Vignette 1 - Understanding the Results of MCIA

Max Mattessich

Joaquin Reyna

Anna Konstorum

9/1/2022

Running MCIA on NCI60 Data and Basic Visualization

NIPALS-MCIA includes a sample multi-omics dataset modified from data collected on the NCI-60 cancer cell lines [CITE: Meng, 2016]. This can be used to illustrate low-dimensional plotting with the global factors.

First, we compute the first 10 global factors for the dataset:

data(NCI60) # this creates the dataset as `data_blocks`
mcia_results <- nipals_multiblock(data_blocks, preprocMethod='colprofile', num_PCs = 10, tol=1e-12)</pre>

Factor Plot

□ mrna○ miRNA△ prot

-2.0

Order 2 Factors

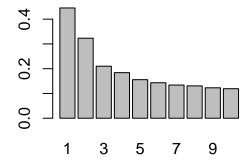
Order 1 Factors

0.0

1.0

-1.0

Global Factor Score Eigenvalues



Global Factor Score Order

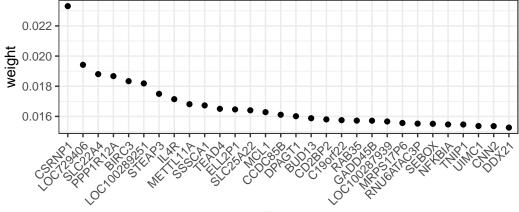
Part 2: Interpreting Global Loadings

Pseudoeigenvalues representing the contribution of each omic to the global factor score

```
## Should make into a function
bs_weights<-as.matrix(data.frame(mcia_results\block_score_weights))
colnames(bs_weights)<-c(1:10)</pre>
ComplexHeatmap::Heatmap(bs_weights, cluster_columns=FALSE, cluster_rows=FALSE,
                         column title side = "bottom", column title = "Factor")
                                               matrix 1
                                     mrna
                                                  0.7
                                                  0.65
                                     miRNA
                                                 0.6
                                     prot
                                                  0.55
                                                  0.5
                                                  0.45
               Factor
```

Scree Plot: Visualizing the top features per factor

```
# make into function
gl<-mcia_results$global_loadings</pre>
omic_list<-gsub("^.*_", "", rownames(gl))</pre>
factor<-4
gl_f<-data.frame(gl[,factor])</pre>
gl_f$omic<-omic_list</pre>
colnames(gl_f)<-c("weight",'omic')</pre>
gl_f_ord<-gl_f[order(gl_f$weight, decreasing=TRUE),]</pre>
# look at mRNA (need to filter since issue with miRNA omic name)
gl_f_ord_mRNA<-gl_f_ord[gl_f_ord$omic=="mrna",]</pre>
omic_name<-sub("_.*", "",rownames(gl_f_ord_mRNA) )</pre>
gl_f_ord_mRNA$omic_name<-sub("_.*", "",rownames(gl_f_ord_mRNA) )</pre>
ggplot(gl_f_ord_mRNA[0:30,], aes(x=factor(omic_name,level=omic_name), y=weight))+
      geom_point()+
      theme_bw()+
      xlab('Feature')+
      theme(axis.text.x=element_text(angle=45, hjust=1))
```



Feature