# Package 'ProteinPCA'

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Type Package
Title Principal Component Analysis (PCA) Tool on Protein Expression
     Data
Version 0.1.0
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Description
     Analysis of protein expression data can be done through Principal Component Analysis (PCA),
     and this R package is designed to streamline the analysis. This package enables users to per-
     form PCA
     and it generates biplot and scree plot for advanced graphical visualization. Optionally, it supports
     grouping/clustering visualization with PCA loadings and confidence ellipses. With this R package,
     researchers can quickly explore complex protein datasets, interpret variance contribu-
     tions, and visualize
     sample clustering through intuitive biplots. For more details, see Jol-
     liffe (2001) <doi:10.1007/b98835>,
     Gabriel (1971) <doi:10.1093/biomet/58.3.453>, Zhang et al. (2024) <doi:10.1038/s41467-024-
     53239-9>, and
     Anandan et al. (2022) <doi:10.1038/s41598-022-07781-5>.
License GPL-3
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```

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pca\_analysis

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#### **Description**

This function performs PCA on protein expression data and produces a biplot using ggplot2. Optionally, it supports grouping/clustering visualization with PCA loadings and confidence ellipses.

### **Arguments**

data	A numeric matrix or data frame of protein expression data. Rows are samples, columns are features.
scale	Logical. Should the data be scaled? Default is TRUE.
center	Logical. Should the data be centered? Default is TRUE.
plot	Logical. Should a PCA biplot be generated? Default is TRUE.
groups	Optional. A factor or character vector specifying group memberships of the samples.

#### Value

A list containing:

pca The PCA object from prcomp.

explained\_variance

The percentage of variance explained by each principal component.

plot A ggplot2 PCA biplot (if plot = TRUE).

#### Author(s)

Paul Angelo C. Manlapaz

#### References

Jolliffe, I. (2001). Principal Component Analysis (2nd ed.). Springer. https://doi.org/10.1007/b98835 Gabriel, K. R. (1971). The biplot graphic display of matrices with application to principal component analysis. Biometrika, 58(3), 453–467. https://doi.org/10.1093/biomet/58.3.453 Zhang, Z., Chen, L., Sun, B., Ruan, Z., Pan, P., Zhang, W., Jiang, X., Zheng, S., Cheng, S., Xian, L., Wang, B., Yang, J., Zhang, B., Xu, P., Zhong, Z., Cheng, L., Ni, H., & Hong, Y. (2024). Identifying septic shock subgroups to tailor fluid strategies through multi-omics integration. Nature Communications, 15(1). https://doi.org/10.1038/s41467-024-53239-9 Anandan, A., Nagireddy, R., Sabarinathan, S.,

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Bhatta, B. B., Mahender, A., Vinothkumar, M., Parameswaran, C., Panneerselvam, P., Subudhi, H., Meher, J., Bose, L. K., & Ali, J. (2022). Multi-trait association study identifies loci associated with tolerance of low phosphorus in Oryza sativa and its wild relatives. Scientific Reports, 12(1). https://doi.org/10.1038/s41598-022-07781-5

# **Examples**

```
set.seed(123)
data_matrix <- matrix(rnorm(100 * 20), nrow = 100, ncol = 20)
rownames(data_matrix) <- paste0('Sample_', 1:100)
colnames(data_matrix) <- paste0('Protein_', 1:20)
groups <- sample(c("Group A", "Group B"), 100, replace = TRUE)
result <- pca_analysis(data_matrix, groups = groups)
print(result$explained_variance)
print(result$plot)</pre>
```

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