

AML7_Dx

jtrincado

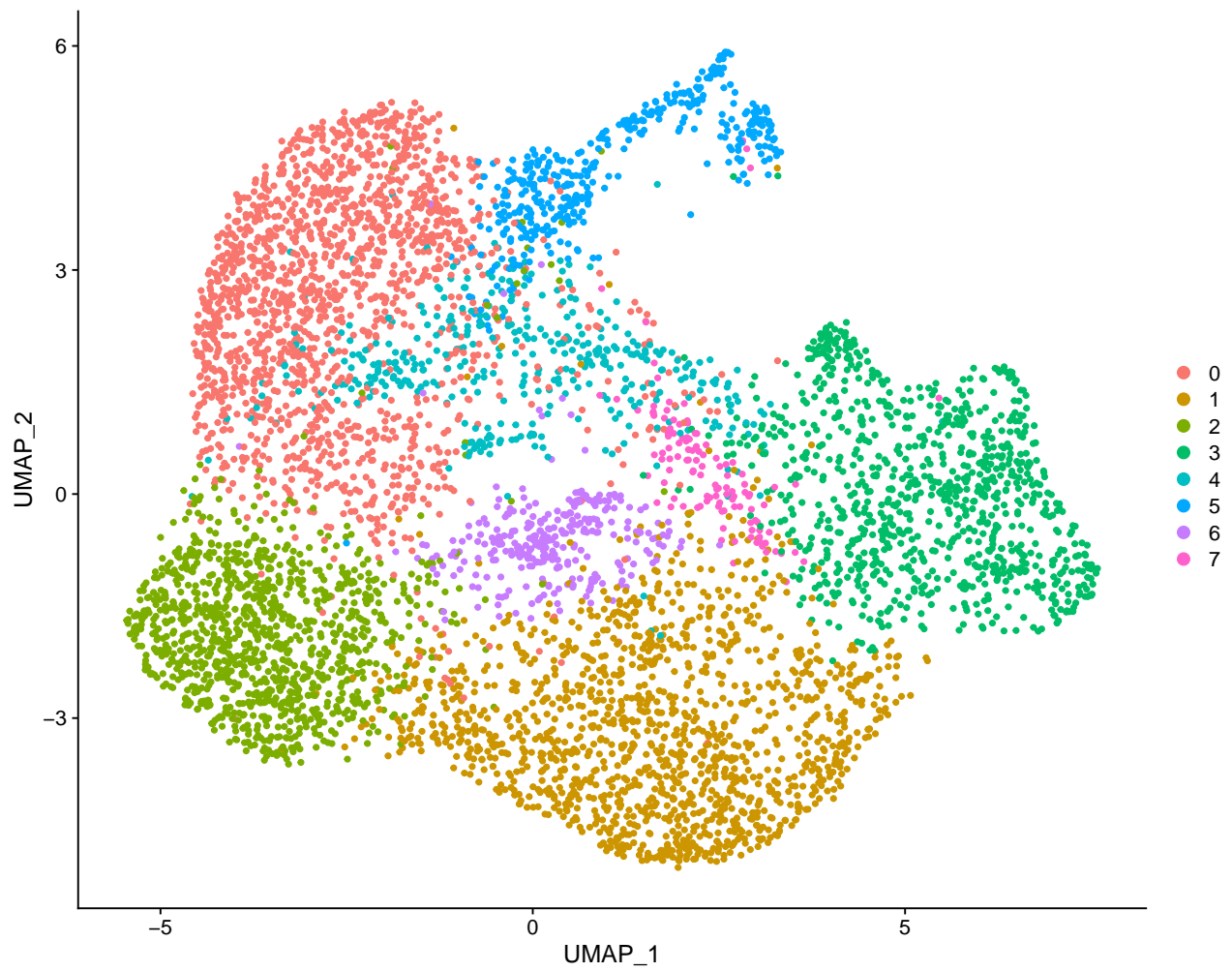
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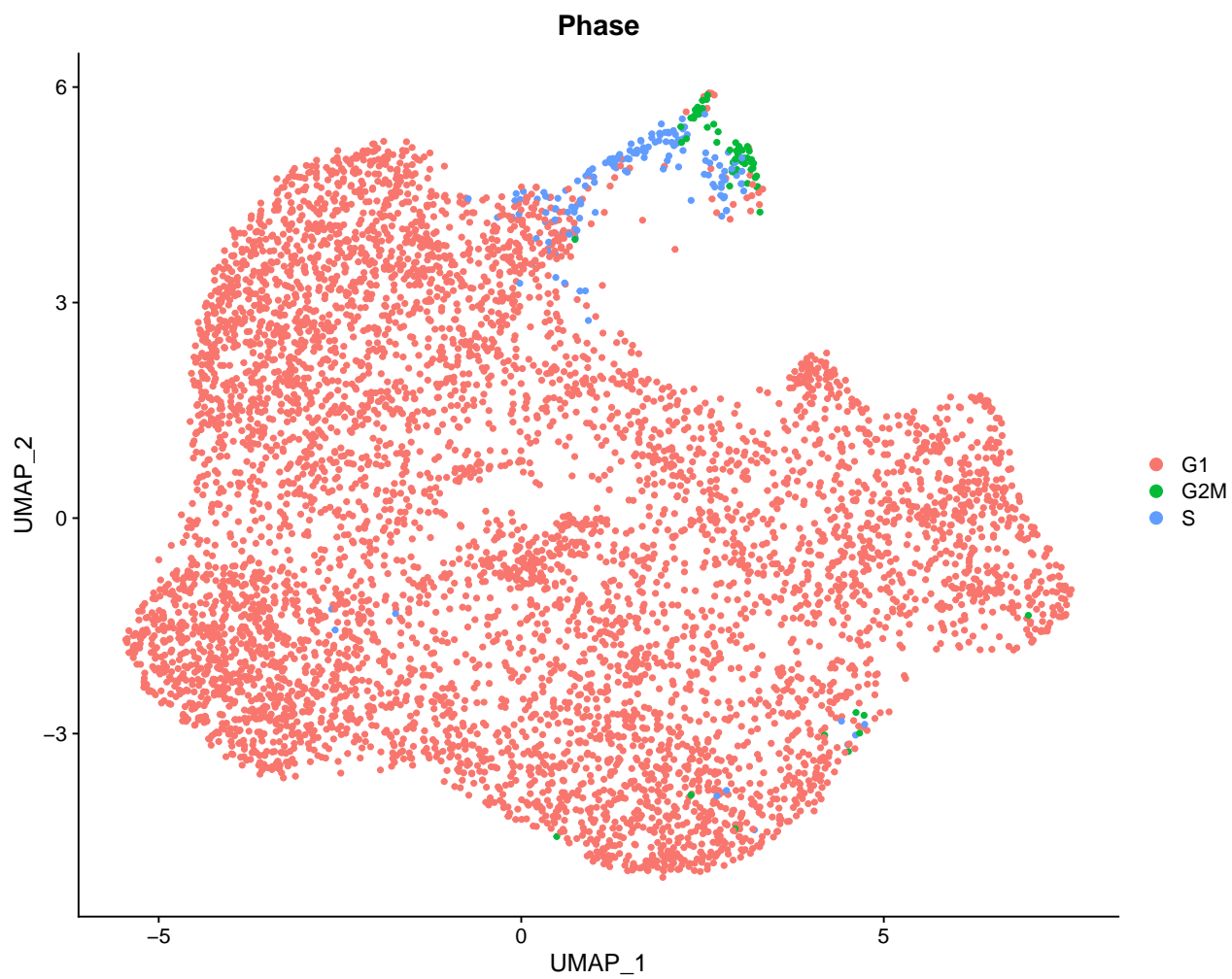
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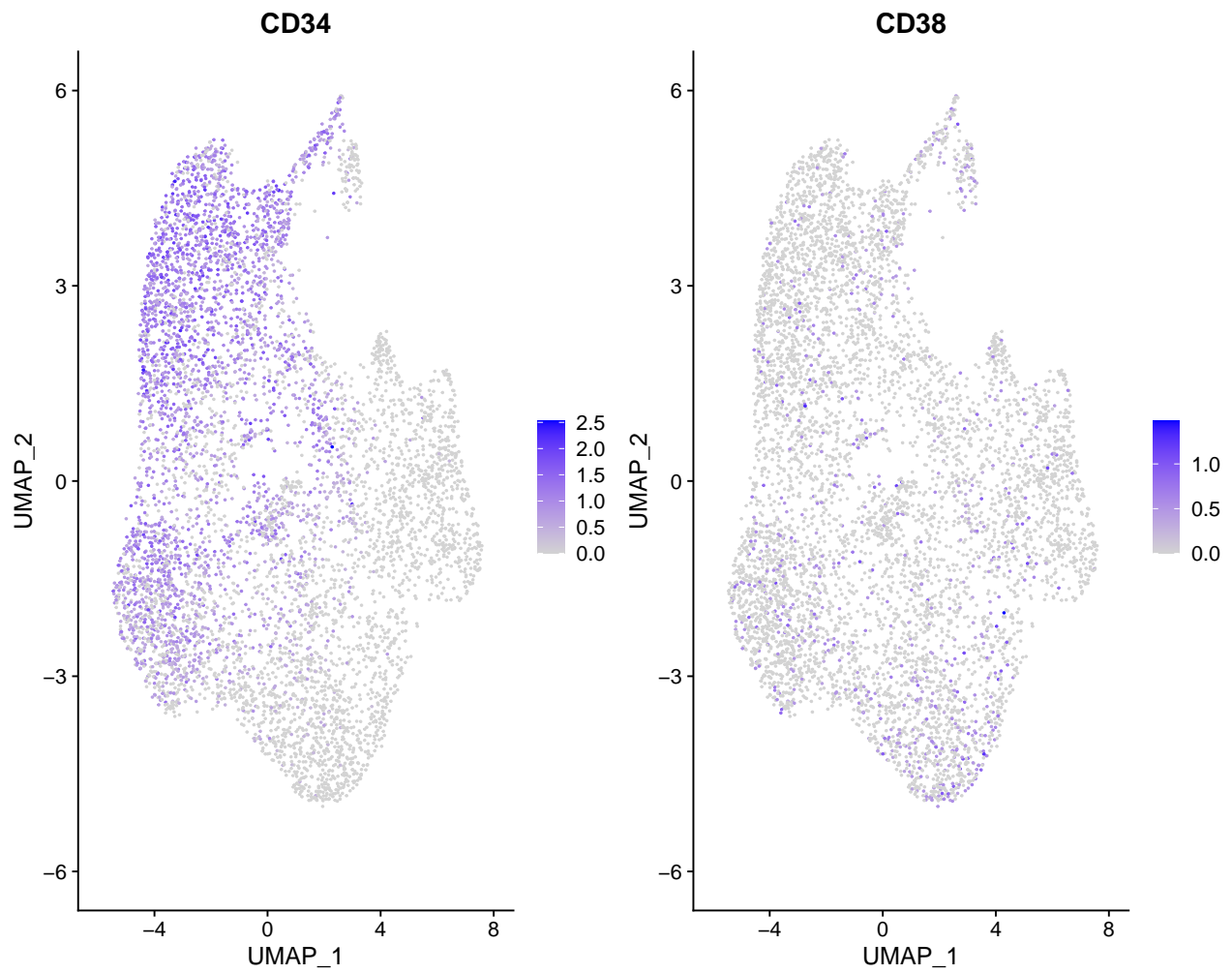
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1. Put together both 34 and 38 libraries. Apply QC and dimensionality reduction.

```
## Modularity Optimizer version 1.3.0 by Ludo Waltman and Nees Jan van Eck
##
## Number of nodes: 6214
## Number of edges: 209221
##
## Running Louvain algorithm...
## Maximum modularity in 10 random starts: 0.8405
## Number of communities: 8
## Elapsed time: 0 seconds
```

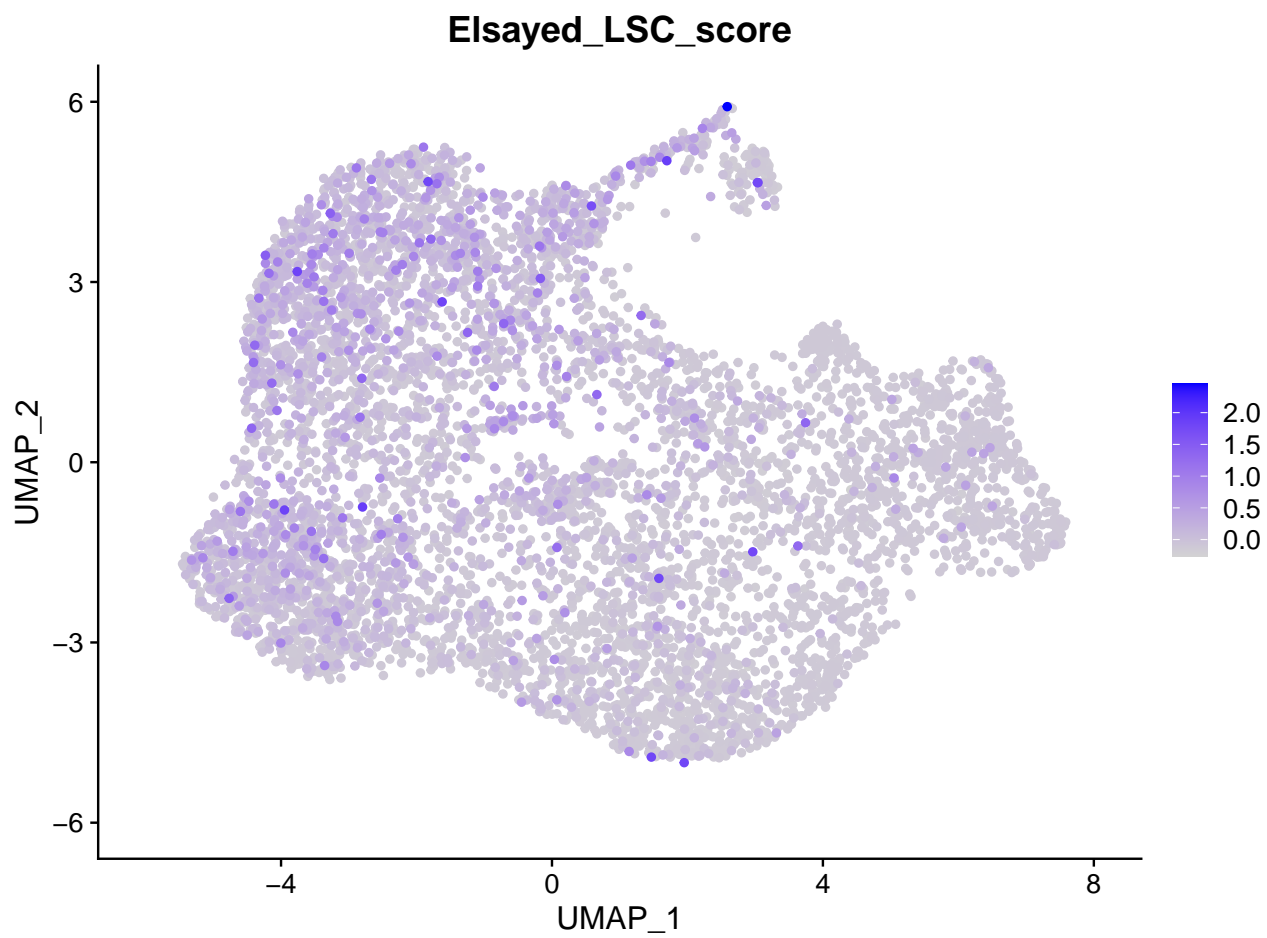


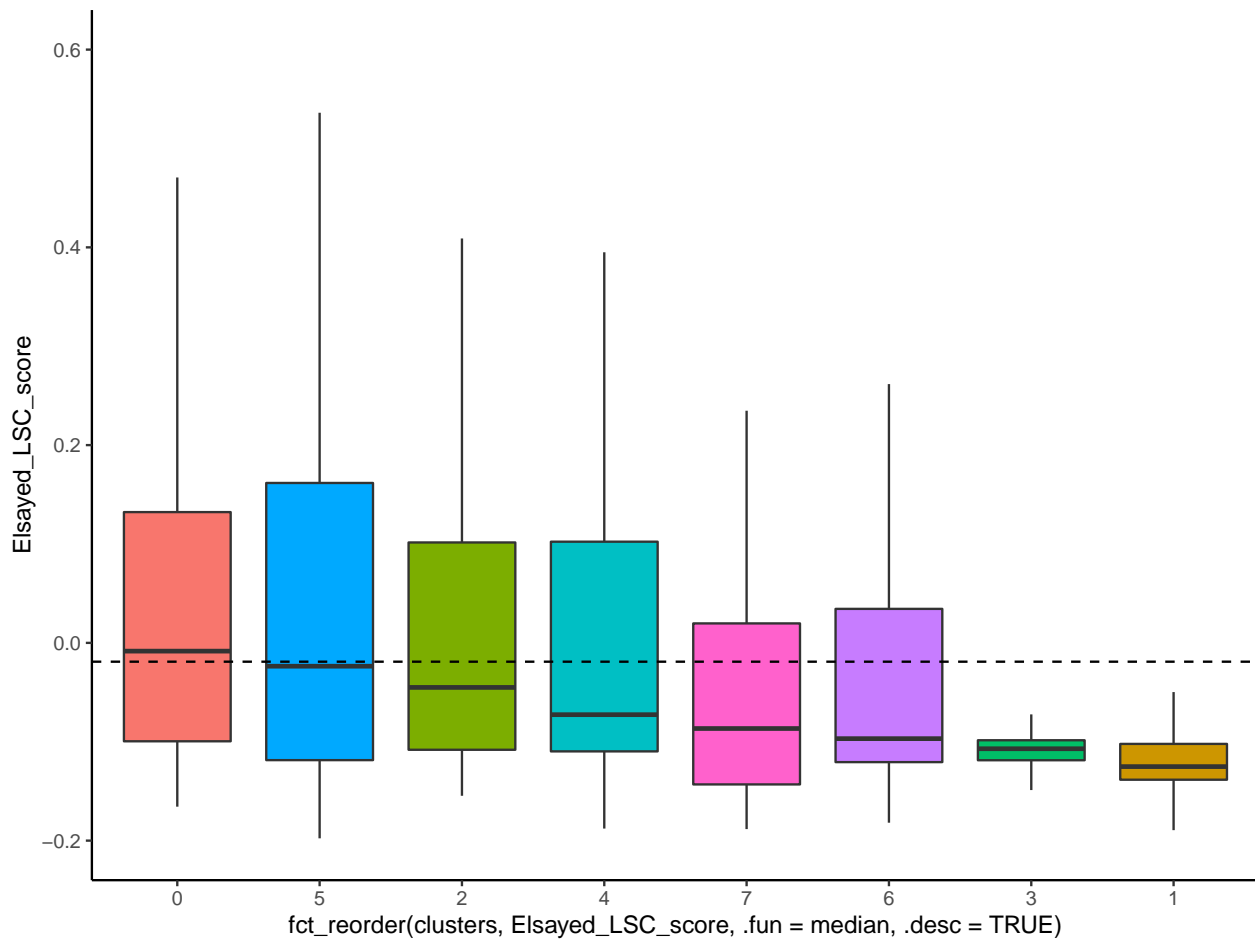




2. Get the LSC6 score

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

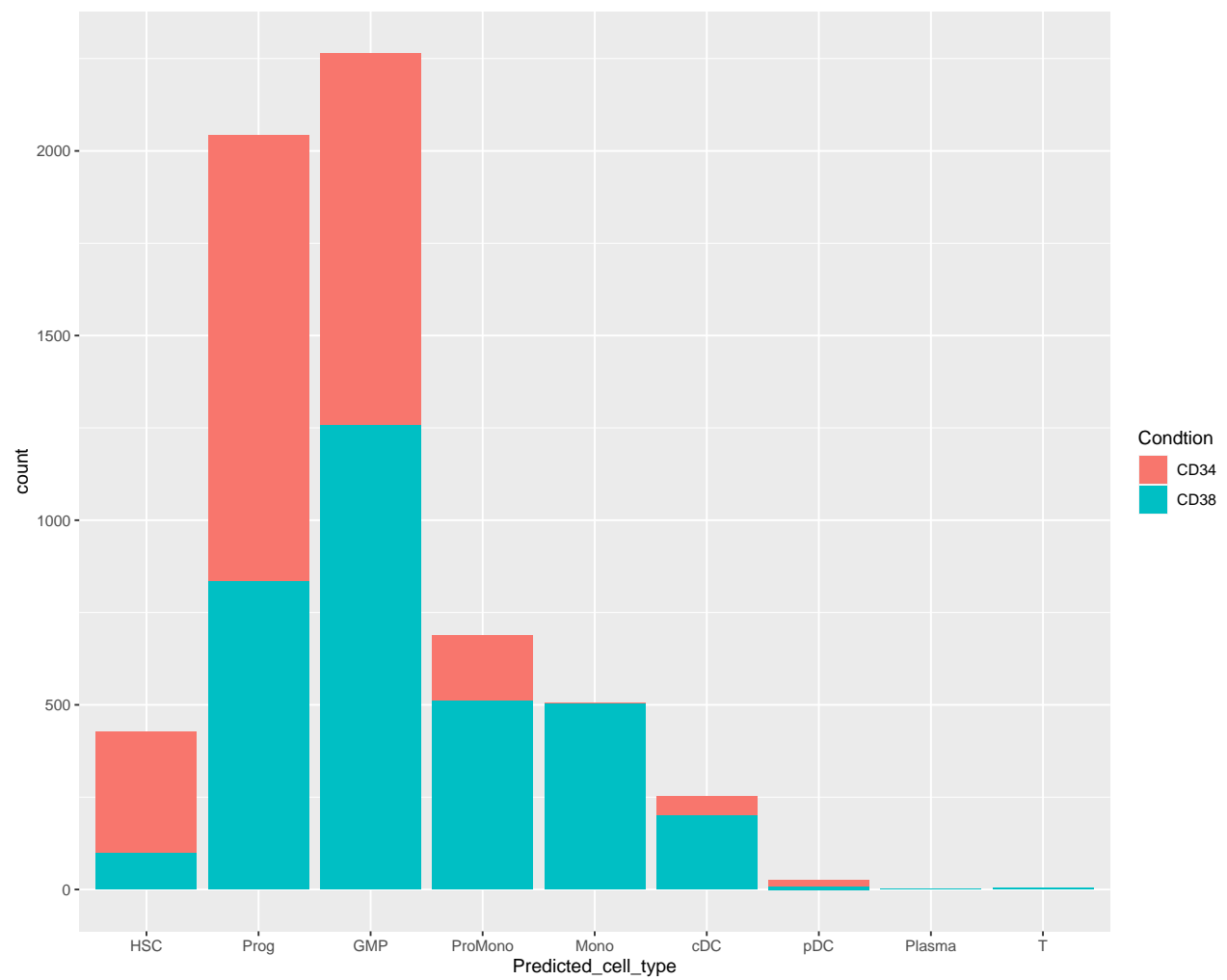


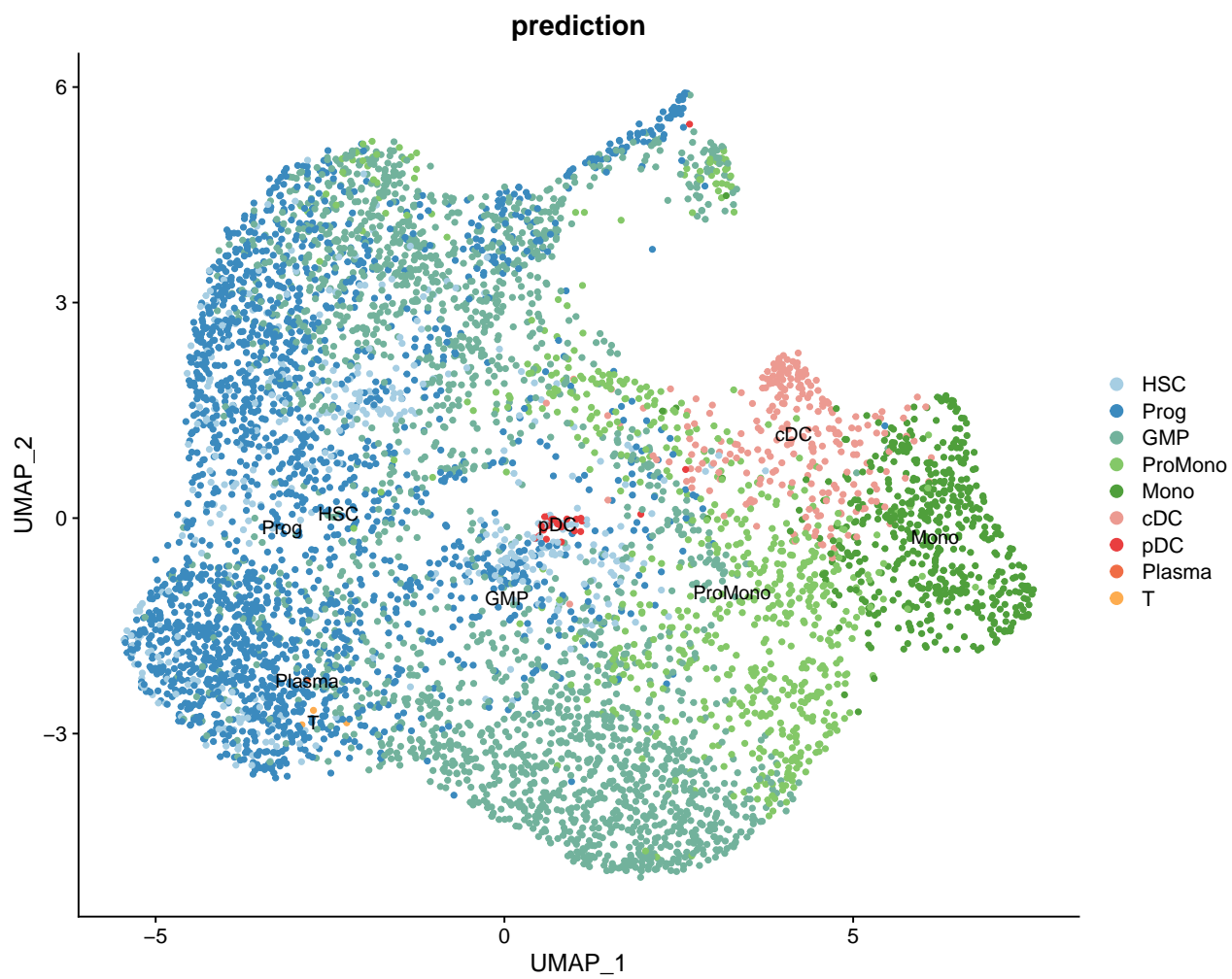


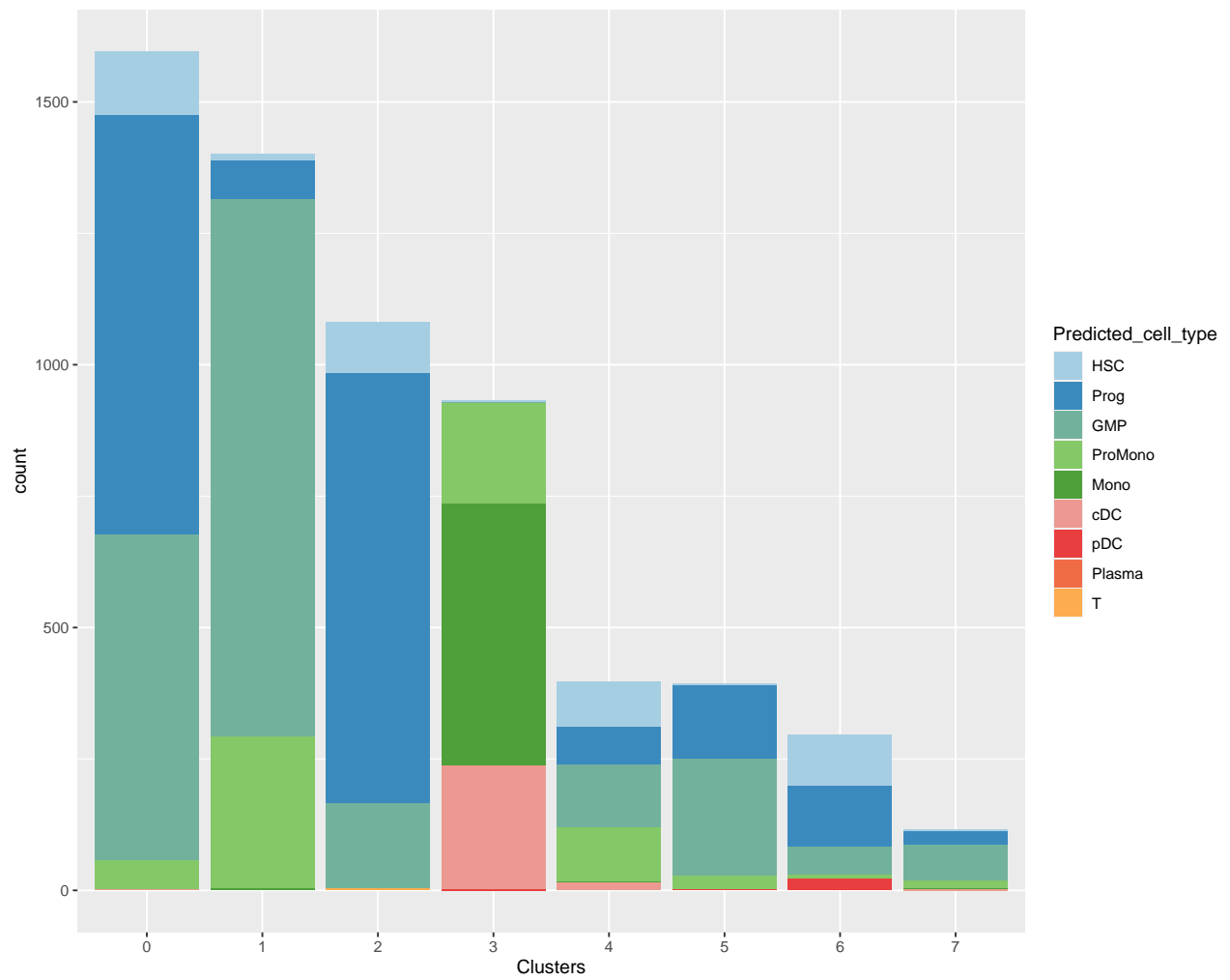
3. Predict the class of the cells using the markers and the expression of the BM cells form Van_Galen paper

```
## Performing PCA on the provided reference using 1821 features as input.
## Projecting cell embeddings
## Finding neighborhoods
## Finding anchors
## Found 2055 anchors
## Filtering anchors
## Retained 1132 anchors
## Finding integration vectors
## Finding integration vector weights
## Predicting cell labels
##
##          HSC Prog  GMP ProMono Mono  cDC  pDC earlyEry lateEry ProB  B Plasma
##  CD34    330 1210 1009    177    5   54   17         0         0    0    0    0
##  CD38     98  834 1256    511  502  199    8         0         0    0    0    1
```

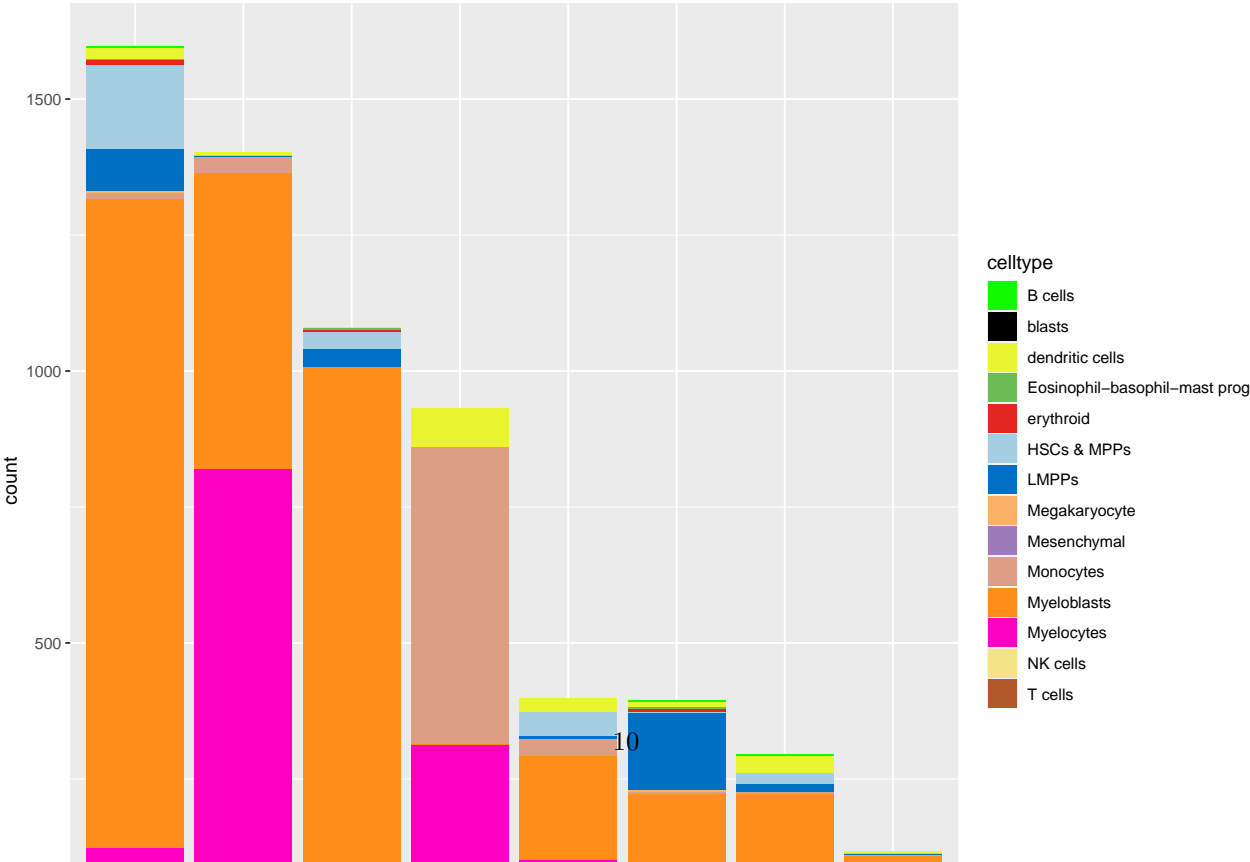
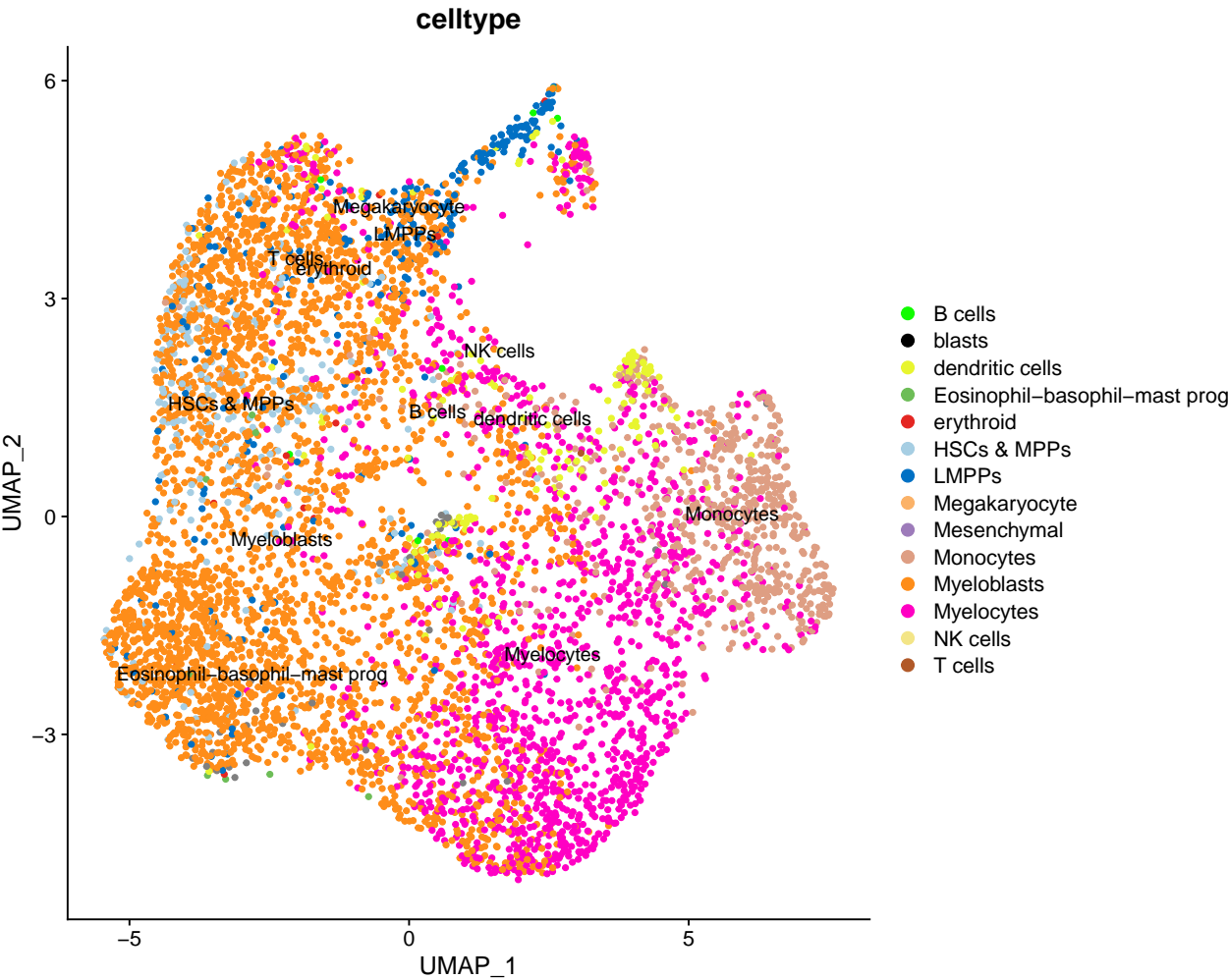
```
##
##      T   CTL   NK
## CD34   0     0    0
## CD38   3     0    0
```







4. Project the predictions from Velten onto our UMAP



Cluster 0 seems the most likely to be enriched in LSC