

## Genomic Data Explorer User Guide (07-10-20)

### Contact:

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URL: <http://doplrshinyweb.mdanderson.edu:8080/>

Login username: ICON

Login password: shinyicon

If the login process is too slow, someone else are probably running the apps. Please be patient and try at a different time. The R Shiny apps are running on a test virtual machine (VM) with limited capability. Infrastructure upgrades are planned.

After logging in, click one of the apps to start. It usually takes 10-20s to start the app.

### DNA Explorer

1. Choose your data source:  
Click one of the three tabs: “[Example](#)”, “[Upload](#)”, and “[Select](#)”. (Currently, “Upload” is not supported)  
There are three types of files – Mutation Annotation Format (MAF), Copy Number Variation (CNV), and Meta Data in “Select”:  
**MAF Data:** [ICON\\_108samples\\_mutect\\_vaf02\\_pindel\\_vaf05.maf](#)  
[ICON\\_108samples\\_mutect\\_vaf02\\_pindel\\_vaf05\\_dropRepeats.maf](#)  
**CNV Data:** [ICON\\_108samples\\_exomecn.txt](#)  
**Meta Data:** [metadata.csv](#)  
In “Example”, the following files are used:  
**MAF Data:** [ICON\\_108samples\\_mutect\\_vaf02\\_pindel\\_vaf05.maf](#)  
**CNV Data:** [ICON\\_108samples\\_exomecn.txt](#)  
**Meta Data:** [metadata.csv](#)
2. Click “Launch” button. Adjust the parameters, and then click “Plot” at the left bottom. The app will start plotting figures for you.
3. Parameters:
  - a. **[Oncoplot](#)** Options:  
Choose to display clinical features, TiTv, CNV, sample names, or whether sort by the annotation.
  - b. Select **[Clinical Features](#)**:  
Choose which annotations to display on the bottom of oncoplot/heatmap.
    - *For example, if you choose [Clinical Features](#) and select [Sex](#), choose [Sort by Annotation](#), and **click “Plot”**, you can view the mutation landscape ([Oncoplot](#)) by comparing male and female groups.*

- c. Gene List:  
Three ways to define genes: 1. Use Top Genes: use the most mutated genes (ranked by mutation frequency by maftools). Use the slide bar to adjust top n mutated genes, and **click "Plot"**; 2. Manual Input: type your favorite genes, split by semicolon or comma, and **click "Plot"**; 3. Upload File: upload your gene list, the file should be plain text file with one gene per row.
- d. Samples List:  
Three ways to define samples: 1. Use Top Samples: use the most mutated samples (ranked by mutation frequency by maftools). Use the slide bar to adjust top n mutated samples, and **click "Plot"**; 2. Manual Input: type your interested sample names, split by semicolon or comma, and **click "Plot"**; 3. Upload File: upload your sample list, the file should be plain text file with one sample per row.
  - *For example, if you upload a list of female patients, you can view the mutation landscape ([Oncoplot](#)) only for the female group.*
- e. Plot Height, Plot Width:  
Adjust the plot size in pixels (px).
- f. Pathway Names:  
Applies to [Oncogenic Signaling Pathways](#). Choose specific pathway name to display pathway-gene association
- g. VAF Column Name:  
Applies to [VAF Plot](#). Specify which column name in MAF file should be used for tumor (i.e., somatic mutation) variant allele frequency (VAF).

## RNA Explorer

1. Choose your data source:  
Click one of the three tabs: "Example", "Upload", and "Select". (Currently, "Upload" is not supported)  
There are two datasets in "Select":  
ICON dataset: "icon\_rnaseq.rds"; Example dataset: "rnaseq.rds".
2. Click "Launch" button. The app will start plotting figures for you. (The first click will take the app 10-15s to load the libraries)
3. Parameters:
  - a. Variable:
    1. Adjust which variable to label the rows and columns of [Heatmap](#); 2. Adjust which variable to color in [PCA](#) plot; 3. Adjust which variable to compare in [Gene Boxplot](#).
  - b. Categorical Palette, Continuous Palette:  
Adjust the palette used for categorical or continuous variables. Can be used in [Heatmap](#), [PCA](#), [Gene Boxplot](#), and [Gene Clustering](#). Please see the last page for palette names and colors.

- c. Reverse Scale Color Direction:  
Only applies to continuous palettes.
- d. Plot Height, Plot Width:  
Adjust the plot size in pixels (px). Maximum 4000px.
- e. Adjusted P-value Cutoff, Log2 Fold Change cutoff:  
Applies to [MA Plot](#), [Volcano Plot](#), and [Table](#).
- f. Adjusted P-value Squash, Log2 Fold Change Squash:  
Applies to [MA Plot](#), [Volcano Plot](#). Used to squash outlier points to acceptable ranges.
- g. Gene List:  
Three ways to define genes: 1. Use Top Genes: use the most significantly differentially expressed genes (ranked by p-values). Use the slide bar to adjust top n genes, and [click "Plot"](#); 2. Manual Input: type your favorite genes, split by semicolon or comma, and [click "Plot"](#); 3. Upload File: upload your gene list, the file should be plain text file with one gene per row.
- h. Pathway List:  
Two ways to define pathways: 1. Use Hallmarks: use 50 hallmark gene sets defined in this link: <https://www.gsea-msigdb.org/gsea/msigdb/collections.jsp>  
2. Upload File: upload your pathway list, the file should be two-column plain text file with the first column as pathway names and the second row as gene names.

