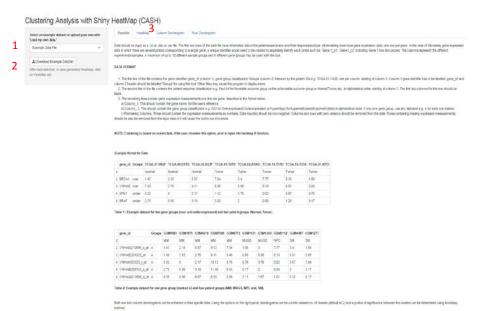
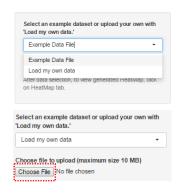
Analyzing data using CASH



1: Select dataset of interest. Using the dropdown, you can choose the example or upload your own. If uploading your own, follow instructions on right of how to format data before inputting.



If loading your own data, click on Choose File and browse to the location of input file and click open. Upload progress bar will initiate.

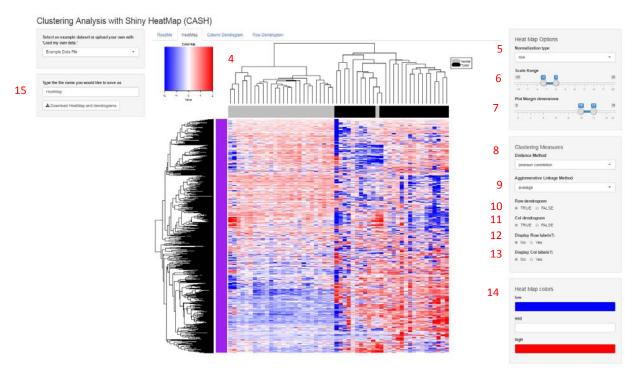


Wait for processing.

2: If using the example dataset (you will need to go straight to 3), but if you wish to download it, the 'Download Example Dataset' button will download the dataset in .csv format as seen below when using Chrome browser.



3: Click on the HeatMap tab in order to view it.



- 4: HeatMap created using 'row' normalization, 'Pearson correlation' distance and 'average' agglomerative linkage method (i.e. default settings). Depending on dataset may take several minutes to load.
- 5: Select a different normalization/scaling you'd like for the data using drop down options. Each time a different type is chosen, the heatmap will be updated.



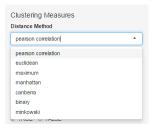
6 (optional): Drag slider to change scale range for the colors. Heatmap will be updated on movement.

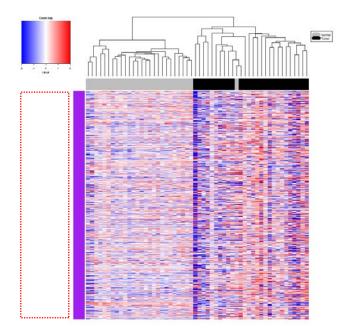


7: Select the Plot margins. If column dendrogram overlaps the legend, increase both margin points and vice –versa until desired.



8, 9: Select Distance method and linkage method of choice using the drop down options. Each selection will display modified heatmap.

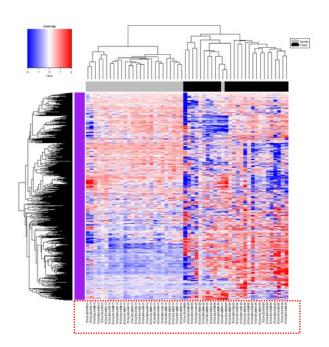




10, 11: Select either to display Row dendrogram or not. If FALSE is chosen, row dendrogram will disappear and data will not be ordered based on means. Same applies to Column dendrogram.

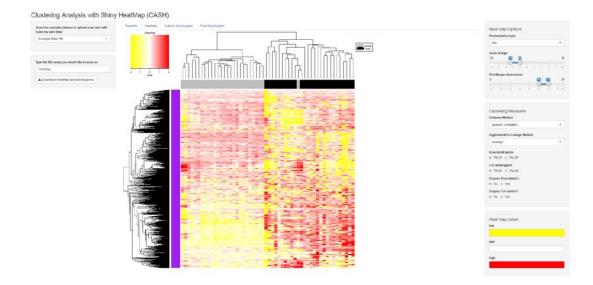
Row dendrogram

TRUE FALSE



12, 13: Select Display Row labels = 'Yes' to see the corresponding CpG sites. Additional slider appears to select, font size. Same applies to Sample labels.



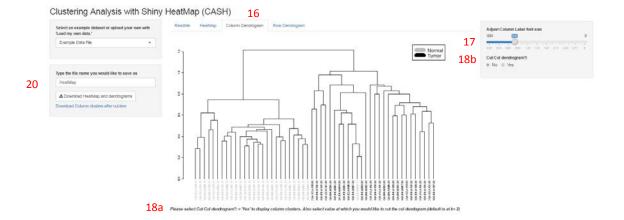


14: Select color scheme.
Red-Black-Green is
typically used for
Expression data and
Blue-White-Red is used
to represent methylation
data. Heatmap will
update as soon as color
is chosen. After choosing
desired color(s), click
anywhere on screen to
come out of color
selection panel.



15: Input file name and click on Download button to save heatmap and the corresponding row and column dendrograms in pdf format as shown below using Chrome browser.







19 Would you want to assess gene set significance in the separation of specimens into two clusters? (Yes/No)

	specificia filo E cidatera : 1
	○ No ● Yes
	Select a dataset or upload your own with 'Load my own data.'
19a	Meth Sampling Data ▼
19h	Sample size for bootstrap:
	1000
	No. of iterations for bootstrap:
19c	1000
19d	Gol Click the button to start sampling using bootstrap method for estimating the p-value. A progress indicator will appear shortly (~approx 10 s.) on too of page indicating
	the status. Once complete, the p-value will be displayed in the main panel.

Assess Gene set significance in separation of

- 16: View in column dendrogram tab
- 17: Slider to adjust font size of the column dendrogram labels
- 18 a, b: a. Option to cut the tree. b. If yes is chosen, user is asked at which position they want to cut the tree (default at 2)

Cut Col Dendrogram at:		
2		

When selected, a table will appear that classifies Samples, their Groups, and their corresponding clusters.

Use the drop down on upper left _____, to display 5/10/All rows of the table.

19: Option to assess gene set significance in separation of the two clusters (Tumor vs Normal). Applicable only when >=2 clusters are available for analysis.

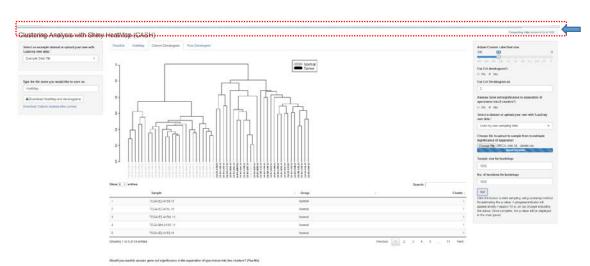
Assess Gene set significance in separation of		
specimens into 2 clusters?:		
No €) Yes	

When 'Yes' is selected, parameters for Monte Carlo pvalue estimation will be made available.

19a: Select Sampling dataset for bootstrap. An example Methylation Sampling data is available or user can input their own (up to 10 MB is allowed).



19b: Choose Sample size of the data for bootstrap. Use a size that does not exceed the original sampling data itself.



19c: Select number of iterations you wish to perform. A good practice is to perform at least 1000 iterations for accuracy of analysis.

19d: Once all options are selected, press 'Go' button to start analysis.

After approximately 10 seconds, a progress indicator and counter (top of page) will appear to track the time remaining for the analysis to get completed.

Computing data Iteration #: 299 of 1000

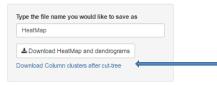
The p-value to test the gene set significance in the separation of specimens into 2 clusters is = 0

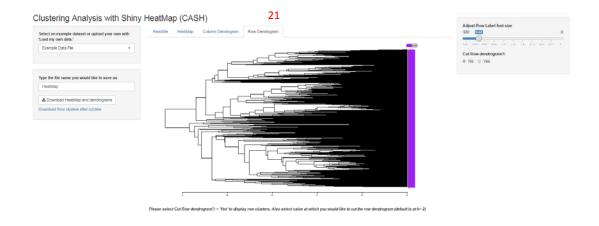
The gene set cluster is statistically significant, i.e., a random sample of CpG probes/gene sets of the same number is Not able to separate the specimens when compared to the CpG probes/gene sets of interest of the same class

p-value results from the boot strap approach for calculation significance of clusters using Fisher's exact test will be displayed under the table along with the interpretation.

20: To download the p-value results as well, input the file name and click on Download button. The heatmap and the corresponding row and column dendrograms followed by the p-value results will be downloaded in pdf format.

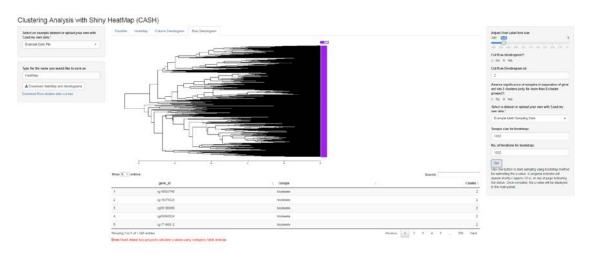
To download the table for the classification of samples by clusters, click on link and the table will be saved as an .csv file.





21: When in Row Dendrogram tab, follow the same steps as when in the column dendrogram tab.

If p-value is calculated here (not applicable here because of a single group), the results will automatically be incorporated in the downloaded pdf file.



If you attempt to calculate p-value where there is just a single group of CpG sites, an error will appear like shown but since analysis was not performed, the result table will not be included for download.

Error: Need atleast two groups to calculate p-values using contigency table anal

Output using the example dataset is available in a separate file titled 'CASH_HeatMap_2016-07-27 13-28-11.pdf'.