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Seer 40th Annual J.P. Morgan Healthcare Conference

Omid Farokhzad, M.D. CEO, Seer

January 10, 2022



Safe Harbor Disclosures

Certain statements in this presentation and the accompanying oral commentary are forward-looking statements within the meaning of the federal securities laws. These statements relate to future events or Seer, Inc. (the "Company")'s future results and involve known and unknown risks, uncertainties and other factors that may cause the actual results, levels of activity, performance or achievements of the Company or its industry to be materially different from those expressed or implied by any forward-looking statements. In some cases, forward-looking statements can be identified by terminology such as "may," "will," "could," "would," "should," "to," "target," "expect," "plan," "anticipate," "intend," "believe," "estimate," "predict," "potential" or other comparable terminology.

All statements other than statements of historical fact could be deemed forward-looking. These forward-looking statements are subject to a number of risks, uncertainties and assumptions, including, among other things: any expectations regarding the Company's projections of market opportunities; statements regarding the Company's business strategy, operations, results of operations, financial needs, and financial condition; statements regarding the Company's long-term expectations; statements that may suggest trends for the Company's business or industry, including expectations that may affect the unmet need and the size of the proteomics market and adjacent markets; statements about the Company's broad commercial release, the center of excellence program and consortium arrangement, statements about the Company's ability to successfully commercialize the Proteograph M Product Suite, demand for the Proteograph Product Suite; the launch of any new or additional products, any expectations or statements regarding customer acquisition in domestic or global markets, including but not limited to Europe and Asia; statements regarding customer adoption of new technologies domestically and globally; the Company's ability to expand life sciences markets through the use of its technology; the discovery of new protein variants and novel biomarkers leading to therapeutic breakthroughs, the scope of protection the Company is able to successfully establish and maintain for intellectual property rights, including its Proteograph Product Suite; projections, assumptions, and estimates of the Company's future performance and the future performance of the markets in which it operates; the Company's expectations regarding its gross margins, and operating income and expenses; any statements of the plans, strategies, and objectives of management for future operations; any statements of expectation or belief regarding future events, opportunities to drive future growth, and potential markets or market size, or technology developments.

While the Company believes these expectations, assumptions, estimates and projections are reasonable, such forward-looking statements are only predictions and involve known and unknown risks and uncertainties, many of which are beyond the Company's control. These and other important factors may cause actual results, performance, or achievements to differ materially from those expressed or implied by these forward-looking statements. The forward-looking statements in this presentation are made only as of the date hereof. For a further description of the risks and uncertainties that could cause actual results to differ from those expressed in these forward-looking statements, as well as risks relating to the business of the Company in general, are described more fully in the Company's filings with the Securities and Exchange Commission ("SEC") and other documents that the Company subsequently files with the SEC from time to time. The Company specifically disclaims any intention to update any forward-looking statements included in this presentation. If one or more of these statements is updated or corrected, investors and others should not conclude that additional updates or corrections will be made.

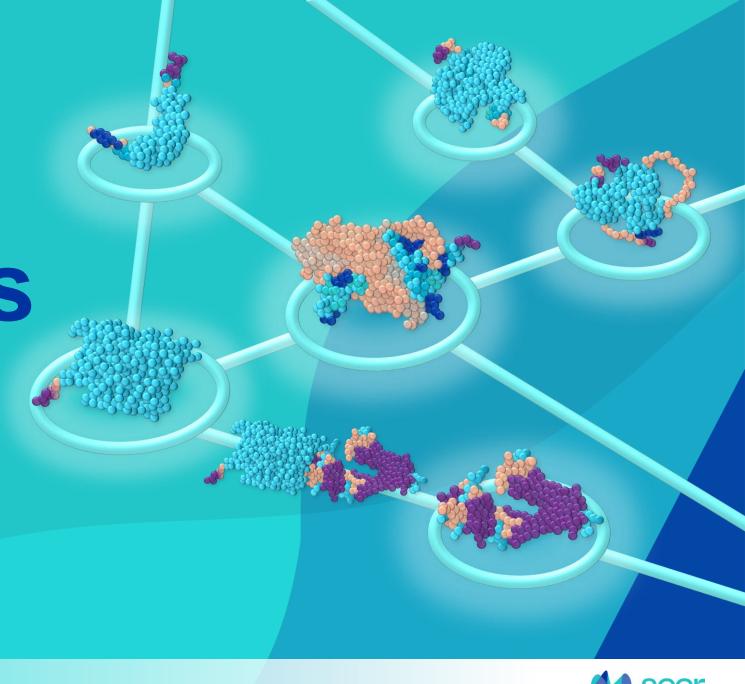
In light of the foregoing, investors are urged not to rely on any forward-looking statement in reaching any conclusion or making any investment decision about any securities of the Company.



We imagine and pioneer new ways to

decode the secrets of the proteome

to improve human health





Seer is leading the proteomics revolution



Leading customers and collaborators

Lighthouse customers
Well designed customer experience



Creating new markets

Industry leading partners
Geographically positioned



Best in class technology

Uniquely enabling technology

Data demonstrating impact

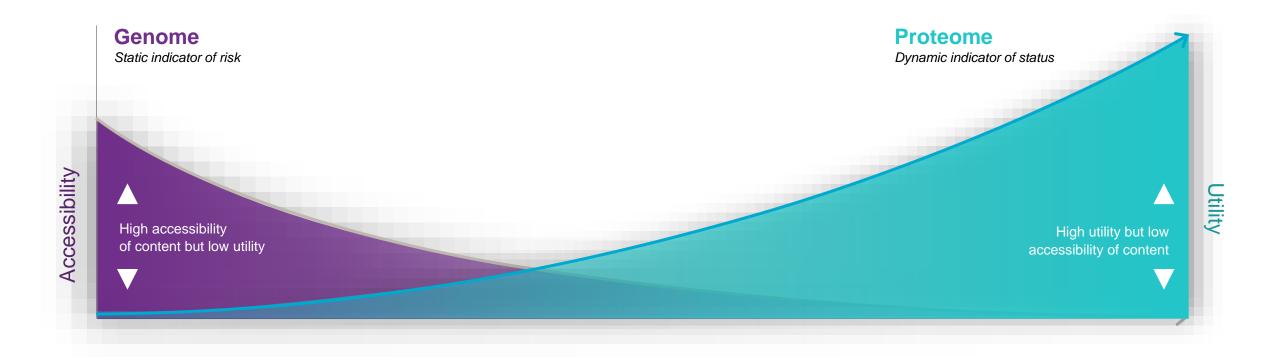


Leading team with multi-omics expertise

Culture of innovation Industry-leading talent



Full characterization of the proteome is essential





10M+ human exomes and 1M+ genomes and counting



~1B genetic variants catalogued



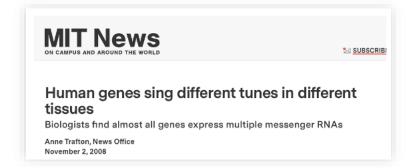
< 0.2% of genetic variants fully characterized

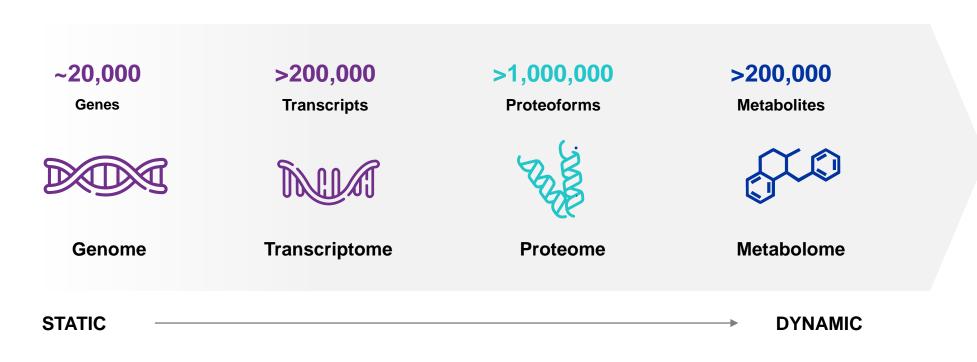


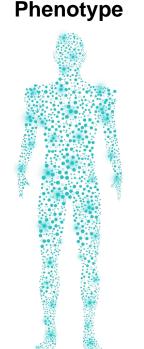
Biology is a dynamic and complex matrix of interactions









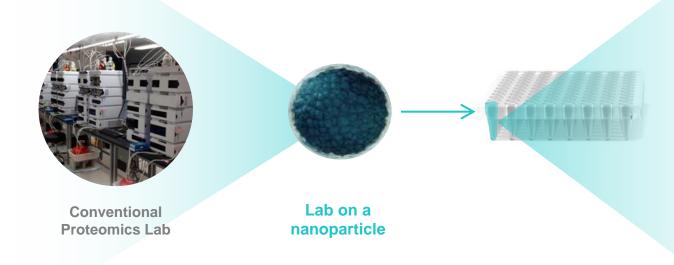




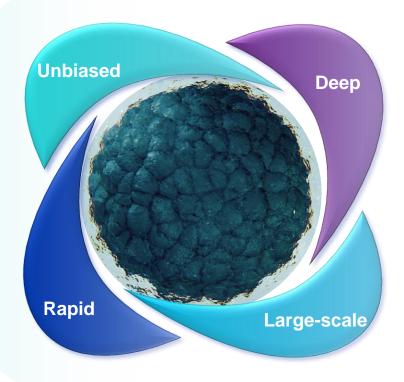
Opening a new frontier with unbiased, deep, rapid proteomics at scale **Proteomics** ~\$32B Genomics



Seer enables unbiased, deep and rapid proteomic analysis at scale

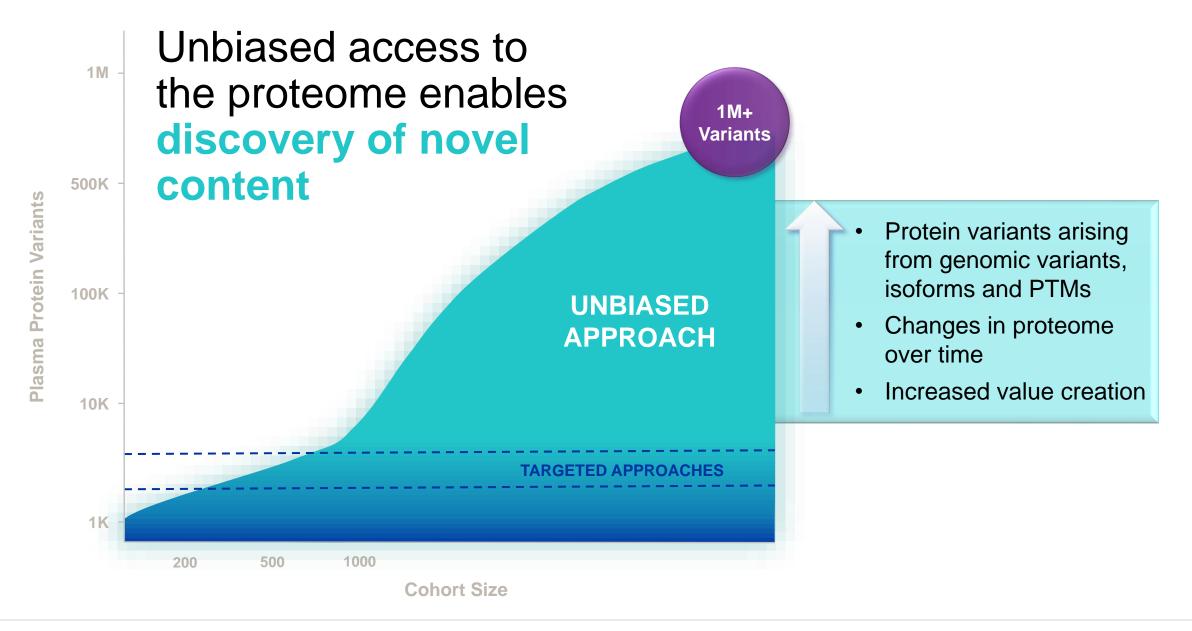


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A New Gateway to the Proteome



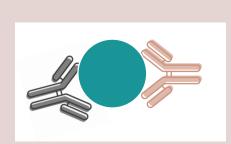




Unbiased approaches identify peptide level insights

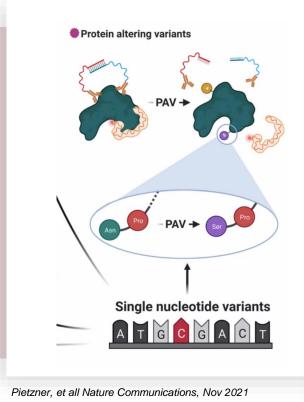
Targeted Approaches

Miss the Vast Majority of Protein Altering Variants



Average human protein: 472aa long

Average epitope: 5-8 aa long

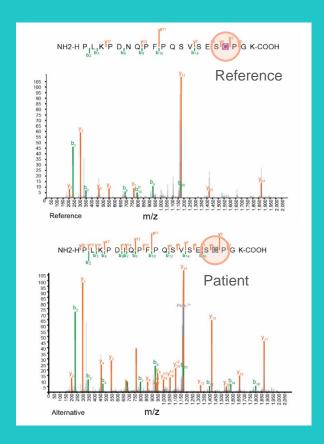


"Protein-phenotype colocalisation ... goes beyond missing target specificity and can be explained by alternative proteoforms induced by genetic variants altering the amino acid sequence of the protein and the effects of alternative splicing"

Pietzner, et al Nature Communications

Unbiased Approaches

Capture Common and Rare Variants





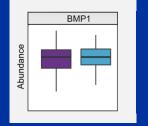
Unbiased approaches identify peptide level insights

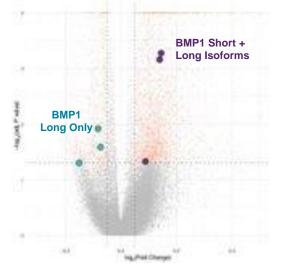


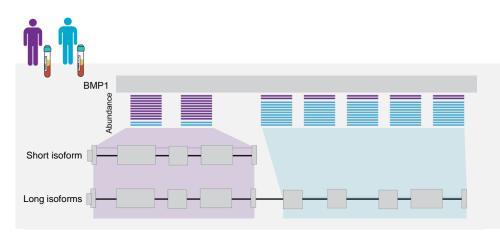
Peptide Level



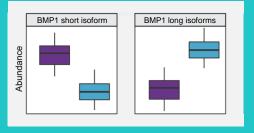
No Difference Detected





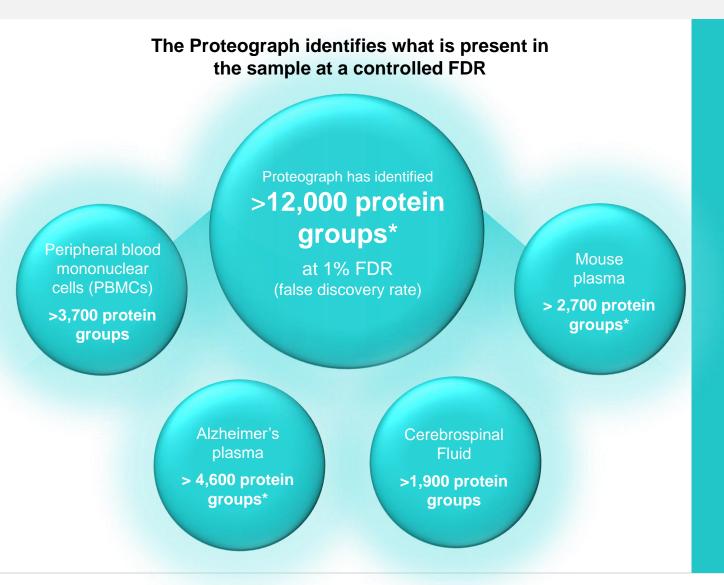


Meaningful Difference Detected

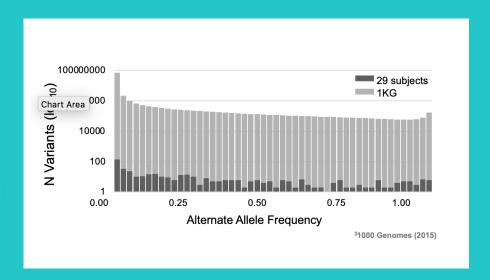




Unbiased approaches identify what is present in a sample



The Proteograph captures protein variants...



...including

Isoforms

PTMs

Alleles (at any frequency in the population)

Small open reading frame (smORFs)



Strong finish to 2021



Completed first two phases of commercial release



25+ unique abstracts



Established industryleading partnerships



Systems sold in US, Korea, China, EMEA



Doubled organization



Announcing Broad Commercial Release of the **Proteograph Product Suite**

Uniquely enabling technology for deep, unbiased proteomics

Streamlined workflow and analysis to enable large-scale studies

Simplified solution accessible to nearly any lab





Positioned to lead the proteomics revolution

Building an ecosystem around unbiased, deep proteomics

Establishing Seer as premier provider in deep proteomics

Empowering leading institutions to perform studies at scale



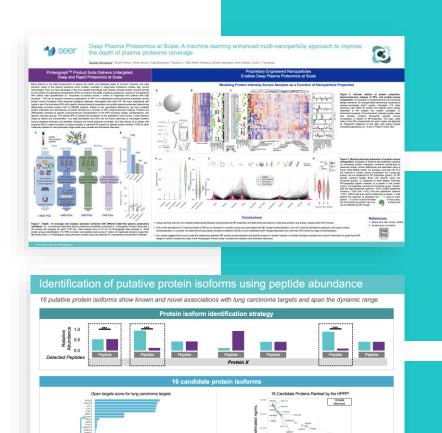
Enabling our customers to derive unique applications and insights

Making deep unbiased proteomics accessible



Technology uniquely positioned to enable deep, unbiased proteomics

Seminal publications demonstrate the power of Seer's proprietary engineered nanoparticles



Deep and precise coverage of the plasma proteome Ferdosi, et al, in press, PNAS

Optimization of protein coronas to increase proteome depth

Proteoform detection with peptide-centric analysis

Manuscript in pre-print*

Hornburg D, et al., 2022. Enhanced competitive protein exchange at the nano-bio interface enables ultra-deep coverage of the human plasma proteome. bioRxiv

Donovan MKR, et al., 2022. Peptide-centric analyses of human plasma enable increased resolution of biological insights into non-small cell lung cancer relative to protein-centric analysis, bioRxiv.







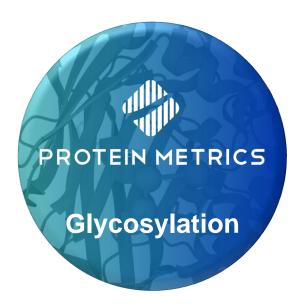


Enabling new applications & insights in customers' hands









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Posters are posted on www.seer.bio/library





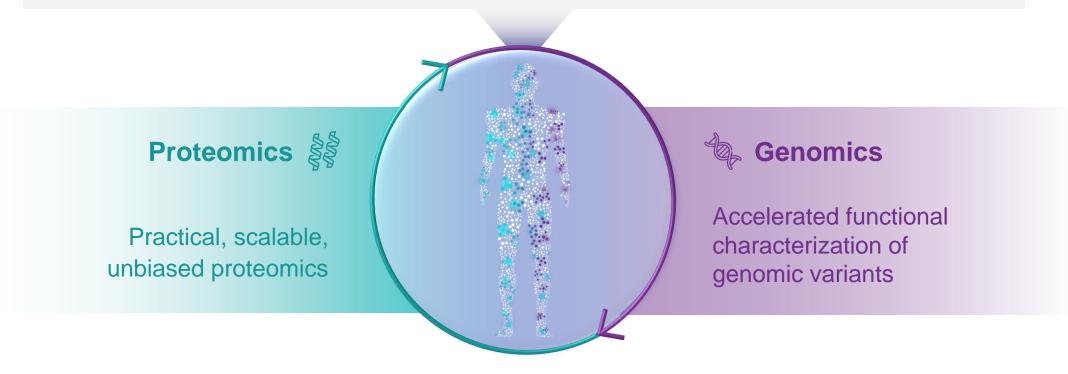




Enabling proteogenomics by impedance matching genomics

Unmet need to functionally characterize genomic variation and deepen our understanding of biology

Oncology Diagnostics Complex Disease Infectious Disease Rare Disease Therapeutics Reproductive Health



PROTEOGENOMICS

Bridge the gap between genomics & proteomics to accelerate our understanding of biology

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Empowering unprecedented studies for unbiased plasma proteomics

Prior to Seer, the largest published deep, unbiased plasma proteomics studies were in the 10s of samples







Prostate Cancer	Disease Multi-omics	Aging	
1,000 samples	2,000 samples	1,500 samples	

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Making deep, unbiased proteomics analysis more accessible

Making it easier to access deep, unbiased, rapid proteomics at scale with Proteograph Analysis Suite

Past

```
t---- Read MS data ----
dat <- readRDS("example_data.rds") %>%
 filter(is.na(mss_controls_applied)) %>%
 filter(mss_biosample_id == "PC3") %>%
 # parse per plate and sample annotations
 rowwise() %>%
 mutate(plate_row = well_location_to_plate_row(mss_well_location),
       plate_col = well_location_to_plate_col(mss_well_location),
        plate_side = ifelse(as.numeric(plate_col) <= 5, "L", "R")) %>%
  mutate(mss_biosample_id = paste(mss_plate_id, mss_biosample_id, plate_row, plate_side, sep = "-")) %>%
 # Only get samples with all 5 NPs
 nest_by(mss_biosample_id) %>%
 mutate(n_nps = n_distinct(data$nps_nanoparticles)) %>%
 ungroup() %>%
 filter(n_nps == 5) %>%
 select(-n_nps) %>%
 unnest(cols = "data") %>%
 # Count the number of panel protein groups per sample
 group_by(mss_sample_id) %>%
 summarize(n_proteins = n_distinct(protein_ids)) %>%
#==== Plot Protein Group Counts per Sample ====
nlt <- gaplot(dat. ges(mss sample id. n proteins)) +
 geom_col(fill = seer_palette[2], color = "black", size = 0.2) +
 geom_hline(yintercept = c(500, 1000, 1500, 2000), linetype = "dashed", color = "gray50") +
 scale_fill_seer(guide = "none") +
 theme_seer() +
  title = "Panel Protein Group Counts Per Sample",
   y = "Number of Panel Protein Groups"
```

80

Advanced Data Analysis



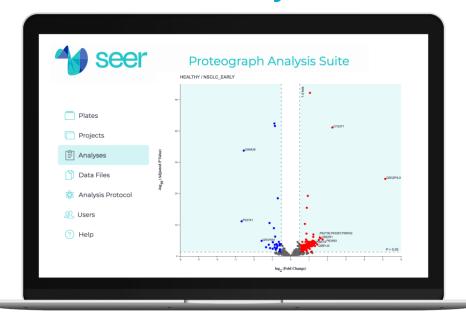


Weeks+



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Today



Recapitulates Nature Communications results



1 user



9 clicks, <1 minute to visualization





Making deep, unbiased proteomics geographically more accessible

Building access to deep, unbiased, rapid proteomics with key partners and Centers of Excellence program

Key partnerships to expand access across geographies and customer groups



Increase access in **Europe** with one of the world's premier service providers







Provide access to research and healthcare customers in **Asia Pacific**





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Making deep, unbiased proteomics more accessible to more labs

Announcing the Proteogenomics Consortium to expand the availability of proteomics to genomics customers







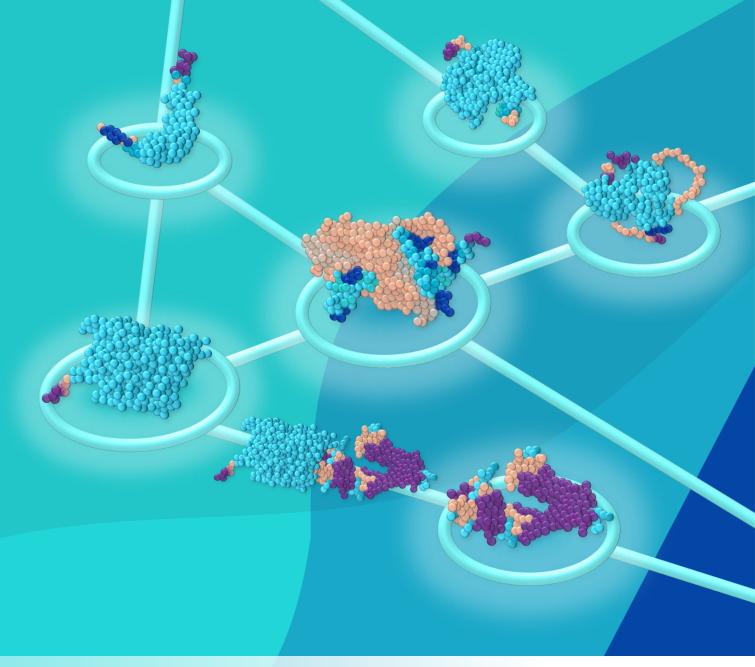


BRINGING PROTEOMICS TO GENOMICS CUSTOMERS



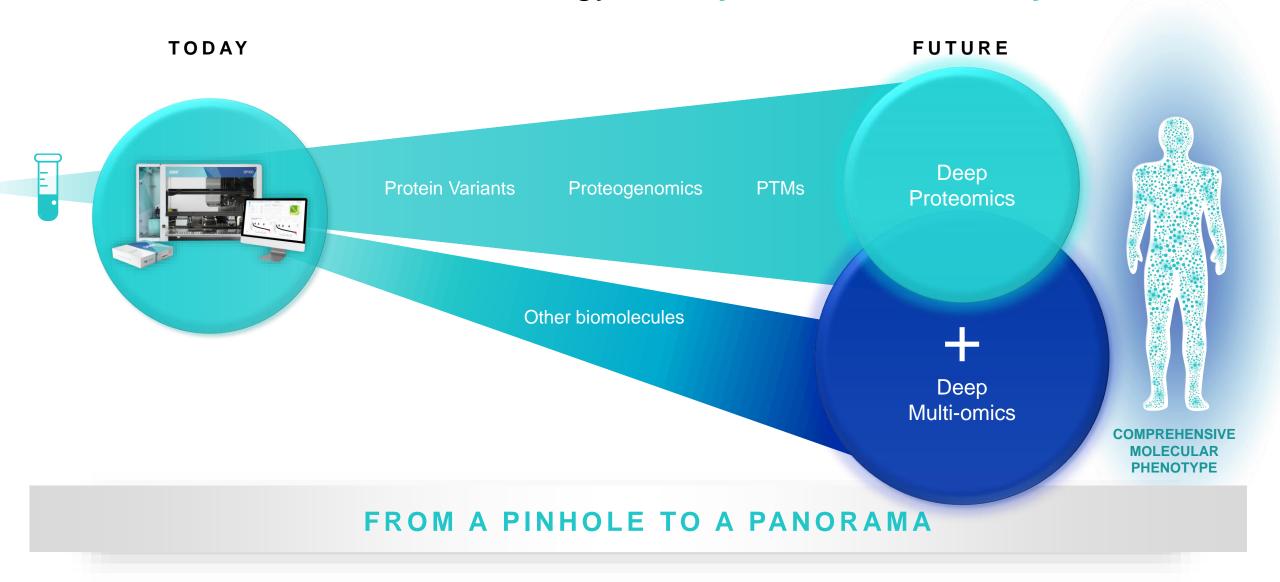


Looking ahead...



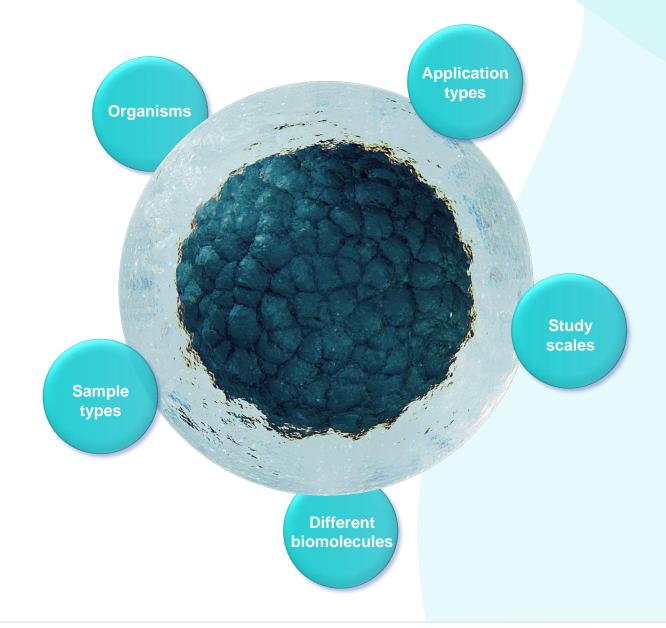


Our vision is to understand biology in a dynamic, broad way





Unconstrained interrogation of biology enabled by proprietary nanoparticles





Continued execution in 2022



Leading customers and collaborators

- ✓ Continue to grow installed base
- ✓ Facilitate the execution of large-scale studies using the Proteograph Product Suite



Best in class technology

- ✓ Further product roadmap to open up a new gateway to biology
- ✓ Pave the way for future clinical validation of our technology
- ✓ Exemplify how deep unbiased proteomics enables discovery of novel content



Creating new markets

- ✓ Partner in novel ways to standardize deep proteomics workflows and insights
- ✓ Broadly enable proteogenomics



Leading team with multi-omics expertise

- ✓ Continue to build a team of world-class talent
- Expand global infrastructure and team to support expanding customer base



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Q & A

Omid Farokhzad, M.D.

