# Volcano Plot using ggplot2

Debojyoti Das

2025-03-02

#### Introduction

This document demonstrates how to generate a volcano plot using ggplot2 by reading a CSV file that contains gene expression data. The dataset must include at least three mandatory columns:

- log2FC (Log2 Fold Change)
- p\_value (P-value for statistical significance)
- Gene\_symbol (or Gene EntrezID or Gene ENSEMBL ID)

Each step is explained in detail, with code chunks for clarity.

#### Installing and Loading Required Libraries

Check if required packages are installed, if not install them

```
if (!requireNamespace("ggplot2", quietly = TRUE)) {
   install.packages("ggplot2")
}

if (!requireNamespace("ggrepel", quietly = TRUE)) {
   install.packages("ggrepel")
}

if (!requireNamespace("dplyr", quietly = TRUE)) {
   install.packages("dplyr")
}
```

Load libraries

```
library(ggplot2)
library(dplyr)

##
## Attaching package: 'dplyr'

## The following objects are masked from 'package:stats':
##
## filter, lag
```

```
## The following objects are masked from 'package:base':
##
## intersect, setdiff, setequal, union
library(ggrepel)
```

#### Setting Up Project Directories

```
project_dir <- "/Users/debda22/Projects/core_facility_projects/ggplot_basics"
data_dir <- paste0(project_dir, "/input_data")
result_dir <- paste0(project_dir, "/results")</pre>
```

Create directories if they don't exist

```
if (!dir.exists(result_dir)) dir.create(result_dir, recursive = TRUE)
```

#### Reading the Input Data

```
input_file <- pasteO(data_dir, "/test_input_file.csv")
data <- read.csv(input_file)</pre>
```

Display first few rows of the dataset

```
head(data)
```

```
Gene_symbol
                   log2FC neg_log10pval log2FC_sq
##
                                                      p_value
## 1
          Gene7 2.267283
                              1.7250171 5.140572 0.0188357482
                              3.8045651 9.166577 0.0001568321
          Gene9 3.027636
## 2
## 3
         Gene11 1.957304
                              0.2616621 3.831041 0.5474417710
## 4
         Gene12 3.429968
                              3.7879732 11.764681 0.0001629396
## 5
         Gene13 -2.083291
                              3.8993947 4.340102 0.0001260681
                              3.9222951 15.877700 0.0001195928
## 6
         Gene18 -3.984683
```

#### Transforming Data for Visualization

Before plotting, we transform the data: - Convert the p\_value column to -log10(p\_value) to emphasize small p-values. - Define upregulated and downregulated genes based on cutoff values. - Reverse the log2FC values for visualization.

We define thresholds for classification: - p\_value cutoff: 0.05 - log2FC cutoff: 1

Define cutoffs

```
pval_cutoff <- 0.05
log2fc_cutoff <- 1</pre>
```

Classify genes into upregulated, downregulated, or Non-significant  $\,$ 

```
data <- data %>%
  mutate(
   logP = -log10(p_value),
   negLog2FC = -log2FC,
  regulation = case_when(
      p_value < pval_cutoff & log2FC > log2fc_cutoff ~ "Upregulated",
      p_value < pval_cutoff & log2FC < -log2fc_cutoff ~ "Downregulated",
      TRUE ~ "Non-significant"
   )
)</pre>
```

#### Selecting Top Genes for Labeling

To highlight important genes, we select the top n genes from the upregulated and downregulated groups.

```
top_n <- 5  # Number of genes to label

top_up <- data %>%
  filter(regulation == "Upregulated") %>%
  arrange(desc(log2FC)) %>%
  head(top_n)

top_down <- data %>%
  filter(regulation == "Downregulated") %>%
  arrange(desc(abs(log2FC))) %>%
  head(top_n)

# Combine top genes
top_genes <- bind_rows(top_up, top_down)</pre>
```

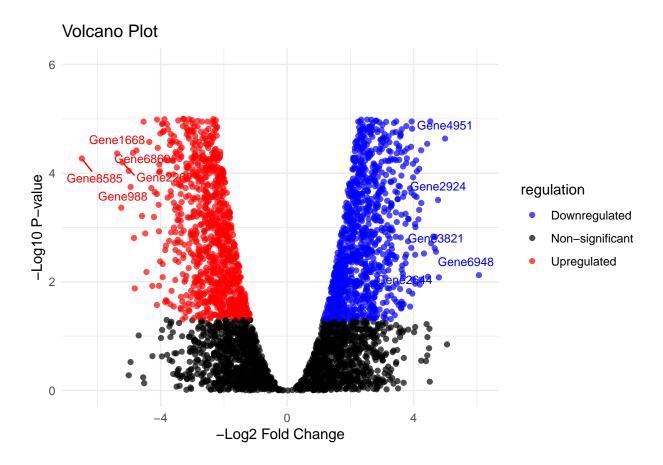
#### Creating the Volcano Plot

We use  $\mathsf{ggplot2}$  to create a volcano plot with color distinctions:

```
volcano <- ggplot(data, aes(x = negLog2FC, y = logP, color = regulation)) +
    geom_point(alpha = 0.7) +
    scale_color_manual(values = c("Non-significant" = "black", "Upregulated" = "red", "Downregulated" = "'
    labs(title = "Volcano Plot", x = "-Log2 Fold Change", y = "-Log10 P-value") +
    theme_minimal() +
    geom_text_repel(
        data = top_genes,
        aes(label = Gene_symbol),
        vjust = -1,
        size = 3,
        show.legend = FALSE
    ) +
    ylim(c(0,6))</pre>
```

Display plot

#### print(volcano)



## Saving the Plot

```
output_file <- pasteO(result_dir, "/volcano_plot.png")
ggsave(output_file, plot = volcano, width = 8, height = 6)</pre>
```

### Conclusion

This document demonstrated how to load, process, and visualize gene expression data using a volcano plot. We added classification for upregulated and downregulated genes, highlighted top genes, and saved the final plot.