AML7_Rx

jtrincado

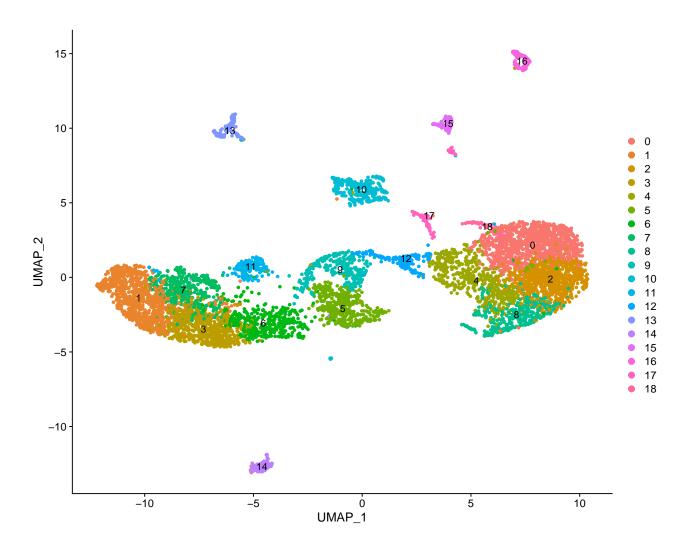
2022-02-09 12:32:52

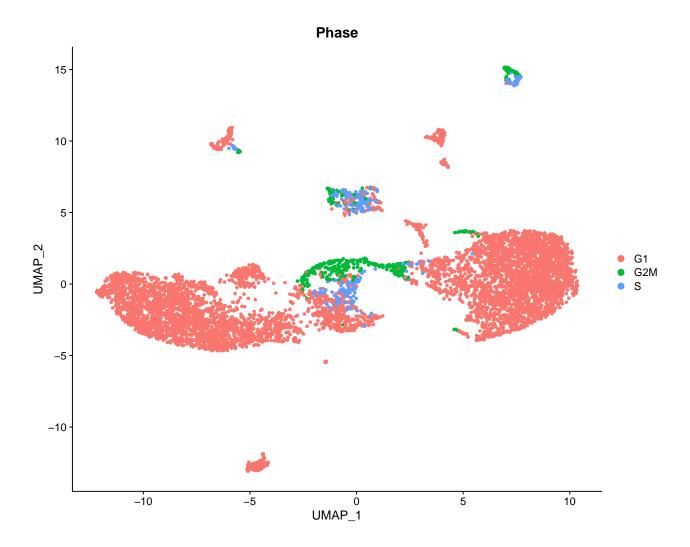
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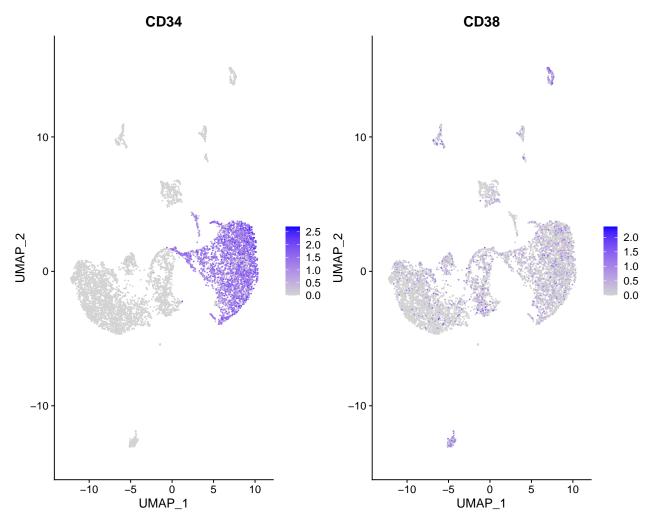
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1. Put together both 34 and 38 libraries. Apply QC and dimensionality reduction.

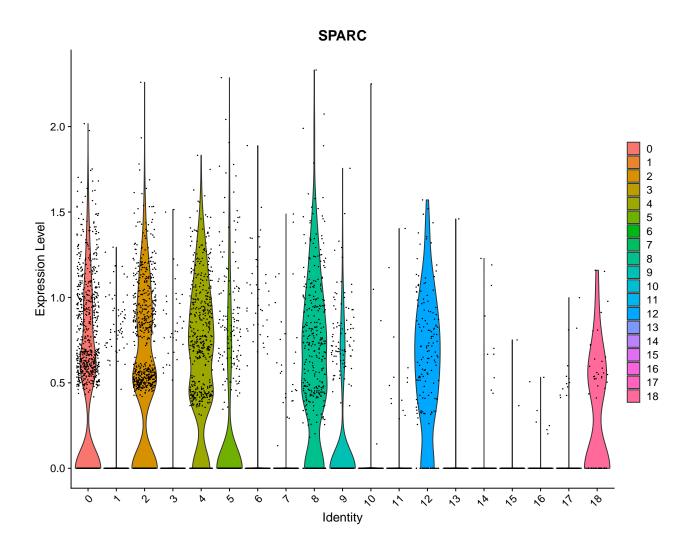
- ## Warning: The default method for RunUMAP has changed from calling Python UMAP via reticulate to the R
- ## To use Python UMAP via reticulate, set umap.method to 'umap-learn' and metric to 'correlation'
- ## This message will be shown once per session





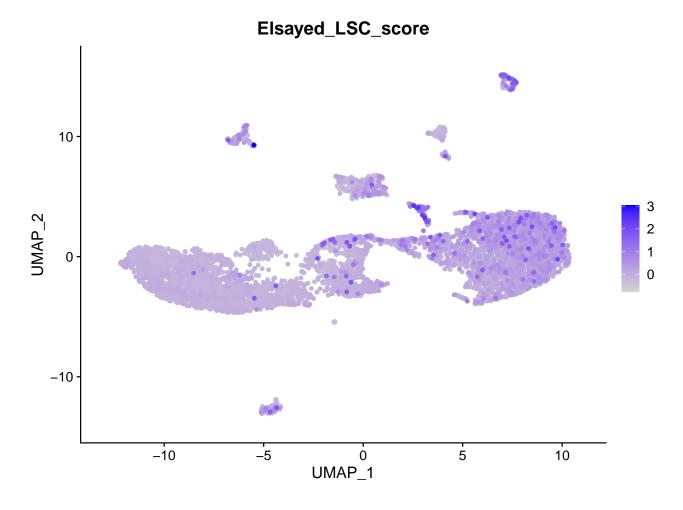


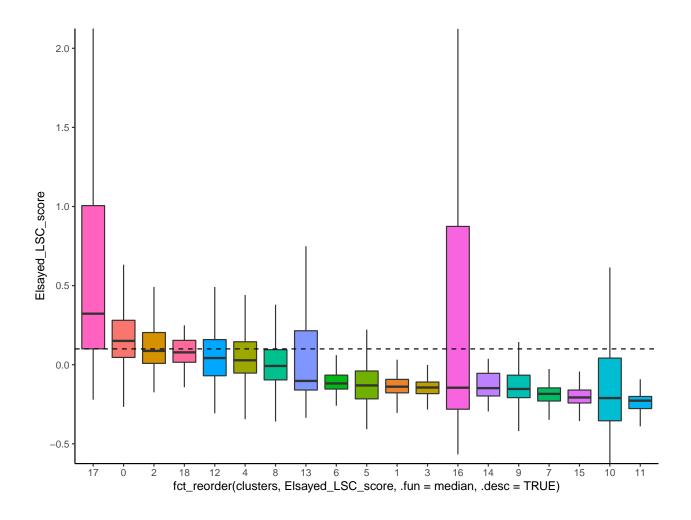
Check for expression of malignant marker for inv16 SPARC



2. Get the LSC6 score

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



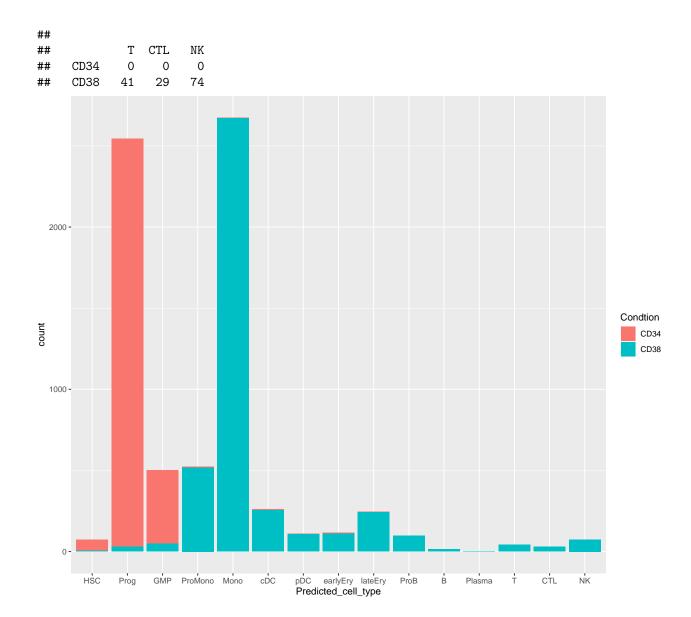


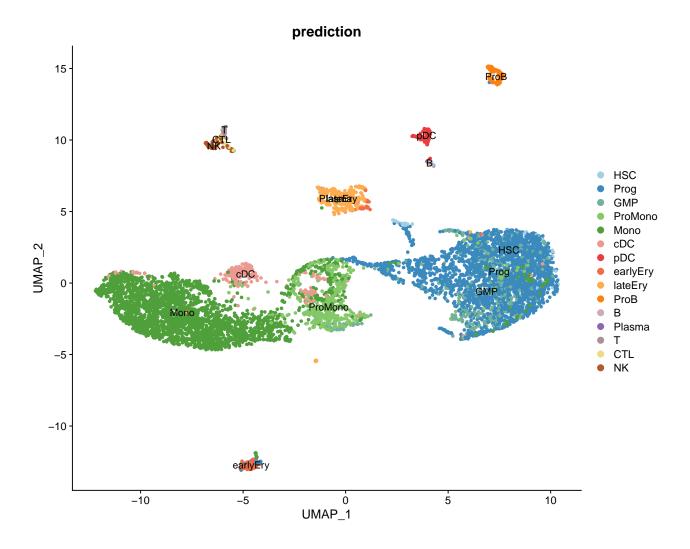
3. Predict the class of the cells using the markers and the expression of the BM cells form Van_Galen paper

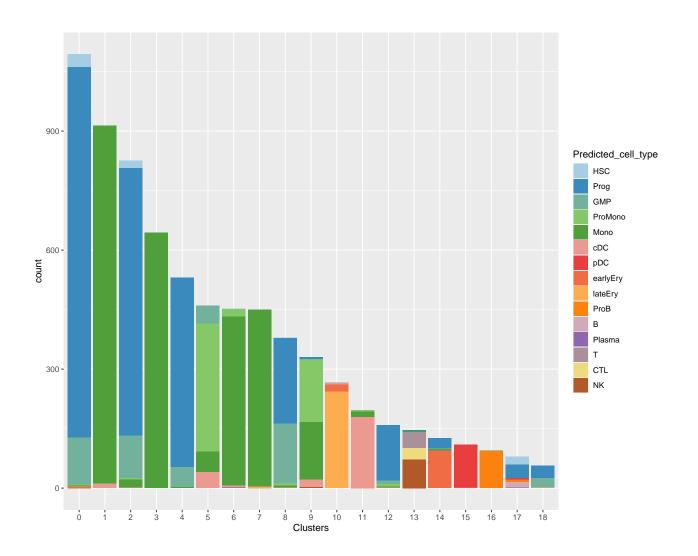
```
## Performing PCA on the provided reference using 1821 features as input.
```

- ## Projecting cell embeddings
- ## Finding neighborhoods
- ## Finding anchors
- ## Found 4361 anchors
- ## Filtering anchors
- ## Retained 3186 anchors
- ## Finding integration vectors
- ## Finding integration vector weights
- ## Predicting cell labels

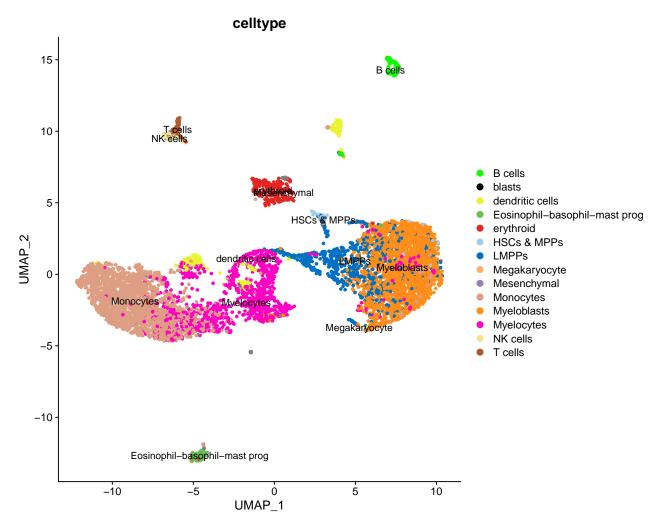
pDC earlyEry lateEry ProB ## HSC Prog GMP ProMono Mono \mathtt{cDC} B Plasma 456 0 ## **CD34** 5 6 0 0 109 CD38 30 48 518 2670 254 106 241 98 15 3

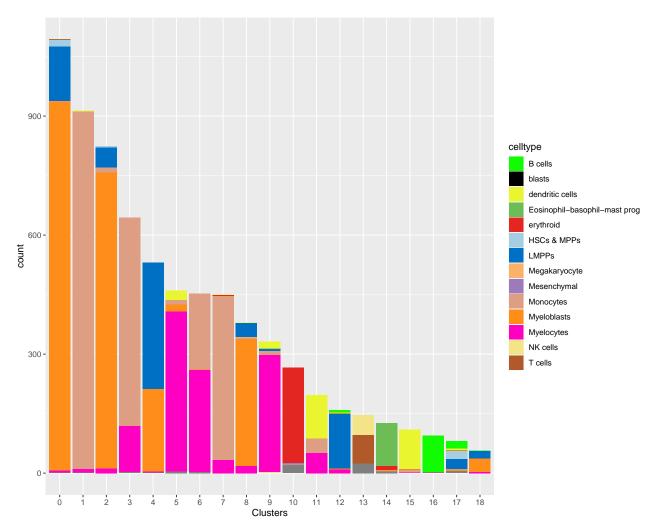






4. Project the predictions from Velten onto our UMAP





Cluster 17 and 0 are among the top according to LSC6 score and they bare a similar proportion of predicted HSCs. However, cluster 0 shows greater expression of malignant marker SPARC. This could suggest cluster 17 are healthy residual cells. In addition, cluster 0 is the one colocalizing with cluster 0 LSC from the paired Dx sample. Therefore, we propose cluster 0 as the one with LSC enrichment