# Accelerate Insights with the Proteograph<sup>™</sup> Analysis Suite















Explore PAS

For Research Use Only. Not for use in diagnostic procedures.

## Get a clearer view of the proteome

The Proteograph<sup>™</sup> Product Suite enables unbiased, deep, and rapid proteomic analysis at scale. Survey thousands of human plasma proteins to discover new insights or quantify protein abundance to uncover biological meaning; Seer's Proteograph Product Suite delivers quantitative, accurate, precise, and reproducible data for proteome studies of any size.

The Proteograph Analysis Suite (PAS) is a dedicated cloud-based software solution for processing, analyzing, and visualizing proteomics data generated by liquid chromatography-mass spectrometry (LC-MS). The integrated search engines power rapid identification and annotation of LC-MS data and the variety of quality control tools ensure the highest confidence in your insights quickly and efficiently.



**Efficient Workflows** 



#### Automated Data Upload

Upload data automatically from the LC-MS to PAS without manual intervention.



# State-of-the-Field Peptide Identification and Protein Quantification

Integrated database search engine and analysis wizard allows automated peptide identification and protein quantification for seamless generation of results. Seer's proprietary human spectral library files are a deep and extensive resource allowing users to identify and quantify proteins in their datasets.



#### Pre-Configured Data Filtering Pipeline

Current best practices for peptide sequence annotation, statistical false discovery filtering, protein assignment, and protein quantification are pre-configured to enable routine analysis without the need for advanced bioinformatics expertise.



#### Visualization Tools for Biological Insight

Results are automatically generated including data tables compatible with downstream analysis tools, and intuitive visualization options to evaluate assay results and performance.



=

## **Analysis Summary and Metrics**

Quickly and easily evaluate experimental results with automatically generated summary and metric plots showing plate map summaries, intensity and coefficient of variation (CV) plots, peptide and protein group counts, protein group overlaps between samples, and sample comparability.

Plate Map Summaries: Quickly assess the results of various experimental metrics across samples to for a high-level overview of experimental performance.

Protein Intensities: Visualize similarity

between samples by comparing the distributions of protein group intensities.

Protein Group Counts

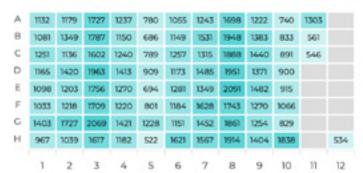


Figure 1: View results for protein groups (shown) and peptide counts, quant mass, miscleavage rate, oxidation ratio and ID rate in a simple and intuitive plate format.

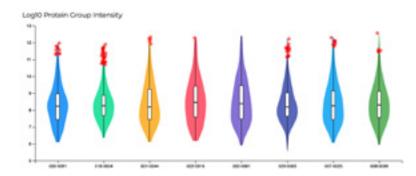


Figure 2: Distributions of protein group intensities and CVs across samples.

Distribution of Detected Proteins in Plasma: Visualize Proteograph's compression of the plasma proteome's

dynamic range by comparing to a deeply covered reference plasma proteome dataset.

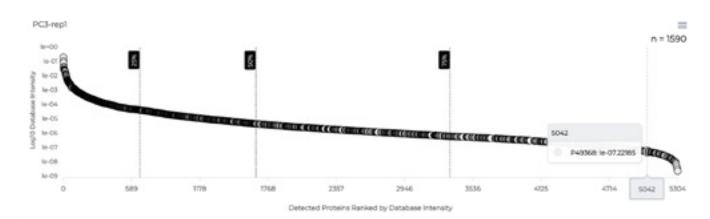
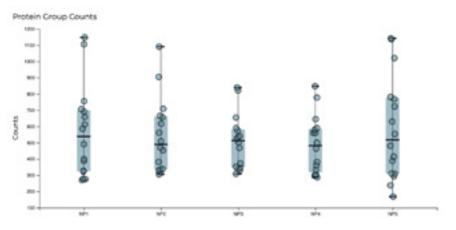
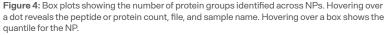


Figure 3: Plots show the dynamic range of identified proteins compared to the deepest reported human plasma proteome index.

**Peptide/Protein Group Counts:** Gain insight to differences in protein groups captured across nanoparticles by examining the number of protein groups identified across NPs.





**Protein Group Overlap:** The Protein Group Overlap Sets section is divided into two bar graphs and a matrix that together show protein group intersections.



Figure 5: (A) Graphs and a matrix show protein group overlaps; Intersection Size bar graph (B) Protein group count bar graph (C) Matrix.

**Sample Comparability**: Displays the degree of statistical correlation between samples based on the Pearson correlation coefficient (PCC), which measures the linear correlation of data.





Figure 6: A color-coded matrix displays sample comparability data using PCC (left) or the Jaccard index (right). Samples on the green end of the spectrum have high correlation, while samples on the red end of the spectrum have low correlation.



## **Quality Control Metrics**

Visualize quality control data in an intuitive, color-coded dashboard. Each chart plots one metric for one process control category. QC charts are organized with the same metric across each row and the control type in each column. The x-axis is labeled with the date of analysis and the y-axis depends on the metric.

												Cartesi		rençi	General		na Can		
MS instru	ment		Color By:	Start Date		End Date				Control Lin		Exclusion Calculate Annetata				a fiel			
- unit	nown 📴 FSNQ0699		Default +		1/2021			3/3/4	100		Preinst	alled	*		D	B	đ		
						,			r		h ,				/				
HS instrument	Control	well	Plate		10	-19	4	4	4	4	1/1	14	4	4	5				
15420699	Process Control	AD	Icarius 2	029ms0234															
F\$N20699	Process Control	,AD	Icarius 2021me0																
P5N20699	MPE Cantrol	сп	loarius 2	2021ma0214	0	0		0	0	0	0	0		0	0				
P5N20699	Mass Spec Control	на	Icarius 2	2021ms0214	0	0		0	0		0		0	0	0				
F5N20699	Digestion Control	811	icarius.)	2027ms0274		0			0										
PSN20690	Digestion Control	811	icarius 2	2021/mai0210		0			0						0				
P5N20699	Mass Spec Control	HIZ	icarius 2	02Pme028		0			0	0	0	0	0	0	0				
						27 Ke	ris												
Process	Control		Digestion Control			=	MPE Control			=		Mass Sp		ass Spec	Control				
540	540 • PLADON		330 • raccare					120 • racises						120					
480			300					80							200				
5		5	2					5						1 Second					
		8	2 140 ······					3+0						240					
3 340	8 360			5-1899					200						200				
1		8	210				8							â					
300	800			100					800						160				
240			150 May 10 2021 3un 20 2021												120	90			
May 10 2021								May 10 2021 3un 20 2021						May 10 2028 Jun 20 2028					

Figure 7: Control Results: (A) Filters for viewing charts for all controls or a selected control type. (B) Toolbar with additional filters and functions. (C) Summary of control data for the selected analysis time frame. (D) QC charts with metrics for each control.



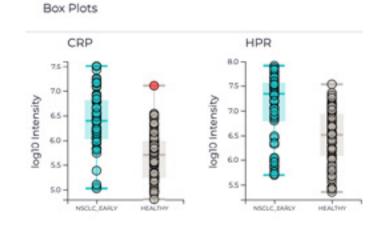
## **Differential Expression Analysis**

Interactive plots enable functional and biological interpretation of results. Seamlessly evaluate expression differences between study samples using Group Analysis or output raw and processed protein expression tables for custom analyses. To support Group Analysis, PAS provides a variety of MS database search engines including MaxQuant<sup>1</sup> for DDA-based analyses, and either EncyclopeDIA<sup>2</sup> or DIA-NN<sup>3</sup> for DIA-based analyses.

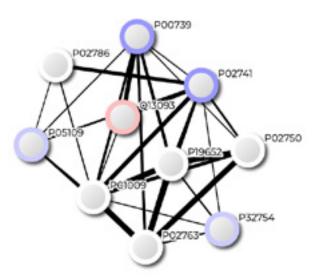
**Sequence Coverage:** Visualize where peptides map relative to the protein sequence



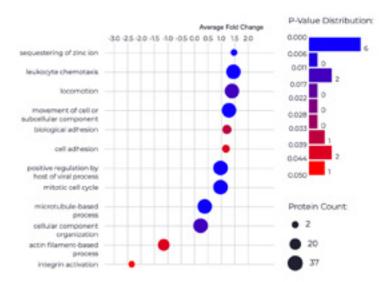
**Intensity Comparison:** View how the intensity of a protein of interest differs between groups



**Protein-Protein-Interactions Comparison:** Build a PPI network using the STRING database<sup>4</sup> to visualize possible protein interactions



**Gene Ontology (GO) Enrichment:** Perform GO enrichment analysis<sup>5</sup> to explore how proteins associated with a group differ functionally<sup>6,7</sup>



## Identify and Explore Variant Peptides with the Proteogenomics Workflow

Proteograph Analysis Suite now includes a Proteogenomics workflow to identify and explore peptide variants arising from allelic variation or other user-defined protein sequence altering variants, through the integration of Proteograph proteomics data with NGS variant information (Figure 8). The workflow offers a scalable and easy-to-use solution to a typically computationally intensive bioinformatics pipeline.

#### Build a custom peptide database

Upload a custom or samplespecific variant call file (VCF) to predict protein altering variants not captured in the canonical reference database. Personalized variant peptides are automatically combined with the canonical reference database to generate a customized database.

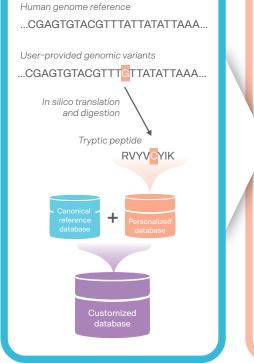
# Perform a search for variant peptides

Using the customized protein sequence database, search your LC-MS/MS Data Dependent Acquisition (DDA) data for variant peptides utilizing MSFragger search algorithms in PAS.

# Browse and explore your variant peptide results

Variant peptide results are summarized in an interactive table and plots. Browse peptide and variant peptide data maps in genomic space at nucleic acid/amino acid resolution (Figure 8). Visualize gene structure, protein domain information, and region information with respect to identified peptides.

# Build custom peptide database



# Perform variant peptide search



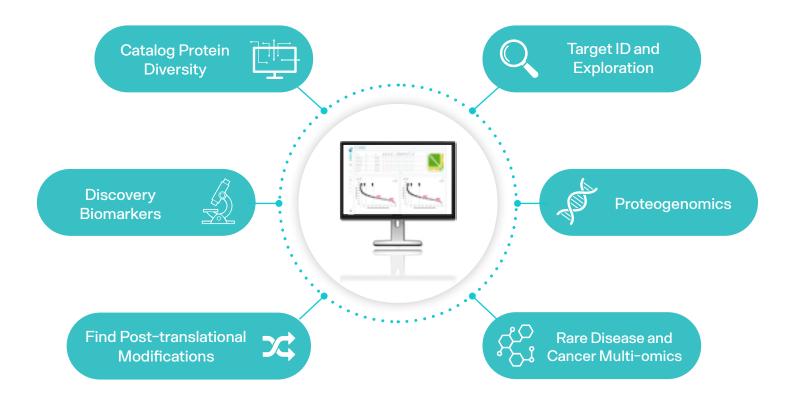
# Browse and explore variant peptide data



Figure 8: Proteograph™ Analysis Suite Proteogenomics Workflow

## Get Started With Proteograph Analysis Suite

Seamless QC and data analysis designed for speed and reproducibility, enabling powerful biological insights.



## Become a PAS User



#### References

- 1. Cox et al., Nat Biotechnol 2008; 1367–1372
- 2. Searle et al., Nat Commun 2018; 5218
- 3. Demichev et al., Nat Methods 2020; 41-44
- 4. Szklarczyk et al. Nucleic Acids Res 2015;43 D447-52
- 5. Faria, D. GOEnrichment, (2017). https://github.com/DanFaria/GOEnrichment
- 6. Ashburner et al., Nat Genet 2000;25(1):25-9.
- 7. The Gene Ontology resource: enriching a GOld mine. Nucleic Acids Res 2021;49(D1):D325-D334.

To learn more about the Proteograph Product Suite or to be kept up to date on recent information – visit our website or follow us on social.



For Research Use Only. Not for use in diagnostic procedures.

Use of the Proteograph Analysis Suite is subject to the terms and conditions contained in Seer's end user license agreement. Use of PAS may incur separate cloud compute and data storage fees. Seer, Proteograph and the Seer logo are trademarks of Seer, Inc. All other marks are the property of their owners.



3200 Bridge Pkwy #102 Redwood City, CA 94065

info@seer.bio | seer.bio