
RBCDE Documentation

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`rbcode.RBC(adata, clus_key='leiden', layer=None, use_raw=False)`

Compute the rank-biserial correlation coefficient for each gene in each cluster. The results can be subsequently turned into a marker list via the helper function `rbcode.filter_markers()`. The primary output is stored as part of either `.var` or `.raw.var`, depending on whether `.raw` data is used.

The rank-biserial correlation coefficient ([Cureton, 1956](#)) can be used as an effect size equivalent of the Wilcoxon test ([Kerby, 2014](#)), which in turn was deemed to perform well on single cell data problems ([Soneson, 2018](#)). Using effect size analyses is recommended for problems with large population sizes ([Sullivan, 2012](#)).

adata [AnnData] Needs per cell normalised data stored somewhere in the object (as either sparse or dense), and the desired clustering/grouping vector included in `.obs`.

clus_key [str, optional (default: "leiden")] The name of the `.obs` column containing the clustering/grouping.

layer [str or None, optional (default: None)] If specified, take the expression data from the matching `.layers` field. Overrides `use_raw` if provided.

use_raw [bool, optional (default: False)] If no `layer` was specified and this is set to `True`, take the data from the `.raw` field of the object. Store results in `.raw.var` to match dimensionality.

`rbcode.filter_markers(adata, thresh=0.5, use_raw=False)`

Filter the rank-biserial correlation coefficients computed with `rbcode.RBC()` to a list of markers for each cluster, provided as a data frame and a Scanpy plotting compatible `var_names` cluster marker dictionary. Returns those two objects, in this order.

adata [AnnData] Needs to have been processed with `rbcode.RBC()`.

thresh [float, optional (default: 0.5)] The threshold value used to call markers. Literature [critical values](#) can be used.

use_raw [bool, optional (default: False)] Set this to `True` if the raw data was used for the computation so that the results can be retrieved from the correct field of the object.

`rbcde.matrix.RBC` (*data*, *clusters*, *vars*)

Compute the rank-biserial correlation coefficient for each gene in each cluster. The results can be subsequently turned into a marker list via the helper function `rbcde.matrix.filter_markers()`. Returns a data frame with the coefficient value for each gene in each cluster.

The rank-biserial correlation coefficient ([Cureton, 1956](#)) can be used as an effect size equivalent of the Wilcoxon test ([Kerby, 2014](#)). Using effect size analyses is recommended for problems with large population sizes ([Sullivan, 2012](#)).

data [`np.array` or `scipy.sparse`] Per cell normalised, if using single cell count data. Variables as rows, observations as columns.

clusters [`np.array` or `list`] A vector of cluster/group assignments for each observation.

vars: np.array or list A vector of variable names, for output generation purposes.

`rbcde.matrix.filter_markers` (*results*, *thresh*=0.5)

Filter the rank-biserial correlation coefficients computed with `rbcde.matrix.RBC()` to a list of markers for each cluster. Returns a data frame of the computed markers.

results [`pd.DataFrame`] The output of `rbcde.matrix.RBC()`.

thresh [`float`, optional (default: 0.5)] The threshold value used to call markers. Literature [critical values](#) can be used.

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