User Manual of CLAM

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1 Introduction

CLAM is an analytical framework for identifying co-expressed gene modules by integrating multi-omics data and known molecular interactions. The source code and the complied tool can be downloaded at https://github.com/free1234hm/CLAM.

2 Preliminaries

- To use CLAM a version of Java 1.5 or later must be installed. If Java 1.5 or later is not currently installed, then it can be downloaded from http://www.java.com.
- CLAM can be executed by double-clicking on 'CLAM.jar', or from a command line change to the CLAM directory and then type: java -mx1024M -jar CLAM.jar. If CLAM reports 'Out of Memory Error', users can increase the -mx parameter.

3 Main Interface

The main interface has three parts: 'Load Data', 'Set Parameters', and 'Search Gene Modules'.

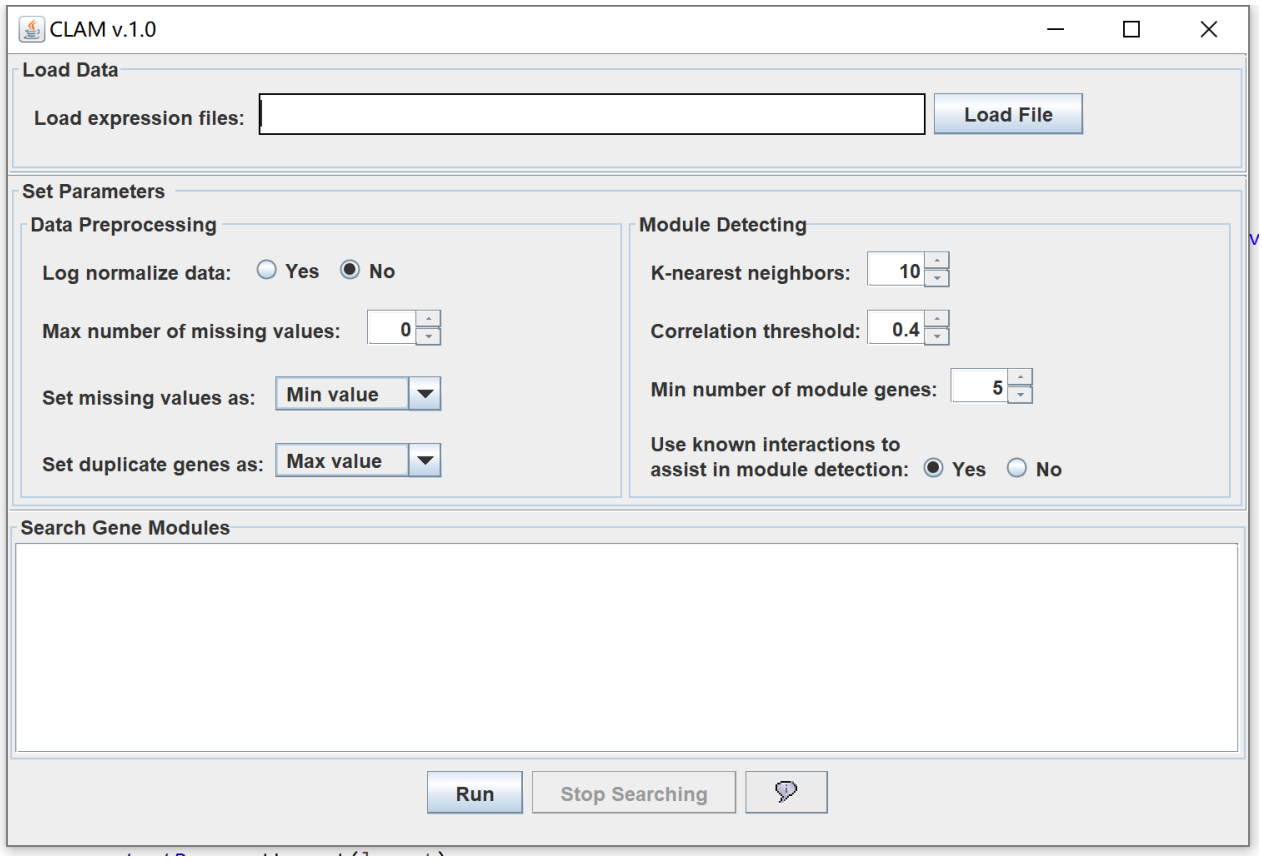

Figure 1. The main interface of CLAM

3.1 Load expression files

The 'Load expression files' field is used to import datasets from different sources. All datasets need to written in same format: the first column is gene or protein names, and the remaining columns contain the measurements in different samples. If a measurement is missing, then the field should be left empty. The first row of the data contains column headers. **Notably, CLAM does not require different omics datasets to share the same genes (proteins) or samples.** A sample expression data file is shown in Figure 2.

Ensembl_ID	TCGA-AA-	TCGA-AA-	TCGA-A6-	TCGA-A6-	TCGA-AA-	TCGA-CK-	TCGA-AA-	TCGA-AA-	TCGA-AU-	TCGA-QG
A1CF	0.92999	1.066218	2.564578	2.242078	1.208603	0.159212	2.189218	0.061351	0.889145	1.183308
A2M	4.529061	4.870139	5.687503	6.662382	5.405043	5.591557	3.445681	5.108467	4.960791	3.854576
A4GALT	2.662687	1.147233	1.104459	2.482948	1.853063	1.383715	0.828363	2.687003	2.098147	1.040627
AAAS	3.833031	3.628329	3.198295	3.443551	4.099529	3.6168	3.774873	3.734673	3.361572	3.774655
AACS	2.145268	2.235108	2.06301	1.917419	2.63819	1.72618	2.344681	1.980136	2.0168	2.221353
AADAT	0.901021	2.117209	2.33651	1.226698	1.584483	3.170597	2.184566	1.468223	2.000801	2.486626
AAGAB	3.384627	4.911665	4.380784	3.860188	3.745479	3.901575	4.193553	4.471359	3.947197	3.8725
AAK1	0.891756	0.938476	2.48266	1.607293	0.550033	1.147723	1.93236	1.247271	1.246245	1.121613
AAMDC	2.905587	2.386935	3.092975	2.742403	2.883851	2.447846	2.444698	3.620468	1.942847	2.423223
AAMP	6.108965	5.606741	5.733432	5.76106	5.797197	5.758501	5.858539	6.147721	5.74096	5.935313
AAR2	4.62933	3.917359	5.064283	4.874728	4.248512	3.939791	4.888974	4.012086	3.845083	3.927044
AARS2	2.320018	2.451048	3.10118	2.902352	3.031804	3.035382	2.926849	3.194291	2.81012	3.086282
AARSD1	0.78421	0.60167	0.864706	0.742812	0.932212	0.947192	1.105827	0.979237	1.248157	1.35347

Figure 2. A sample of expression data file when viewed in Microsoft Excel.

3.2 Set Parameters

Through the parameters on the 'Data Preprocessing' panel the user can adjust the criteria for filtering genes. If a gene is filtered, then it will be excluded from further analysis. Assuming that the expression

vector of a gene is $\{v_1, v_2, \ldots, v_n\}$.

- Log normalize data—transforms the vector to $\{\log_2 v_1, \ldots, \log_2 v_n\}$.
- Maximum number of missing values—a gene will be filtered if the number of missing values in all samples exceeds this parameter.
- Set missing value as—the missing value of a gene can be set as 'min value' (default), 'mean value' or 'zero'.
- Set duplicated genes as—combine the duplicate expression profiles of a gene based on the 'mean value' or 'max value' (default).
- K-nearest neighbors—search the $\boldsymbol{k}$ nearest neighbors for each gene. The higher $\boldsymbol{k}$ is, the fewer cluster centers will be identified; as a consequence, fewer clusters will be generated.
- Correlation threshold—among the genes assigned to an expression pattern (p_1), genes showing absolute correlation coefficients higher and lower than this threshold (0.4 as default) with p_1 are divided into two groups, which are further applied to construct two new expression patterns. This parameter influences the final module number.
- Min number of module genes—the minimum (5 as default) number of genes in a module.
- Use known interactions to assist in module detection—if 'Yes' is selected, the files that currently is present in the 'Interaction data' directory of the CLAM directory will be used to assist in module detection.

3.3 Search Gene Modules

The text box here displays the running progress of CLAM after pressing the 'Run' button. After the analysis process is finished, CLAM will create a new folder named 'Results' in its directory to save the analysis results, including the table of final modules and the functional enrichment analysis results.