

# Correlation analysis

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## What is Correlation analysis?

*Correlation analysis* is a statistical test that lets you rank features by their correlation with numeric attributes using Pearson (linear), Spearman (rank), or Kendall (tau) correlation.

## Running Correlation analysis

We recommend normalizing you data prior to running *Correlation analysis*, but it can be invoked on any counts data node.

- Click the counts data node
- Click the **Statistics** section in the toolbox
- Click **Correlation**
- Choose the method to use for correlation analysis (Figure 1)

Method to use for correlation analysis	
<input checked="" type="radio"/> Feature many-to-one correlation Rank features by their correlation with one or more quantitative attributes.	<input type="radio"/> Correlation across assays Correlate features in one data node vs features in another data node.

Figure 6. Choose the method to use for correlation analysis

## Feature many-to-one correlation

When multiple numeric factors are added, the correlation analysis will perform each factor with a feature in the data node independently. If you are interested in particular features, use the **Search features** box to add one or more.

- Select the factors and interactions to include in the statistical test (Figure 2).


### Select factor(s) for analysis

**Numeric factors**

☐ # Cells
 ☒ Age in years
 ☐ Expressed genes
 ☐ Total count

**Features**

Search features...

Add factors 

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### Selected factor(s)


Factor	Delete
Age in years	

Figure 7. Select the factors and interactions to include

- Click **Next**
- It is optional to apply a lowest coverage filter or configure the advanced settings
- Click **Finish** to run

*Correlation analysis* produces a *Correlation* data node; double-click to open the task report (Figure 3) which is similar to the [ANOVA/LIMMA-trend/LIMMA-voom](#) and [GSA](#) task reports and includes a table with features on rows and statistical results on columns.


Feature list									
Results: 2120		Optional columns							
Filter <a href="#">Clear all</a>					Age in years				
<input type="checkbox"/> Feature ID		View	Feature ID ↑↓	Ensembl ID ↑↓	Gene name ↑↓	P-value ↑↓	FDR step up ↑↓	Partial correlation ↑↓	
<input type="checkbox"/> Ensembl ID		1	MYO1F	MYO1F	MYO1F	0	0	0.21	
<input type="checkbox"/> Gene name		2	MAL	MAL	MAL	0	0	-0.19	
<input type="checkbox"/> P-value		3	CST7	CST7	CST7	0	0	0.34	
<input type="checkbox"/> FDR step up		4	B2M	B2M	B2M	0	0	0.32	
<input type="checkbox"/> Low expressed		5	CD63	CD63	CD63	0	0	0.18	
<input type="checkbox"/> Partial correlation		6	FGFBP2	FGFBP2	FGFBP2	0	0	0.32	
<a href="#">Save filter</a>		7	ID2	ID2	ID2	0	0	0.25	
Saved filters 		8	TMSB4X	TMSB4X	TMSB4X	0	0	0.23	
(No saved filters available)									
<a href="#">Generate filtered node</a>									
<a href="#">Save as managed list</a>									

Figure 8. Correlation analysis task report

Each numeric attribute includes p-value, adjusted p-value columns (FDR step up and/or Storey q-value if included), and a partial correlation value. Each interaction will have p-value and adjusted p-value columns (FDR step up and/or Storey q-value if included).

Each feature includes  [chromosome view](#),  [dot plot](#),  [correlation plot](#), and extra details  buttons in the *View* column.

## Correlation analysis advanced options

### Low value filter

*Low-value filter* allows you to specify criteria to exclude features that do not meet the requirements for the calculation. If there is a filter feature task performed in the upstream analysis, the default of this filter is set to **None**, otherwise, the default is **Lowest average coverage** is set to **1**.

*Lowest average coverage*: the computation will exclude a feature if its geometric mean across all samples is below the specified value

*Lowest maximum coverage*: the computation will exclude a feature if its maximum across all samples is below the specified value

*Minimum coverage*: the computation will exclude a feature if its sum across all samples is below the specified value

*None*: include all features in the computation

### Multiple test correction

Multiple test correction can be performed on the p-values of each comparison, with **FDR step-up** being the default. If you check the *Storey q-value*, an extra column with q-values will be added to the report.

### Use only reliable estimation results

There are situations when a model estimation procedure does not fail outright but still encounters some difficulties. In this case, it can even generate p-value and fold change on the comparisons, but they are not reliable, i.e. they can be misleading. Therefore, the default of *Use only reliable estimation results* is set **Yes**.

### Correlation type

Sets the type of correlation used to calculate the correlation coefficient and p-value. Options are *Pearson (linear)*, *Spearman (rank)*, *Kendall (tau)*. Default is **Pearson (linear)**.

## Correlation across assays

*Correlation across assays* should be used to perform correlation analysis across different modalities (e.g. ATAC-Seq enriched regions vs. RNA-Seq expression) for multiomics data analysis.

- Select the data node to be compared to the node that the task has been invoked from using the **Select data node** button
- Modify any parameters (Figure 4)
- Click **Finish**

Select data

Select data node

Clear selection

Correlation and similarity measures

Features to correlate

☒ Features within same chromosome  
Restrict features to chromosome location.
☐ All features in one data node vs all features in the other data node  
All combinations

Correlation method

☒ Pearson
☐ Spearman

Report correlation pairs

☐ P-value < 
☐ abs(Correlation coefficient) >

Figure 9. Correlation across assays can be performed with multiomic data

## Correlation across assays analysis options

### Correlation and similarity measures

*Features within same chromosome:* this option will restrict feature comparison to the chromosome location

*All features in one data node vs all features in the other data node:* this option will perform the comparison using all combinations without location constraint

$$r_{xy} = \frac{\sum_i (x_i - \bar{x})(y_i - \bar{y})}{\sqrt{\sum_i (x_i - \bar{x})^2} \sqrt{\sum_i (y_i - \bar{y})^2}}$$

*Pearson:* linear correlation:

$$r_s = \frac{\sum_i (R_i - \bar{R})(S_i - \bar{S})}{\sqrt{\sum_i (R_i - \bar{R})^2} \sqrt{\sum_i (S_i - \bar{S})^2}}$$

*Spearman:* rank correlation:

### Report correlation pairs

*P-value:* select a cut-off value for significance and only those pairs that meet the criteria will be reported

*abs(Correlation coefficient):* select a cutoff for reporting the absolute value of the correlation coefficient (represented by the symbol r) where a perfect relationship is 1 and no relationship is 0

*Correlation across assays* produces a *Correlation pair list* data node; double-click to open the table (Figure 5). The table can be sorted and filtered using the column titles.

Optional columns

View	Feature 1 ID ↑↓	Feature 2 ID ↑↓	Chromosome ↑↓	Pearson's r ↑↓	P-value ↑↓
	FCRL1	chr1:38474653-38475376	1	0.50	0
	MS4A1	chr11:60455254-60456156	11	0.56	0
	IRAK3	chr12:119988567-119989449	12	-0.47	1.47E-292
	BLK	chr8:11545456-11546358	8	0.47	2.75E-287
	LRRK2	chr12:119988567-119989449	12	-0.45	1.1E-263
	SYK	chr9:107489482-107490363	9	0.43	3.59E-241
	CD22	chr19:17776260-17777083	19	0.43	1.03E-239
	ADAM28	chr8:11545456-11546358	8	0.42	1.11E-231

Figure 10. Correlation across assays table

Click [View correlation plot](#) to open the correlation plot for each comparison. Scroll to the bottom of the table to [Download table](#) download the full table report.

Additional Assistance

If you need additional assistance, please visit [our support page](#) to submit a help ticket or find phone numbers for regional support.

Your Rating: Results: 12 rates