

# plot\_function

2024-08-08

```
# Define a custom theme
custom_theme <- theme(
  panel.background = element_rect(fill = '#333333'),
  plot.title = element_text(hjust = 0.5, color = 'white', size = 16),
  axis.title = element_text(size = 14, color = 'white'),
  axis.text = element_text(size = 12, color = 'white'),
  plot.background = element_rect(fill = '#333333'),
  panel.grid.major = element_line(color = "gray", size = 0.1),
  legend.position = 'top',
  legend.background = element_rect(fill = "#333333"),
  legend.text = element_text(size = 12, color = 'white'),
  legend.title = element_text(size = 12, color = 'white'),
  legend.key = element_rect(fill = "#333333")
)

# Define the function to create plots
create_plots <- function(meta_data_path, gene_data_path, gene_names,
                          continuous_covariate, categorical_covariate1,
                          categorical_covariate2) {

  # Load the datasets
  genes_data <- read_csv(gene_data_path, show_col_types = FALSE)
  metadata <- read_csv(meta_data_path, show_col_types = FALSE)

  # Check if the data is read properly
  if (is.null(genes_data) || is.null(metadata)) {
    stop("Failed to read the datasets")
  }

  # Rename the column `...1` to `Genes` if it exists
  if ("...1" %in% colnames(genes_data)) {
    genes_data <- genes_data %>% rename(Genes = `...1`)
  } else {
    stop("Column '...1' not found in gene data")
  }

  # Set gene names as row names
  rownames(genes_data) <- genes_data$Genes

  for (gene_name in gene_names) {
    gene_name <- as.character(gene_name)
    #print(paste("Processing gene:", gene_name))
    if (!(gene_name %in% rownames(genes_data))) {
      warning(paste("Gene name", gene_name, "not found in the genes data"))
      next
    }
  }
}
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}

# Select the gene row and transpose
gene <- genes_data[gene_name, ]
transposed_gene <- as.data.frame(t(gene))
colnames(transposed_gene) <- gene_name
# Combine two datasets with participant_id and gene
transposed_gene_data <- cbind(participant_id = rownames(transposed_gene), transposed_gene)
# Merge the dataset
data_frame <- merge(metadata, transposed_gene_data, by = "participant_id")
# Ensure the gene expression values are numeric
data_frame[[gene_name]] <- as.numeric(data_frame[[gene_name]])
# Filter out NA values for the continuous covariate
data_frame[[continuous_covariate]] <- as.numeric(data_frame[[continuous_covariate]])
data_frame <- data_frame %>% filter(!is.na(!sym(continuous_covariate)))
# Filter out NA values for categorical covariate
data_frame <- data_frame %>% filter(!(sym(categorical_covariate1) != 'unknown'))
filtered_data <- data_frame %>% filter(!(sym(categorical_covariate2) != 'unknown'))
# Histogram for gene expression
gene_histogram <- ggplot(filtered_data, aes(x = !sym(gene_name))) +
  geom_histogram(binwidth = 0.05, color = "#1dddeb", fill = "#1dddeb", alpha = 0.7) +
  labs(x = gene_name,
       y = "Number",
       title = paste("Histogram of",
                     gene_name,
                     "Expression")) +
  custom_theme

# Save the histogram
ggsave(paste0("plots/", gene_name, "_histogram.png"),
       plot = gene_histogram, width = 16, height = 9)
# Print the histogram
print(gene_histogram)

# Scatter Plot with rainbow colors
scatter_plot <- ggplot(filtered_data,
                       aes(x = !sym(continuous_covariate),
                           y = !sym(gene_name),
                           color = !sym(gene_name))) +
  geom_point(size = 2) +
  geom_rug(color = "white") + # Add a rug plot
  scale_color_gradientn(colors = rainbow(7)) + # Apply rainbow colors
  labs(x = continuous_covariate, y = gene_name, title = paste("Scatter Plot of",
                                                             gene_name,
                                                             "Expression vs",
                                                             continuous_covariate)) +
  custom_theme

# Save the scatter plot
ggsave(paste0("plots/", gene_name, "_scatter_plot.png"),
       plot = scatter_plot, width = 16, height = 9)
# Print the scatter plot
print(scatter_plot)

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# Define labels for the legend
labels <- unique(filtered_data[[categorical_covariate2]])
# Create the palette dynamically based on the labels
color <- c("#66c2a5", "#fc8d62")
palette <- setNames(color, labels)
# https://www.statology.org/setnames-r/
# Boxplot
boxplot <- ggplot(filtered_data, aes(x = !!sym(categorical_covariate1),
                                     y = !!sym(gene_name),
                                     fill = !!sym(categorical_covariate2))) +
  geom_boxplot(outlier.color = "red", color = "white") +
  labs(x = categorical_covariate1, y = gene_name, fill = categorical_covariate2,
       title = paste("Boxplot of", gene_name, "Expression by",
                     categorical_covariate1, "and", categorical_covariate2)) +
  scale_fill_manual(values = palette, labels = labels) +
  custom_theme

# Save the boxplot
ggsave(paste0("plots/", gene_name, "_boxplot.png"),
       plot = boxplot, width = 16, height = 9)
# Print the boxplot
print(boxplot)
}

create_plots('dataset/QBS103_GSE157103_series_matrix.csv',
             'dataset/QBS103_GSE157103_genes.csv',
             c("A1BG", "A2M", "AANAT"),
             "age", "sex",
             "disease_status")

```









