

# AML10\_Dx

jtrincado

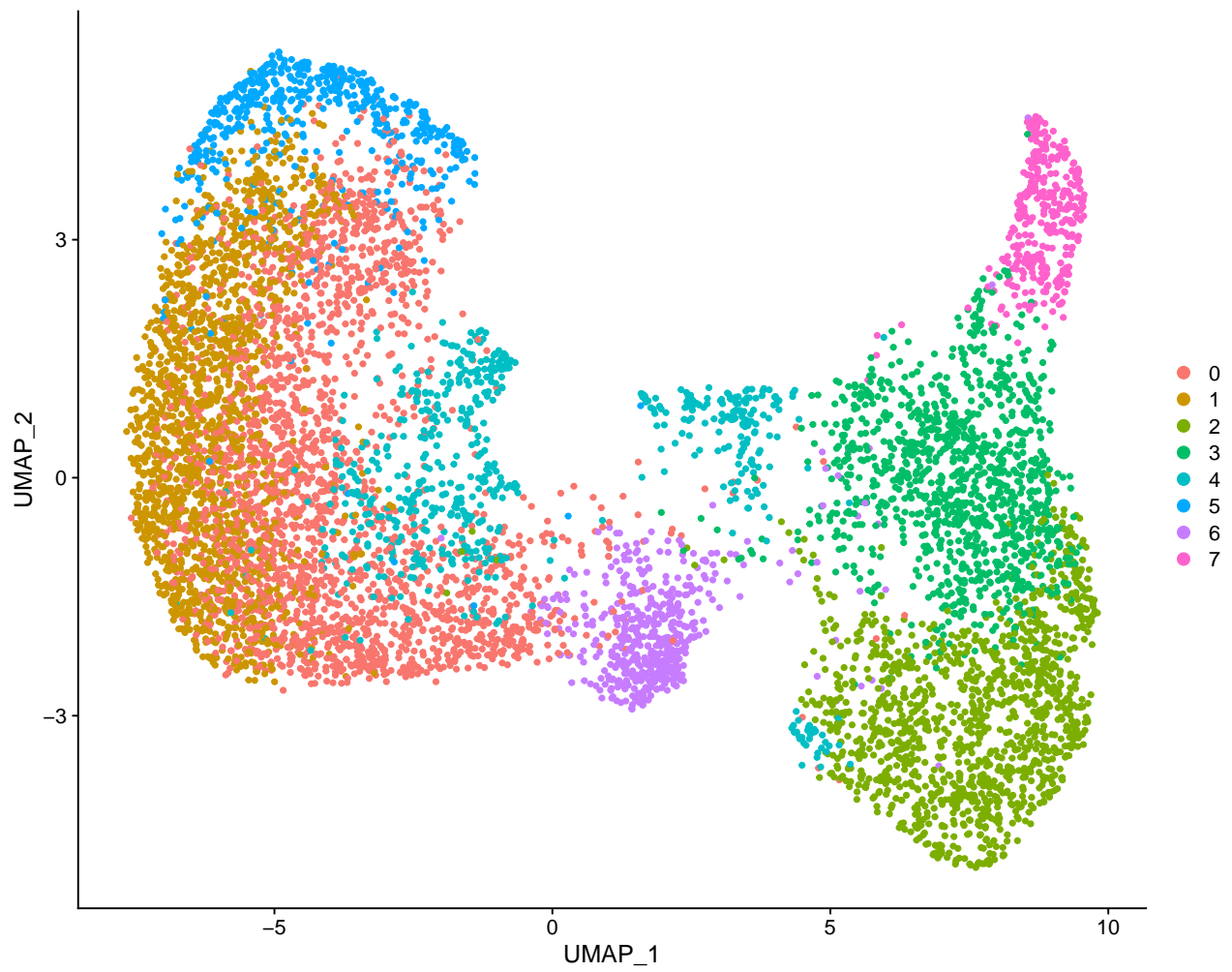
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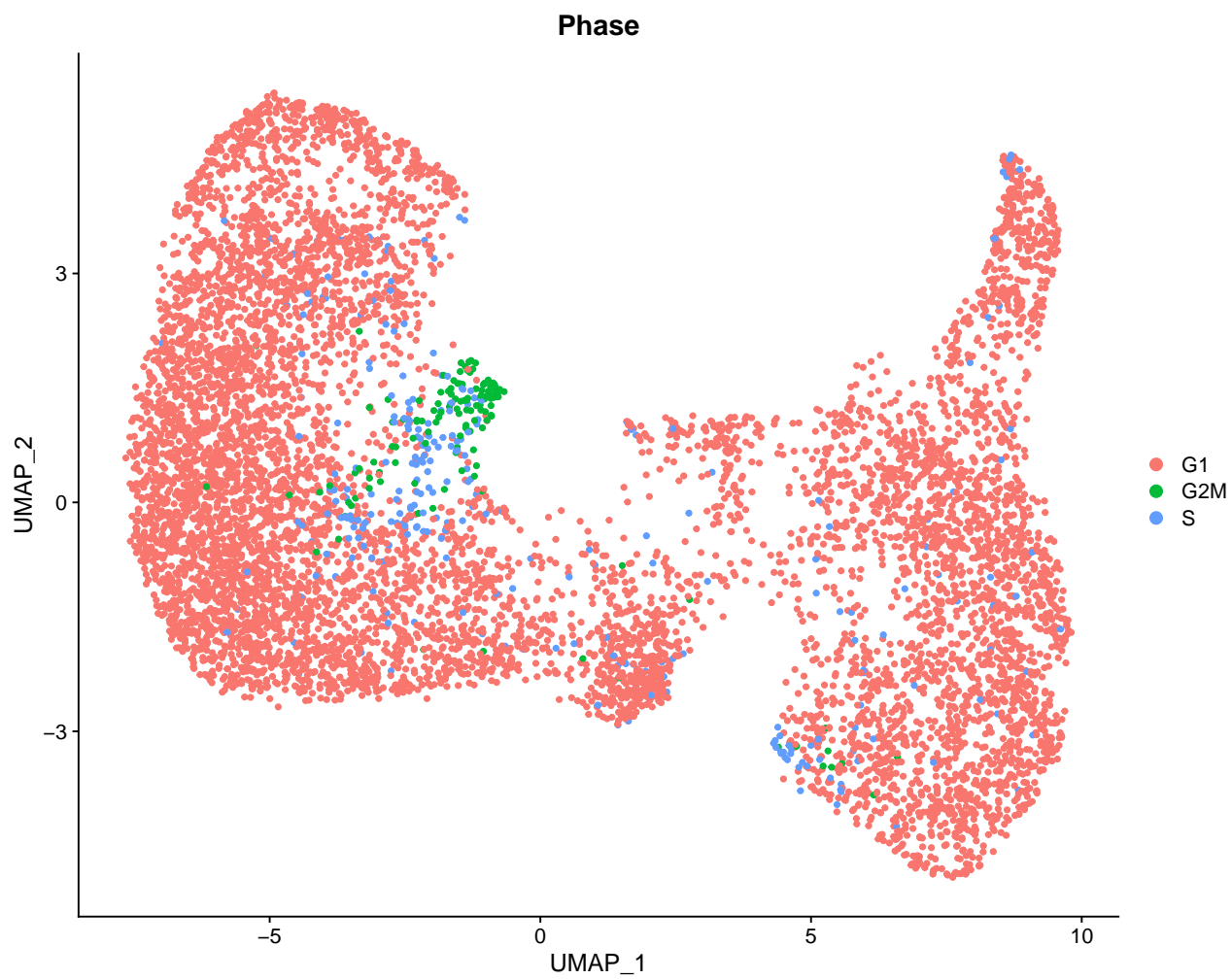
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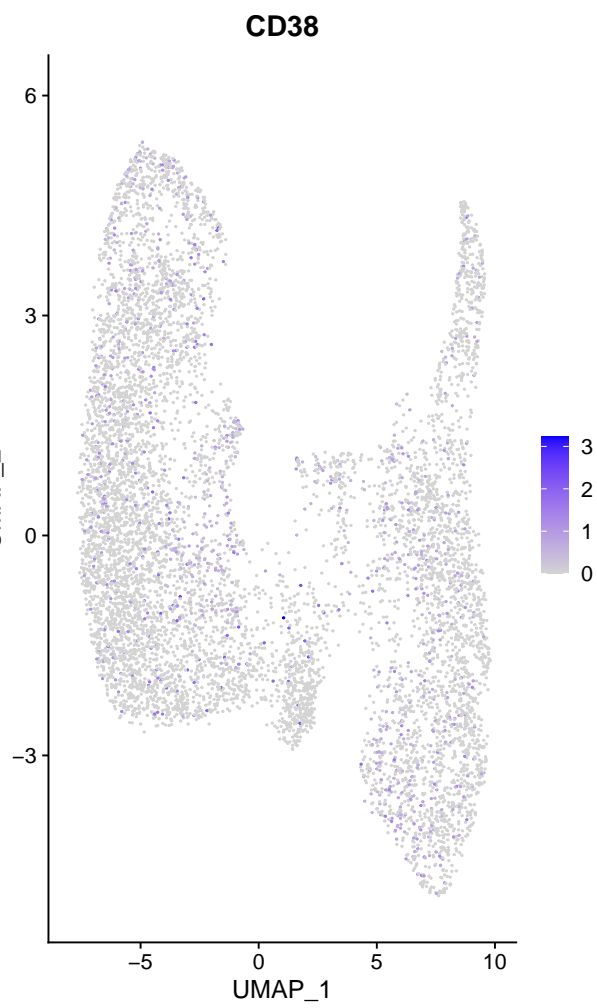
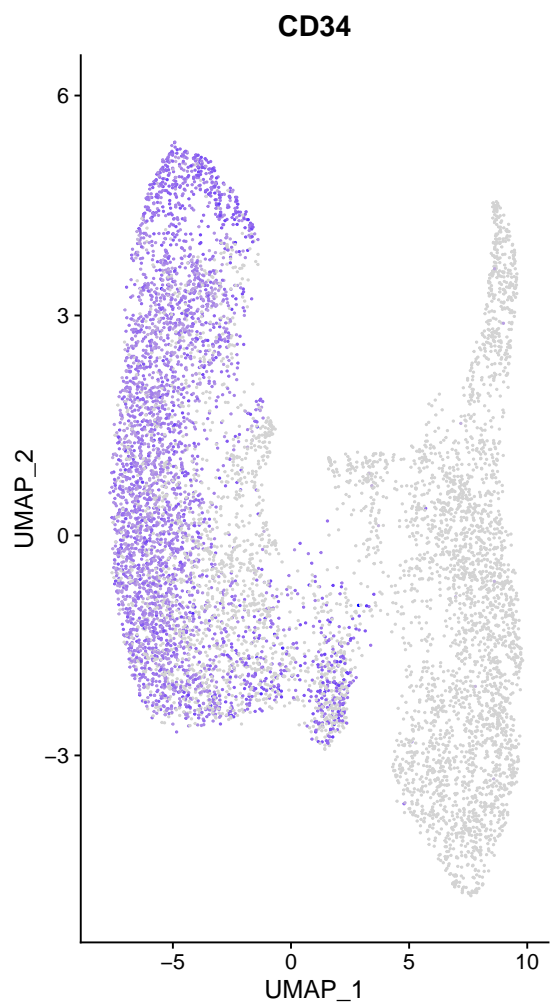
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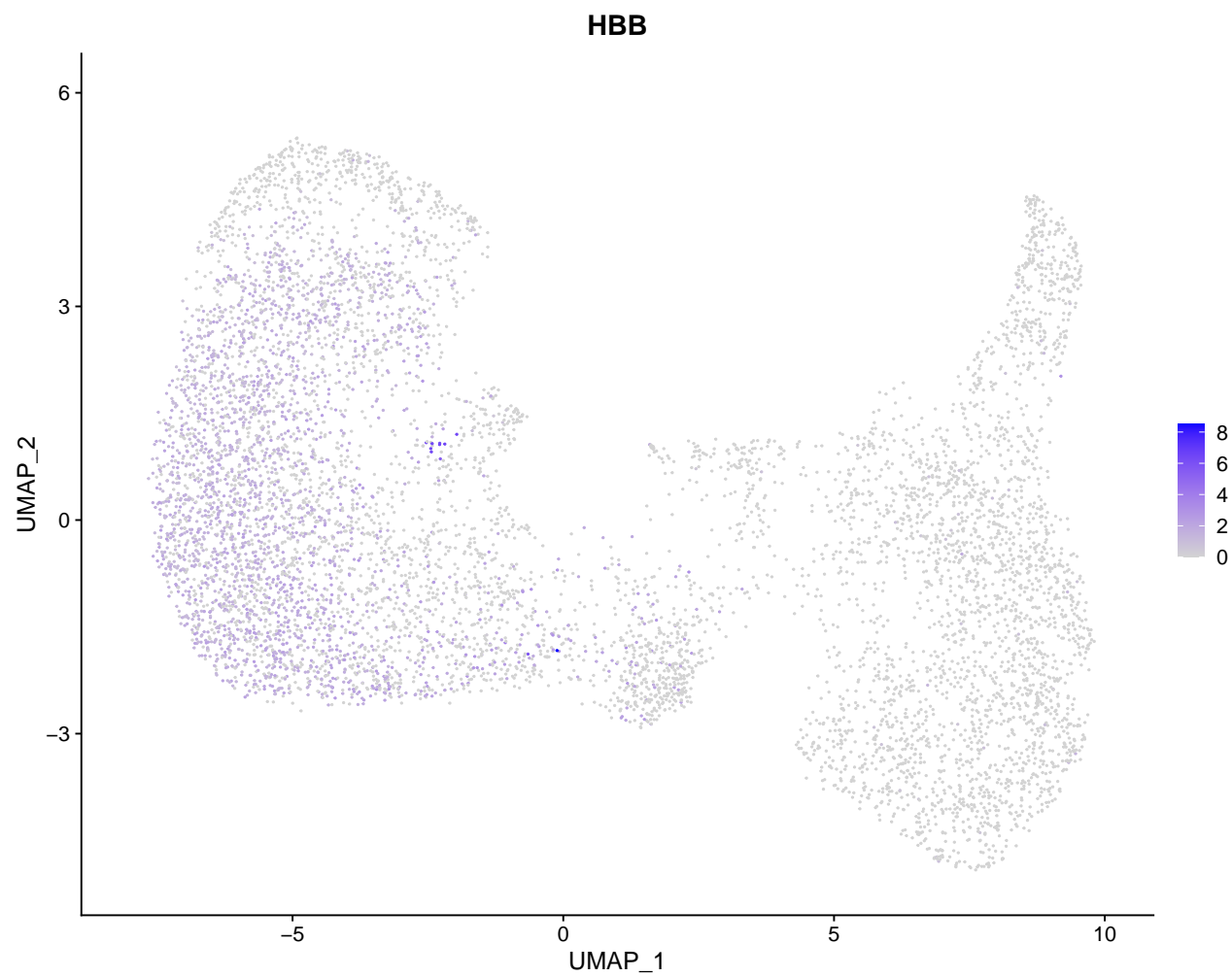
## 1. Put together both 34 and 38 libraries. Apply QC and dimensionality reduction.

```
## CD34_AAACCCAAGCGTGAGT-1 CD34_AAACCCACAAGCGAAC-1 CD34_AAACCCACACCTCAGG-1
##                               1                               0                               6
## CD34_AAACCCACATTGACTG-1 CD34_AAACCCATCTGTCGTC-1
##                               0                               0
## Levels: 0 1 2 3 4 5 6 7
```



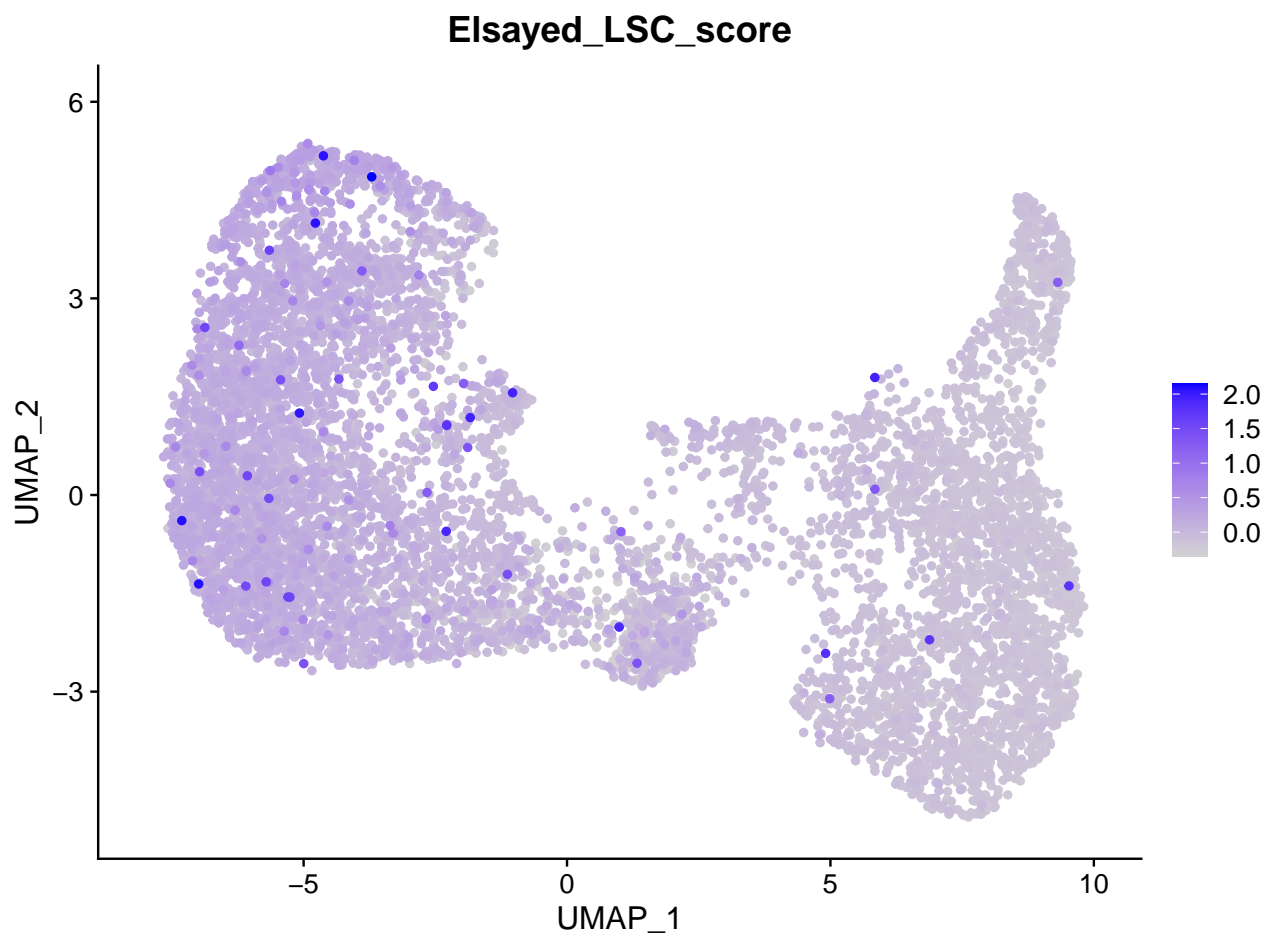


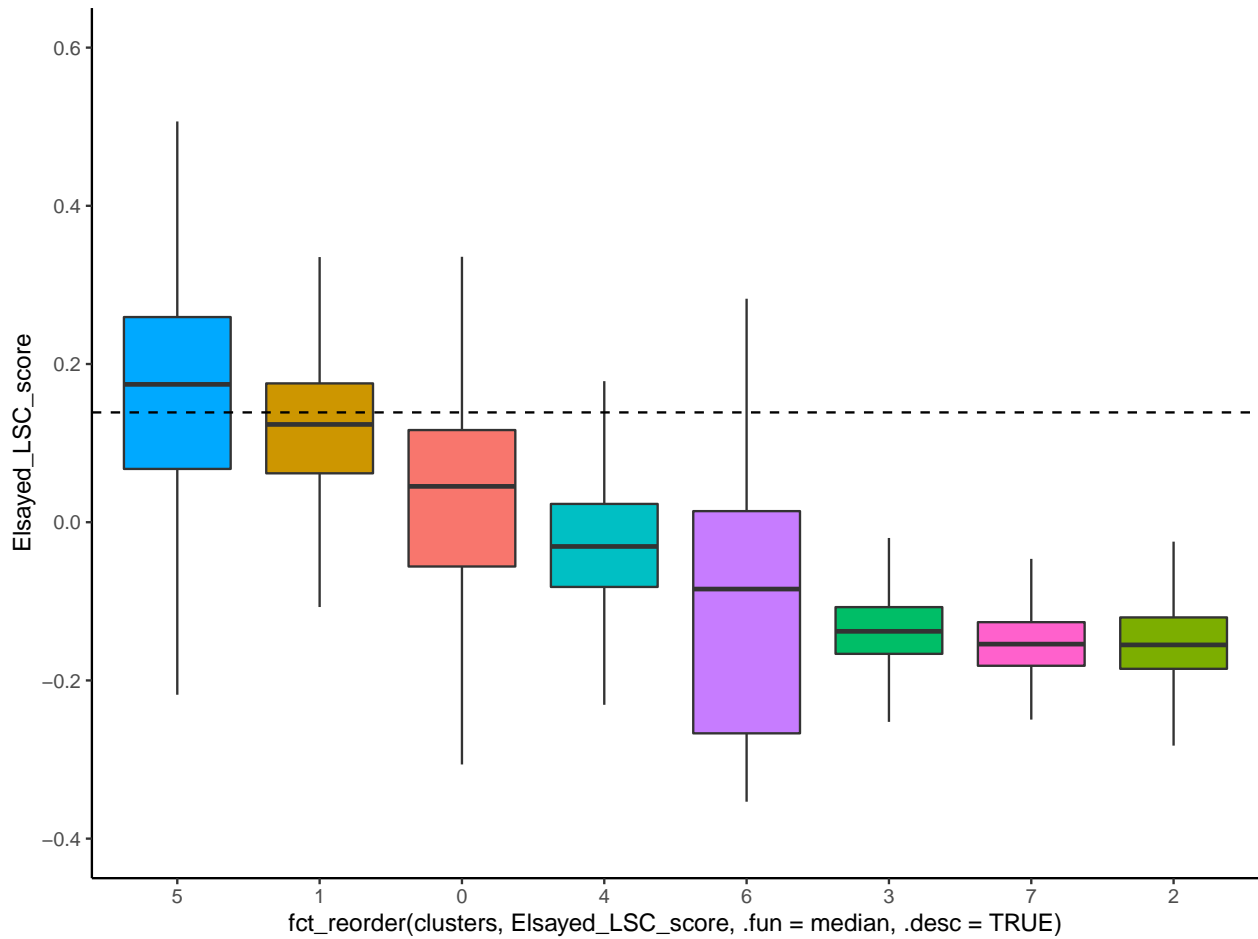




## 2. Get the LSC6 score

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

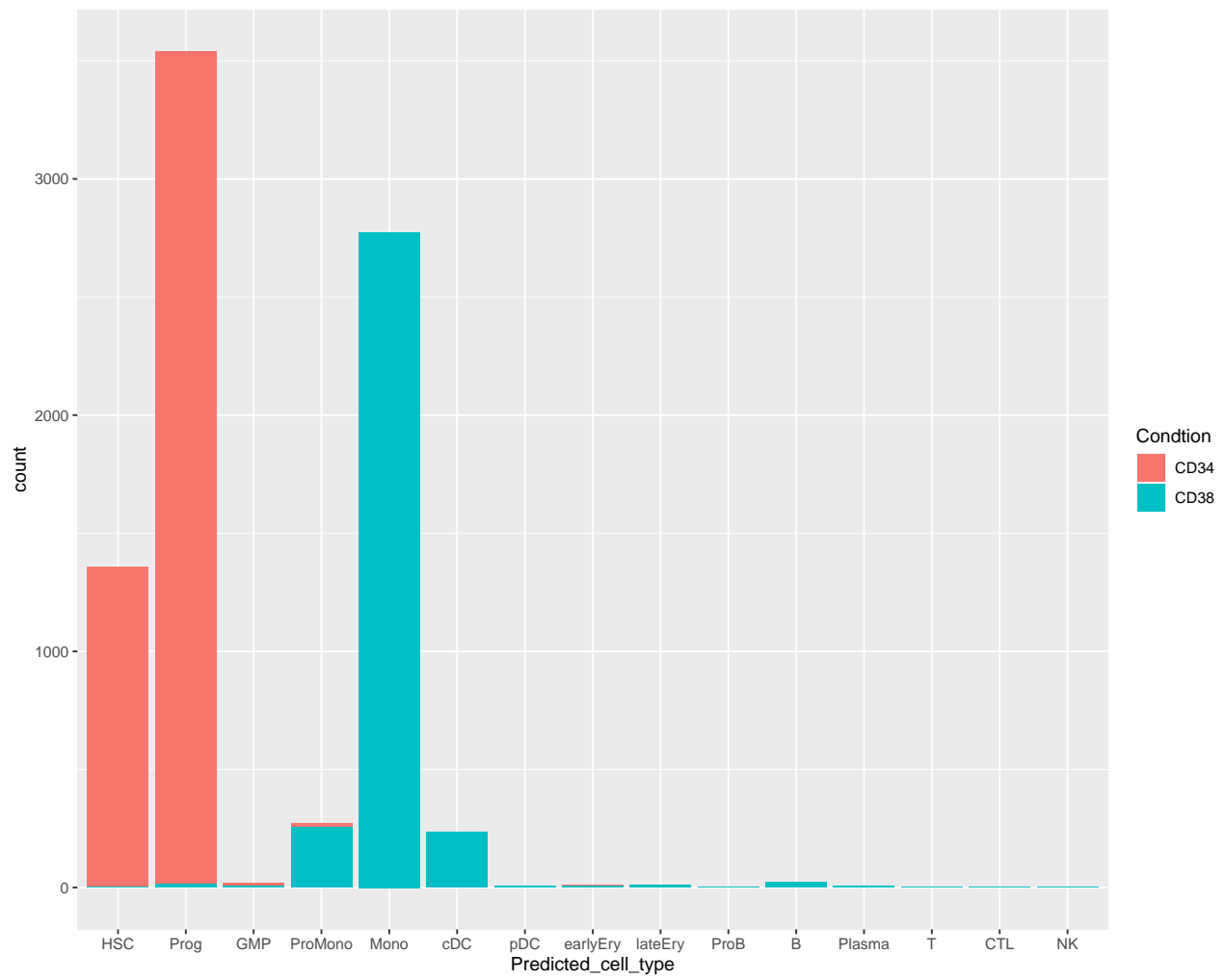




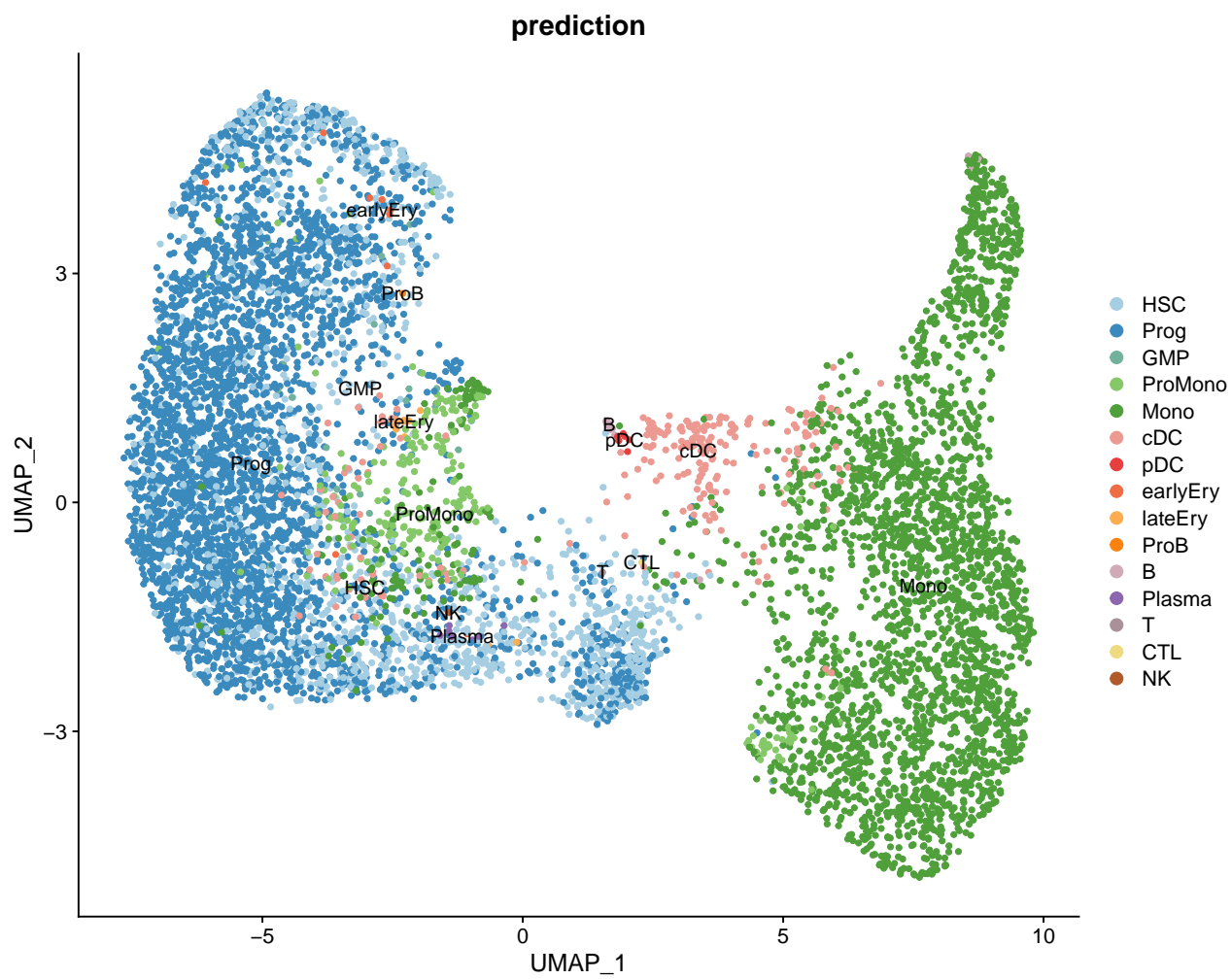
### 3. Predict the class of the cells using the markers and the expression of the BM cells form Van\_Galen paper

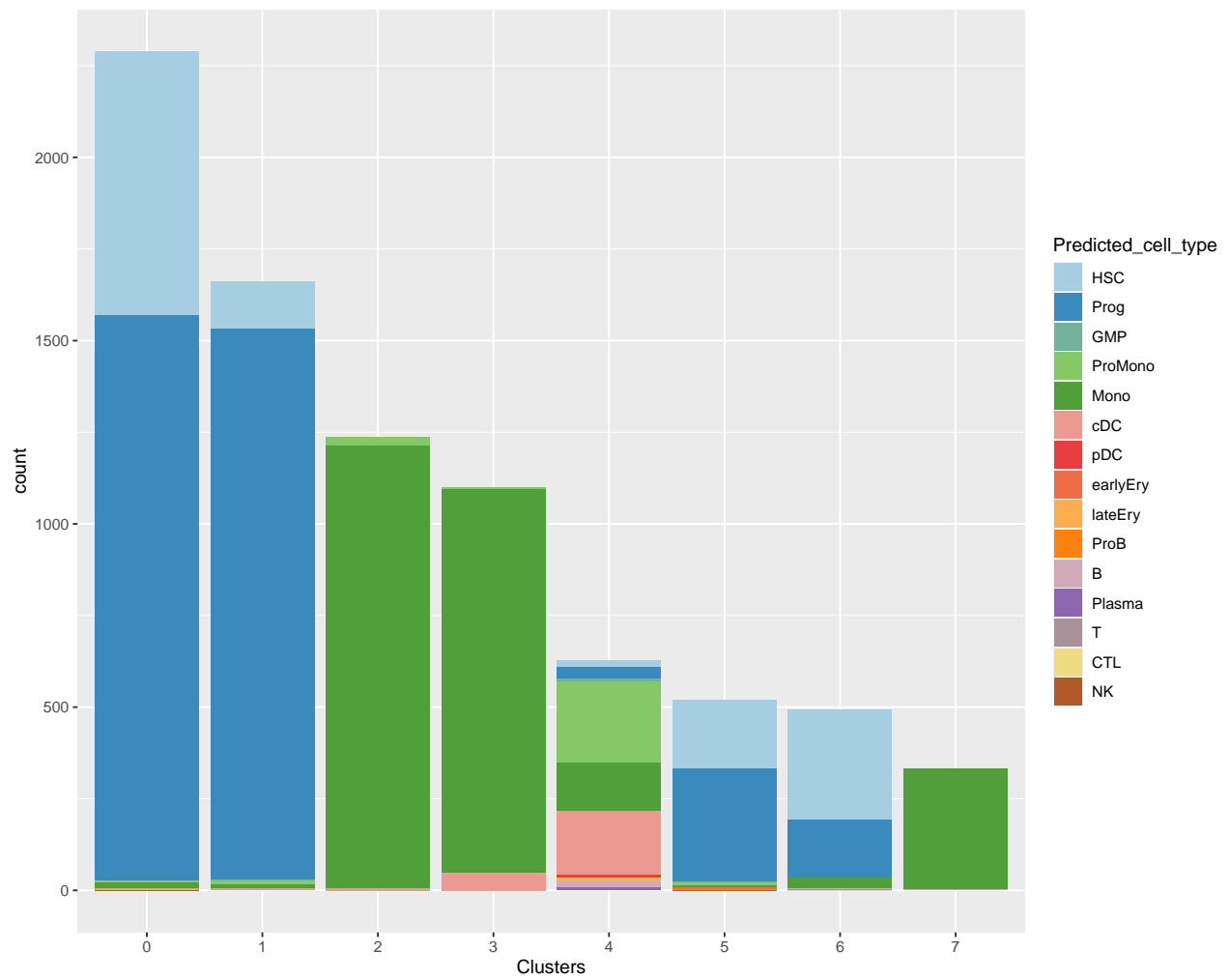
```
## Performing PCA on the provided reference using 1821 features as input.
## Projecting cell embeddings
## Finding neighborhoods
## Finding anchors
## Found 2785 anchors
## Filtering anchors
## Retained 1482 anchors
## Finding integration vectors
## Finding integration vector weights
## Predicting cell labels
##
##          HSC Prog  GMP ProMono Mono  cDC  pDC earlyEry lateEry ProB  B Plasma
##  CD34 1353 3527   14    18    0    1    0          1      2    0    0    0
##  CD38   3   14    7   254 2774 235    7          8      9    1   22    8
```

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

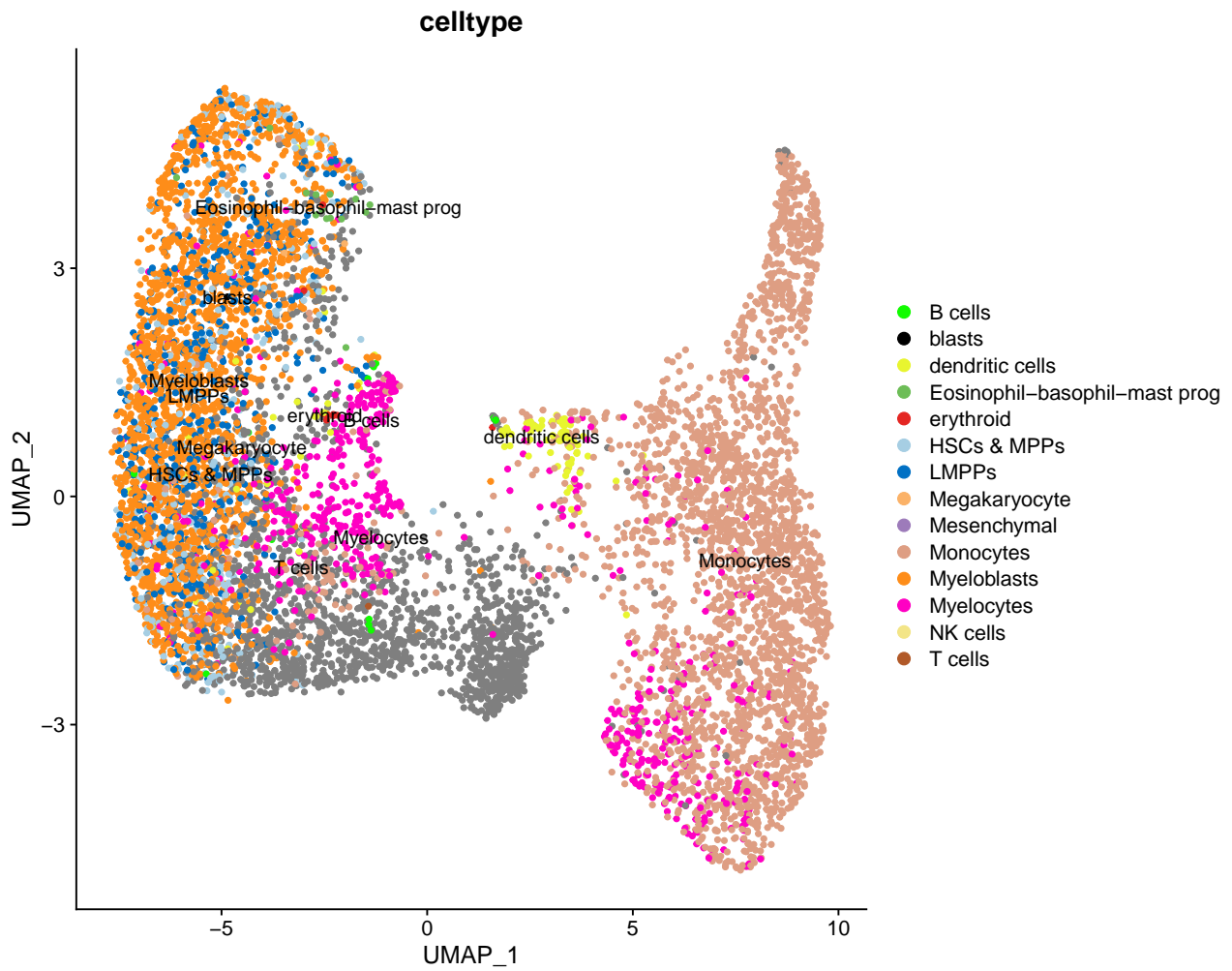


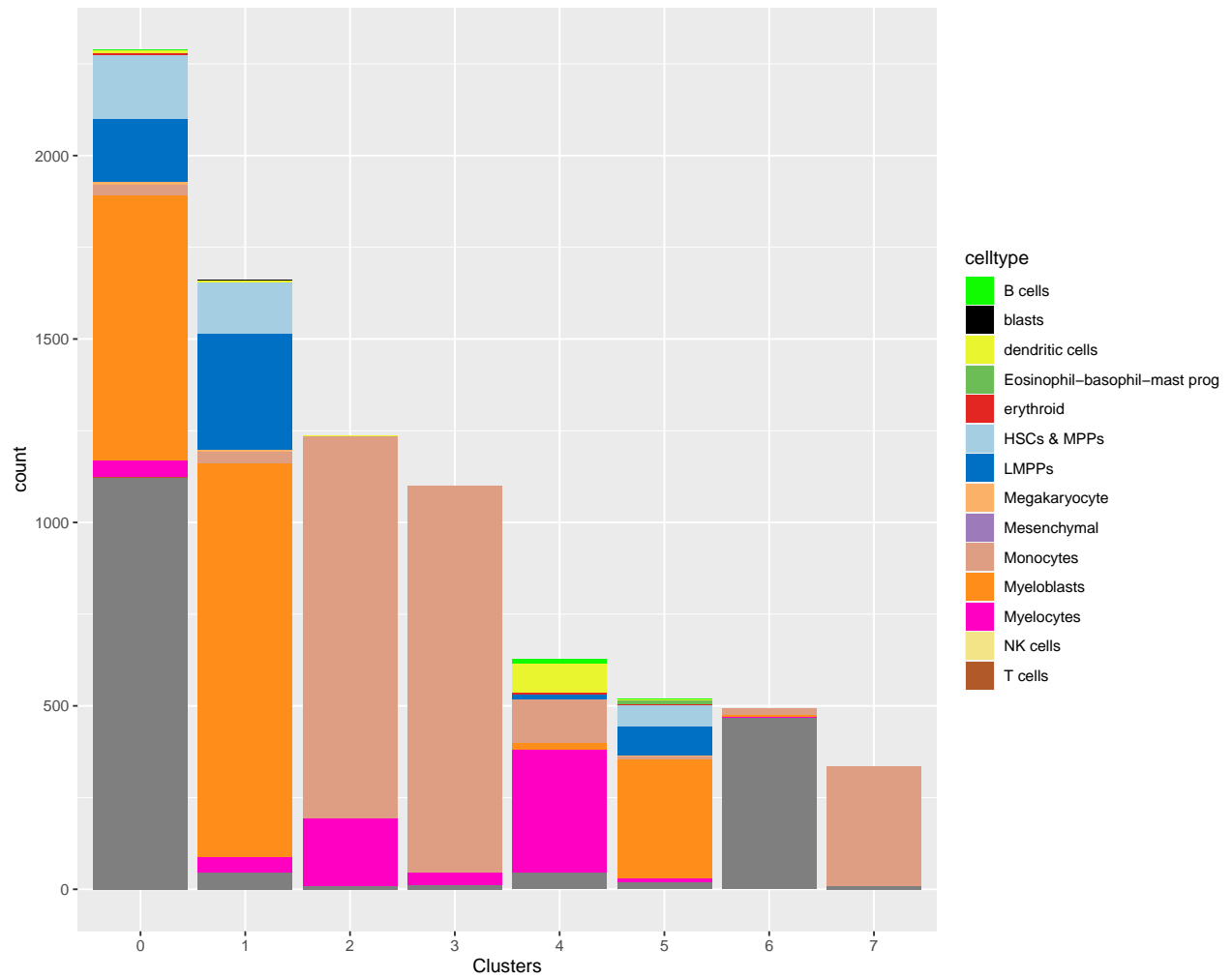






#### 4. Project the predictions from Velten onto our UMAP





Cluster 5 seems the most likely to be enriched in LSC