

AML10_Rx

jtrincado

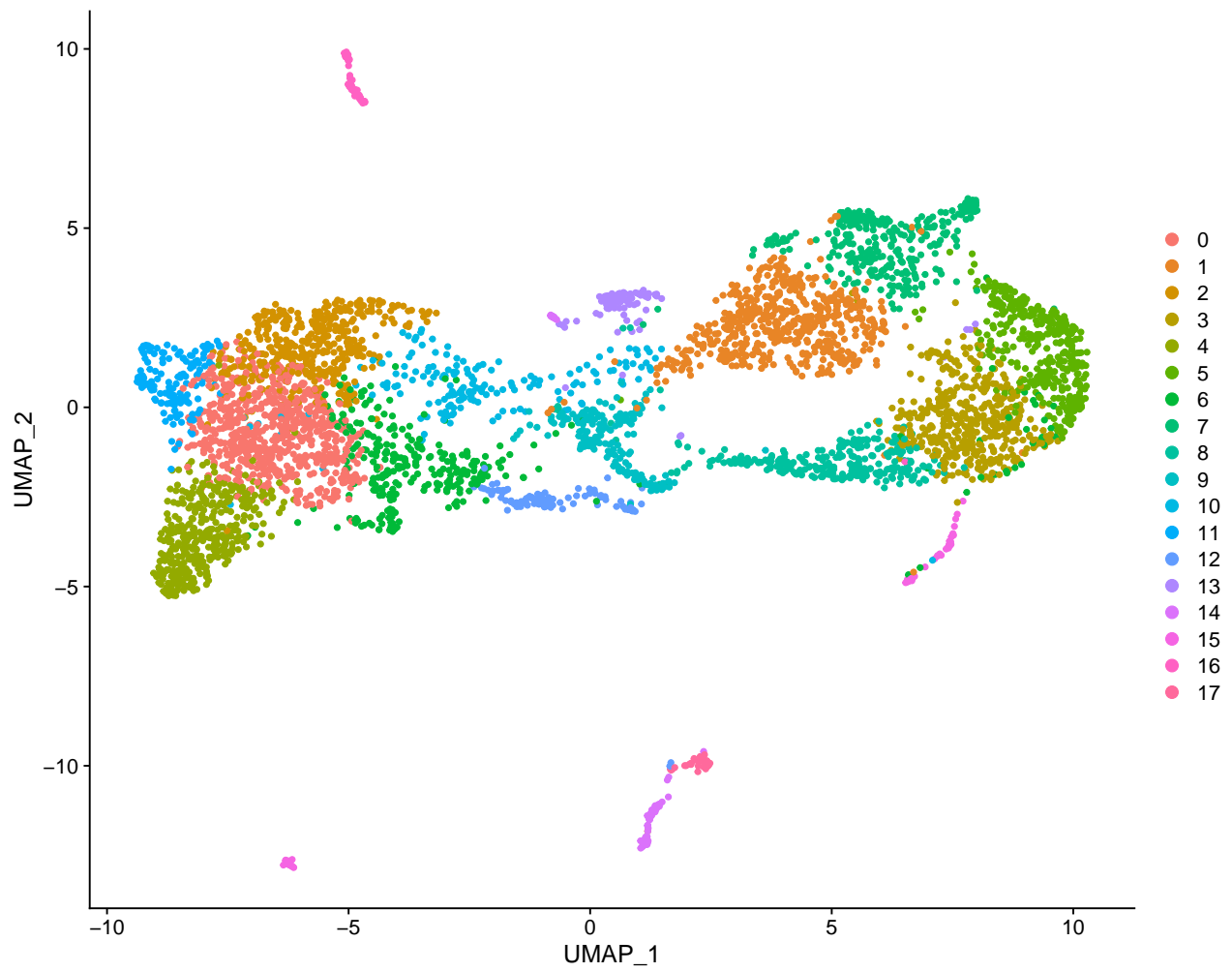
2022-02-09 10:58:54

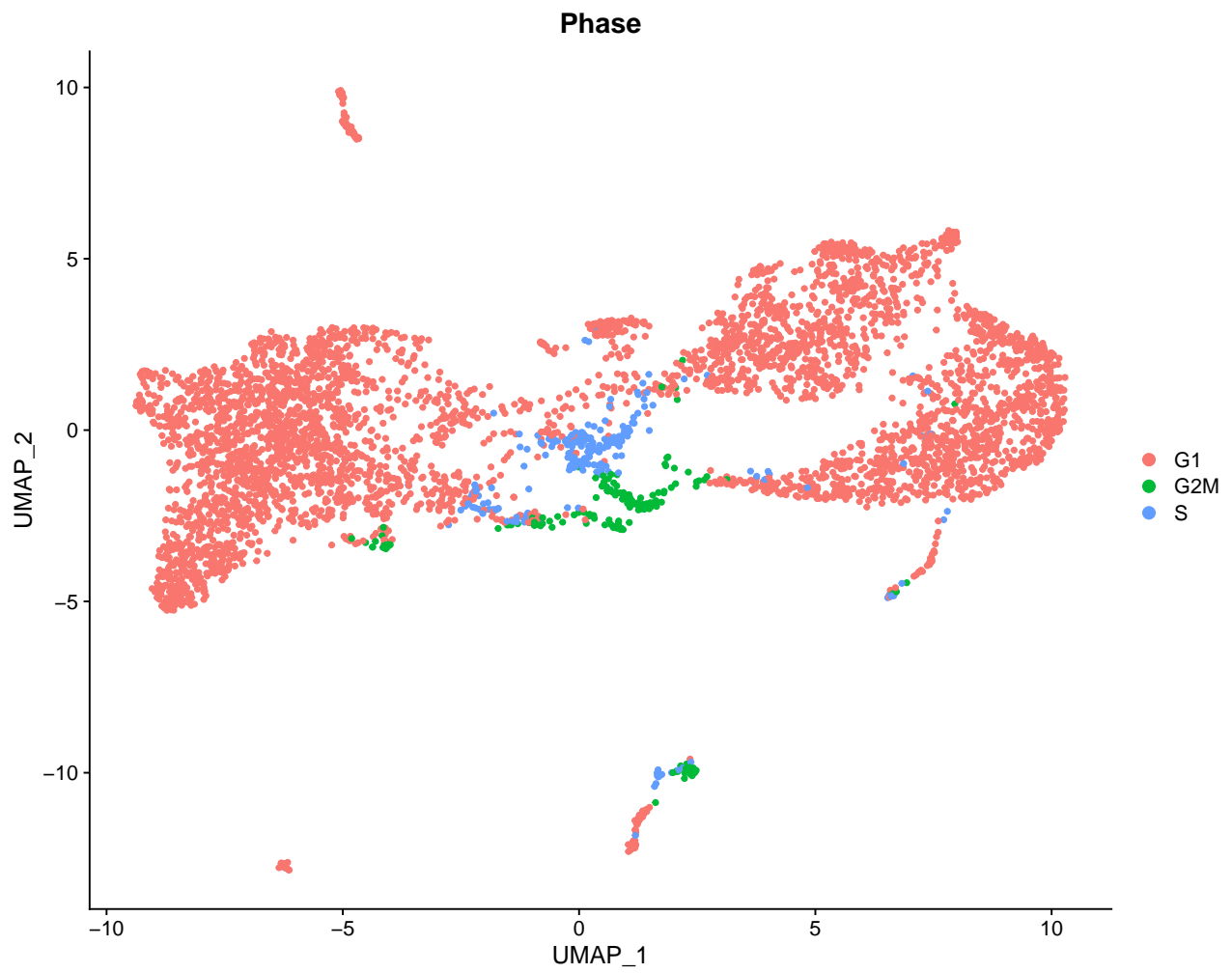
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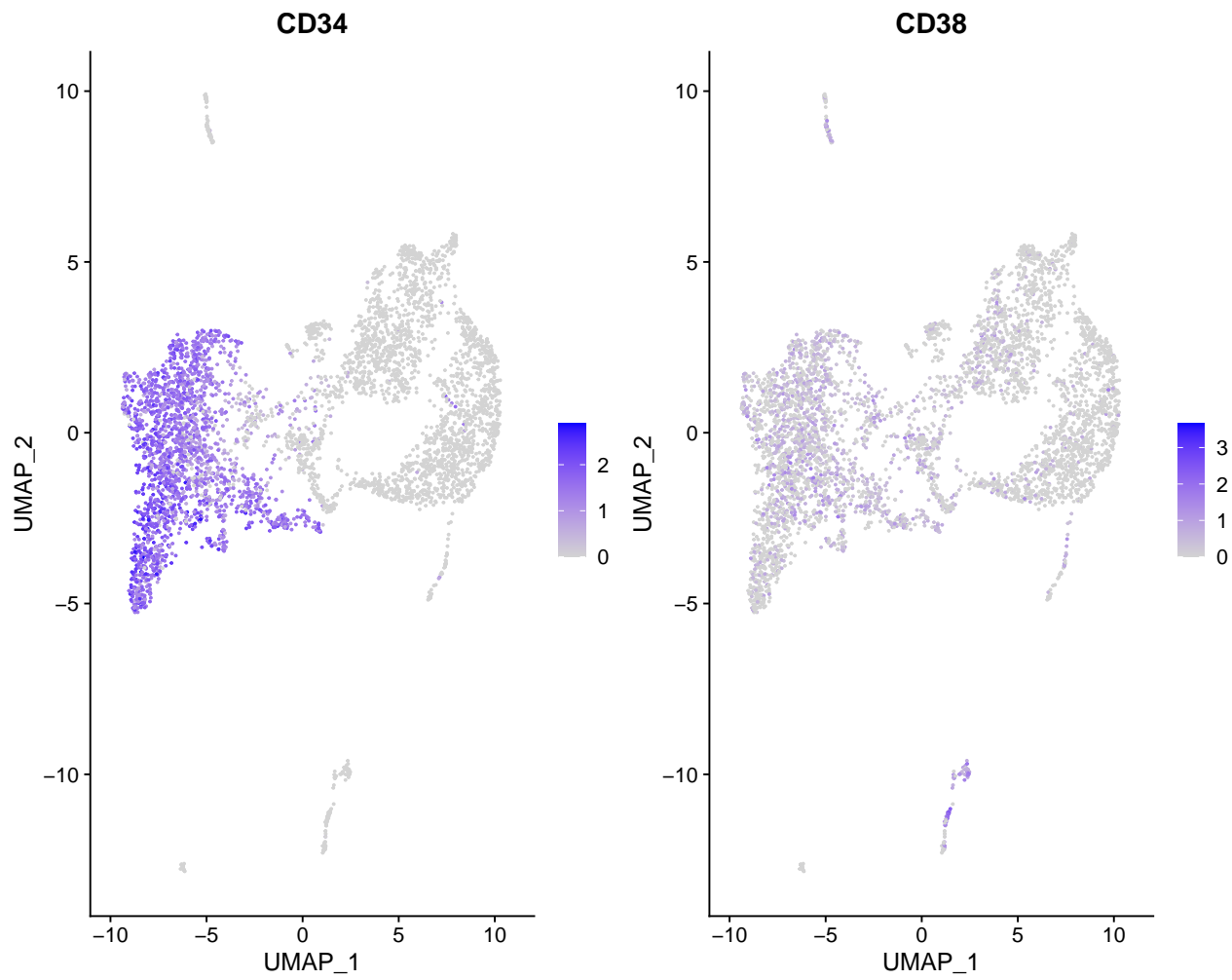
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1. Put together both 34 and 38 libraries. Apply QC and dimensionality reduction.

```
## Warning: The default method for RunUMAP has changed from calling Python UMAP via reticulate to the R  
## To use Python UMAP via reticulate, set umap.method to 'umap-learn' and metric to 'correlation'  
## This message will be shown once per session
```

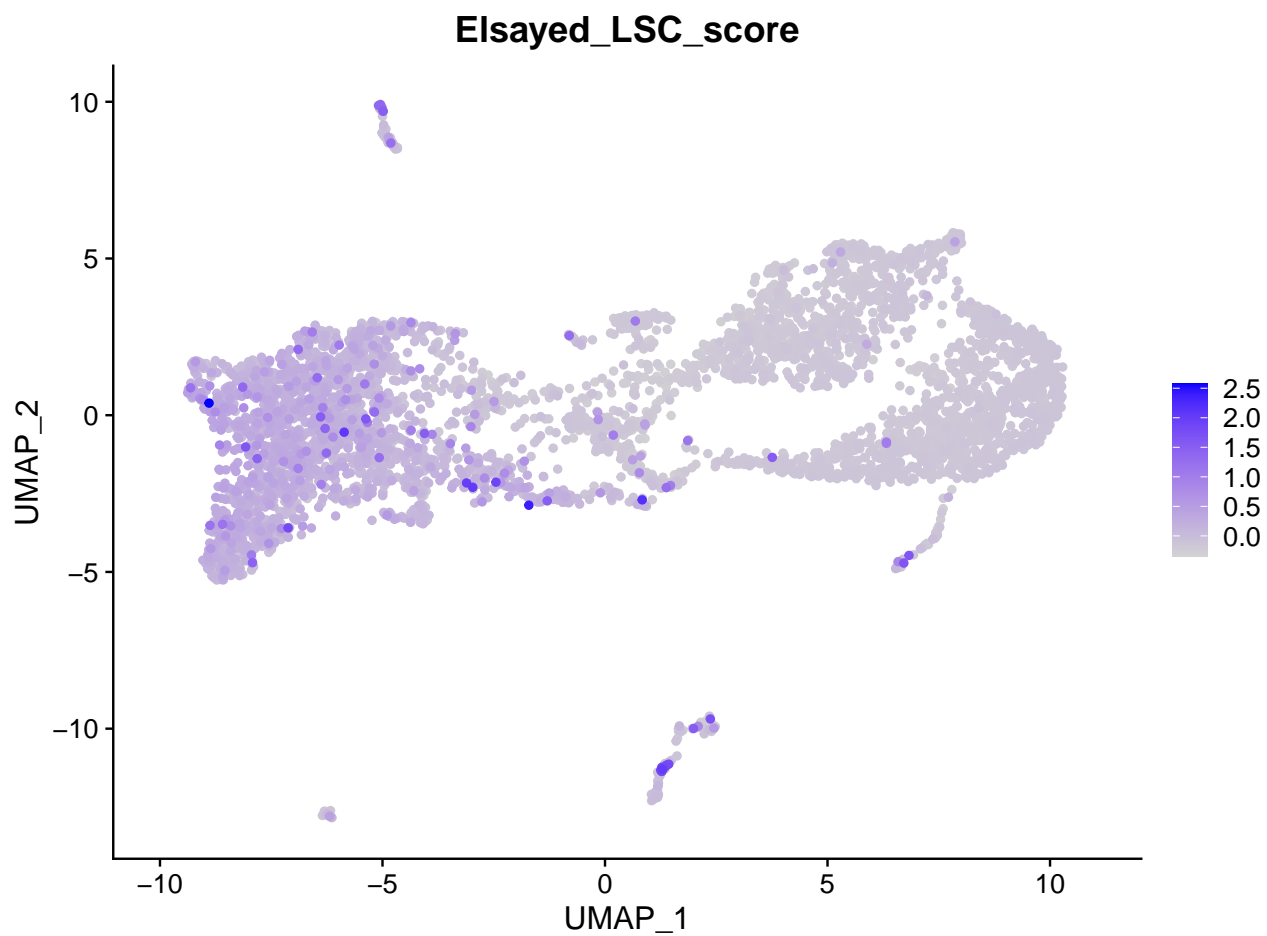


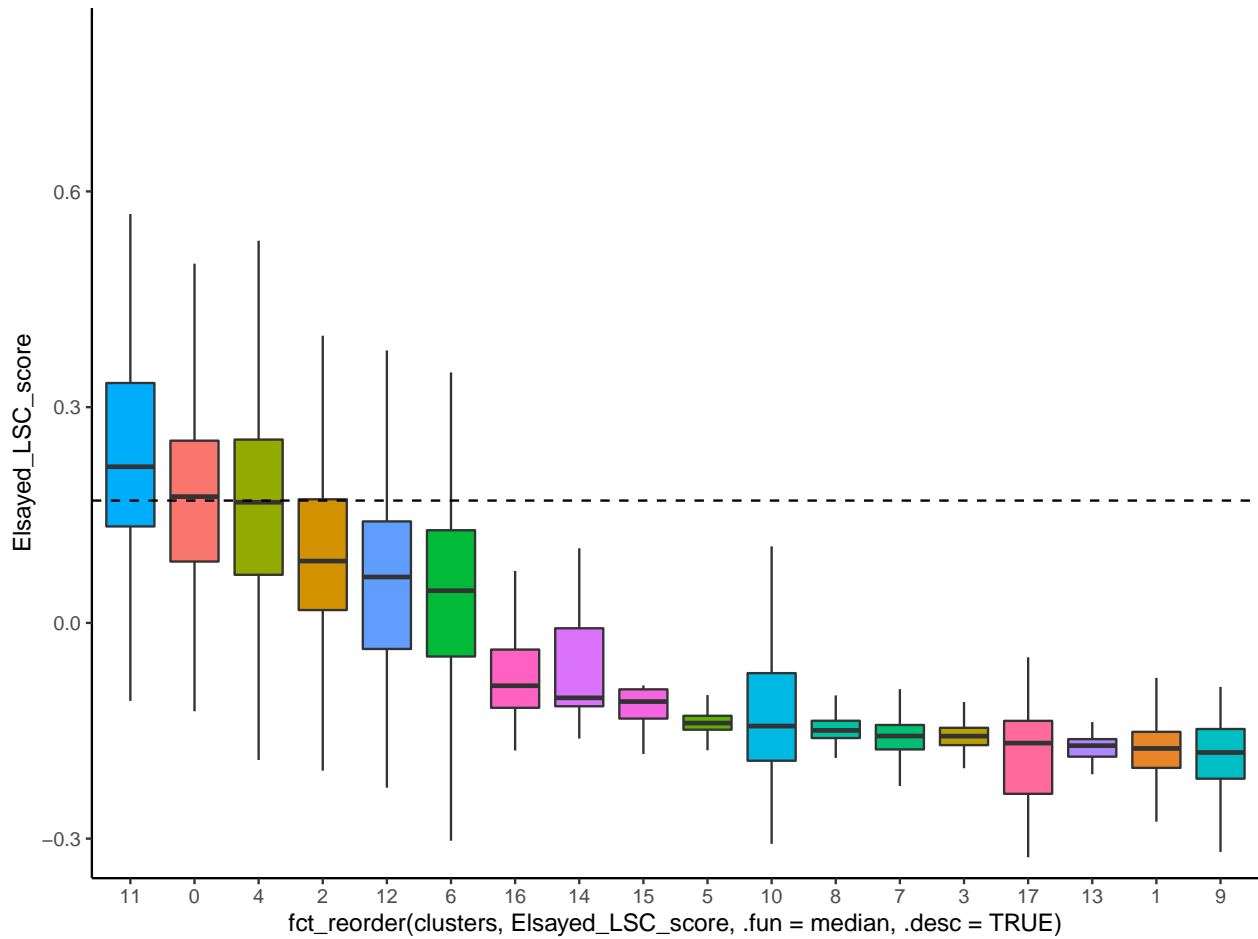




2. Get the LSC6 score

```
## [1] "CD34" "SPINK2" "SOCS2" "FAM30A" "ADGRG1" "DNMT3B"
```

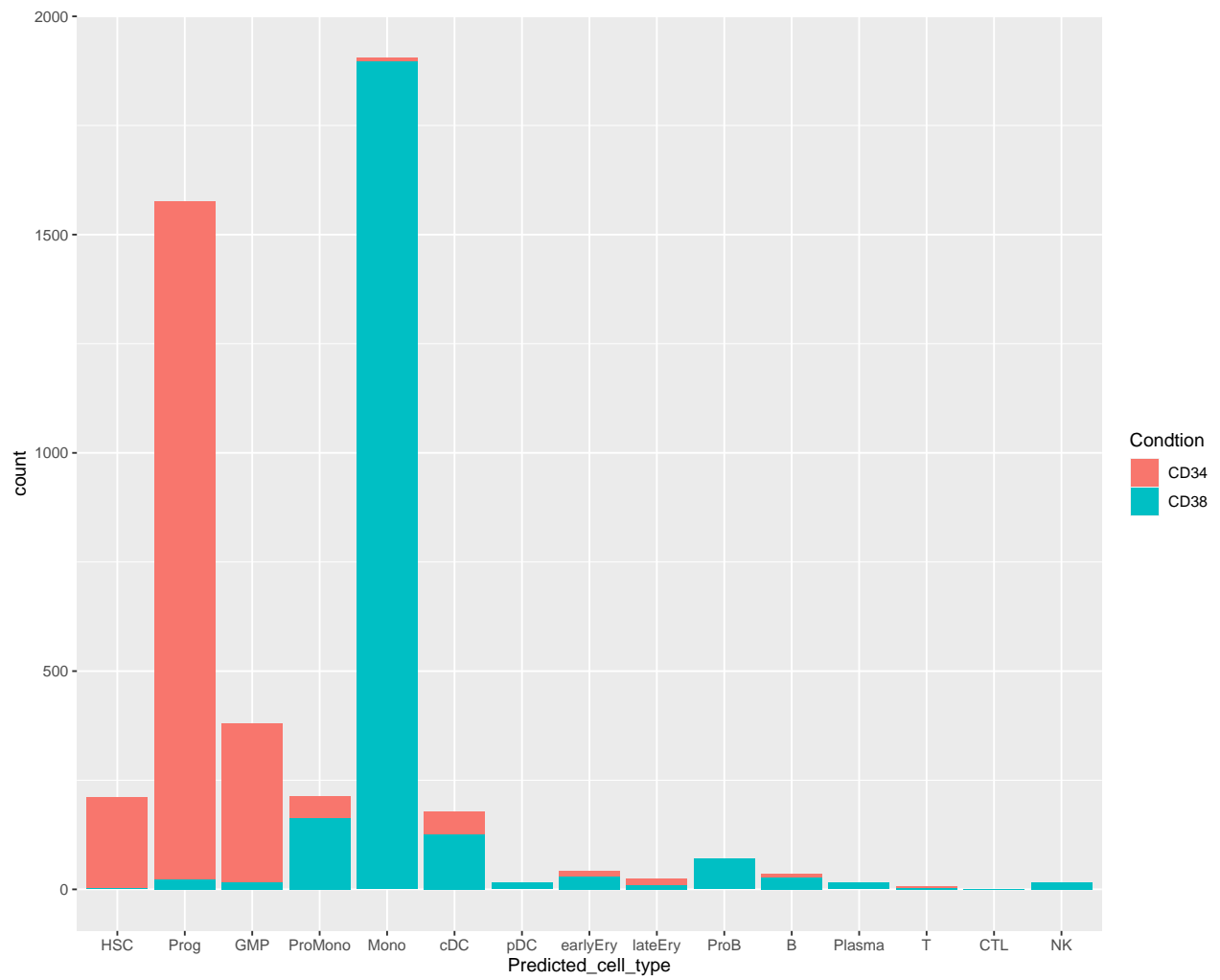


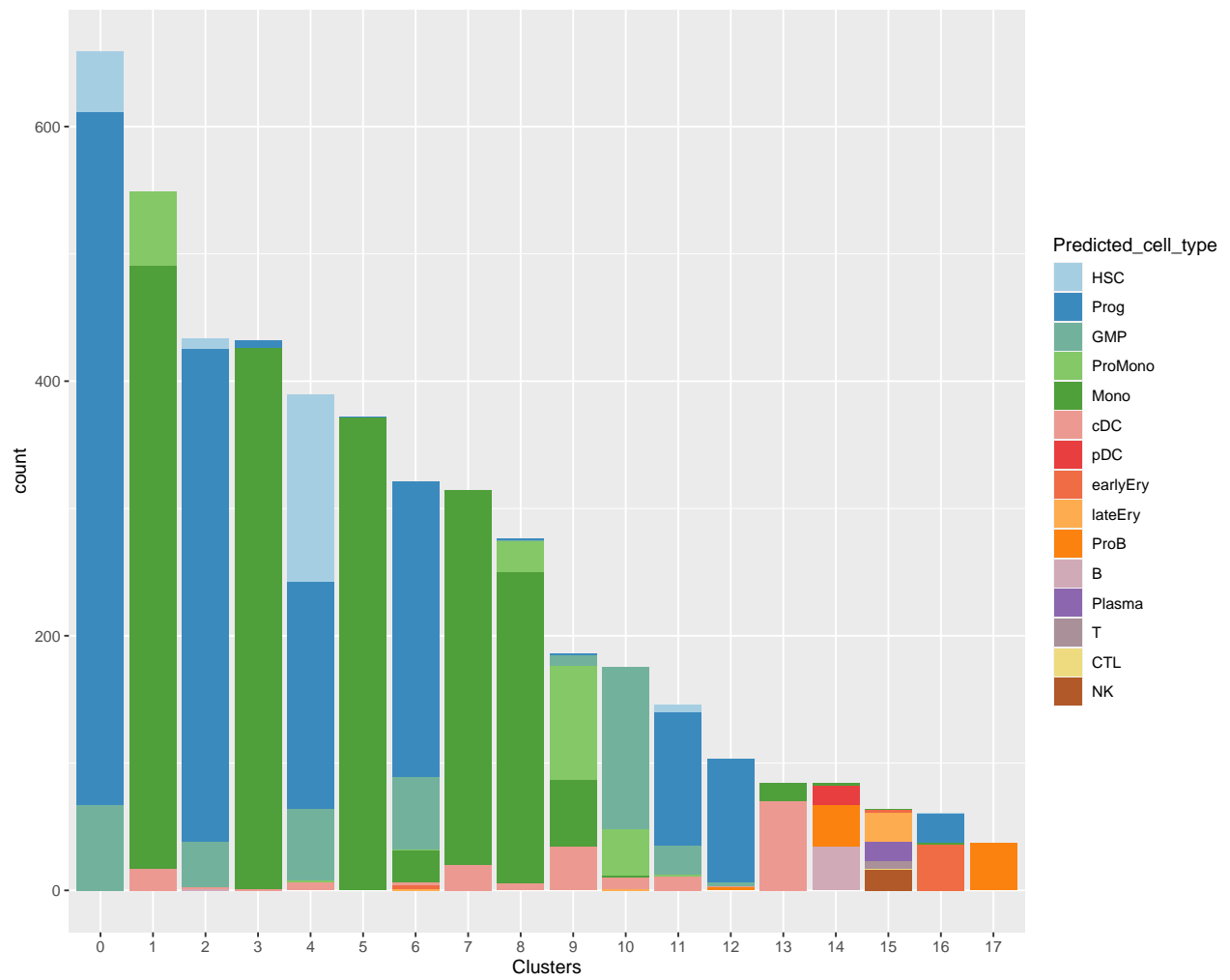


3. Predict the class of the cells using the markers and the expression of the BM cells form Van_Galen paper

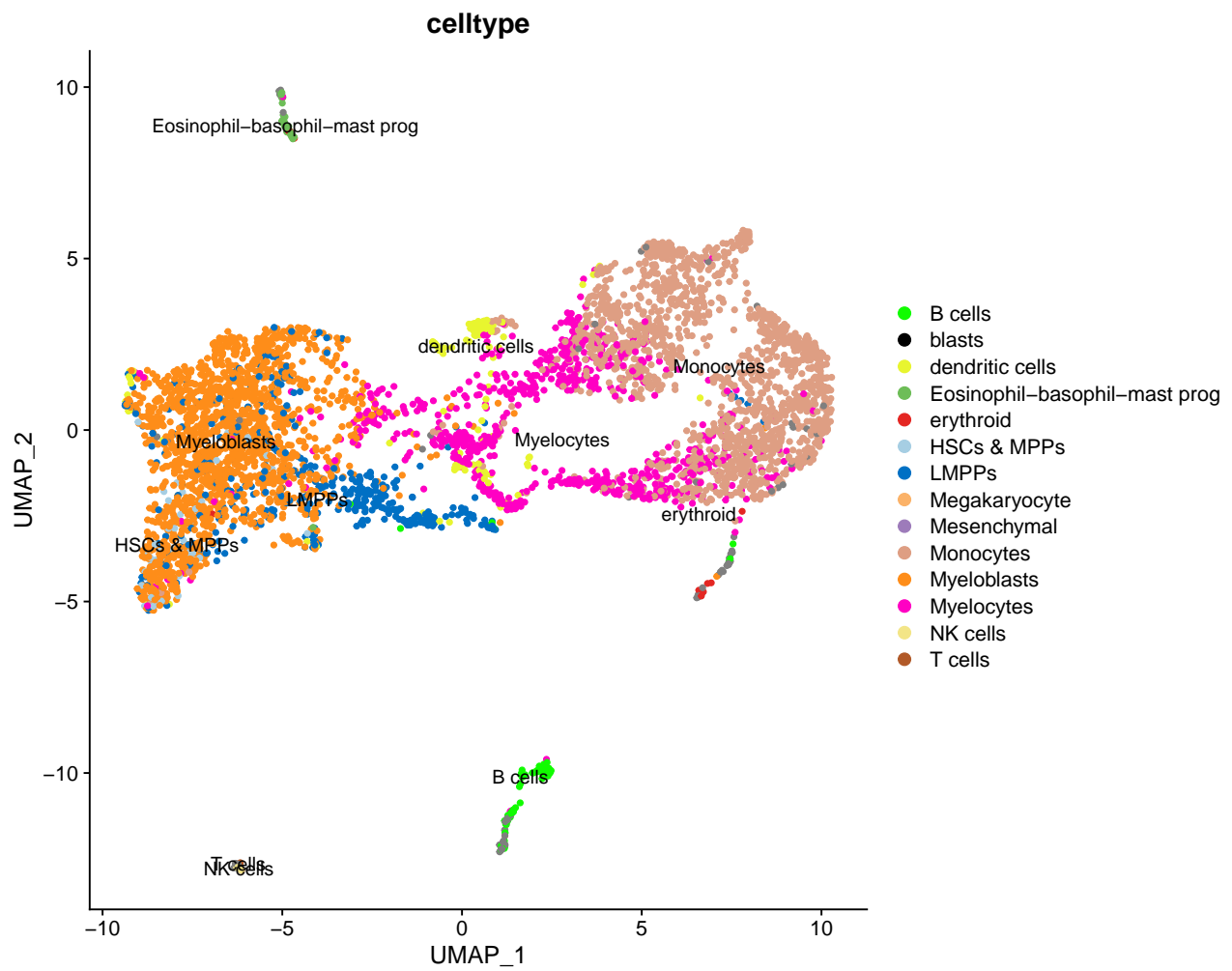
```
## Performing PCA on the provided reference using 1821 features as input.
## Projecting cell embeddings
## Finding neighborhoods
## Finding anchors
## Found 3632 anchors
## Filtering anchors
## Retained 2622 anchors
## Finding integration vectors
## Finding integration vector weights
## Predicting cell labels
##
##           HSC Prog  GMP ProMono Mono  cDC  pDC earlyEry lateEry ProB  B Plasma
##   CD34   210 1552  363    49   9   51   0    12    16   2    7    0
##   CD38    2   23   16   164 1896 126  15    29    9   70   28   15
```

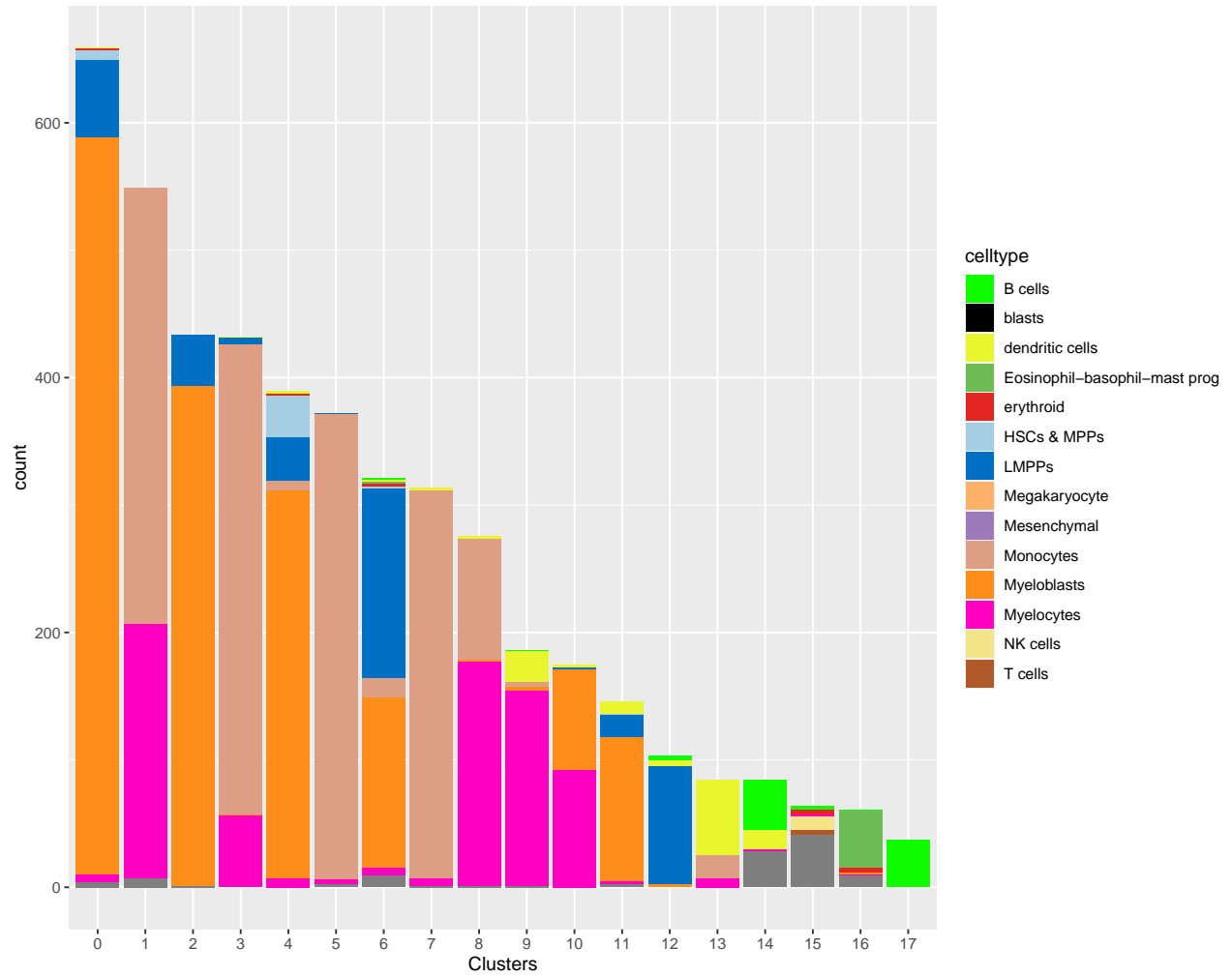
```
##
##           T   CTL   NK
##   CD34    3     0     0
##   CD38    3     1    16
```





4. Project the predictions from Velten onto our UMAP





Cluster 11 and 0 are the ones above the D10 threshold. Nevertheless, cluster 4 has more HSC predicted cells. In addition, is this cluster 4 the ones that colocalizes with cluster 5 LSC from the paired Dx sample. Therefore, we propose cluster 4 as the more likely to have enrichment of LSC