Unsupervised Multiblock analyses

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- · Analysis of complete dataset
 - Question 1: based on lipids and genes data, do we observe clusters of samples?
 - Question 2: how do both blocks contribute to each dimension?
 - Question 3: observe the samples distributions in the space of the common dimensions, what are the main sources of variation?
 - Question 4: which genes and lipids are responsible of the samples differences?
- · Repeat analysis for wt samples only
- · Repeat analysis for ppar samples only

Nutrimouse dataset

The data sets come from a nutrigenomic study in the mouse (Martin et al., 2007) in which the effects of five regimens with contrasted fatty acid compositions on liver lipids and hepatic gene expression in mice were considered. Two sets of variables were acquired on forty mice:

gene: expressions of 120 genes measured in liver cells, selected (among about 30,000) as potentially relevant in the context of the nutrition study. These expressions come from a nylon macroarray with radioactive labelling;

lipid: concentrations (in percentages) of 21 hepatic fatty acids measured by gas chromatography.

Biological units (mice) were cross-classified according to two factors experimental design (4 replicates):

Genotype: 2-levels factor, wild-type (WT) and PPARα -/- (PPAR).

Diet: 5-levels factor. Oils used for experimental diets preparation were corn and colza oils (50/50) for a reference diet (REF), hydrogenated coconut oil for a saturated fatty acid diet (COC), sunflower oil for an Omega6 fatty acid-rich diet (SUN), linseed oil for an Omega3-rich diet (LIN) and corn/colza/enriched fish oils for the FISH diet (43/43/14).

HIDE

```
data("nutrimouse")
genes <- nutrimouse$gene
lipids <- nutrimouse$lipid
metadata <- data.frame(genotype = nutrimouse$genotype, diet = nutrimouse$diet)
metadata$sample_name <- paste0(rownames(metadata), "_", metadata$genotype, "_", metadata$di
et)
rownames(genes) <- metadata$sample_name
rownames(lipids) <- metadata$sample_name</pre>
```

Analysis of complete dataset

Question 1: based on lipids and genes data, do we observe clusters of samples?

Prepare dataset - concatenate genes and lipids dataframes - define the number of variables of both block

Run ComDim analysis - use ComDim() from MBAnalysis package

HIDE

```
# prepare dataset
ComDim_data <- cbind.data.frame(genes, lipids)
n_group <- c(dim(genes)[[2]], dim(lipids)[[2]])
# run analysis
ComDim_res <- ComDim(X = ComDim_data, group = n_group)</pre>
```

Question 2: how do both blocks contribute to each dimension?

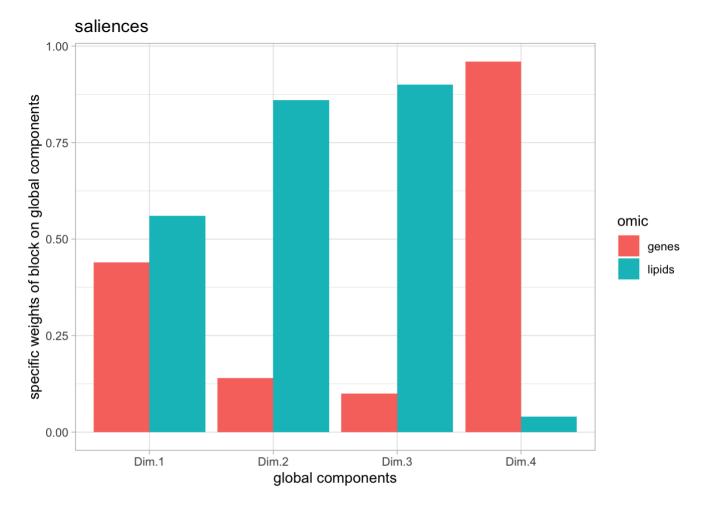
- · plot saliences
- plot block contributions (contrib)

```
# saliences

saliences <- ComDim_res$saliences
rownames(saliences) <- c("genes", "lipids")
saliences <- as.data.frame(t(saliences[,1:4]))
saliences$Dim <- rownames(saliences)

saliences <- melt(saliences)

ggplot(saliences, aes(x=Dim, y=value, fill=variable)) +
    geom_bar(stat = "identity", position=position_dodge()) +
    theme_light() +
    labs(x = "global components", y = "specific weights of block on global components", fill
= "omic",
    title = "saliences")</pre>
```



```
# block contributions

contributions <- ComDim_res$contrib

rownames(contributions) <- c("genes", "lipids")

contributions <- as.data.frame(t(contributions[,1:4]))

contributions$Dim <- rownames(contributions)

contributions <- melt(contributions)

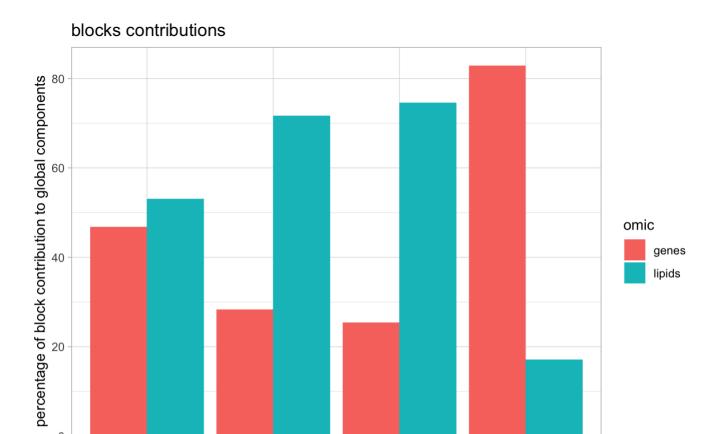
ggplot(contributions, aes(x=Dim, y=value, fill=variable)) +

geom_bar(stat = "identity", position=position_dodge()) +

theme_light() +

labs(x = "global components", y = "percentage of block contribution to global component
s", fill = "omic",

title = "blocks contributions")</pre>
```



Question 3: observe the samples distributions in the space of the common dimensions, what are the main sources of variation?

global components

Dim.3

Dim.4

• plot scores (T) on Dim.1 vs Dim.2 with percentages of explained variance on axes

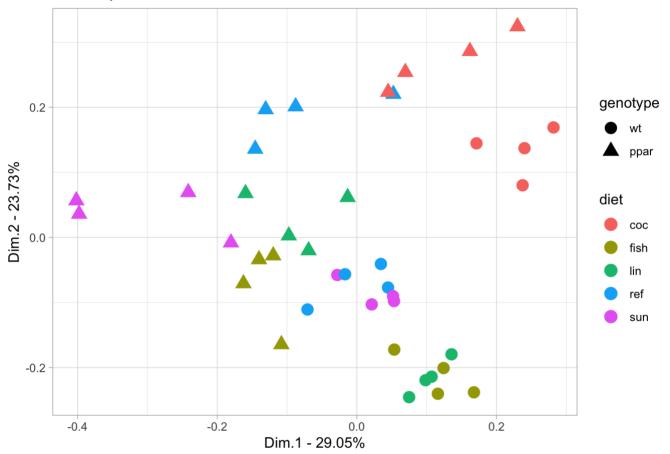
Dim.1

• plot scores (T) on Dim.3 vs Dim.4 with percentages of explained variance on axes

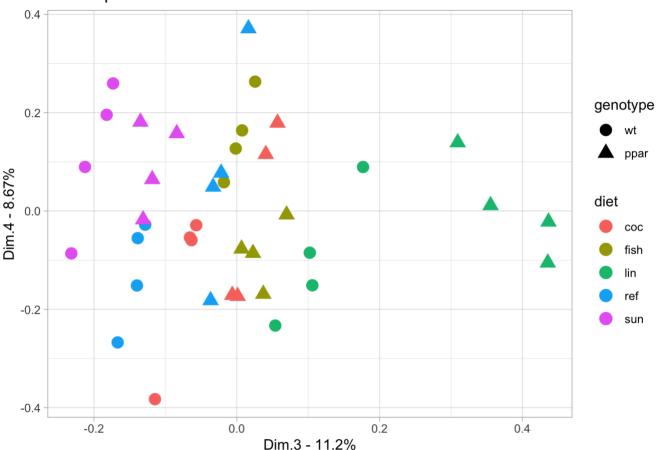
```
scores <- data.frame(metadata, ComDim_res$T)

ggplot(scores, aes(x=Dim.1, y=Dim.2, col=diet, shape = genotype)) +
    geom_point(size=4) +
    labs(x=paste0("Dim.1 - ", ComDim_res$cumexplained[1,"%explX"], "%"),
        y=paste0("Dim.2 - ", ComDim_res$cumexplained[2,"%explX"], "%"),
        title = "scores plots on Dim.1 Dim.2") +
    theme_light()</pre>
```

scores plots on Dim.1 Dim.2



scores plots on Dim.3 Dim.4



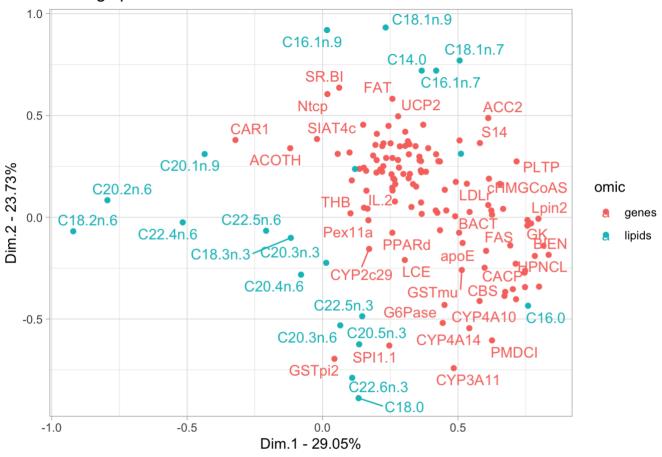
Question 4: which genes and lipids are responsible of the samples differences?

- plot scores (globalcor) on Dim.1 vs Dim.2 with percentages of explained variance on axes
- · plot scores (globalcor) on Dim.3 vs Dim.4 with percentages of explained variance on axes

```
loadings <- data.frame(ComDim_res$globalcor)
loadings$omic <- c(rep("genes", dim(genes)[[2]]), rep("lipids", dim(lipids)[[2]]))
loadings$variable <- rownames(loadings)

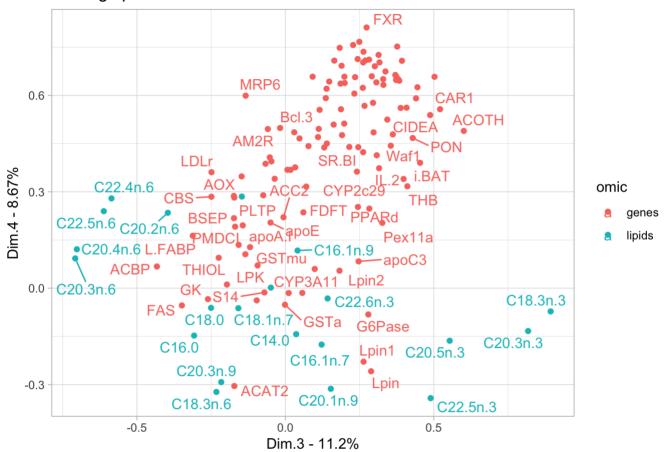
ggplot(loadings, aes(x=X1, y=X2, col=omic, label=variable)) +
    geom_point() +
    geom_text_repel() +
    labs(x=paste0("Dim.1 - ", ComDim_res$cumexplained[1,"%explX"], "%"),
        y=paste0("Dim.2 - ", ComDim_res$cumexplained[2,"%explX"], "%"),
        title = "loadings plots on Dim.1 Dim.2") +
    theme_light()</pre>
```

loadings plots on Dim.1 Dim.2



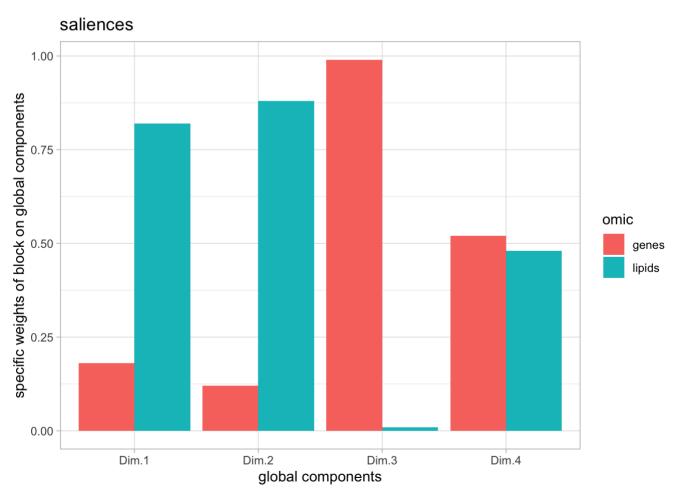
```
ggplot(loadings, aes(x=X3, y=X4, col=omic, label=variable)) +
  geom_point() +
  geom_text_repel() +
  labs(x=paste0("Dim.3 - ", ComDim_res$cumexplained[3,"%explX"], "%"),
       y=paste0("Dim.4 - ", ComDim_res$cumexplained[4,"%explX"], "%"),
       title = "loadings plots on Dim.3 Dim.4") +
  theme_light()
```

loadings plots on Dim.3 Dim.4



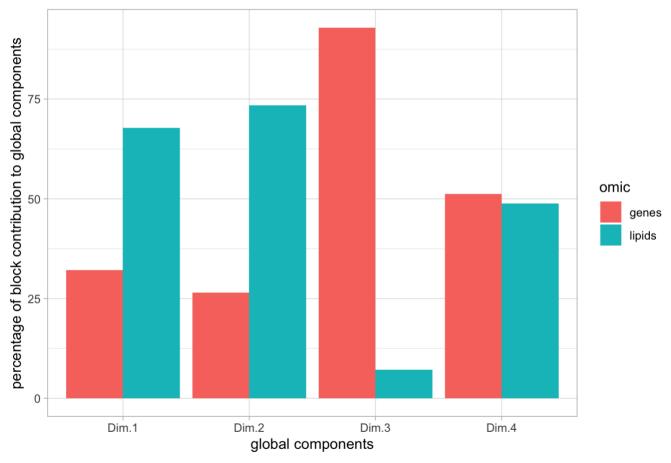
Repeat analysis for wt samples only

```
# prepare dataset
wt samples <- metadata$sample name[metadata$genotype == "wt"]</pre>
wt_ComDim_data <- cbind.data.frame(genes[wt_samples,], lipids[wt_samples,])</pre>
n_group <- c(dim(genes)[[2]], dim(lipids)[[2]])</pre>
# run analysis
wt_ComDim_res <- ComDim(X = wt_ComDim_data, group = n_group, option = "uniform", plotgraph</pre>
= F)
# saliences
wt saliences <- wt ComDim res$saliences
rownames(wt saliences) <- c("genes", "lipids")</pre>
wt saliences <- as.data.frame(t(wt saliences[,1:4]))</pre>
wt_saliences$Dim <- rownames(wt_saliences)</pre>
wt saliences <- melt(wt saliences)</pre>
ggplot(wt_saliences, aes(x=Dim, y=value, fill=variable)) +
  geom bar(stat = "identity", position=position dodge()) +
 theme light() +
  labs(x = "global components", y = "specific weights of block on global components", fill
= "omic",
       title = "saliences")
```



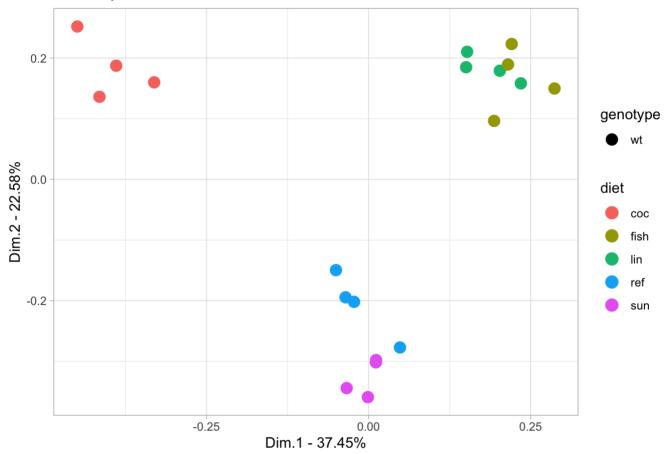
```
# block contributions
wt_contributions <- wt_ComDim_res$contrib
rownames(wt_contributions) <- c("genes", "lipids")
wt_contributions <- as.data.frame(t(wt_contributions[,1:4]))
wt_contributions$Dim <- rownames(wt_contributions)
wt_contributions <- melt(wt_contributions)
ggplot(wt_contributions, aes(x=Dim, y=value, fill=variable)) +
    geom_bar(stat = "identity", position=position_dodge()) +
    theme_light() +
    labs(x = "global components", y = "percentage of block contribution to global component
s", fill = "omic",
    title = "blocks contributions")</pre>
```

blocks contributions



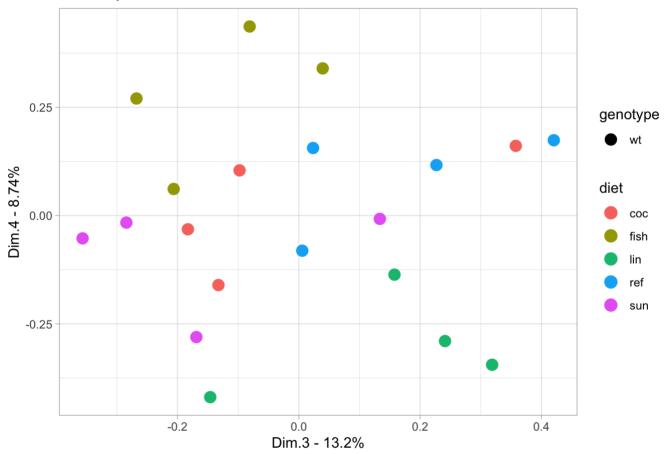
```
# scores plots
wt_scores <- data.frame(metadata[metadata$genotype == "wt",], wt_ComDim_res$T)
ggplot(wt_scores, aes(x=Dim.1, y=Dim.2, col=diet, shape = genotype)) +
    geom_point(size=4) +
    labs(x=paste0("Dim.1 - ", wt_ComDim_res$cumexplained[1,"%explX"], "%"),
        y=paste0("Dim.2 - ", wt_ComDim_res$cumexplained[2,"%explX"], "%"),
        title = "scores plots on Dim.1 Dim.2") +
theme_light()</pre>
```

scores plots on Dim.1 Dim.2



```
ggplot(wt_scores, aes(x=Dim.3, y=Dim.4, col=diet, shape = genotype)) +
  geom_point(size=4) +
  labs(x=paste0("Dim.3 - ", wt_ComDim_res$cumexplained[3,"%explX"], "%"),
      y=paste0("Dim.4 - ", wt_ComDim_res$cumexplained[4,"%explX"], "%"),
      title = "scores plots on Dim.3 Dim.4") +
  theme_light()
```

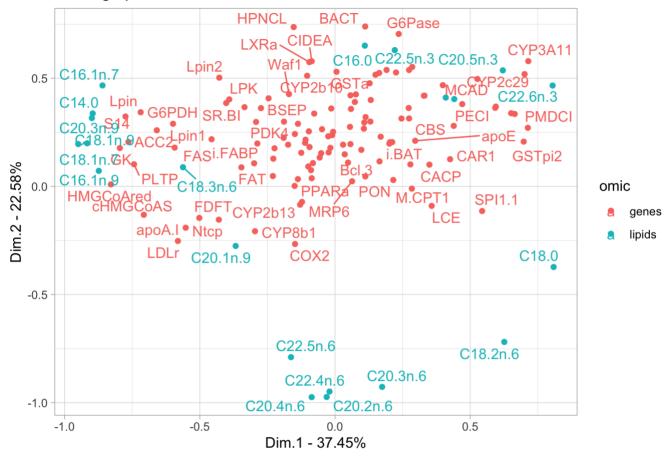
scores plots on Dim.3 Dim.4



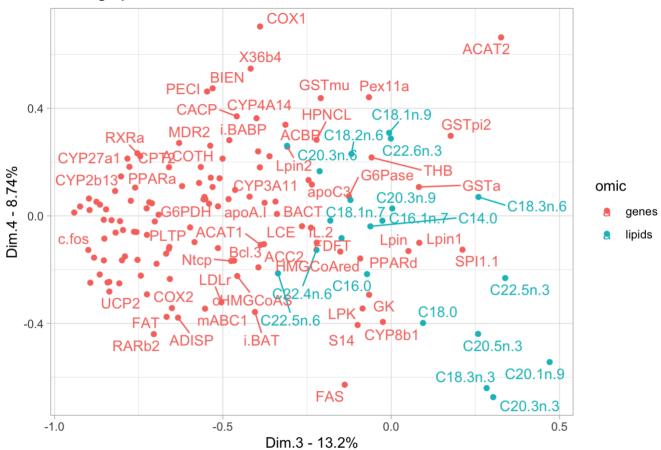
```
# loadings plots
wt_loadings <- data.frame(wt_ComDim_res$globalcor)
wt_loadings$omic <- c(rep("genes", dim(genes)[[2]]), rep("lipids", dim(lipids)[[2]]))
wt_loadings$variable <- rownames(wt_loadings)
ggplot(wt_loadings, aes(x=X1, y=X2, col=omic, label=variable)) +
    geom_point() +
    geom_text_repel() +
    labs(x=paste0("Dim.1 - ", wt_ComDim_res$cumexplained[1,"%explX"], "%"),
        y=paste0("Dim.2 - ", wt_ComDim_res$cumexplained[2,"%explX"], "%"),
        title = "loadings plots on Dim.1 Dim.2") +
    theme_light()</pre>
```

loadings plots on Dim.1 Dim.2

theme_light()

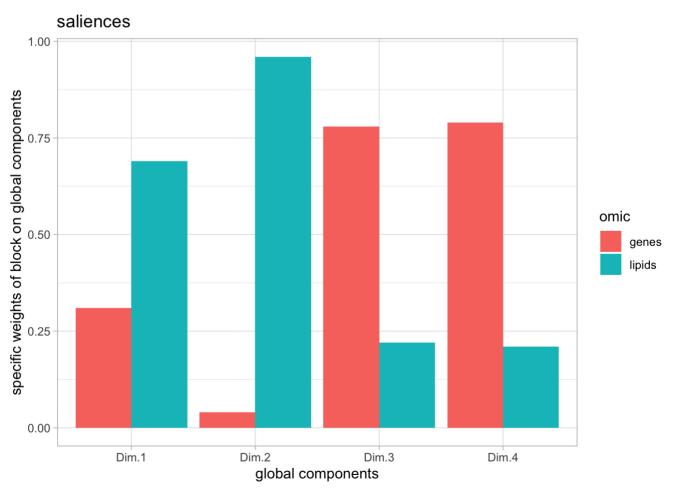


loadings plots on Dim.3 Dim.4



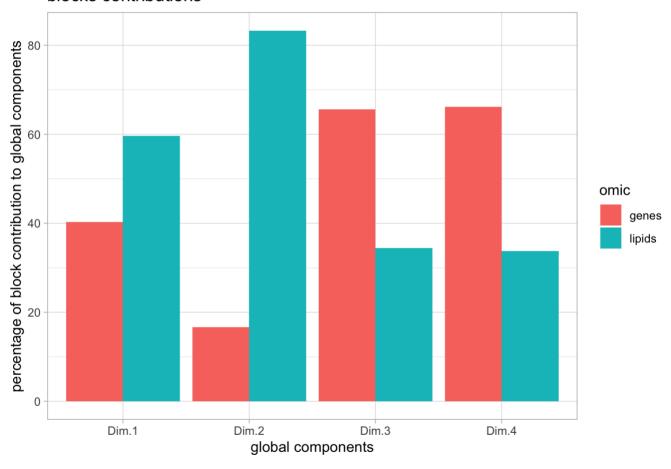
Repeat analysis for ppar samples only

```
# prepare dataset
ppar samples <- metadata$sample name[metadata$genotype == "ppar"]</pre>
ppar_ComDim_data <- cbind.data.frame(genes[ppar_samples,], lipids[ppar_samples,])</pre>
n_group <- c(dim(genes)[[2]], dim(lipids)[[2]])</pre>
# run analysis
ppar_ComDim_res <- ComDim(X = ppar_ComDim_data, group = n_group, option = "uniform", plotgr</pre>
aph = F)
# saliences
ppar_saliences <- ppar_ComDim_res$saliences</pre>
rownames(ppar_saliences) <- c("genes", "lipids")</pre>
ppar_saliences <- as.data.frame(t(ppar_saliences[,1:4]))</pre>
ppar_saliences$Dim <- rownames(ppar_saliences)</pre>
ppar_saliences <- melt(ppar_saliences)</pre>
ggplot(ppar_saliences, aes(x=Dim, y=value, fill=variable)) +
  geom bar(stat = "identity", position=position dodge()) +
  theme light() +
  labs(x = "global components", y = "specific weights of block on global components", fill
= "omic",
       title = "saliences")
```



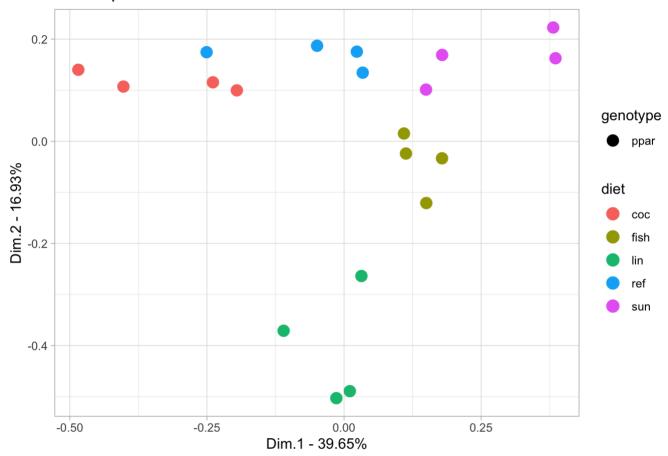
```
# block contributions
ppar_contributions <- ppar_ComDim_res$contrib
rownames(ppar_contributions) <- c("genes", "lipids")
ppar_contributions <- as.data.frame(t(ppar_contributions[,1:4]))
ppar_contributions$Dim <- rownames(ppar_contributions)
ppar_contributions <- melt(ppar_contributions)
ggplot(ppar_contributions, aes(x=Dim, y=value, fill=variable)) +
    geom_bar(stat = "identity", position=position_dodge()) +
    theme_light() +
    labs(x = "global components", y = "percentage of block contribution to global component
s", fill = "omic",
    title = "blocks contributions")</pre>
```

blocks contributions



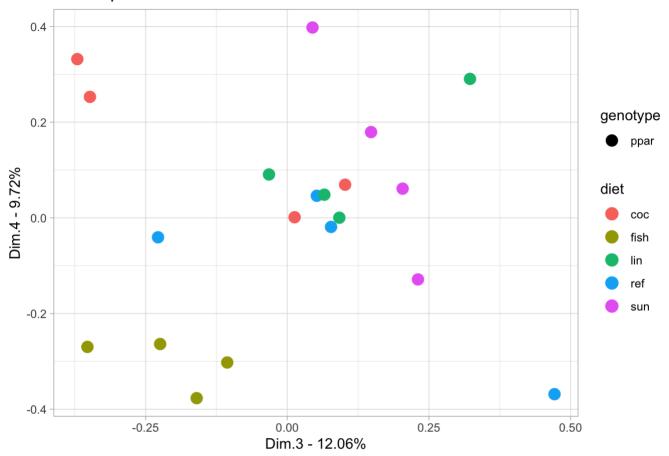
```
# scores plots
ppar_scores <- data.frame(metadata[metadata$genotype == "ppar",], ppar_ComDim_res$T)
ggplot(ppar_scores, aes(x=Dim.1, y=Dim.2, col=diet, shape = genotype)) +
    geom_point(size=4) +
    labs(x=paste0("Dim.1 - ", ppar_ComDim_res$cumexplained[1,"%explX"], "%"),
        y=paste0("Dim.2 - ", ppar_ComDim_res$cumexplained[2,"%explX"], "%"),
        title = "scores plots on Dim.1 Dim.2") +
    theme_light()</pre>
```

scores plots on Dim.1 Dim.2



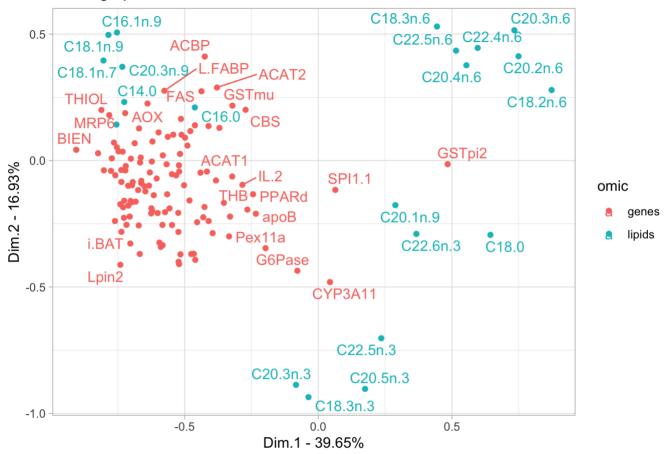
```
ggplot(ppar_scores, aes(x=Dim.3, y=Dim.4, col=diet, shape = genotype)) +
  geom_point(size=4) +
  labs(x=paste0("Dim.3 - ", ppar_ComDim_res$cumexplained[3,"%explX"], "%"),
       y=paste0("Dim.4 - ", ppar_ComDim_res$cumexplained[4,"%explX"], "%"),
       title = "scores plots on Dim.3 Dim.4") +
  theme_light()
```

scores plots on Dim.3 Dim.4



```
# loadings plots
ppar_loadings <- data.frame(ppar_ComDim_res$globalcor)
ppar_loadings$omic <- c(rep("genes", dim(genes)[[2]]), rep("lipids", dim(lipids)[[2]]))
ppar_loadings$variable <- rownames(ppar_loadings)
ggplot(ppar_loadings, aes(x=X1, y=X2, col=omic, label=variable)) +
    geom_point() +
    geom_text_repel() +
    labs(x=paste0("Dim.1 - ", ppar_ComDim_res$cumexplained[1,"%explX"], "%"),
        y=paste0("Dim.2 - ", ppar_ComDim_res$cumexplained[2,"%explX"], "%"),
        title = "loadings plots on Dim.1 Dim.2") +
    theme_light()</pre>
```

loadings plots on Dim.1 Dim.2



loadings plots on Dim.3 Dim.4

