

OMICs data analysis in Qlucore: Plots in focus

Yana Stackpole, PhD Training, Application Support



Visualize and Explore

- QC (outliers, mislabeled samples)
- Make observations identify structures, patterns
- Generate new hypotheses
- Browse the genome



for Legend	• • •		Q	acore '	Test D	lata S	et 1.1							
				Г			_			7				
- 1							1			-				
	Standard Extended		[_										
- 1	Input All active variables 50/50 va								L .	_				
	Input: All active variables 50(50 va	1			⊥				1	Т	l	Т		
1	Filter by Variance (d/ame)													T
н	0 0													Treatment
						_	_	_	_	_	_	_		
	Filter by Two Group Comparison													
														ID_13
	Treatment 📴 🛄 Placebo													10_15
н	Eliminated factors													
														ID_21
		-												
	p = 0.0295 C q = 0.015598 C													
	tu ≥ 2.5373 R ≥ 0.62582													ID_40
	Filter by Fold Change													
		5306	5307	5305	5308	5302	5303	5301	5304	5312	5311	ä	5310	
	1.5	ð	1	S	8	Ň	ω	Ħ	¥	N	÷	ö	9	
- 1														

Analysis

- t-test, ANOVA, Regressions, R scripts. Open API to R, Batch exec
- Variant calling
- Easy generation and export of reports, status, and plots
- Save your session, share

Biological Insight

- GSEA using MSigDB, reactome, or custom gene sets
- GO Browser
- GO enrichment
- NDEx

AI - Classify and Predict

- Build classifiers
- kNN, SVM, RT
- Predict sample class, outcome, etc.



Enjoy fantastic computing speed on a laptop to boost your discovery and scientific creativity

Benchmark examples (static). Compared to R	Times faster
ANOVA (22k var. + 130 samples)	2800
t-test (two-groups, selected from 22k var. + 130 samples)	1000
Kruskal-Wallis (22k var. + 130 samples)	900
Mann-Whitney U-test	480
(two groups, 30k var. + 5k samples)	
ANOVA (30k var. + 5000 samples)	180
PCA calculations (30k var. + 150 samples)	77
UMAP (22k var. + 130 samples)	13

The speed enables a more flexible workflow – generating better results faster.

Details at: https://glucore.com/calculation-benchmarks

"This tool might literally save you years of your life"

Prof. Ulrich Steidl at Albert Einstein College of Medicine



Plot options

	Window	License	Help	
			Qlucore Omic <mark>s Explorer 3.9</mark>	
oolbox			Launch Data Met	hod Optic
Multi. Move Color List Info Mark Flip Clear Annot. Label Corr.	Cent.		Plot Mode Mode PCA Co Co <	Normalizatio
Samples Variables Log Variables sts [+ 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1			PCA SampleHistogram VariableTableTalusPCA VariableBar SampleVolcanoScreeScatter SampleBox Sample2D t-SNEPie Chart SampleScatter VariableBox Variable2D UMAPPie Chart Variable	
Acute Lymphoblaemia_ Collapse		100	Line Sample Violin Sample Kaplan-Meier Fusions Circular	
anova subtype top 100		100 100	Line Variable Violin Variable Silhouette Genome Histogram Sample Heat Venn	
high variance_top 100	100	100		
T ALL vs others, top 100	100	100		
HALLMARK_P53_PATHWAY	200	193		



Venn Diagram

John Venn introduced it in an 1880 paper entitled:

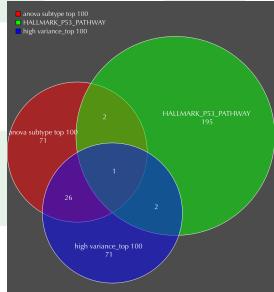
<u>"On the Diagrammatic and Mechanical Representation of Propositions and Reasonings</u>"

Math: Venn diagrams are used for sets, intersections, and unions

Statistics and probability: Venn diagrams are used to express the relationships between two or more data sets in an easily understood way – efficient connections between things

Simple + Powerful, especially:

- to highlight similarities and differences, and
- to compare and contrast the characteristics of different sets.





When?

- Venn's initial rationale for it: "Some of X is Y and some of Y is X." If *this* is what you're trying to communicate, then Venn.
- "These sets have more (or less) in common than you might think."
- "We're talking about a *very specific* subset."



Examples of Sets/Lists

- 1. Whole data sets
- 2. Data subsets:
- Statistically significant results
- Variance-based lists
- Pathways, GO lists
- Any lists

Qlucore Omics Explorer File	Windo	w Lice
•••		
Toolbox		
Multi. O Move O Color O List		ent. 🔘 🕽
Clear Info Mark Flip		Val. 🔿 Y
Annot. Label Cor	F.	
Samples Variables Log	NGS	
Variables		
Lists		Q
Search	0	0
Acute Lymphoa_ Collapsed	13266	13266
anova subtype top 100	100	100
high variance_top 100	100	100
🗌 💋 T ALL vs others, top 100	100	100
HALLMARK_P53_PATHWAY	200	193
from Venn anova+P53+high var	1	1
IDs 📶 🗸		<i>L</i>

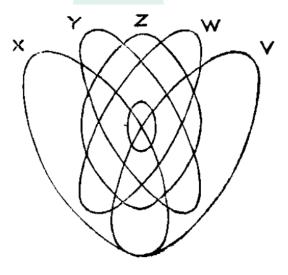


Venn Diagram

How many sets (terms, gene lists, data sets/subsets) can be used?

See example with 5 sets/terms below (original paper)

I and w, but are no part of 2). It must be admitted that such a diagram is not quite so simple to draw as one might wish it to be; but then we must remember what are the alternatives before any one who wishes to grapple effectively with five terms and all the thirty-two possibilities which they yield. He must either write down or in some

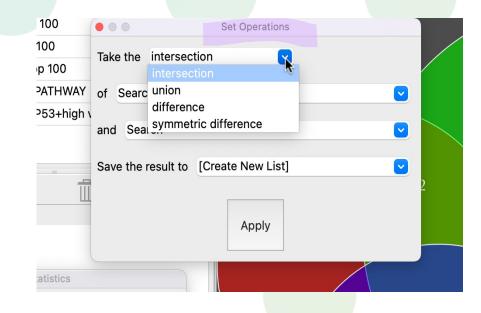




Handling sets two at a time?

Yes! Not visual, manual, tedious, more operator error prone:

- Intersection of A and B
- Difference A vs B (unique to A)
- Symmetric Difference (unique to A and B merged)
- Merge sets without duplicates





What makes the Qlucore software unique?

- Instant visual feedback
- Easy: to learn to use to remember how to use
- Automated access to public data as GEO, TCGA
- Easy import of other public data in a matrix format like GREIN
- Share analyses, save your sessions to resume later \bigcirc



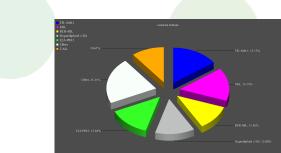
Pie Chart – part-to-whole relationship in your data

- Displays **relative proportions** of multiple classes of data (%, frequency).
- Size of the circle can be made **proportional** to the total quantity it represents. Hence, can serve as a visual QC check of calculation accuracy.
- Summarizes a large data set in visual form overview of classes.
- Visually simpler than other types of graphs.

1st known Pie – 1801 Scottish writer on political economy <u>William Playfair</u> (1759-1823), known as the inventor of statistical graphs.

"Making an appeal to the eye when proportion and magnitude are concerned, is the best and readiest method of conveying a distinct idea"

The pie chart and <u>bar chart</u> were described. Some believe that he also invented the <u>line</u> plot



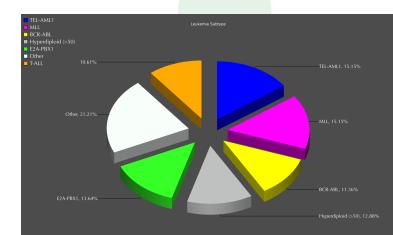


Pie Chart –

part-to-whole relationship in your data

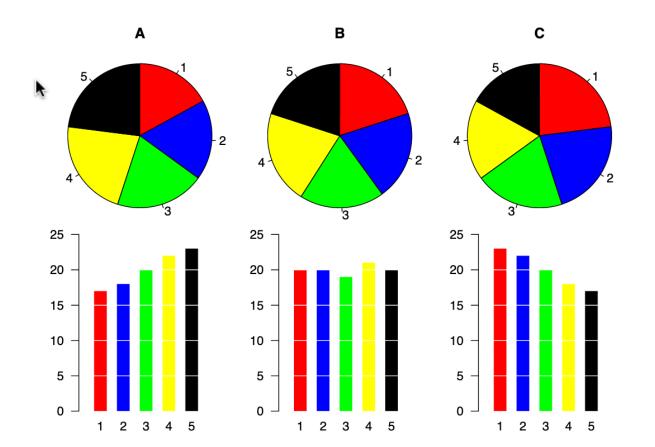
Keep in mind:

- Too similar slices (unless you need to show that, but really no need in a plot). Simple bar plot is better then.
- Exploded plot may distort proportions, but helps with many slices.
- Too many slices.
- Use of **color**.





Pie vs Histogram "evil pie"





DEMO



Next steps Experience your data in depth:

 Join free fully supported trial -- download and install the trial version (<u>Mac or Win version - links</u>), and then update the license key in Qlucore – License – Activation Key with the key from us

Book a session with us – get help setting up your analysis.
 <u>https://calendly.com/yana-stackpole/30min</u>

3. Link to join our Discord channel with instructional videos and 30 min webinars:

https://discord.gg/zu8h8FJx

Reach out any time <u>Yana.Stackpole@Qlucore.com</u>



System requirements Base module

FOR WINDOWS

- Windows 7, Windows 8 or Windows 10
- 512 MB of RAM memory
- A graphical card with support of at least Open GL 2.1
- 5 GB of free hard disk space
- The program takes full advantage of processors with multiple cores and computers with multiple processors.

FOR MAC

- Max OS X 10.15 or 10.14
- 512 MB of RAM memory
- A graphical card with support of at least Open GL 2.1
- 5 GB of free hard disk space