#### UNITED STATES SECURITIES AND EXCHANGE COMMISSION

Washington, D.C. 20549

#### FORM 8-K

CURRENT REPORT Pursuant to Section 13 or 15(d) of the Securities Exchange Act of 1934

Date of Report (Date of earliest event reported): March 8, 2022

### **QUANTUM-SI INCORPORATED**

(Exact name of registrant as specified in its charter)

Delaware

(State or other jurisdiction of incorporation)

001-39486

(Commission File Number)

**85-1388175** (IRS Employer Identification No.)

530 Old Whitfield Street Guilford, Connecticut

(Address of principal executive offices)

**06437** (Zip Code)

Registrant's telephone number, including area code: (203) 458-7100

N/A

(Former name or former address, if changed since last report)

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions:

□ Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)

□ Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)

□ Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))

□ Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

Securities registered pursuant to Section 12(b) of the Act:

Title of each class	Trading Symbol(s)	Name of each exchange on which registered		
Class A common stock, par value \$0.0001 per share	QSI	The Nasdaq Stock Market LLC		
Redeemable warrants, each whole warrant exercisable for	QSIAW	The Nasdaq Stock Market LLC		
one share of Class A common stock, each at an exercise				
price of \$11.50 per share				

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (§230.405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§240.12b-2 of this chapter).

Emerging growth company  $\Box$ 

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act.

#### Item 7.01 Regulation FD Disclosure.

From time to time, Quantum-Si Incorporated (the "Company") presents and/or distributes slides and presentations to the investment community to provide updates and summaries of its business. On March 8, 2022, the Company gave a presentation at the 42<sup>nd</sup> Annual Cowen Health Care Conference. The presentation slides and a replay of the webcast is available on the "Investors" section of the Company's website at https://ir.quantum-si.com. This presentation is also furnished as Exhibit 99.1 to this Current Report on Form 8-K.

The information in this Item 7.01 is being furnished and shall not be deemed "filed" for purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the "Exchange Act"), or otherwise subject to the liabilities of that Section, nor shall it be deemed incorporated by reference into any registration statement or other filing under the Securities Act of 1933, as amended, or the Exchange Act, except as shall be expressly set forth by specific reference in such filing.

#### Item 9.01. Financial Statements and Exhibits.

(d) Exhibits.

Exhibit No.	Description
<u>99.1</u>	Corporate Presentation of Quantum-Si Incorporated dated March 8, 2022.
104	Cover Page Interactive Data File (embedded within the Inline XBRL document).

#### SIGNATURES

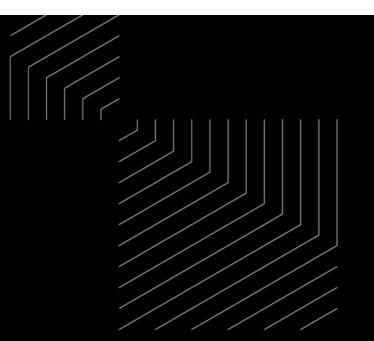
Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

#### QUANTUM-SI INCORPORATED

By: /s/ Claudia Drayton Name: Claudia Drayton Title: Chief Financial Officer

Date: March 8, 2022

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## Quantum-Si

Investor Update

March 8 2022

### Disclaimer

This presentation includes "forward-looking statements" within the meaning of the "safe harbor" provisions of the United States Private Securities Litigation Reform Act of 1995. The actual results of the Company may differ from its expectations, estimates, and projections and, consequently, you should not rely on these forward-looking statements as predictions offuture events. Words such as "expect," "estimate," "project," "budget, "forecast, "fanticipate," "intend," "plan," "may," "will," "could," "should," "believes," "predicts, "potential, "continue," and similar expressions (or the negative versions of such words or expressions) are intended to i dentify such forward-looking statements. The se forward-looking statements include, without limitation, the Company's expectations with respect to future performance and development and commercialization of products and services. These forward-looking statements involve significant risks and uncertainties that could cause the actual results to differ materially from those discussed in the forward-looking statements. Most of these factors are outside the Company's control and are difficult to predict. Factors that may cause such differences indude, but are not limited to: the impact of COVID-19 on the Company's business the inability to maintain the listing of the Company's Class A common stock on The Nasdaq Stock Market; the ability to recognize the anticipated benefits of the recently completed business combination, which may be affected by, among other things, competition and the ability of the Company to grow and manage growth profitably and retain its key employees; our ongoing leadership transition; changes in applicable laws or regulations; the ability of the Company to raise financing in the future, the success, cost and timing of the Company's product development and commercialization activities, the potential attributes and benefits of the Company's products and services; the Company's ability to obtain and maintain regulatory approval for its products, and any related restrictions and limitations of any approved product; the Company's ability to identify, in-license or acquire additional technology; the Company's ability to maintain its existing lease, license, manufacture and supply agreements; the Company's ability to compete with other companies currently marketing or engaged in the development or commercialization of products and services that the Company is developing the size and growth potential of the markets for the Company's future products and services, and its ability to serve those markets, either alone or in partnership with others; the pricing of the Company's products and servi cesfollowing anticipated commercial launch; the Company's estimates regarding future expenses, future revenue, capital requirements and needs for additional financing; the Company's financial performance; and other risks and uncertainties discussed in the "Risk Factors" section of the Company's periodic reports filed with the U.S. Securities and Exchange Commission (SEC), and risks described in other filings the Company may make with the SEC in the future. The Company cautions that the foregoing list of factors is not exclusive. The Company cautions readers not to place undue reliance upon any forward-looking statements, which speak only as of the date made. The Company does not undertake or accept any obligation or undertaking to release publicly any updates or revisions to any forward-looking statements to reflect any change in its expectations or any change in events, conditions, or circumstances on which any such statement is based.

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# "I start each company to improve the life of somebody I love."

Dr. Jonathan M. Rothberg Founder & Chairman Interim CEO Quantum-Si

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## We build ecosystems to

Digitize Medicine Apply Deep Learning Enabled AI Democratize Healthcare



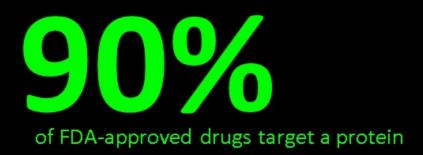
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454 SEQUENCING RainDance interent States Hyperfine M Liminal Sciences TESSERACT A ITherapeutics Detect PROTEIN

## We harness the power of semiconductor technology for

## Simplicity Speed Scale

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Most diseases are linked to dysfunctional proteins, as humans share 99.9% of DNA sequence. 85% of the human proteome is currently undrugged<sup>2</sup>, potential for game changing drug development.

Protein modifications are real-time indicators of health and disease, making them ideal markers for disease, drug response and health.

Source(s): 1. The Hurman Proteome Tissue Atlas - Druggs ble Proteome, 2015, The Hurman Protein Atlas Project 2. A Quest to Drug the Undruggs ble, June 2015, Chemical & Engineering News

# 150,000

research papers found

thousands of protein biomarkers

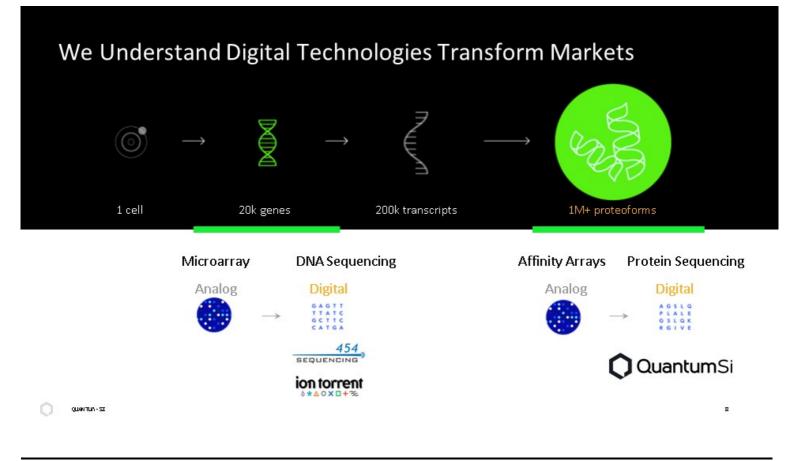
yet less than

are routinely used in clinic<sup>1</sup>

w.nature.com/articles/469156a

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Current tools limit the use of protein biomarkers. Routine tests for Serum & Cerebrospinal Fluid (CSF) are constrained by number of analytes they can look at, sensitivity, and specificity.



## Analog vs Digital Approaches to Proteomics

#### **Analog Affinity-based Approaches**

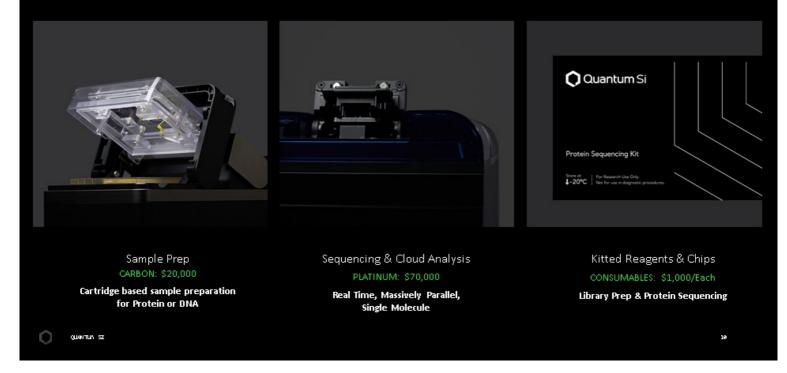
#### Identify known proteins

#### **Digital Sequencing-based Approaches**

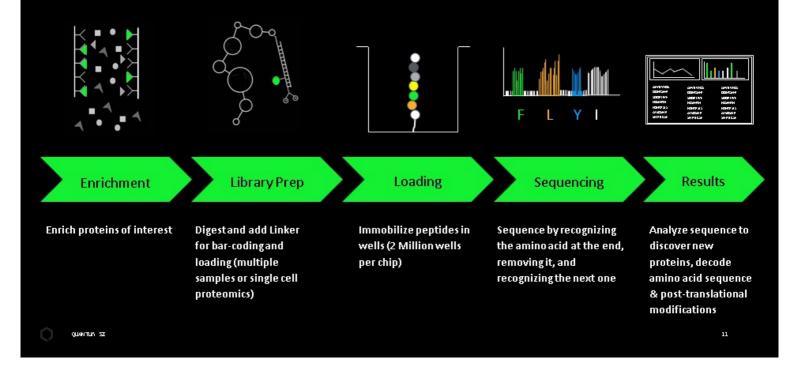
#### Decode Novel Sequences

	SomaLogic	Olink	Nautilus	Quantum-Si	Encodia	Erisyon
Detection Method	Aptamers	Antibodies	Aptamers / Antibodies	Direct Real-Time	N-terminal binding+ Edman degradation with NGS readout	Side chain labeling+ Edman degradation with s canning
Instrument Costs	\$\$ - \$\$\$ (NGS Optional)	S - SSS (NGS Optional)	\$\$\$		55 - 555 (NGS Required)	55
Run Costs	55 - 555 (NGS Optional)	5 - 555 (NGS Optional)	55 - 555		55 - 555 (NGS Required)	55
AA Sequencing	NO	NO	NO	YES	LIMITED	LIMITED
Read Length Scaling	N/A	N/A	N/A	HIGH	LOW	LOW
PTM Detection	? (Affinity Reagent)	<b>?</b> (Affinity Reagent)	<b>?</b> (Affinity Reagent)	SCALABLE		
Notes	Notes *Can't differentiate between proteoforms unless they create a specific affinity reagent			Kinetics for amino acids & PTMs	NGS erases quantitative information	Harsh acidic environment limits utility
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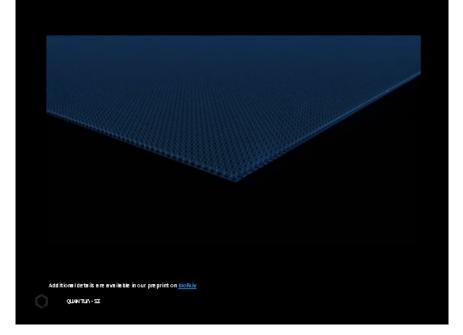
## Q-Si End-to-End Proteomics Solution



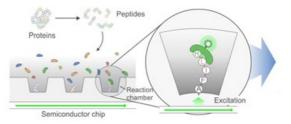
## Workflow for Q-Si Next-Gen Protein Sequencing™



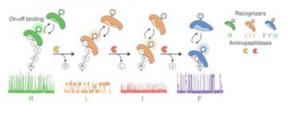
## World's First Massively Parallel Next-Gen Protein Sequencing<sup>™</sup>



#### Library Preparation and Loading

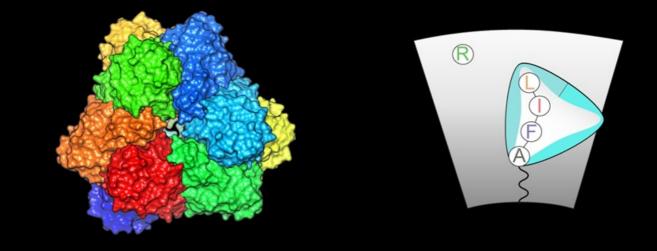


#### **Protein Sequencing**



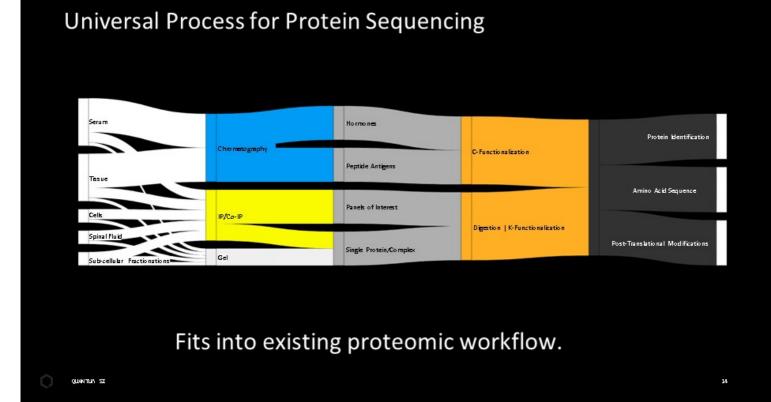
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## The Magic Enabling Broad Coverage of the Proteome

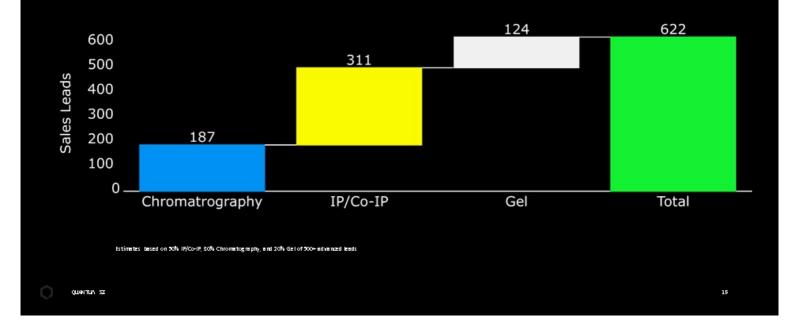


While each protein is unique, the enzyme machines we engineer (aminopeptidases) have evolved to make all peptides behave the same in our system!

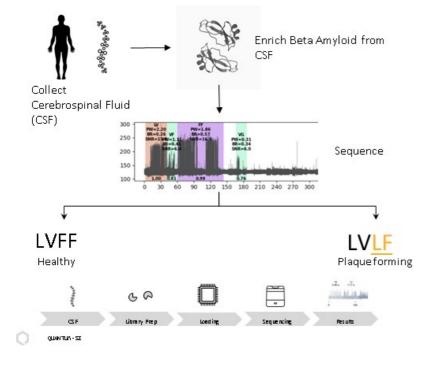
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## 500+ Advanced Leads by Workflow



## Alzheimer's Risk Assessment



## How can you test for early onset Disease?

#### Biological & Clinical Challenge:

- Less than 1% of Alzheimer's caused by an inherited single gene.
- Somatic mutations mutations accumulated over a lifetime.

#### Technical Challenge:

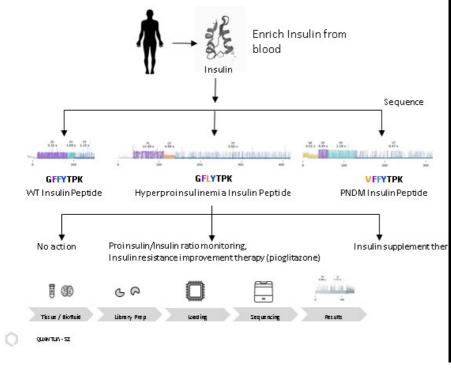
- Source of mutation not known.
- Mass spec is expensive, inconsistent, and often not sensitive enough.

#### Solution:

 Sequencing of the peptides to identify changes in amino acid sequence.

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### Metabolic Disease



## How can you identify modification of critical peptide hormones?

#### **Biological & Clinical Challenge:**

 Heterogeneous populations of variants.

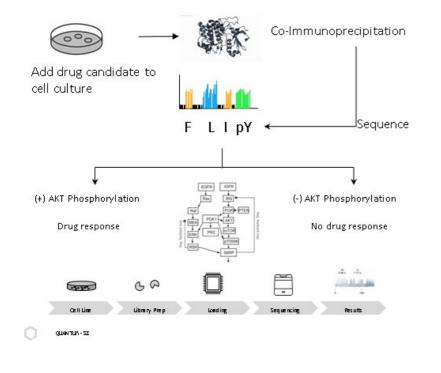
#### Technical Challenge:

- Accuracy of detection for small mass differences is not consistent by mass spec.
- Sensitivity challenging for less abundant modifications.

#### Solution:

 Immunoprecipitation and Sequencing of the peptides to identify point mutations.

## **Drug Development**



## How do you identify proteins that interact with target proteins of interest?

#### **Biological & Clinical Challenge:**

- New proteins in my pathway?
- How does the complex change in disease?

#### Technical Challenge:

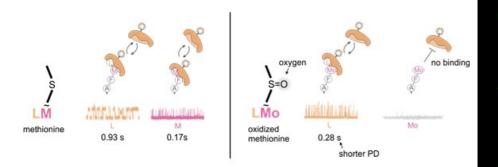
 Routine, robust, scalable, sensitive tools to discover new proteins and post-translational modifications.

#### Solution:

- Peptide sequencing to discover new proteins.
- Comparisons between samples to identify new post-translational modifications.

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### Single Molecule, Single Atom Detection



No a priori knowledge needed to detect new biological markers.

The oxidation of the penultimate residue is detected by a reduction in the average pulse duration of the N-terminal recognizer (as well as by the blocking of recognition of methionine when it becomes the N-terminal amino acid, as sequencing proceeds).

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#### 1,000,000+ Protein Variations!

#### **Biological & Clinical Challenge:**

- A protein's modifications determine its function.
- What biomarkers can we discover?

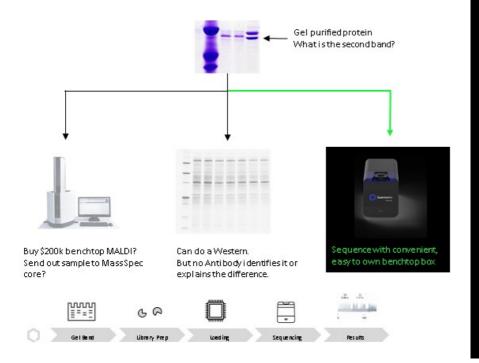
#### **Technical Challenge:**

 Impossible technical challenge to generate affinity reagents to ALL PTMs in context (over 1 million).

#### Solution:

- Q-Si detects modifications without a priori knowledge.
- Powerful new method for comparing disease & treatment states to find biomarkers.
- Kinetics enable the detection of posttranslational modifications; oxidation, phosphorylation, glycosylation (in the penultimate amino acids).

## Proteomics Core on Your Benchtop



## Sequence Proteins Like we Sequence DNA

#### **Biological & Clinical Challenge:**

- What is this protein?
- How is it modified?

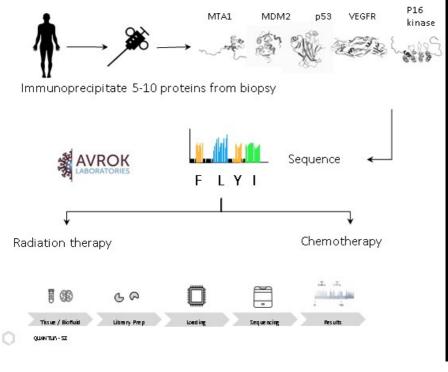
#### Technical Challenge:

- Weeks waiting for answers from a mass spec core facility.
- Antibodies don't provide new insights.

#### Solution:

- Discover new proteins of interest.
- Identify new protein variations and post-translational modification.

## **Future of Therapy Selection**



#### **Profiling Cancers to Guide Therapy**

#### **Clinical Challenge:**

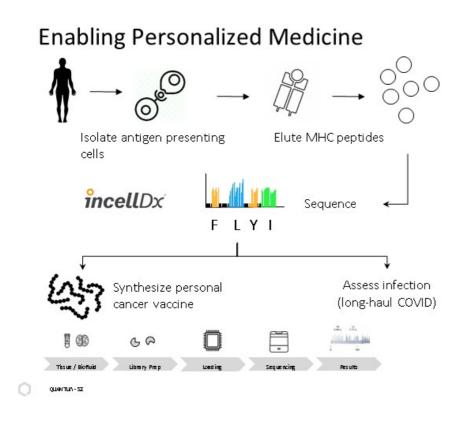
 Survival depends on early correct therapy selection and modifications of treatment regimens.

#### Technical Challenge:

- Genetic tests detect chromosomal aberrations, not protein alterations.
- Disease instability requires frequent testing & new understanding,

#### Solution:

 Q-Si Sequencing enables biomarkers select therapy.



#### Identify Antigens for Personalized Immunotherapy or to Understand Infection

#### **Clinical Challenge:**

- Highly diverse peptides of unknown origin.
- Need to identify modifications.

#### Technical Challenge:

- Peptides missed by mass spec
- Relevant neoantigens or antigenic pathogens relatively low abundant.

#### Solution:

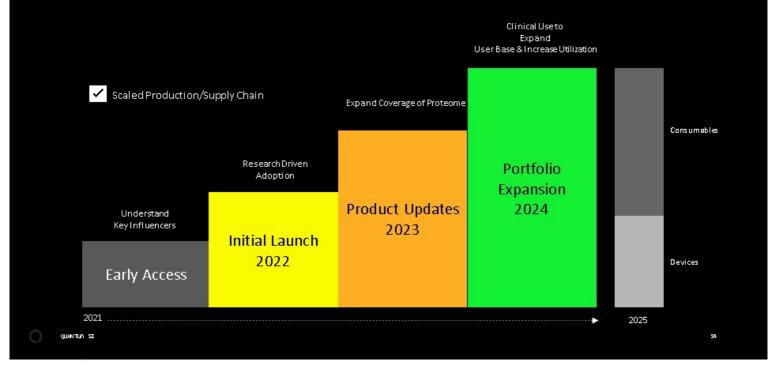
- Q-Si sequencing to identify antigen targets.
- Understanding of new Covid Variants.

## Goals for Commercial Launch

~70% loading of proteome after library prep into 5 to 25 amino acid long peptides 200,000+ reads per run, with 10 to 20 reads for each high confidence call 5 to 50+ proteins over 3 or 4 logs concentration range

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## Roadmap for Customer Adoption & Growth



# ~\$470 M

to fund work through 2024

## Experienced

team of



With team members that invented and commercialized the first Next Generation DNA Sequencing and put DNA sequencing on a Semiconductor chip, we are well positioned to launch the World's First Next-Generation Protein Sequencing

## **Catalyst for Success**

Oncology drove Next Generation DNA Sequencing

Immunology, Immuno-oncology & Infectious Disease will drive Next-Gen Protein Sequencing™

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The vision to bring Moore's Law to DNA Sequencing

The team to bring Next-Gen Protein Sequencing™ to the world

**Quantum**Si

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