

# AML9\_Rx

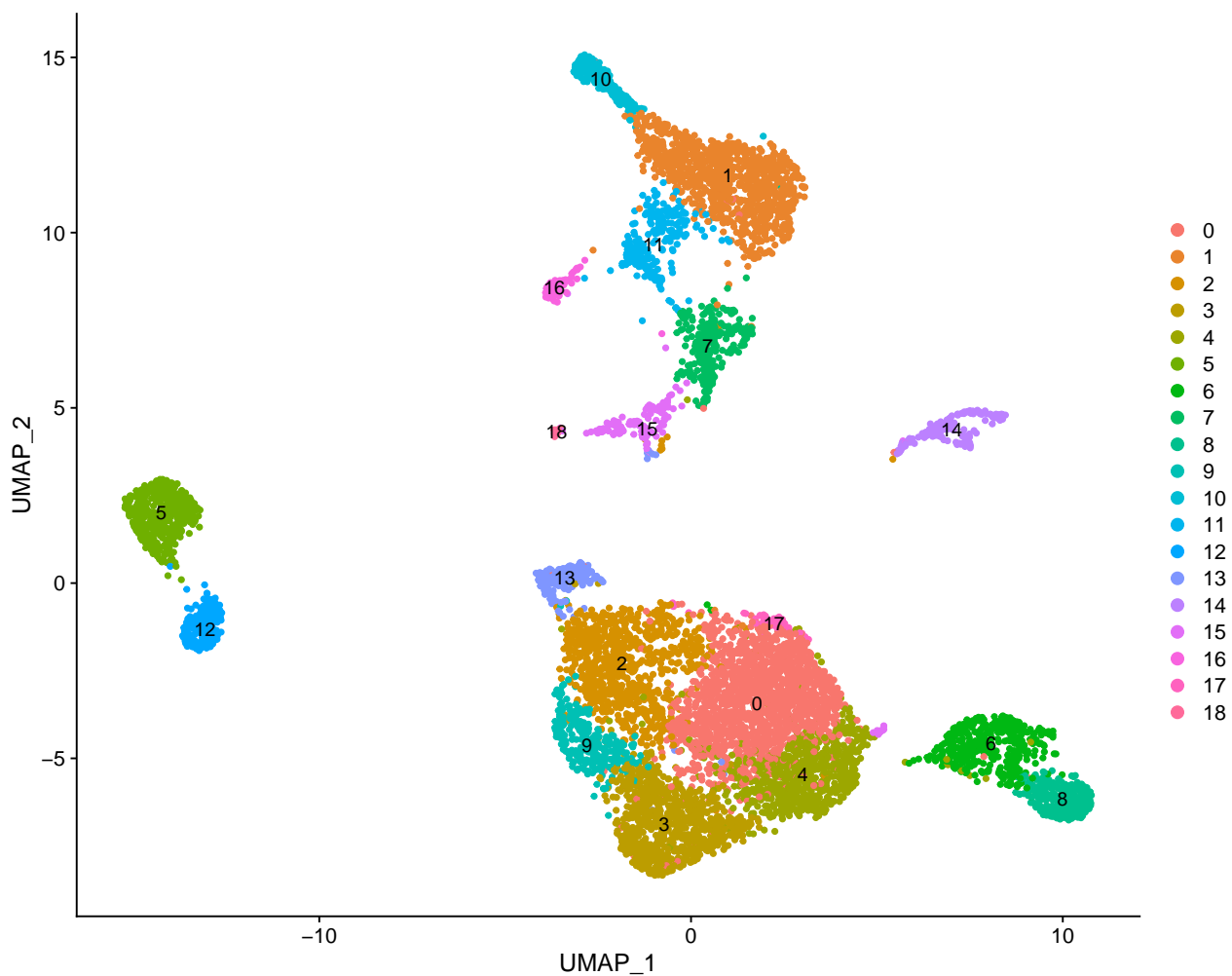
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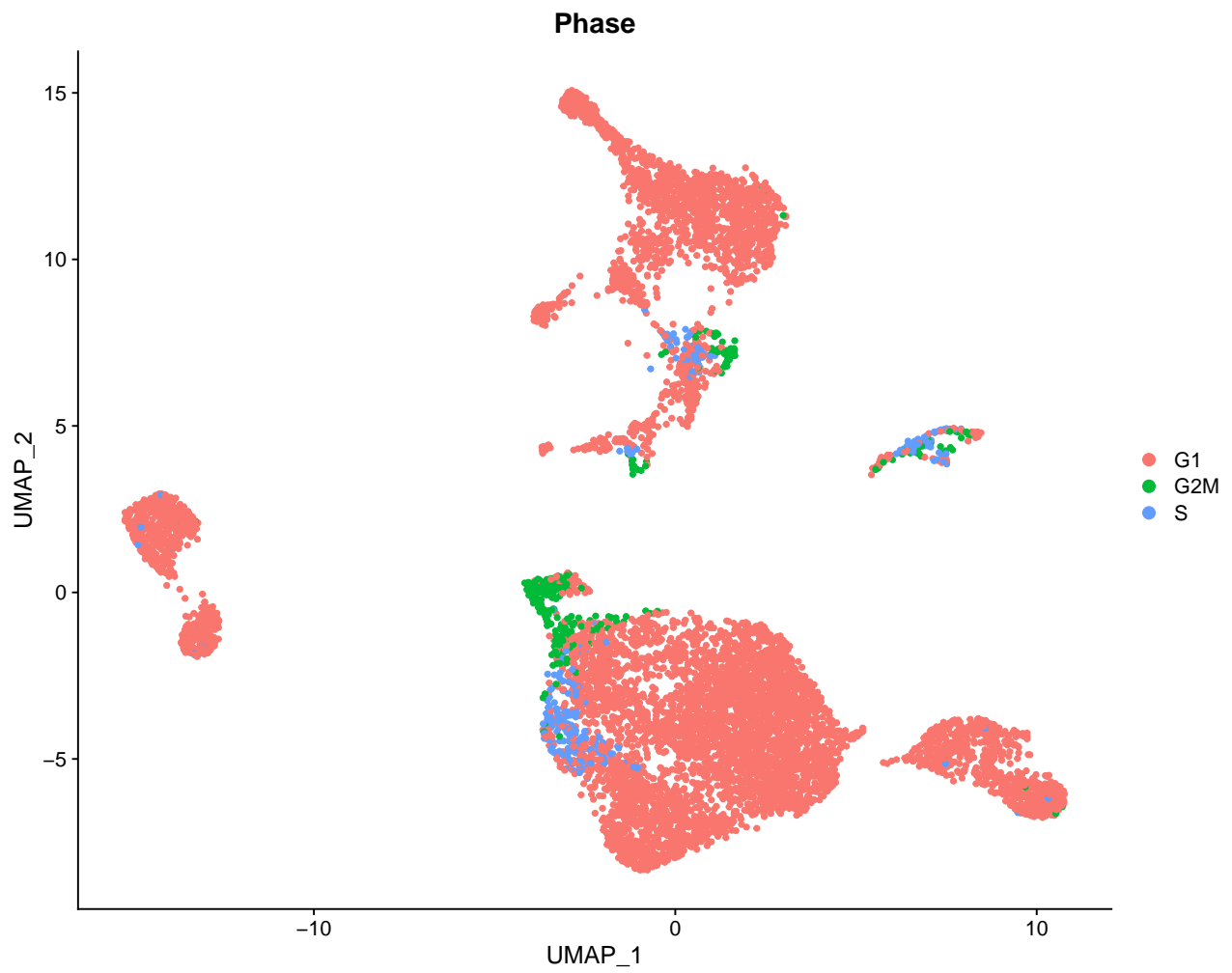
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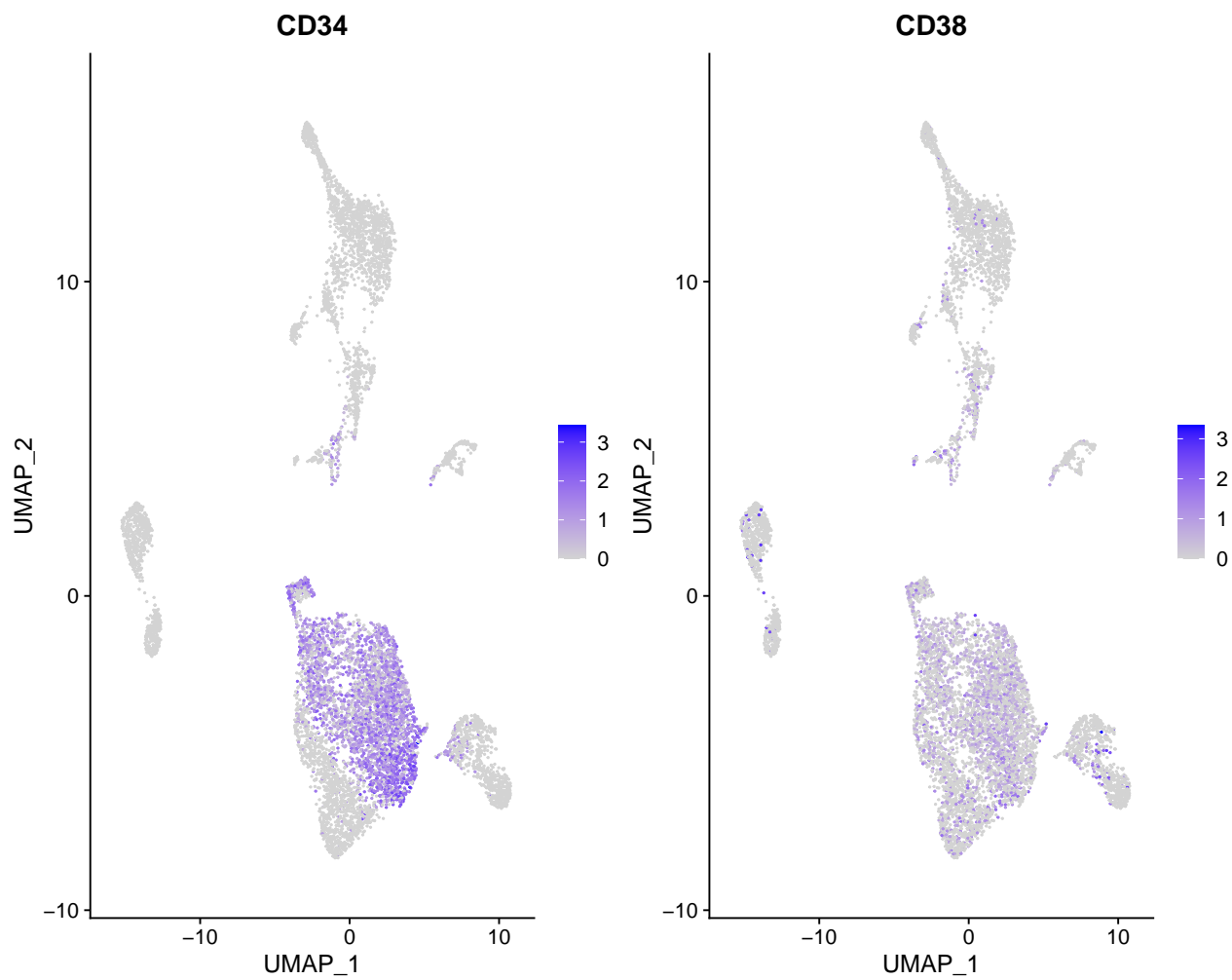
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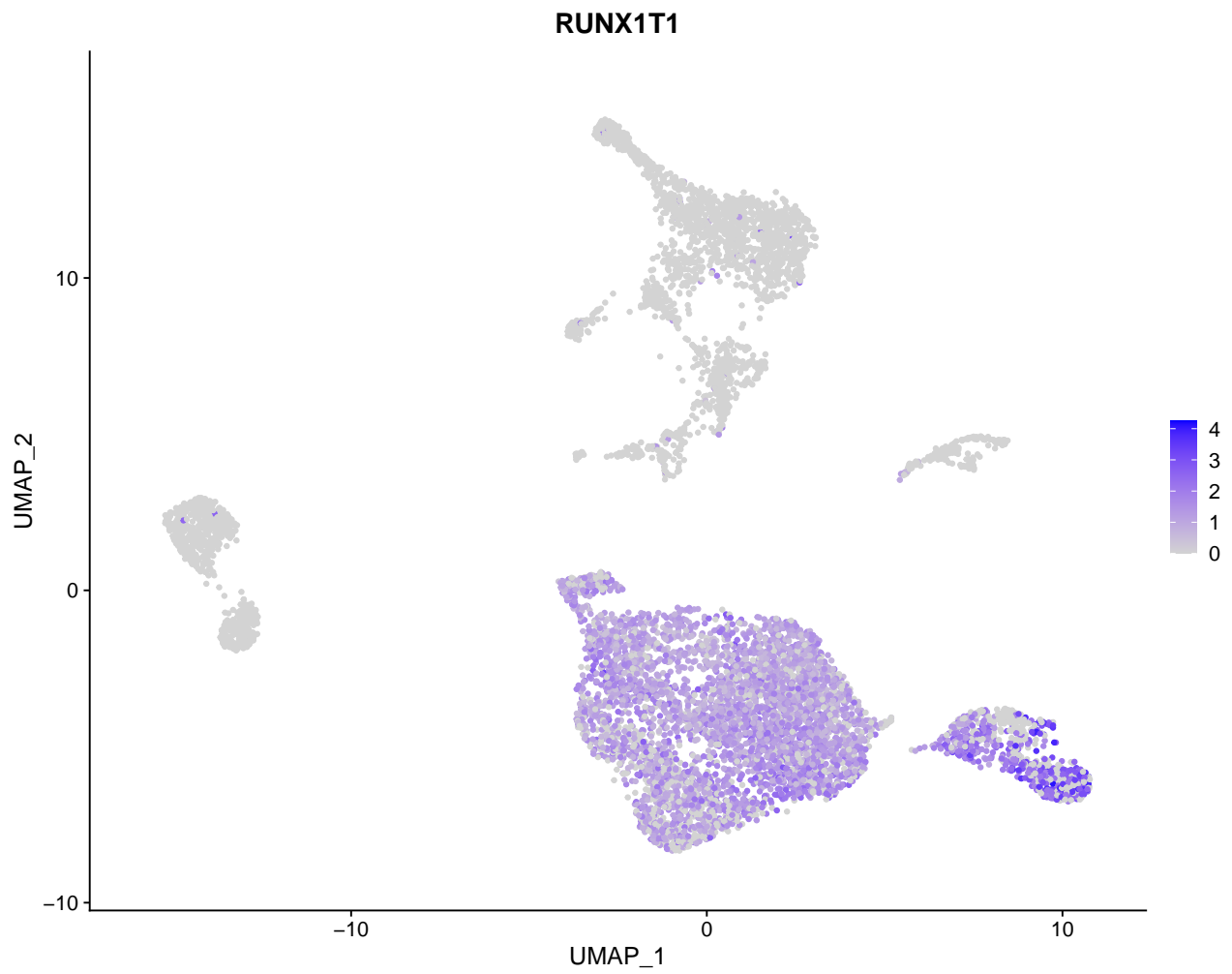
1. Put together both 34 and 38 libraries. Apply QC and dimensionality reduction.





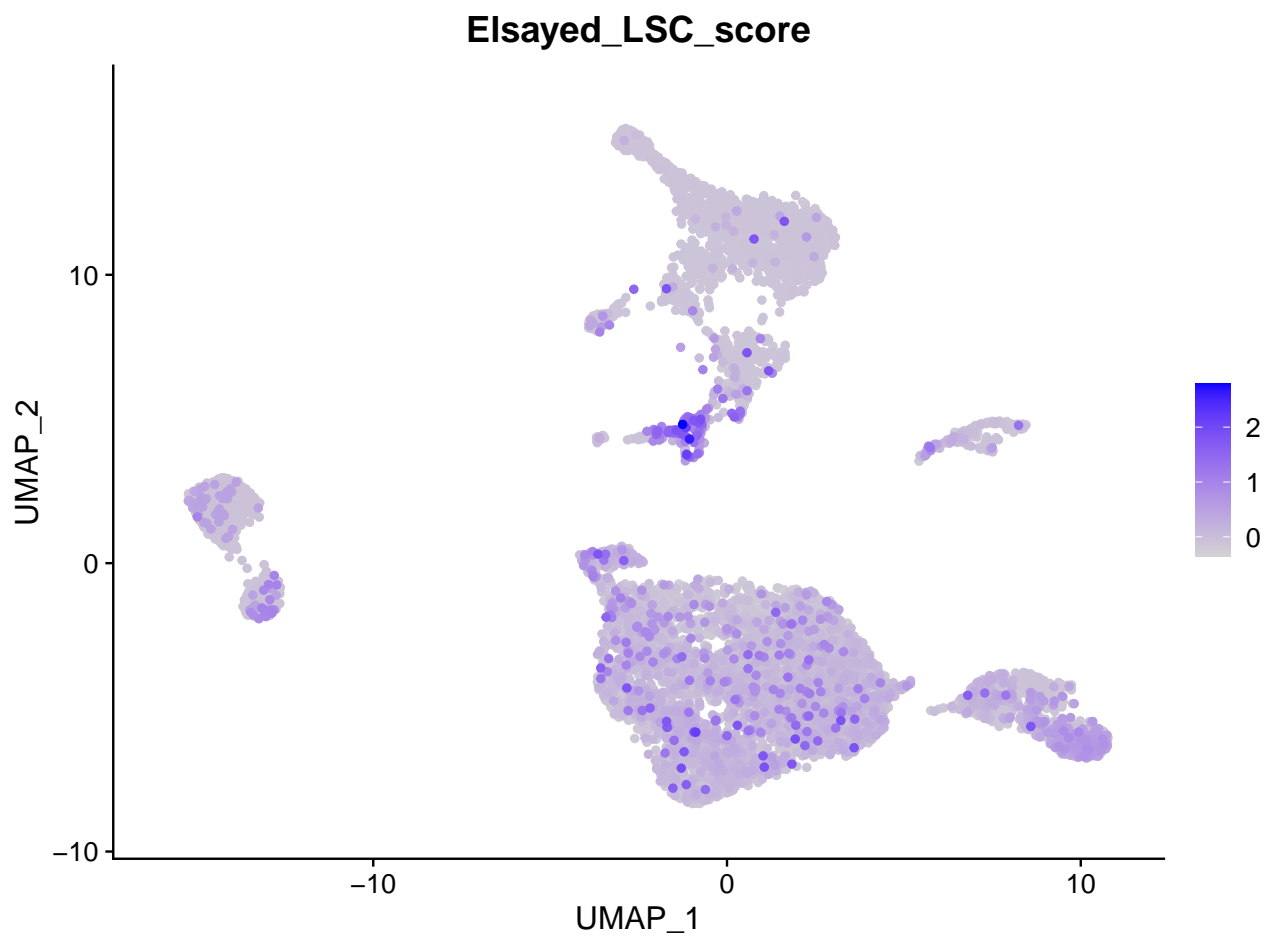


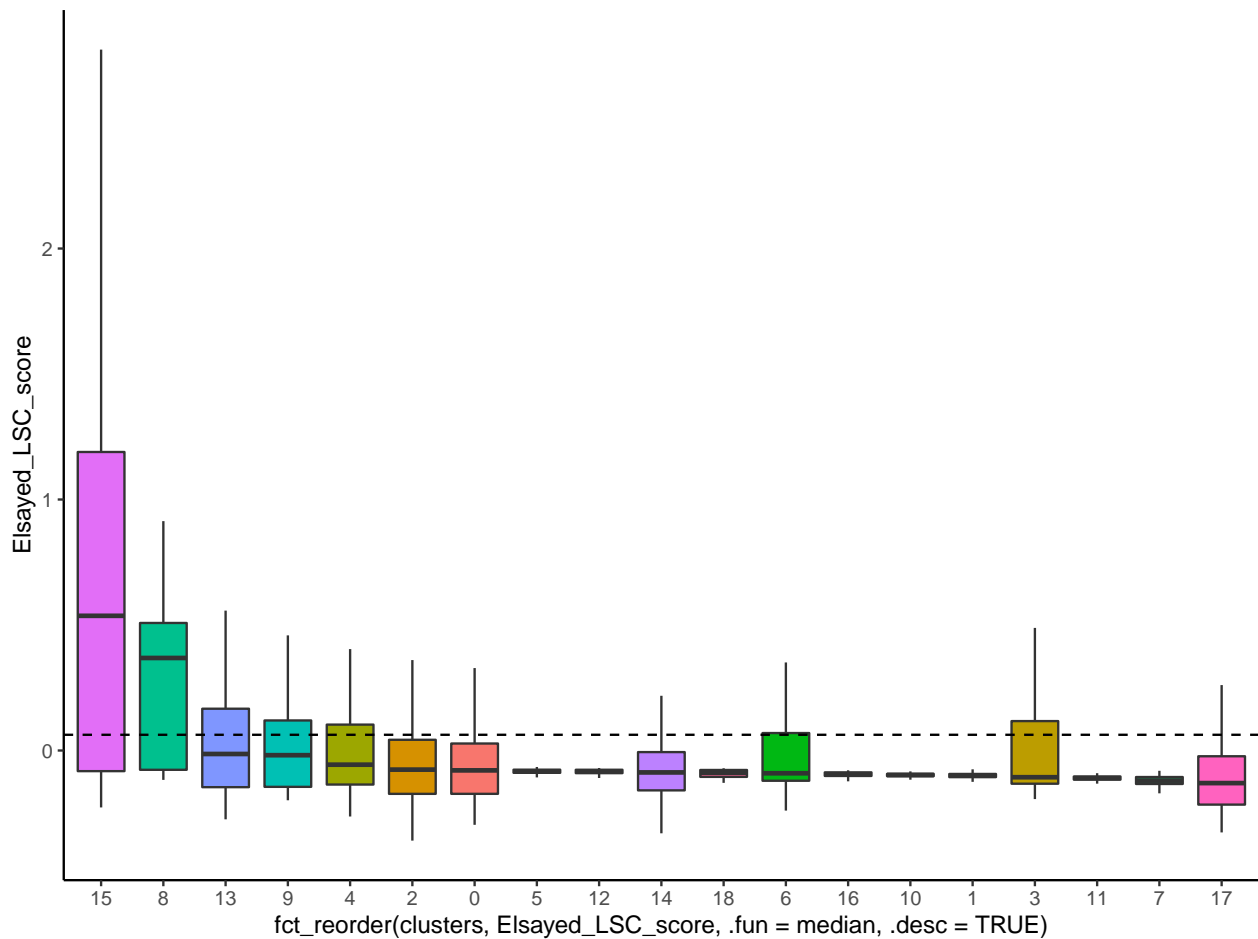
## Check for expression of malignant marker for t(8;21) RUNX1T1



## 2. Get the LSC6 score

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

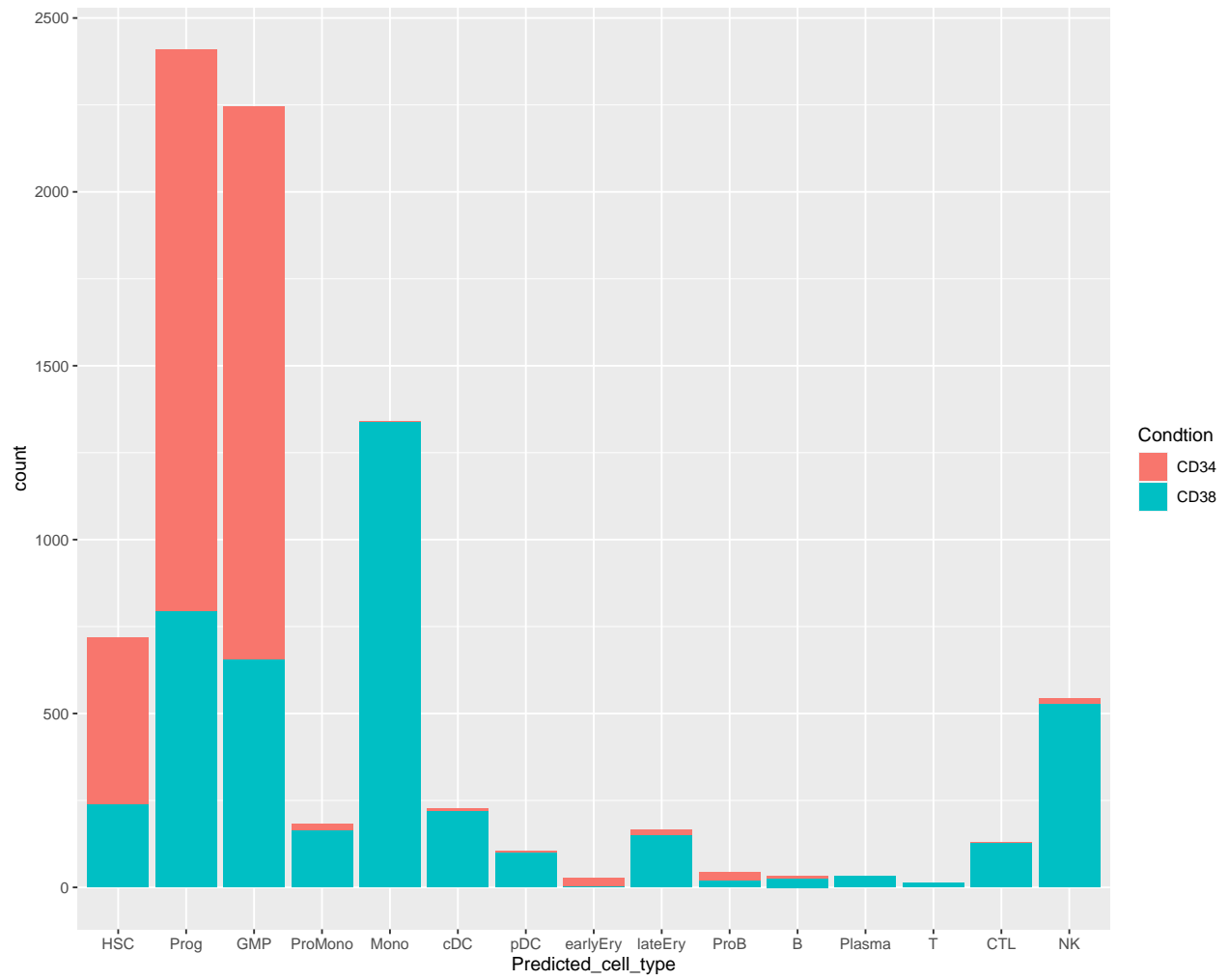




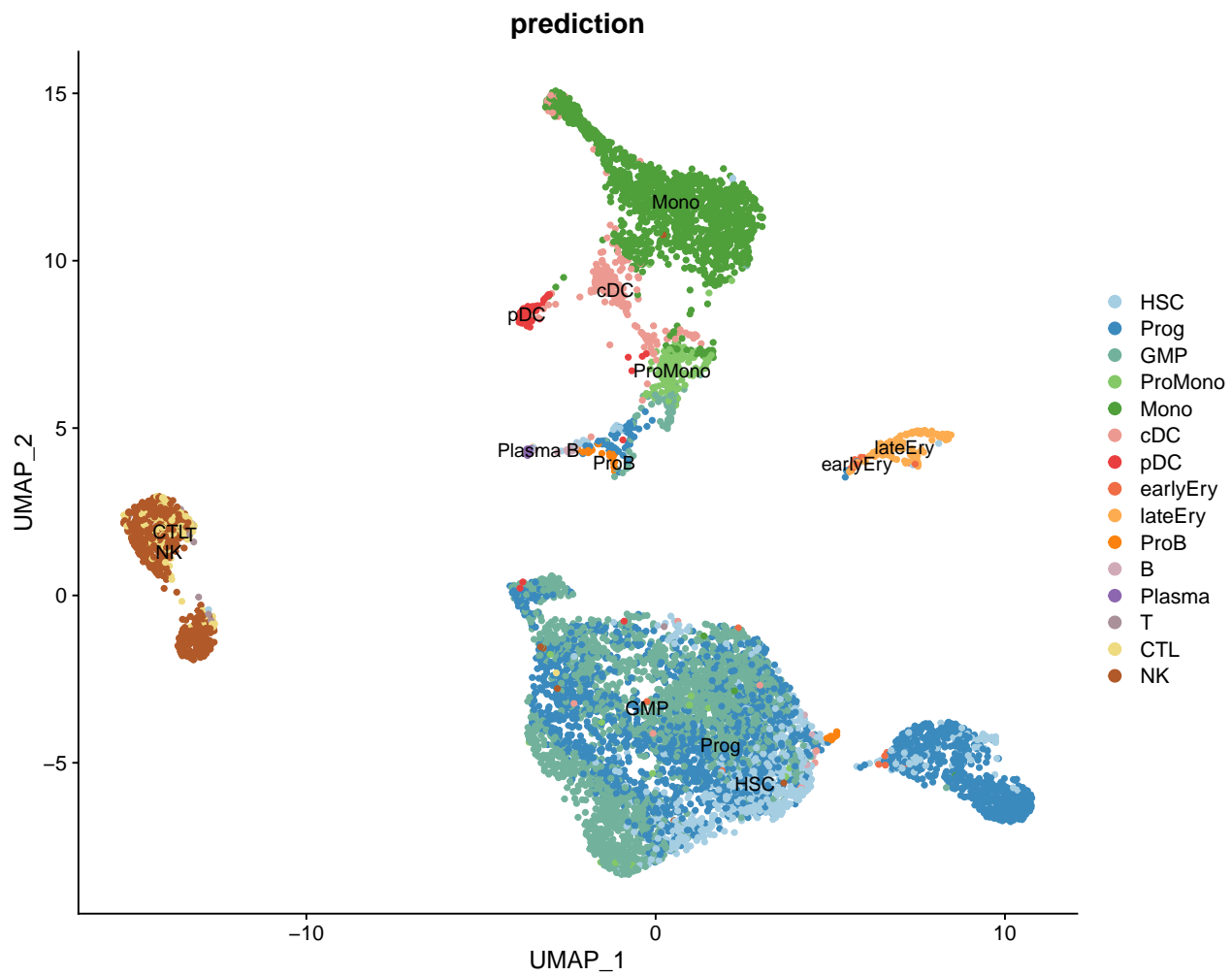
### 3. Predict the class of the cells using the markers and the expression of the BM cells form Van\_Galen paper

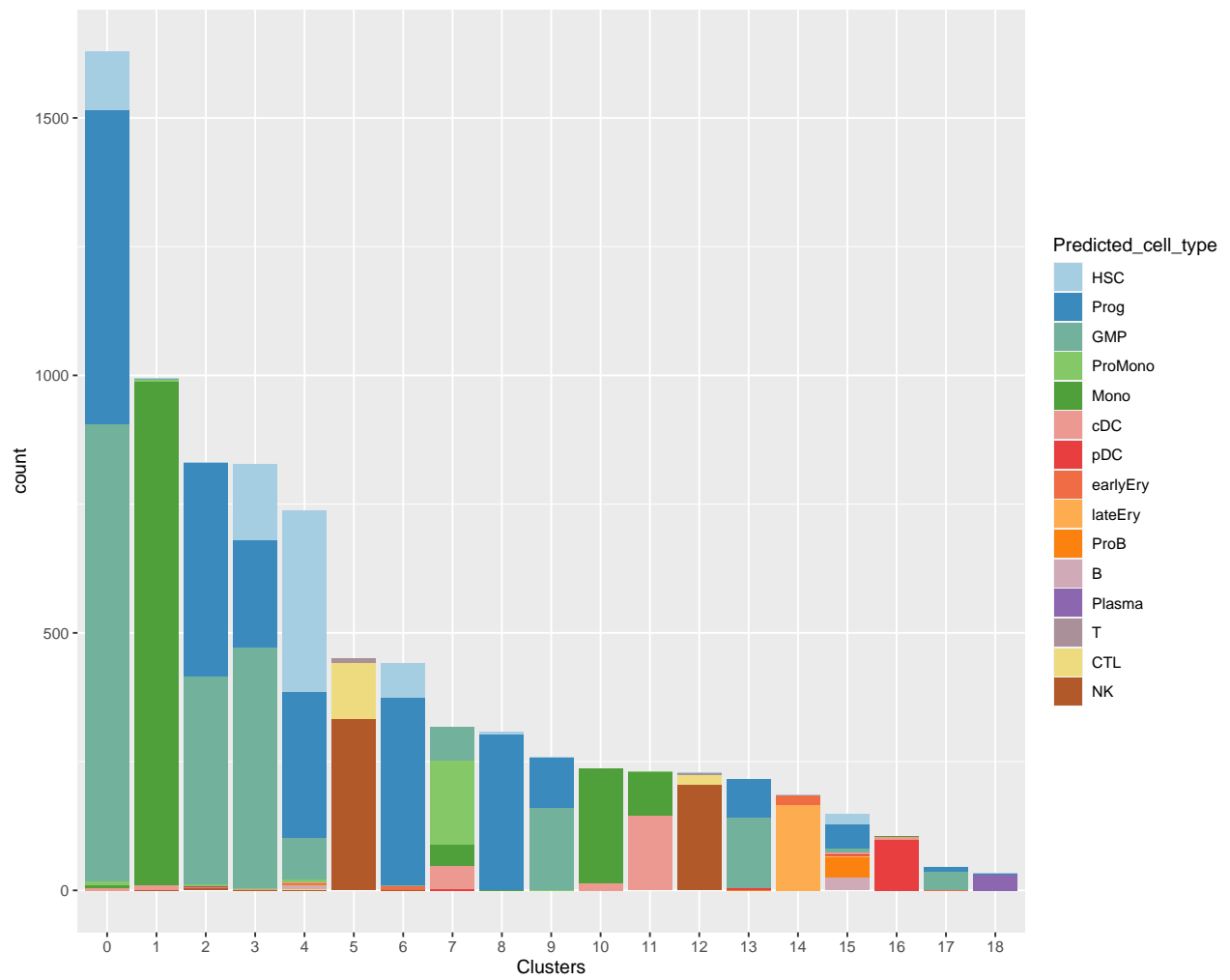
```
## Performing PCA on the provided reference using 1821 features as input.
## Projecting cell embeddings
## Finding neighborhoods
## Finding anchors
## Found 3841 anchors
## Filtering anchors
## Retained 2763 anchors
## Finding integration vectors
## Finding integration vector weights
## Predicting cell labels
##
##           HSC Prog  GMP ProMono Mono  cDC  pDC earlyEry lateEry ProB  B Plasma
##   CD34  482 1618 1593    22    1    9    8      25    18   26    8    0
##   CD38  238  792  654   162 1338 218  98      2    149   18   25   31
```

##		T	CTL	NK
##	CD34	1	5	16
##	CD38	13	126	527

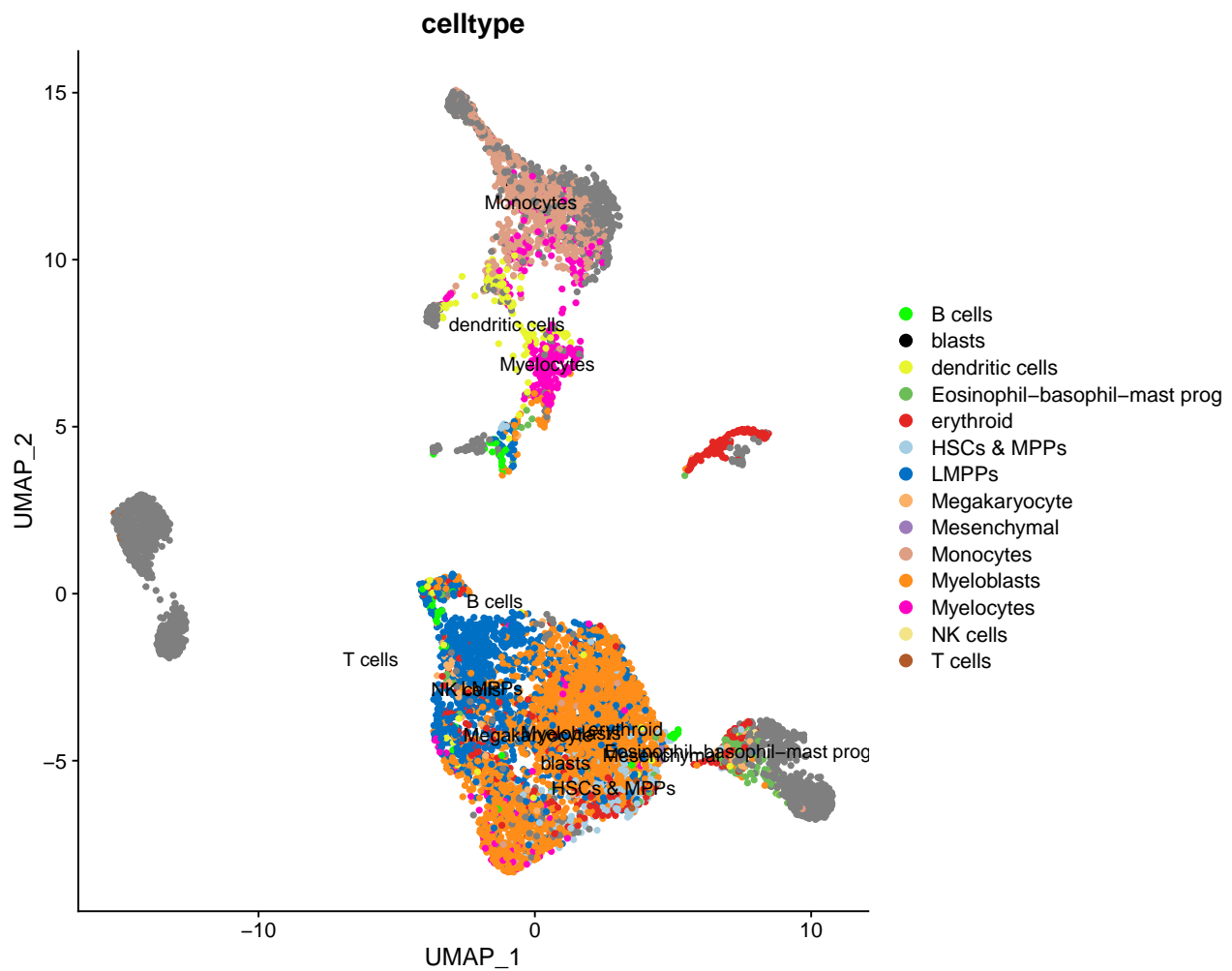


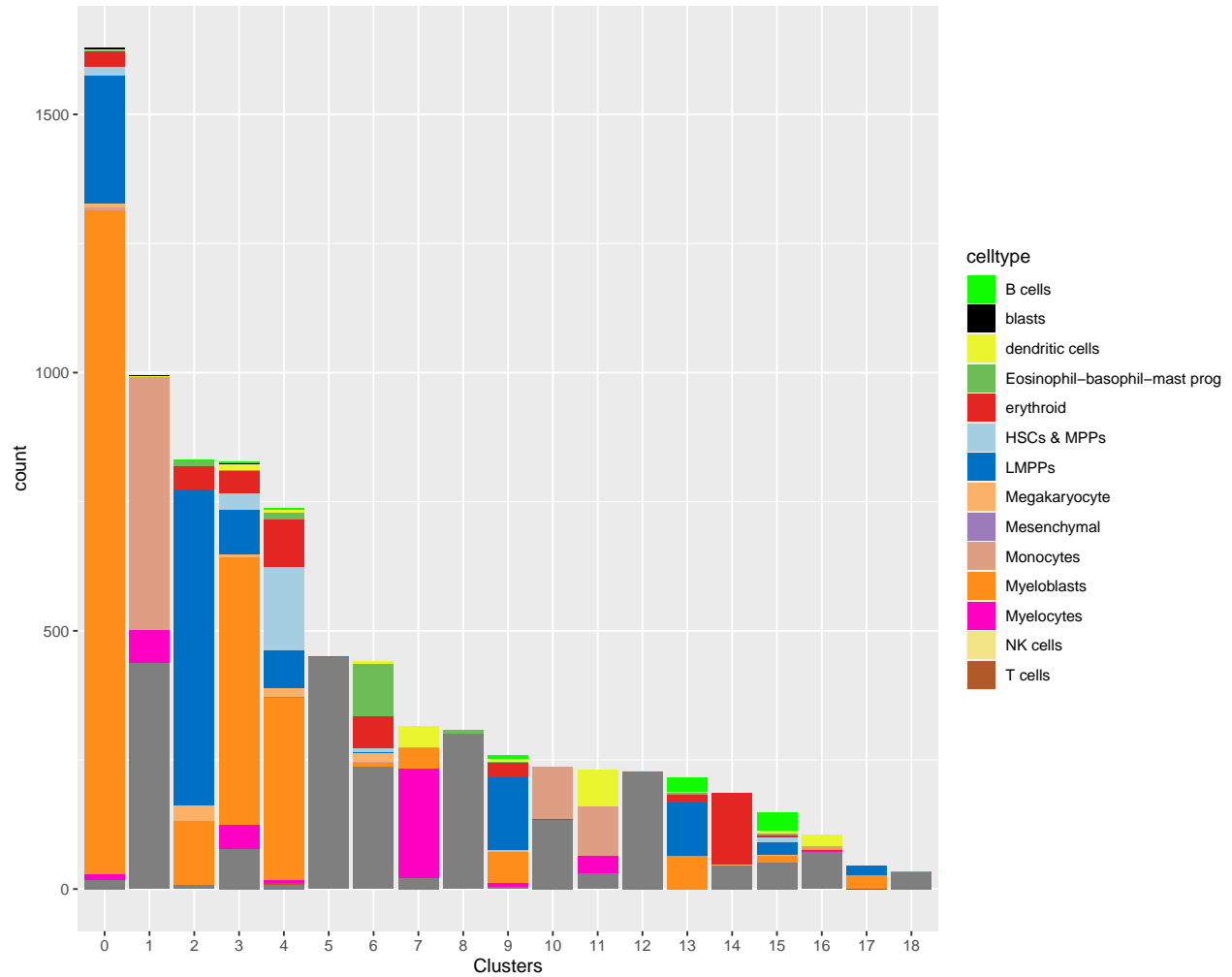






#### 4. Project the predictions from Velten onto our UMAP





Cluster 15 seems to be healthy residual cells, non expressing tumoral marker RUNX1T1. Clusters 8, 13 and 9 they express RUNX1T1, but have not predicted any HSC from both in-silico predictions. Cluster 4, spite of would be under the threshold, is way more enriched in HSC predicted cells. In addition, this is the cluster that colocalize with cluster LSC 2 from the paired Dx. Therefore we would consider this cluster as the more likely enriched on LSC