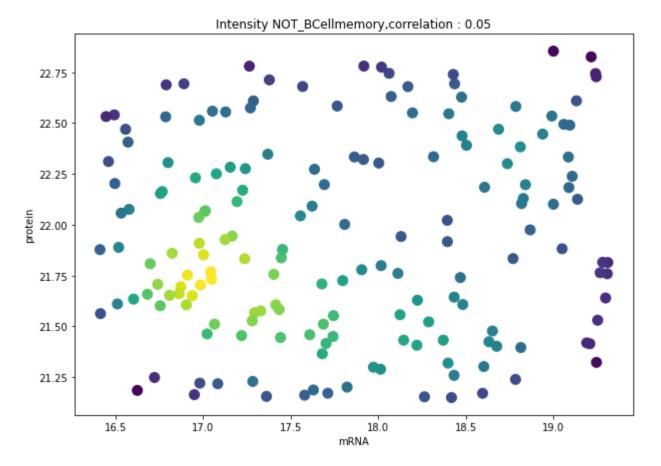
#### **▼** 3.1.2 med values + taking only percantile 0.5 to 0.9

Here we kept only high values (mostly remove zeros from proteins)

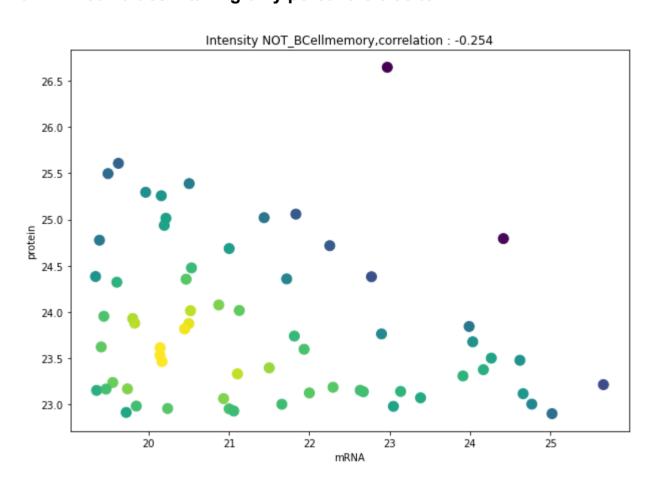


## ■ 3.1.3 med values + taking only percantile 0.95 to 0.8

4



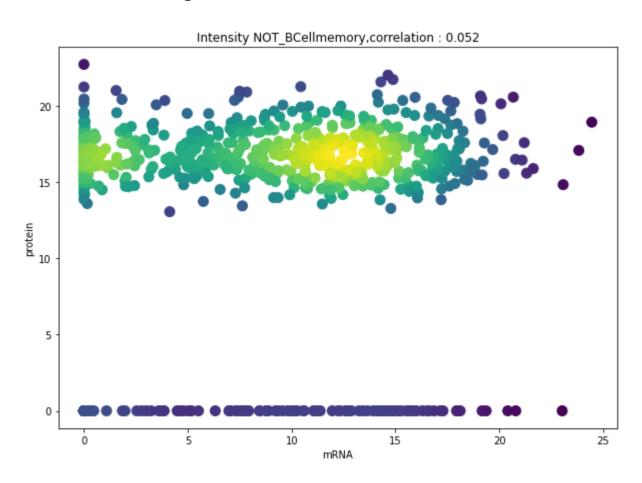
## 3.1.4 med values + taking only percantile 0.95 to 1



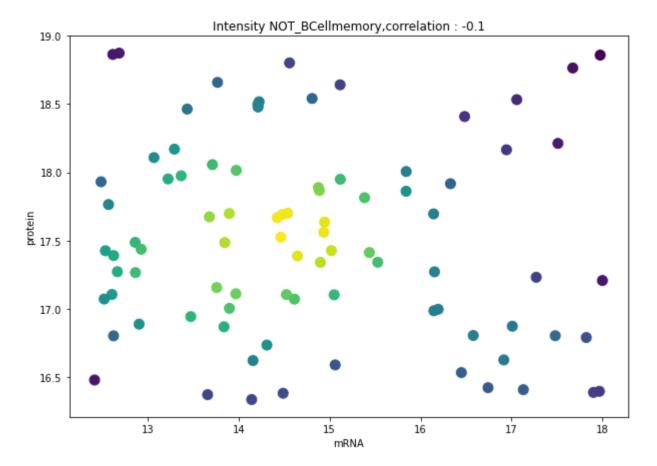
## **▼** 3.2 remove high STD proteins

Here we removed proteins we are "not sure about". aka - proteins where the std between versions is bigger then 0.8

## ▼ 3.2.1 med value of genes



### **▼** 3.2.2 med values + taking only percantile 0.5 to 0.9



## 3.2.3 med values + taking only percantile 0.99 to 0.7

