

## SurvivalGene Documentation

**Description:** Select genes correlated with time-to-event clinical outcome

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#### Summary

This module evaluates correlation of each gene's expression level with time-to-event clinical outcome such as time to tumor recurrence, disease progression, or death using Cox score [1,2] or Cox regression coefficient. Significance of the correlation is assessed based on standard random permutation test.

In contrast to gene selection based on class comparison (e.g., comparison between two sample groups using t-test), the correlation with time-to-event outcome is more affected by measurement in each single sample. That is, an outlier measurement in a single poor quality sample could falsely yield a high correlation. To avoid selecting such false positives, it is recommended to further refine the gene list by running LoocvSurvival module (this module takes an output file of SurvivalGene module. "SurvivalGene emp.stat.txt", as an input).

#### References

[1] N Engl J Med. 2008 Nov 6;359(19):1995-2004

[2] PLoS Med. 2006 Jan;3(1):e13

#### Parameters

Name	Description
input filename gct	Gene expression dataset (.gct)
input filename clinical	Clinical data (tab-delimited .txt, sample oder should be the same with .gct)
output file	Prefix for output file name, Default: SurvivalGene
time field	Field name for time variable, Default: time



censor field	Field name for censoring information (0: censor, 1: event) , Default: status	
statistic selection	Statistic to use, Choice: Cox.score, Cox.regression.coefficient, Default: Cox.score	
trim percent 2 side	Proportion of samples with outlier outcome to trim (2-side, e.g., "0.05"), Default: 0	
nperm	Number of sample name permutations to compute empirical distribution of statistic (must be >2 for Shapiro-Wilk normality test), Default: 1000	
rnd seed	Random seed, Default: 56438219	
emp stat dist	Output mean & SD of empirical distribution of statistic used for LoocvSurvival module? Perform Shapiro-Wilk normality test? (for Cox score only), Default: No	

#### Input Files

1. input filename gct

Normalized gene expression dataset in GenePattern GCT file format.

2. input filename clinical

Clinical dataset including time-to-clinical event variable and censoring information. Tab-delimited text. First column should be sample identified in the same order with the gene expression dataset (.gct).

#### **Output Files**

1. SurvivalGene.txt

Gene list with statistic (Cox score or Cox regression coefficient), p-value, Benjamini-Hochberg false discovery rate (BH.FDR), and Bonferroni-corrected p-value. Tabdelimited text.

 SurvivalGene\_emp.stat.txt (optional) Gene list with mean and sample standard deviation for Cox score. Last column is p-Shapiro-Wilk test p-value (test of normality for Cox score).



### Example Data

"SurvivalGene\_example\_gene\_expression\_data.gct": Gene expression data "SurvivalGene\_example\_clinical\_data.txt": Clinical data

# **Platform Dependencies**

Module type:	Gene List Selection
CPU type:	Any
OS:	Any
Language:	R