

# Pseudotime for clusters 3.0 and 3.1

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April 8, 2020

## Load library

```
library(monocle3,verbose = FALSE)
library(dplyr,verbose = FALSE)
```

## Read the data

```
## Loads a sparse matrix RawCountsPseudotime
load("RawCountsPseudotime.rda")
dim(RawCountsPseudotime)
```

```
## [1] 20271 3127
```

## Load cluster data

```
## Loads a factor variable ClusterPseudotime containing cluster identities
load("ClusterPseudotime.rda")
## Change the name of variable to remind
DataClusters <- ClusterPseudotime

table(DataClusters)
```

```
## DataClusters
##      0.0      0.1      1.0      1.1     10.0     10.1     10.2
##      238      213      410      30       0       0       34
## 11.0_1_2    11.3    11.4    12.0    12.1    13.0    14.0
##       0       0       0       0       0       0       0
##      16.0     16.1     17.0     17.1     19.0     19.1      2.0
##      160      42      169      36       0       0     383
##      20.0     22.0     22.1    22.2_3    22.4     26.0     26.1
##       0       0       0       0       0       0       0
##      26.2    27.0_3     27.1     27.2    27.NA 28.0_2_3_4     28.1
##       0       0       0       0       0       0       0
##      29.0     29.1      3.0      3.1     30.0     31.0     33.0
##       0       0     253     119       0       0      90
##      34.0     35.0     35.1     36.0     37.0     37.1     38.0
##       0       0       0       0       0       0       0
##      39.0      4.0      4.1     40.0     41.0     42.0     42.1
##       0      206     142       0       0      40      27
```

```
##      43.0      44.0      45.0      46.0      48.0      49.0      5.0
##      0        0        0        0        0        0        0
##      5.1      50.0      51.0      51.1      52.0      53.0      54.0
##      0        0        0        0        0        0        0
##      55.0      56.0      57.0      59.0      6.0      60.0      61.0
##      0        43        0        0        0        0        0
##      62.0      63.0      7.0      8.0      8.1      8.2      8.3
##      0        0        0        0        77       27       15
##      10.0.0  10.0.1_4  10.0.2_3  10.0.5  10.1.0  10.1.1  8.0.0
##      45        59        57        9        33       21      103
##      8.0.1
##      46
```

```
length(DataClusters)
```

```
## [1] 3127
```

## Convert DataClusters to a matrix format for input to Monocle

```
DataCluster.ID <- matrix(as.numeric(levels(DataClusters))[DataClusters], ncol = 1)
```

```
## Warning in matrix(as.numeric(levels(DataClusters))[DataClusters], ncol = 1): NAs
## introduced by coercion
```

```
rownames(DataCluster.ID) <- names(DataClusters)
```

```
colnames(DataCluster.ID) <- "Cluster.IDs"
```

```
DataCluster.ID[1:10,]
```

```
## cele-001-008.GATCAGTCAT cele-001-027.ACTCCGCCAA cele-001-042.TTCCTAGACC
##      0        0        0
## cele-001-046.TTCTACGCCA cele-001-047.TTCGCTGCCT cele-001-047.ATGGAAGCAT
##      0        0        0
## cele-001-064.AAGCTGACCT cele-001-065.GCCATCAACT cele-001-068.ACGGCAACCA
##      0        0        0
## cele-001-071.GTCATTGCGC
##      0
```

## Generate matrix of gene short names for Monocle

```
geneNames <- matrix(rownames(RawCountsPseudotime), ncol = 1)
```

```
rownames(geneNames) <- rownames(RawCountsPseudotime)
```

```
colnames(geneNames) <- "gene_short_name"
```

```
head(geneNames)
```

```
##      gene_short_name
## aap-1 "aap-1"
## aat-1 "aat-1"
## aat-2 "aat-2"
## aat-3 "aat-3"
## aat-4 "aat-4"
## aat-5 "aat-5"
```

## Initiate Monocle object

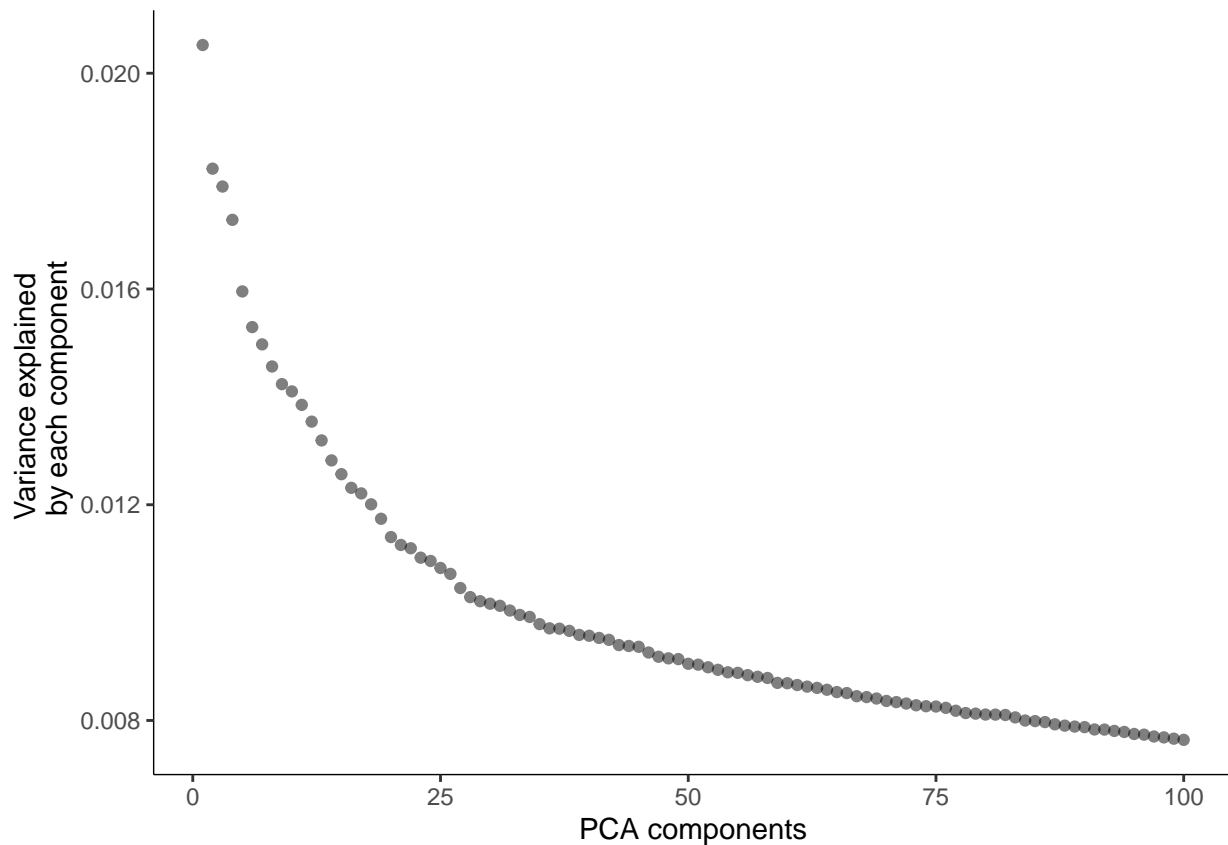
```
cds <- new_cell_data_set(expression_data = RawCountsPseudotime,  
                        cell_metadata = DataCluster.ID,  
                        gene_metadata = geneNames)  
  
cds <- cds[,names(ClusterPseudotime[ClusterPseudotime %in% c("3.0","3.1")])]  
  
colData(cds)
```

```
## DataFrame with 372 rows and 2 columns  
##               Cluster.IDs      Size_Factor  
##               <numeric>      <numeric>  
## cele-001-002.ACGACCAATA      3 0.512985911598468  
## cele-001-003.GGATTCTATC      3 0.625482822036728  
## cele-001-003.TCCAGAAGGT      3 0.764978990980171  
## cele-001-013.GATATGGTCT      3 0.787478373067823  
## cele-001-030.CGTCCGTCCT      3 0.503986158763407  
## ...  
## cele-010-072.CCTGACGTTC      3.1 1.50745859987269  
## cele-010-081.AAGTCTTCCG      3.1 1.69645340940897  
## cele-010-088.TGCCAGATGG      3.1 2.12394166907436  
## cele-010-090.CAGGCGCCAT      3.1 2.78542350245133  
## cele-010-092.GAAGTCCGTC      3.1 1.16996786855791  
## Column Cluster.IDs contains the original DataCluster IDs  
colData(cds)$Cluster.IDs <- factor(colData(cds)$Cluster.IDs)  
colData(cds)
```

```
## DataFrame with 372 rows and 2 columns  
##               Cluster.IDs      Size_Factor  
##               <factor>      <numeric>  
## cele-001-002.ACGACCAATA      3 0.512985911598468  
## cele-001-003.GGATTCTATC      3 0.625482822036728  
## cele-001-003.TCCAGAAGGT      3 0.764978990980171  
## cele-001-013.GATATGGTCT      3 0.787478373067823  
## cele-001-030.CGTCCGTCCT      3 0.503986158763407  
## ...  
## cele-010-072.CCTGACGTTC      3.1 1.50745859987269  
## cele-010-081.AAGTCTTCCG      3.1 1.69645340940897  
## cele-010-088.TGCCAGATGG      3.1 2.12394166907436  
## cele-010-090.CAGGCGCCAT      3.1 2.78542350245133  
## cele-010-092.GAAGTCCGTC      3.1 1.16996786855791
```

## Step 1: Normalize and pre-process the data

```
cds <- preprocess_cds(cds, num_dim = 100)  
plot_pc_variance_explained(cds)
```



```
cds <- preprocess_cds(cds, num_dim = 20)
```

## Step 2: Reduce the dimensions using UMAP

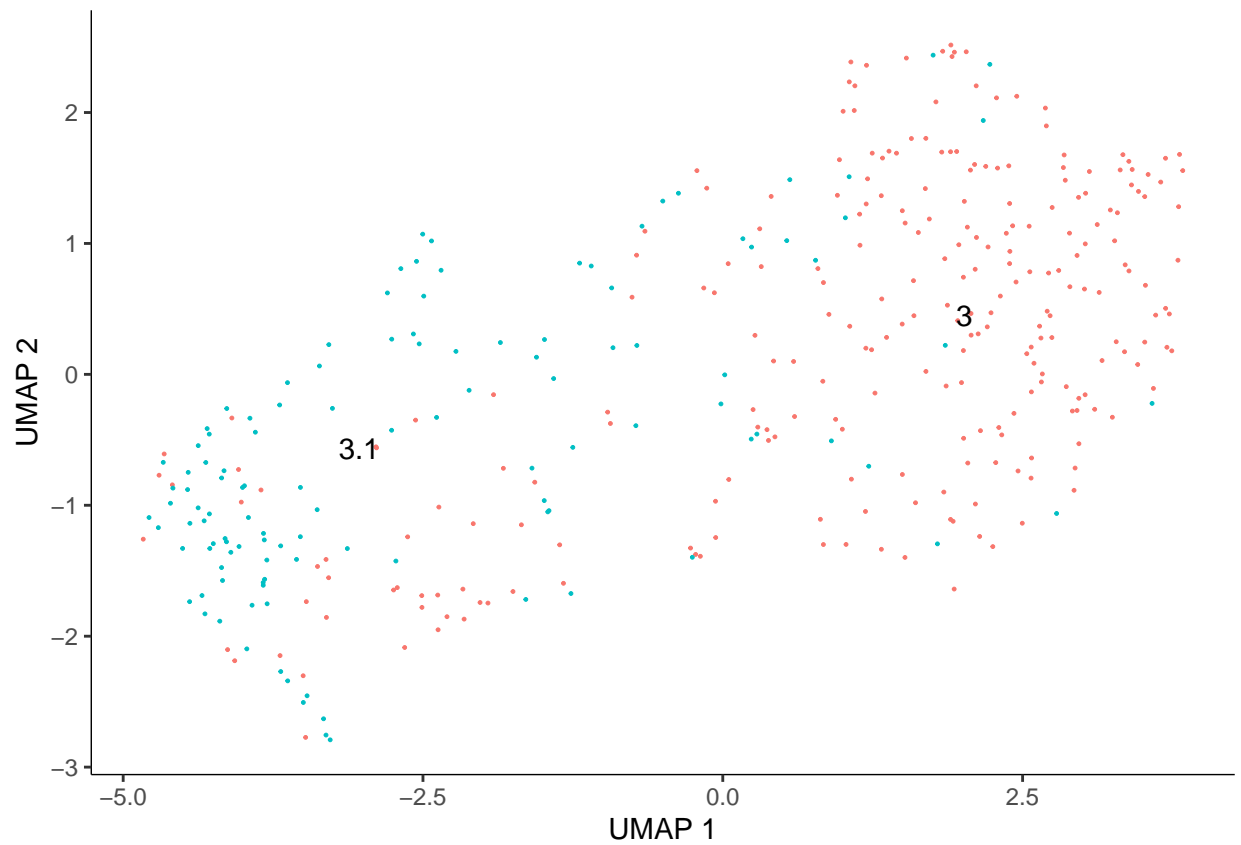
```
cds <- reduce_dimension(cds, umap.min_dist = 0.1, cores = 8)
```

```
## No preprocess_method specified, using preprocess_method = 'PCA'
```

```
## Note: reduce_dimension will produce slightly different output each time you run it unless you set 'umap.random_seed'
```

```
plot_cells(cds, color_cells_by = "Cluster.IDs", group_label_size = 4, cell_size = 0.5)
```

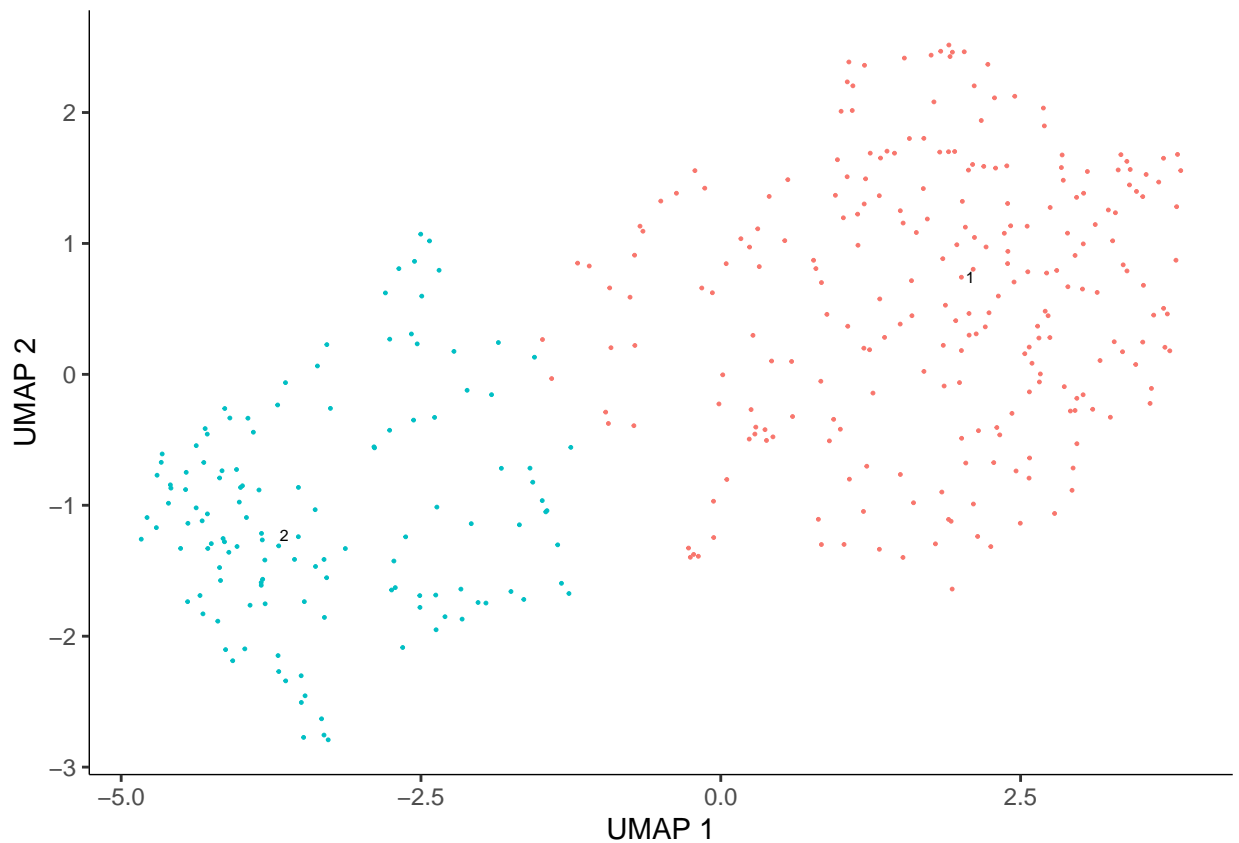
```
## No trajectory to plot. Has learn_graph() been called yet?
```



##Step 3: Cluster the cells

```
# cds = cluster_cells(cds, resolution=0.02)
cds = cluster_cells(cds, resolution=0.01)
plot_cells(cds, cell_size = 0.5)
```

## No trajectory to plot. Has learn\_graph() been called yet?



#### Step 4: Learn a graph and order cells

```
cds <- learn_graph(cds)
```

```
##
```

```
|
|
|
|=====| 100%
```

```
## With Shiny
```

```
# cds <- order_cells(cds)
```

```
# #
```

```
# save(file = "cdsRoots.rda", cds) #Save object state after selection of roots
```

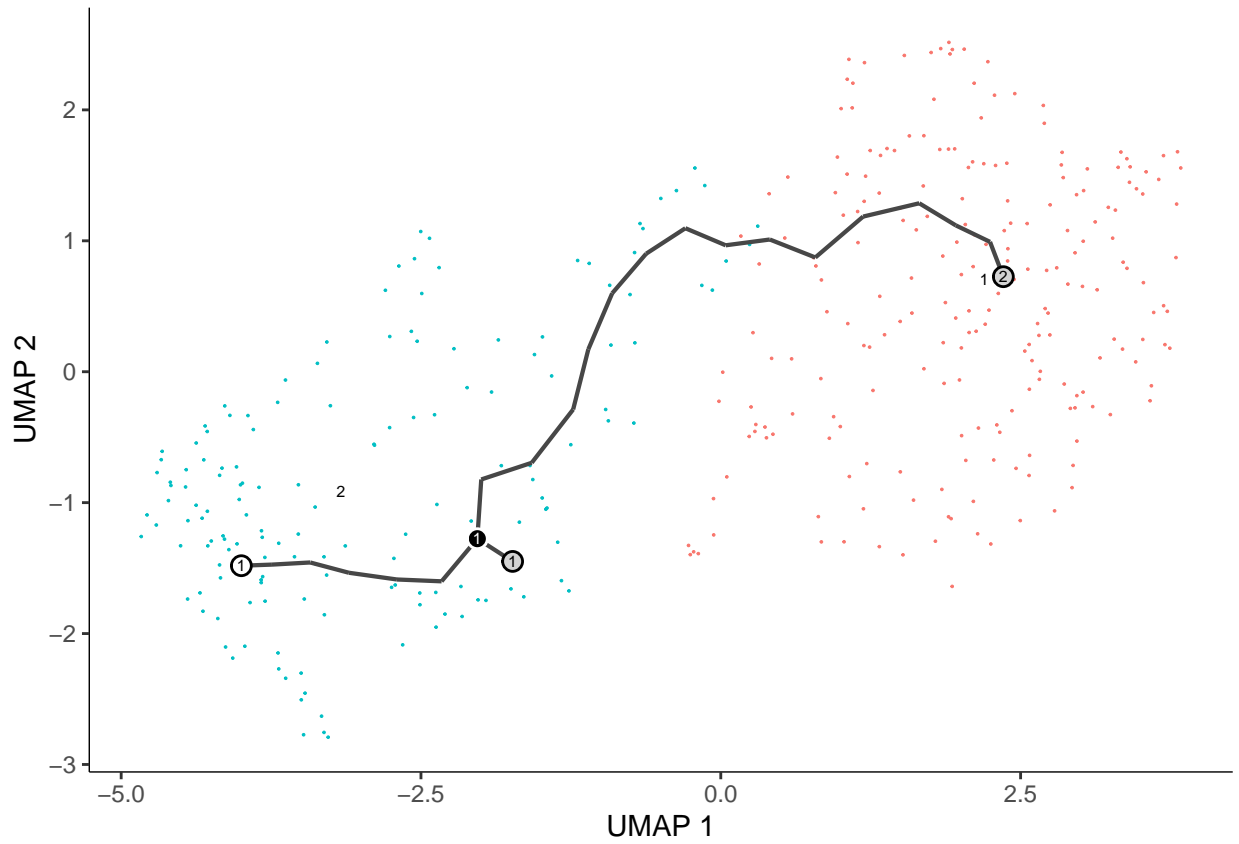
```
## Without Shiny
```

```
load("cdsRoots.rda") #Load object
```

```
print(cds@principal_graph_aux$UMAP$root_pr_nodes)
```

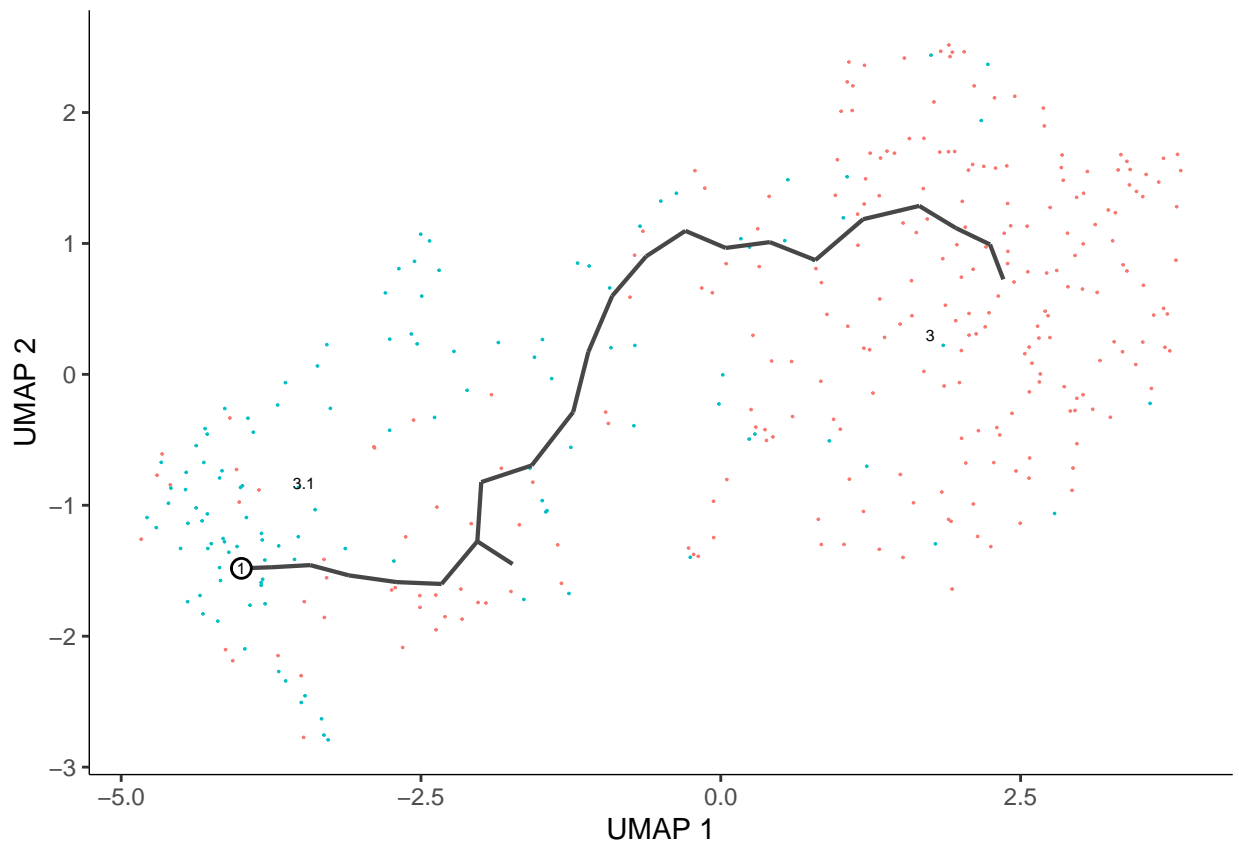
```
## [1] "Y_22"
```

```
plot_cells(cds)
```



Plot the pseudotime graph

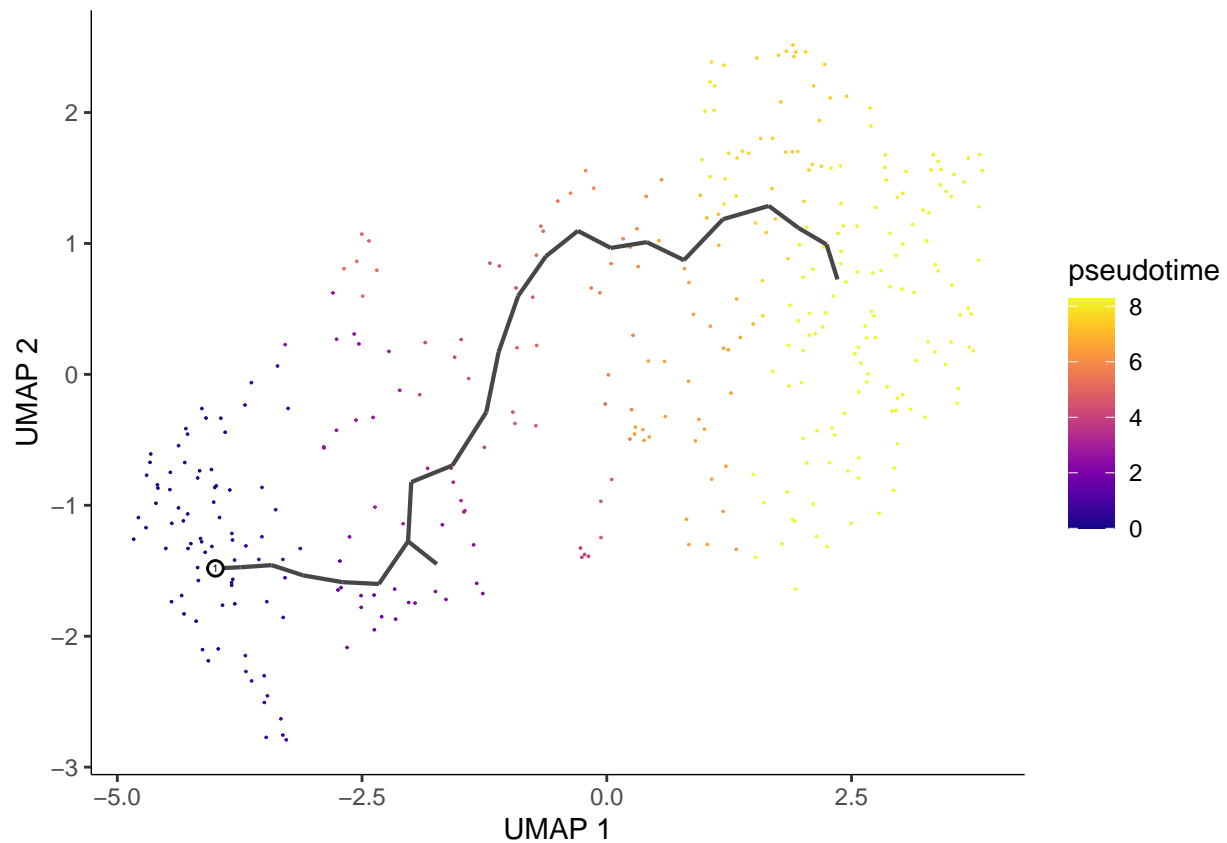
```
plot_cells(cds,  
            color_cells_by = "Cluster.IDs",  
            label_groups_by_cluster=FALSE,  
            label_leaves=FALSE,  
            label_branch_points=FALSE)
```



### Plot the pseudotime graph

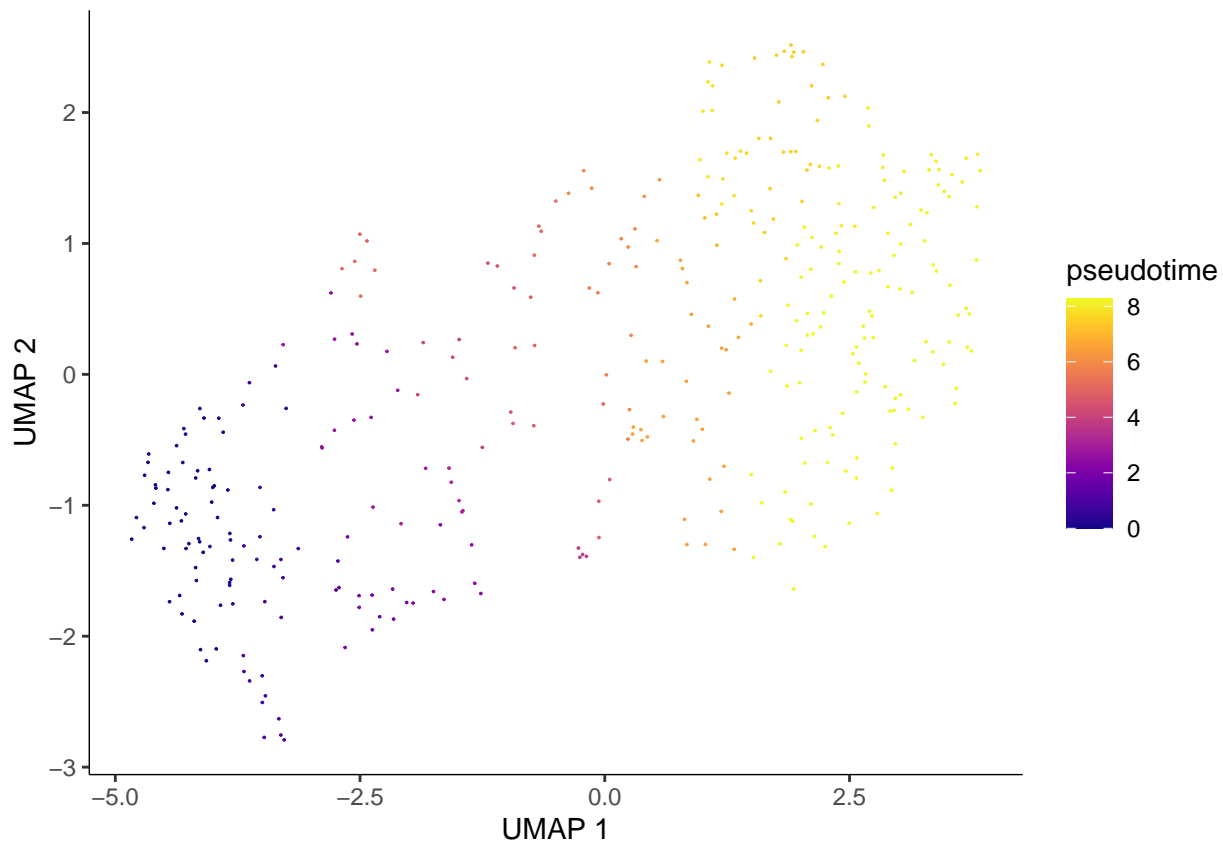
```
plot_cells(cds,
           color_cells_by = "pseudotime",
           label_cell_groups=FALSE,
           label_leaves=FALSE,
           label_branch_points=FALSE,
           graph_label_size=1.5)
```





Plot the pseudotime graph without trajectory

```
plot_cells(cds,
            color_cells_by = "pseudotime",
            label_cell_groups=FALSE,
            label_leaves=FALSE,
            label_branch_points=FALSE,
            graph_label_size=1.5,
            show_trajectory_graph = FALSE)
```



## Correlation Analysis

### Cluster 3

```
cluster3.cellNames <- rownames(pData(cds))[pData(cds)$Cluster.IDs %in% c(3, 3.1)]

cds_3 <- cds[,cluster3.cellNames]

cds3_pg <- graph_test(cds_3, neighbor_graph="principal_graph", cores=4, verbose = F)
cds3_genes <- cds3_pg %>%
  filter(q_value < 0.05) %>%
  arrange(morans_I) %>%
  select(gene_short_name)
```

```
cds3_genes$gene_short_name
```

##	[1]	T07A9.10	alh-11	inx-10	npp-3	stg-2	dgn-1
##	[7]	B0035.15	dpt-1	Y105E8A.2	Y76B12C.3	ubc-22	ddx-35
##	[13]	K03H1.11	hot-8	parg-1	lin-61	F01E11.18	R10E4.1
##	[19]	lmtr-3	T26C11.4	F56D1.2	nlp-6	K06H7.2	mes-2
##	[25]	haf-2	W03D8.3	glrx-22	F46F11.8	agt-1	acd-3
##	[31]	F54D5.12	C46H3.2	cebp-1	gly-20	pfd-6	srp-7
##	[37]	mms-19	dhhc-8	nft-1	riok-3	gab-1	F40F11.3
##	[43]	F25F8.1	F54D5.7	col-17	atg-4.2	pcaf-1	nape-1
##	[49]	slx-1	Y57A10A.26	sym-2	T10B11.7	W09C5.1	K05F1.5

```

## [55] mup-4      lin-15A    pyc-1      acl-1      C53B4.2    R10E11.6
## [61] Y66D12A.9  pfd-3      Y39A1A.24  T14B4.8    Y105E8A.13 gfi-2
## [67] F13B12.6   nuo-1      B0416.5    K07C5.6    C18A3.2    sqv-8
## [73] snap-1     spat-2     lin-66     rps-27     Y105E8A.11 ribo-1
## [79] hsp-3      mev-1      pan-1      gbb-2      rpl-7      C24A3.2
## [85] lin-39     dip-2      xbp-1      Y54F10AM.5 cyn-16     F21F3.6
## [91] rps-16     glb-25     pam-1      F31C3.3    nra-1      ekl-5
## [97] rpl-39     rpl-11.2   tag-80     Y71H2AM.5  lmn-1      gdi-1
## [103] klp-7      unc-115    sacy-1     ZC395.10   dnj-5      hmr-1
## [109] ser-4      F32B4.5    des-2      rbm-28     cul-1      chc-1
## [115] lin-10     inf-1      F46F11.10  F52C9.3    hsp-1      let-716
## [121] B0252.8    psd-1      set-15     gars-1     cnnm-1     atp-5
## [127] rpl-5      ceh-44     sur-6      C08B6.8    Y53F4B.18  ZK1307.8
## [133] ZK673.2    fkh-9      let-504     sas-4      fln-1      brp-1
## [139] C03A3.1    T04F8.2    fbx1-1     F14F9.5    eak-7      M03E7.1
## [145] odr-2      ftt-2      rpl-27     hmg-11     vha-1      ztf-13
## [151] top-2      imp-2      usp-46     let-70     tba-2      nlr-1
## [157] gopc-1     F29B9.11   lgc-49     his-35     exp-2      gop-3
## [163] rgef-1     F55C12.4   lin-53     elf-1      tag-196    dnj-18
## [169] nkb-1      emb-5      rpl-30     pab-1      ZC434.7    idhg-1
## [175] rpl-13     somi-1     aldo-1     cmd-1      R106.5     acr-14
## [181] nsf-1      rpl-26     T27C4.1    C09G1.5    ubl-1      trap-4
## [187] hex-2      rpt-6      ras-1      Y94H6A.8   mlc-4      lpd-7
## [193] C26D10.6   rps-25     F44E7.4    mrpl-24     lin-52     clr-1
## [199] K08F4.1    vig-1      Y119C1B.3  gon-1      rpl-16     rps-9
## [205] mthf-1     daf-21     lfe-2      tmc-1      ensa-1     F36A2.7
## [211] ncx-1      W09D10.1   ykt-6      rpl-34     inst-1     T20F5.7
## [217] Y105E8A.3  R02E12.5   smc-3      cyn-3      ama-1      madd-4
## [223] C25H3.11   rpl-6      W05B5.1    atp-2      nono-1     cct-8
## [229] sup-1      hipr-1     his-72     npp-7      cct-4      mig-22
## [235] rps-8      mel-11     Y66D12A.10 unc-13     K10B2.4    anc-1
## [241] frpr-4     rpl-3      F02A9.1    F28E10.1   mai-1      vha-14
## [247] F12F6.1    cct-1      brf-1      mma-1      mex-3      uig-1
## [253] rpy-1      C36B7.5    rps-0      rpl-43     lin-25     oaz-1
## [259] tbc-9      Y57G11C.9  flp-13     rpl-29     set-20     vha-13
## [265] vha-2      eif-3.E    Y97E10C.1  rpl-38     rps-14     rpl-25.2
## [271] prpf-4     C27A7.5    tbb-2      ser-2      rps-12     F40F8.1
## [277] F19G12.2   skr-1      spe-15     tin-44     eef-2      Y39B6A.36
## [283] nlp-9      Y55F3AM.5  unc-40     T25F10.3   pas-4      mom-5
## [289] ZC190.4    F26H9.2    R13A5.9    trap-1     unc-83     hse-5
## [295] unc-25     chdp-1     his-37     rpl-22     R12B2.6    tba-1
## [301] T19D12.6   rps-1      rpl-32     Y110A2AR.3 rpl-9      T10C6.6
## [307] rpl-41.2   W05E10.5   rpc-2      pcca-1     act-4      eef-1A.1
## [313] clic-1     rps-13     igcm-4     marc-4     iff-2      rps-11
## [319] rpl-10     rpn-12     gpb-1      patr-1     rla-1      K05B2.2
## [325] snrp-200   nlp-21     Y37E3.8    aqp-7      cdh-8      rsp-5
## [331] sir-2.2    C53H9.2    Y44A6D.2   far-1      rla-0      rpl-18
## [337] rpl-4      B0041.5    baf-1
## 20271 Levels: 2L52.1 2RSSE.1 4R79.2 6R55.2 aagr-1 aagr-2 aagr-3 aagr-4 ... zyx-1

```

```
# Plot a few genes
```

```

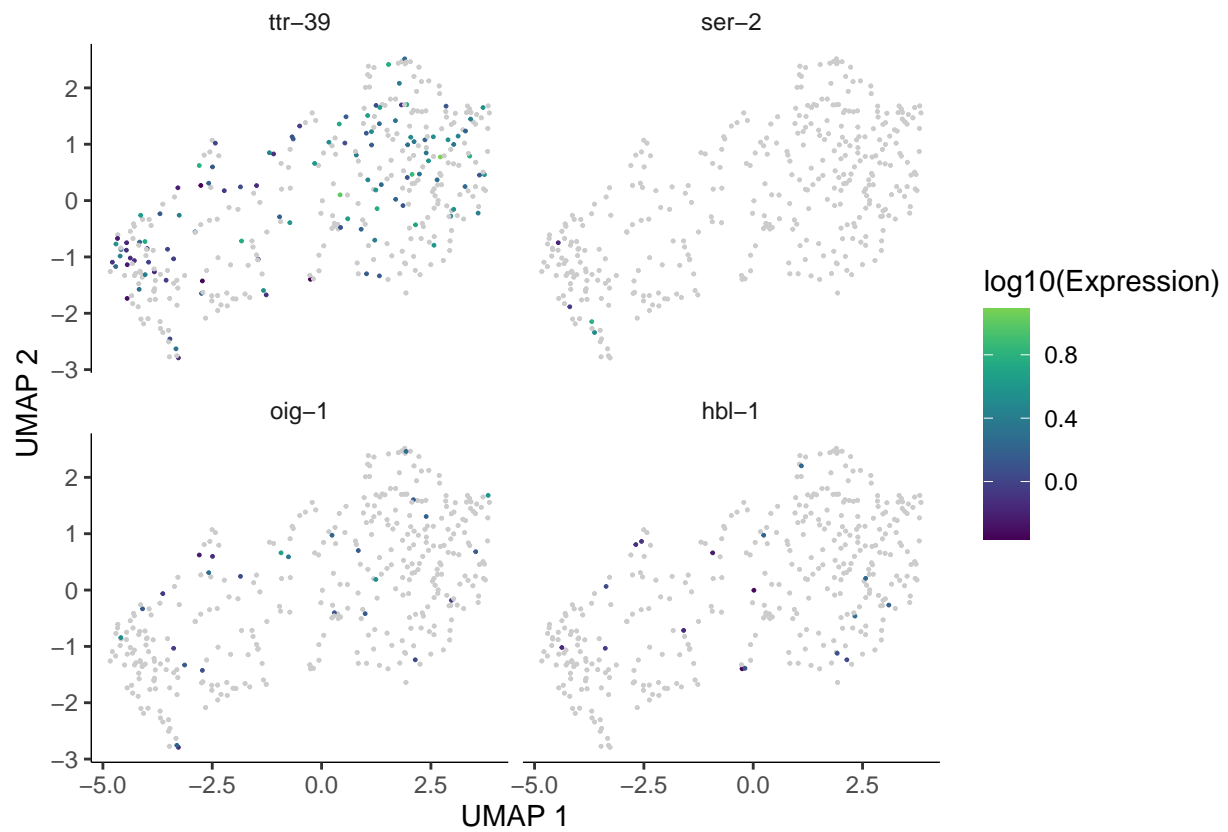
plot_cells(cds_3, genes = c("ttr-39", "ser-2", "oig-1", "hbl-1"),
  show_trajectory_graph=FALSE,
  label_cell_groups=FALSE,

```

```

label_leaves=FALSE,
cell_size = .5
)

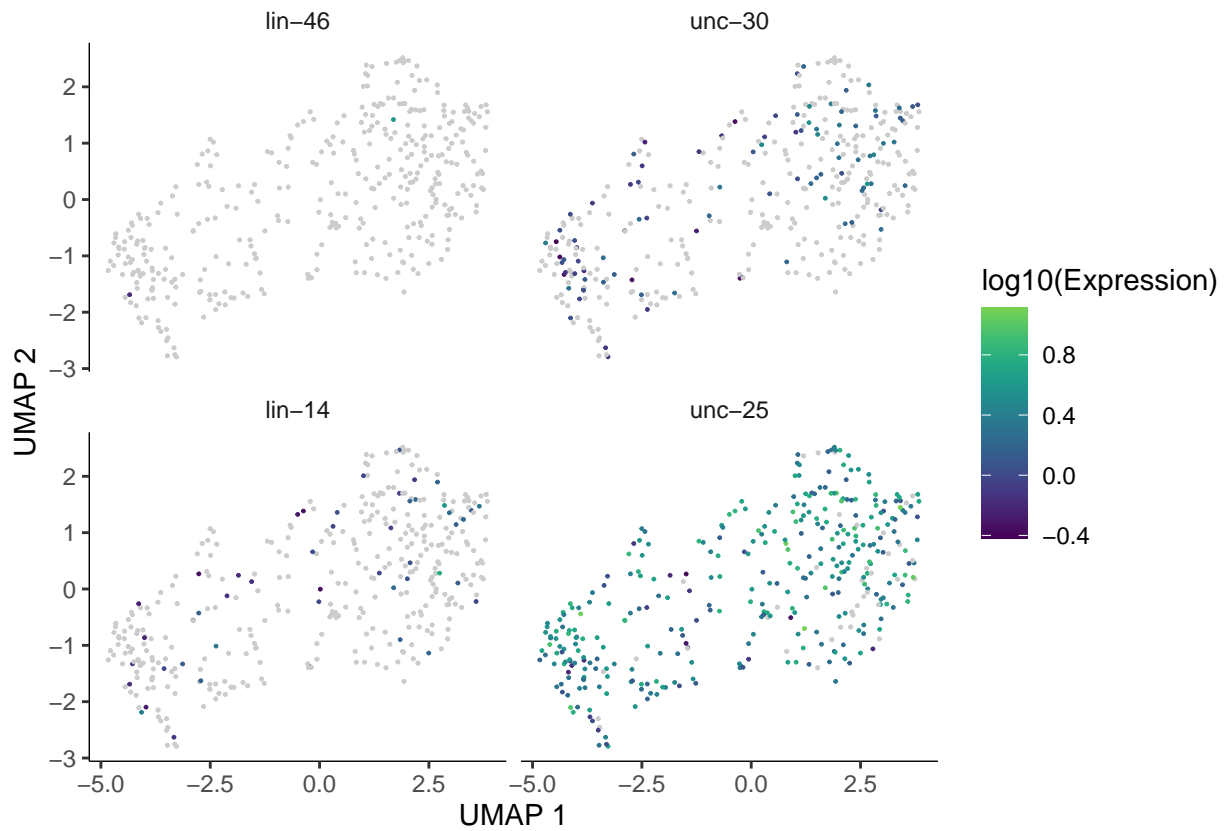
```



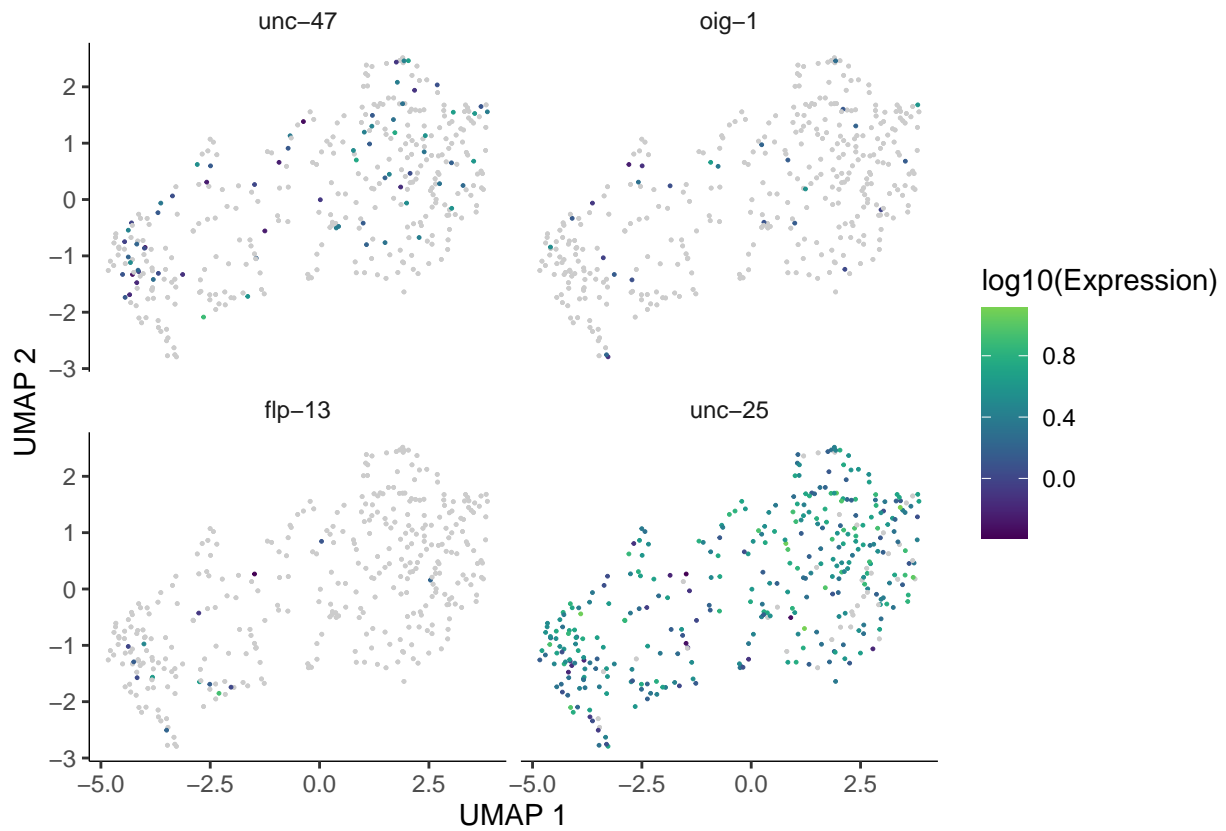
```

# Plot a few genes
plot_cells(cds_3, genes= c("lin-46", "unc-30", "lin-14","unc-25"),
  show_trajectory_graph=FALSE,
  label_cell_groups=FALSE,
  label_leaves=FALSE,
  cell_size = .5
)

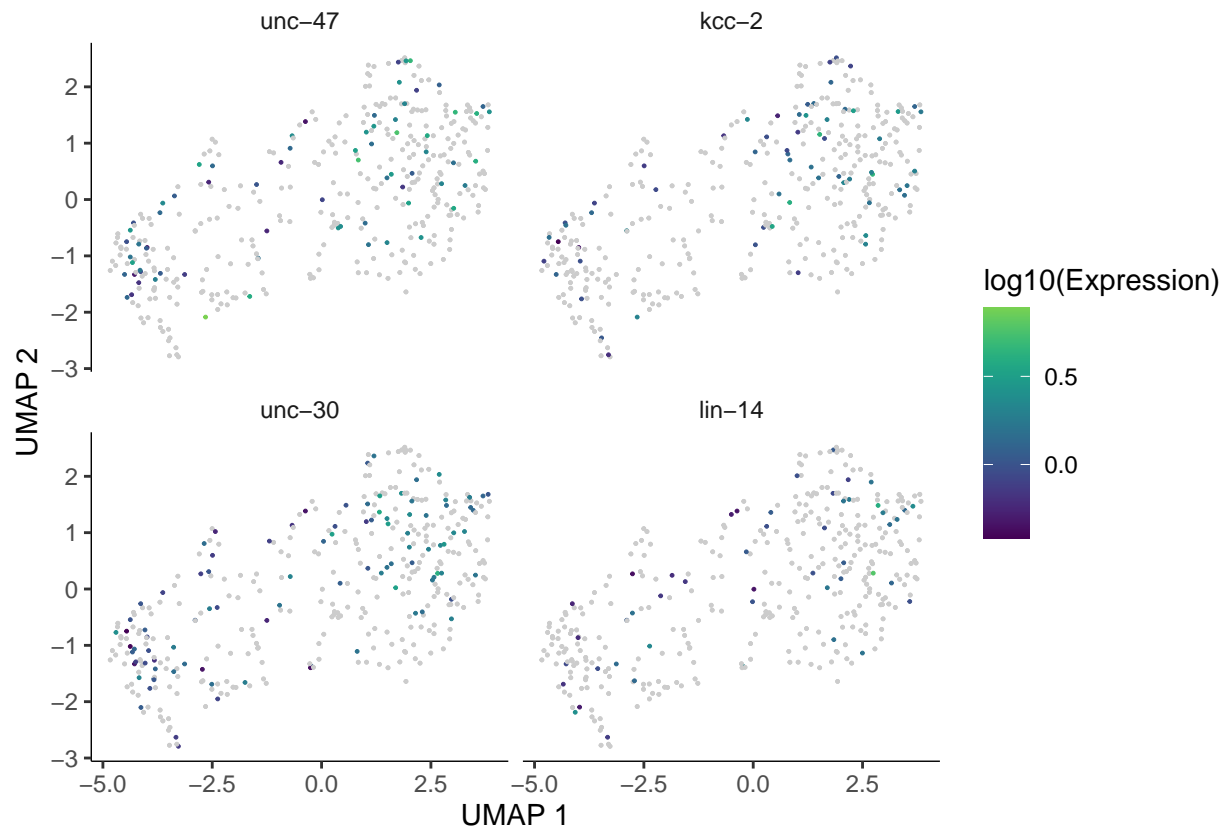
```



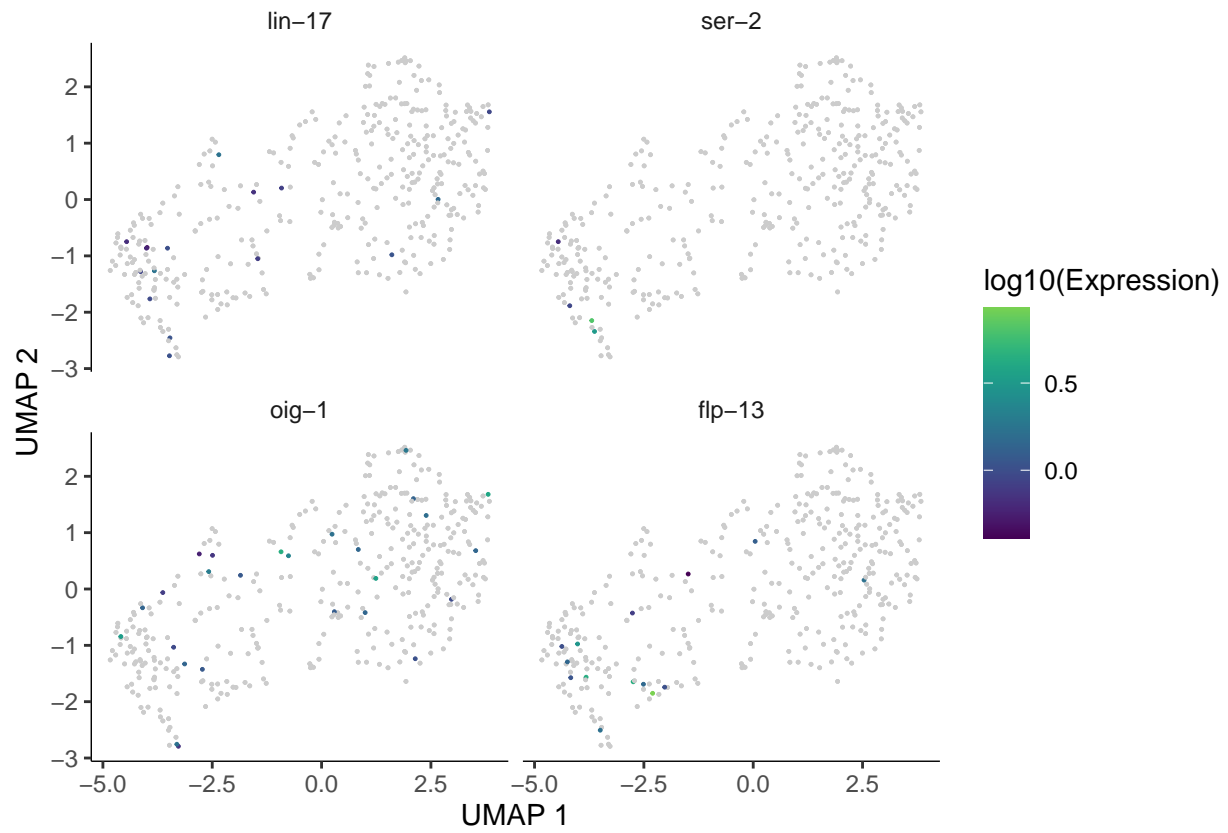
```
# Plot a few genes
plot_cells(cds_3, genes= c( "unc-47", "oig-1", "flp-13", "unc-25"),
  show_trajectory_graph=FALSE,
  label_cell_groups=FALSE,
  label_leaves=FALSE,
  cell_size = .5
)
```



```
# Plot a few genes
plot_cells(cds_3, genes= c("unc-47", "kcc-2", "unc-30", "lin-14"),
  show_trajectory_graph=FALSE,
  label_cell_groups=FALSE,
  label_leaves=FALSE,
  cell_size = .5
)
```



```
# Plot a few genes
plot_cells(cds_3, genes= c("lin-17", "ser-2", "oig-1", "flp-13"),
  show_trajectory_graph=FALSE,
  label_cell_groups=FALSE,
  label_leaves=FALSE,
  cell_size = .5
)
```



```
cds_3_lineage_cds <- cds_3[rowData(cds_3)$gene_short_name %in% c("unc-30", "unc-47", "kcc-2", "oig-1",
plot_genes_in_pseudotime(cds_3_lineage_cds,
min_expr=0.05)
```



