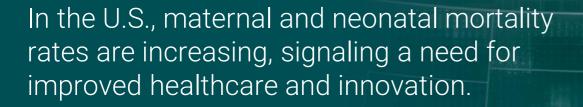


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This presentation contains forward-looking statements that involve substantial risks and uncertainties. All statements, other than statements of historical facts, contained in this presentation, including statements regarding our strategy, future operations, future financial position, future revenue, projected costs, prospects, plans and objectives of management, are forward-looking statements. The words "anticipate," "believe," "estimate," "expect," "intend," "may," "plan," "predict," "project," "target," "potential," "will," "would," "could," "should," "continue" and similar expressions are intended to identify forward-looking statements, although not all forward-looking statements contain these identifying words. We may not actually achieve the plans, intentions or expectations disclosed in our forward-looking statements, and you should not place undue reliance on our forward-looking statements. Actual results or events could differ materially from the plans, intentions and expectations disclosed in the forward-looking statements we make.

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One-third of pregnant women experience complications, often without warning, posing significant

health risks for both mother and child.



Vision

Sera Prognostics aims to be a global leader in highvalue women's health diagnostics, delivering pivotal information to physicians and mothers to improve the health of pregnant women and newborns, and to simultaneously improve the economics of pregnancy care.

Mission

Pregnancy complications occur frequently, often without warning. In the U.S., this is the reality for one-third of pregnancies. Sera Prognostics is transforming the pregnancy journey with innovative products aimed at identifying complications early and accurately. We provide personalized insights into the risk of preterm birth and other pregnancy complications, empowering all pregnant women and their caregivers with innovative products aimed at delivering better outcomes for mothers and babies.





The Pregnancy Company®

PREGNANCY OUTCOMES

Provides information to healthcare providers to guide personalized interventions

SPECIALTY LAB

Focuses on obstetric test offerings and analytics products

EMPOWERMENT

Aims to empower physicians and expectant mothers with actionable insights

RESEARCH-DRIVEN

Committed to driving advancements in maternal-fetal medicine through rigorous research and clinical trials

INNOVATIVE TESTS & PRODUCTS

Utilizes cutting-edge technologies such as proteomics and Al

GLOBAL IMPACT

Works towards shaping the future of pregnancy care worldwide



SOCIETAL IMPACT

Focusing on neonatal and maternal outcomes can yield numerous benefits for all members of society

Families: Decrease the burden linked to pregnancy complications

Payers: Improve outcomes and the economics of care delivery

Communities: Address growing health disparities in maternal care

Healthcare Providers: Provide tools to address unmet needs in pregnancy care

Self-insured Employers: Enhance employer healthcare for employees and their families

Policy: Support government programs targeting improvements in maternal healthcare



Improved pregnancy outcomes start with better information



Utilize proteomic testing early in the pregnancy to identify high-risk pregnancies

Target individuals identified as high-risk for tailored preventive treatments

Achieve healthier outcomes by reducing morbidity, mortality, and hospitalization time for newborns



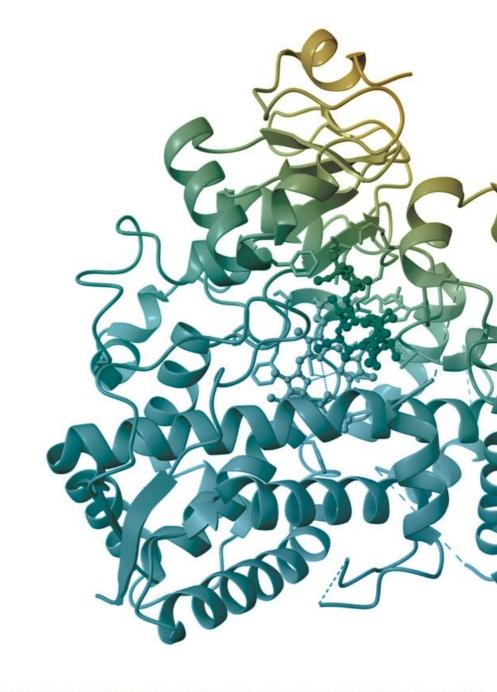
Answers provided by **proteomics**.

Given the limitations of current screening practices for pregnancy complications, scientists at Sera discovered a biomarker prediction that fills a critical diagnostic gap.

Why Proteomics?

Personalized insights for each pregnancy. Proteomics provide insights into the biology of each pregnancy and go beyond genetic inheritance.

Early detection and intervention. Proteins reflect the functions of specific cells allowing for early detection and intervention before symptoms arise.





Sera's proprietary data assets and technology platform generate powerful insights

Biobank of Blood Samples

20,000 pregnant women

Clinical Outcomes Data

Multiple clinical utility studies

Demographic & Geographic Diversity

Broad sample collection across U.S. population

Global Dataset

North America, Europe, Asia and Africa

Advanced Mass Spectrometry

300+ proteins analyzed

Artificial Intelligence & Machine Learning

Discovering important predictors



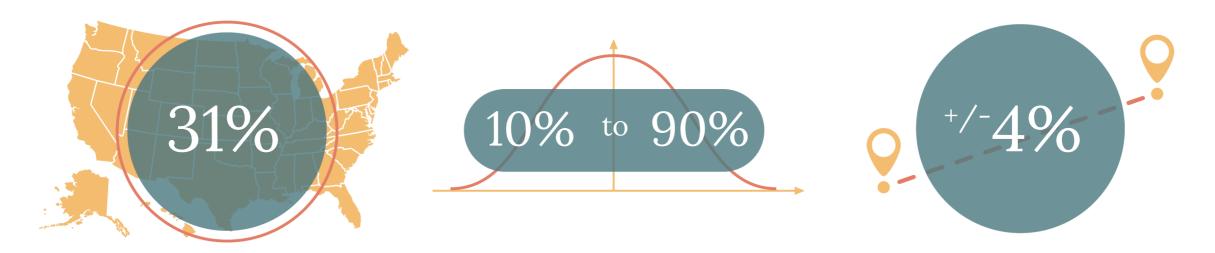


POWERFUL INSIGHTS:

- **Early risk prediction** such as preterm birth or preeclampsia, before symptoms manifest.
- Improving health outcomes by identifying high-risk individuals and implementing targeted interventions
- Shared decision-making by empowering patients to make informed decisions regarding prenatal care
- Reduced healthcare costs through improvements in the health of the baby and reducing the need for the NICU
- Research advancements to help understanding the biology of pregnancy



Sera's technology platform transforms national averages into personalized insights



The national risk of a complicated pregnancy is 31%.

The probability of pregnancy complications range from 10 - 90%.

Utilizing proteomic markers and data analytics, Sera can identify a mother's pregnancy journey inclusive of her complication risk within a 4% margin.







NEAR TERM EXECUTION

Accelerate PreTRM Adoption & Revenue

Addressing the Problem of Preterm Birth

1 in 1 babies are born too soon¹

Preterm birth contributes to 34.2% of newborn deaths²

Causes numerous medical issues requiring more hospital time and pediatric visits^{3,4}



References: 1. Osterman MJK, Hamilton BE, Martin, J.A., Driscoll AKV, C.P. Births: Final Data for 2020. Division of Vital Statistics, National Center for Health Statistics, U.S. Centers for Disease Control and Prevention. 2022 Feb 7 (https://www.cdc.gov/nchs/nvss/births.html. 2. Callaghan WM, et al. The contribution of preterm birth to infant mortality rates in the United States. Pediatrics. 2006 Oct;118(4):1566-73. 3. Howson CP, et al. Born Too Soon: Preterm birth matters. Reprod Health 10, S1 (2013). 4. Crump C, et al. Prevalence of Survival Without Major Comorbidities Among Adults Born Prematurely. JAMA. 2019 Oct 22;322(16):1580-1588.



Reducing the health impacts of an early delivery with the PreTRM® Test

Challenges facing risk assessment for preterm birth

Up to half of all pregnant women who deliver prematurely have no known risk factors^{1,2}

Knowing the Risk Enables Informed Action

PreTRM's sensitivity of 88% allows healthcare providers to detect higher-risk patients who would have been missed with current screening tools³

Reduce the impacts of preterm birth

Recent studies demonstrated an 18% reduction in severe neonatal morbidity and mortality when higher-risk pregnancies identified by the PreTRM Test received risk-reduction interventions compared to standard care⁴



A strong foundation of evidence

CLINICAL STUDIES

ANALYTICAL VALIDATION

CLINICAL VALIDATION CLINICAL

ECONOMIC UTILITY

CLINICAL MASS SPECTROMETRY / Bradford et al., 2017

 Achieved robust analytical validation of protein biomarkers for risk of spontaneous preterm birth



PAPR / Saade et al., 2016

AJOG editor's choice establishing PreTRM® analytical validation



ACCORDANT THRESHOLD / Burchard et al., 2021

 Validated PreTRM threshold for clinical decision-making for risk of spontaneous preterm birth (sPTB) and adverse maternal and neonatal outcomes in cohorts with diverse racial and ethnic backgrounds



AJP REPORTS / Caughey et al., 2016

 Empirical data demonstrating the clinical and cost impact of prognostic test for early detection of preterm birth







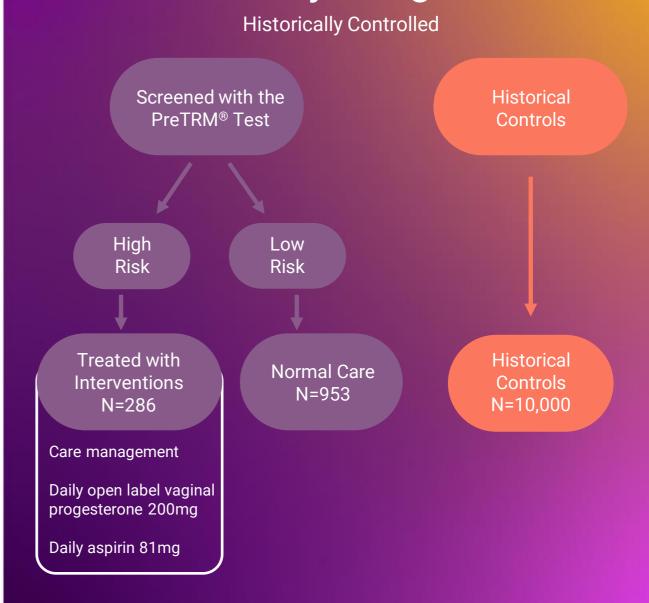
Building on the foundation with new clinical utility evidence

CLINICAL STUDIES	ANALYTICAL VALIDATION	CLINICAL VALIDATION	CLINICAL UTILITY	ECONOMIC UTILITY
COST-EFFECTIVENESS OF A PROTEOMIC TEST FOR PRETERM BIRTH PREDICTION / Grabner et al., 2021				
Showed both improved neonatal outcomes and reduced immediate and long-term treatment costs associated with premature birth, when compared to routine care				
PREVENT / Branch et al., 2021 Study demonstrates positive impact of the company's PreTRM® test-and-treat strategy on improving neonatal healthcare				
ACCORDANT CU (TREETOP) / Burchard et al., 2022 Combines real-world observational data with simulation to project significant potential improvements in neonatal outcomes among racially and ethnically diverse populations				
PREDICTION AND PREVENTION OF PTB / Combs et al., 2023 Study concludes that screening with the PreTRM® Test followed by care management intervention and LDA prolonged pregnancy and reduced adverse outcomes				
AVERT / Hoffman et al., 2023 PreTRM® test-and-treat strategy demonstrates statistically and clinically significant improvement in neonatal health outcomes and hospital length-of-stay				
PRIME / Iriye et al., ongoing Further investigates the value of implementing the PreTRM® test-and-treat strategy to reduce both adverse singleton pregnancy outcomes and overall healthcare costs				
			6	SERA PROGNOSTI

The AVERT PRETERM Trial studied the health impacts of the PreTRM® test-and-treat Strategy

Neonatal outcomes after proteomic biomarker guided intervention: the AVERT PRETERM TRIAL

- The study took place from June 2018 -September 2020 at ChristianaCare Hospital (Newark, DE)
- The proportion of Black women in both arms was 26.5%, reflecting the racial diversity in the study site's patient population

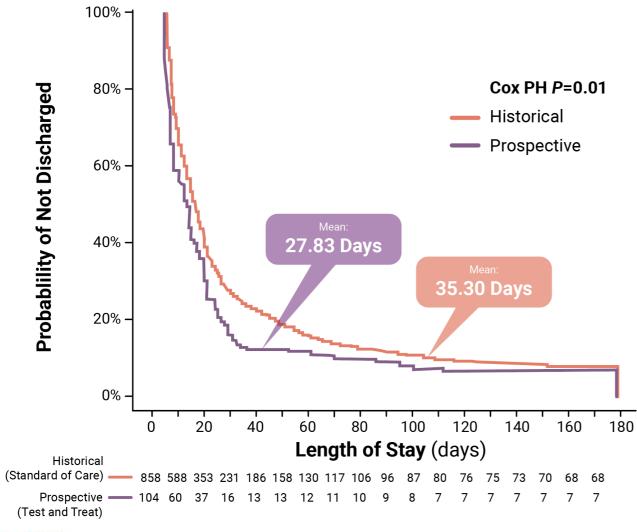


Study Design



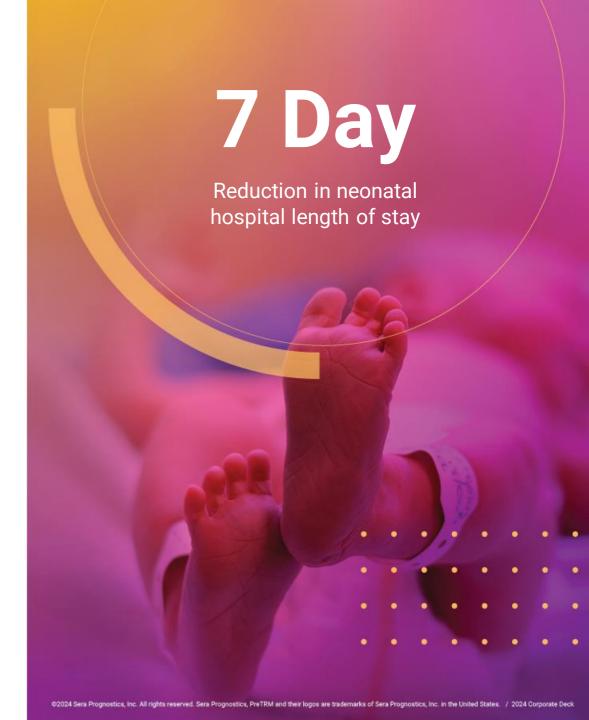
AVERT Study Results

Reduce the time the baby spends in the hospital on average by one week

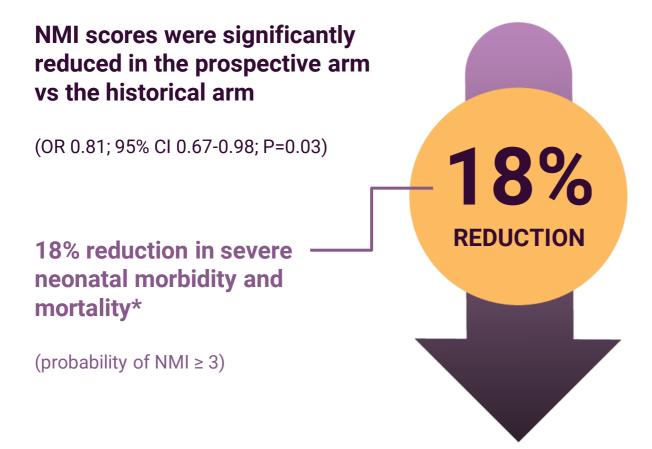




Reference: Matthew K. Hoffman, Carrie Kitto, Zugui Zhang, et al. Neonatal outcomes after proteomic biomarker-guided intervention: the AVERT PRETERM TRIAL. medRxiv 2023.09.13.23295503; doi: https://doi.org/10.1101/2023.09.13.23295503



Severe morbidity and mortality rates were significantly reduced





Reference: Matthew K. Hoffman, Carrie Kitto, Zugui Zhang, et al. Neonatal outcomes after proteomic biomarker-guided intervention: the AVERT PRETERM TRIAL. medRxiv 2023.09.13.23295503; doi: https://doi.org/10.1101/2023.09.13.23295503

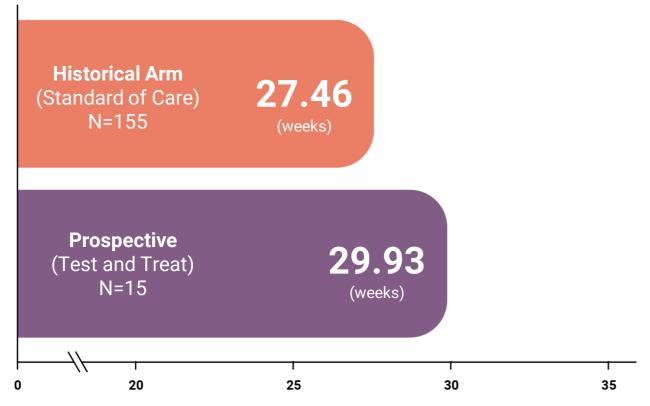
*Severe neonatal morbidity and mortality are defined as Neonatal Composite Morbidity & Mortality Index (NMI) ≥3

- 1-4 score given (4 = infant mortality)
- The score increases by 1 point for each additional diagnosis of:
 - Respiratory distress syndrome
 - Bronchopulmonary dysplasia
 - Intraventricular hemorrhage grade III or IV
 - All stages of necrotizing enterocolitis
 - Periventricular leukomalacia
 - Proven severe sepsis
- The scale uses NICU stays to determine index scores if the length of stay gives a higher score than concomitant diagnoses: 1-4 days give a score of 1, 5-20 days a score of 2 and >20 days a score of 3.1

Increase gestational age at birth to improve outcomes for those at risk of earliest delivery

Gestational Age at Birth

<32 Weeks



Mean Gestational Age at Birth (weeks)



Increased Gestational Age at Birth on Average by 2.48 Weeks ©2024 Sera Prognostics, Inc. All rights reserved. Sera Prognostics, PreTRM and their logos are trademarks of Sera Prognostics, Inc. in the United States. / 2024 Corporate Deck

Reference: Matthew K. Hoffman, Carrie Kitto, Zugui Zhang, et al. Neonatal outcomes after proteomic biomarker-guided intervention: the AVERT PRETERM TRIAL. medRxiv 2023.09.13.23295503; doi: https://doi.org/10.1101/2023.09.13.23295503

AVERT Study Results

Reduce the time the neonate spends in the hospital on average by one month for those at risk of earliest delivery

Neonatal length of hospital stay (days) for babies born <32 weeks' gestation 100% Cox PH P =0.046 Remaining in the Hospital Prospective **Proportion of Neonates** 60% 97.23 Days 40% 100 120 140 **Neonatal Hospital Length of Stay (days)** Historical (Standard of Care) — 155 155 147 131 121 114 98 Prospective (Test and Treat) — 15 15 15 11 8 8



Reference: Matthew K. Hoffman, Carrie Kitto, Zugui Zhang, et al. Neonatal outcomes after proteomic biomarker-guided intervention: the AVERT PRETERM TRIAL. medRxiv 2023.09.13.23295503; doi: https://doi.org/10.1101/2023.09.13.23295503

28 Day

Reduction in neonatal length of hospital stay for patients born before 32 weeks' gestation

Advancing PreTRM®'s Evidence Development for Enhanced Pregnancy Outcomes

<u>Prematurity Risk Assessment Combined With Clinical Interventions for Improving Neonatal outcoMEs (PRIME)</u>

A pioneering clinical trial assessing the efficacy of the PreTRM Test and preventive interventions in lowering the occurrence of adverse pregnancy outcomes.



Achievement of a primary endpoint resulted in the early termination of the study.



Published Elevance Cost-Effectiveness Model

Elevance Claims Data

Analysis of >40,000 mothers and babies within commercially insured Elevance membership

- Evaluated screening with PreTRM along with proactive interventions given to PreTRM-higher risk patients vs. standard care
- Demonstrated robust clinical and economic impacts of the PreTRM test-and-treat strategy

Results

- 20% reduction in preterm birth rates
- \$1,608 gross savings, excluding a \$745 PreTRM list price per pregnant woman (amortized over all pregnancies including non-tested)
- 10% reduction in neonatal intensive care admissions
- 7% reduction in overall hospital length-of-stay
- 33% reduction in births <32 weeks
- The test-and-treat strategy is dominant with respect to cost savings across all conservative probabilistic sensitivity analyses and scenarios







Pregnancy Product Road Map





More than just the PreTRM® Test

2023 2024 2025 2026 2027

PRETRM TEST

PREDICTIVE ANALYTICS PRODUCT

TIME-TO-BIRTH TEST

PREGNANCY RISK PREDICTION PANEL









Developing a portfolio to provide better answers to life changing questions

Am I at risk for a premature birth?

PreTRM® Test

To provide a woman and her physician a risk assessment for spontaneous preterm birth

What do pregnancies like mine look like?

Predictive Analytics Product

To show a woman the most common outcomes in pregnancies of other women with similar clinical and demographic characteristics

When will my baby be born?

Time-to-Birth Test

To provide a more accurate estimate of when a mother's baby will be delivered for the purposes of planning maternity leave, travel, visitors, etc. This is a lifestyle and wellness test rather than a clinical test.

Am I at risk for the common pregnancy complications?

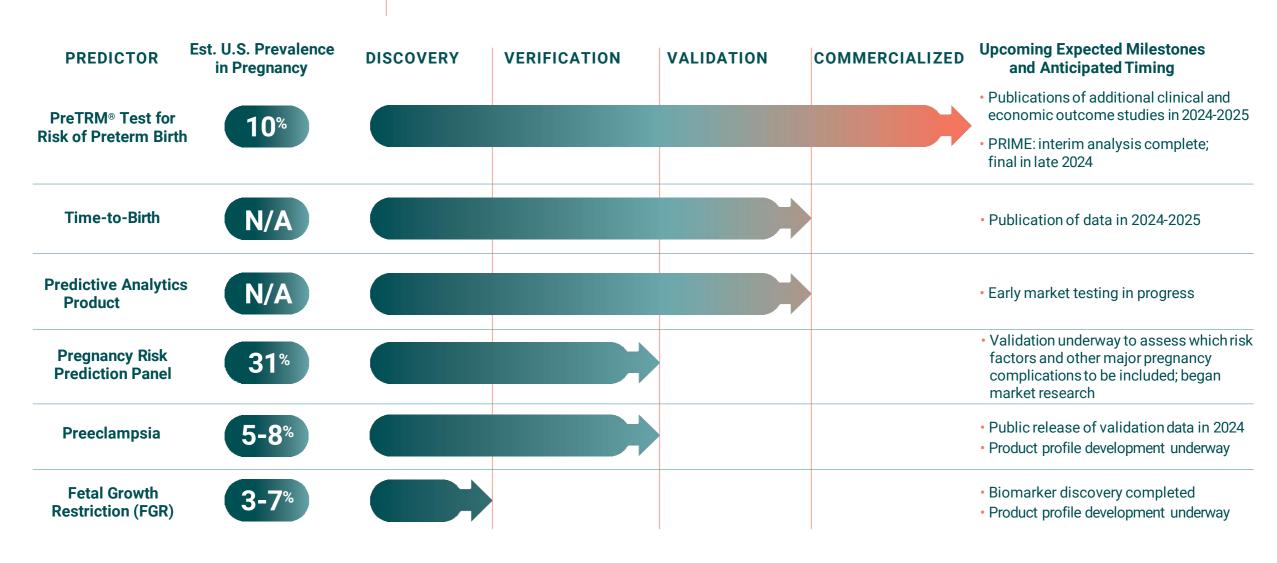
Pregnancy Risk Prediction Panel Test

To provide a woman and her physician a risk assessment for multiple pregnancy complications assessing both High Risk (rule-in) and Low Risk (rule-out) results in a single test





Robust Biomarker Pregnancy Pipeline





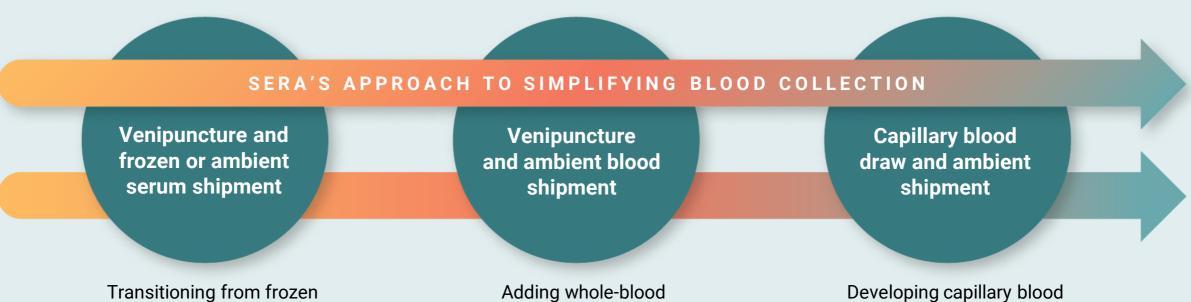
Technology Road Map





Transitioning analyses for cost-effectiveness, faster turnaround times, and enhanced market accessibility

- Diagnostic tests rely on biological samples, but there are current limitations on collection and transport
- Sera is shifting to ambient conditions for sample transport to enhance the patient and provider experience and improve access to testing



Transitioning from frozen shipments to ambient conditions for sample transport

Completed

Adding whole-blood sample types to test menus, alongside serum

Ongoing

Developing capillary blood draw capabilities without the need for venipuncture

Ongoing



Transitioning analyses for cost-effectiveness, faster turnaround times, and enhanced market accessibility

- Sera utilizes proteomic methods for discovering, verifying, and validating pregnancy tests.
- The PreTRM® test, a prime example, is commercially launched using mass spectrometry, enabling swift market access and evidence accumulation.
- Immunoassays may offer cost-effectiveness, rapid turnaround times, and enhanced market accessibility via point-of-care testing or kit distribution.
- Antibodies are developed for PreTRM analytes to facilitate immunoassay creation.
- Immunoassay methods at Sera include affinity-capture mass spectrometry (AC-MS) and enzyme-linked immunosorbent assays (e.g., ELISA, ECLIA), with the latter typically requiring two antibodies per analyte.
- AC-MS serves as a transitional tool to ELISA/ECLIA platforms.
- In some instances, Sera directly transitions from proteomics discovery to commercialization via an ELISA format.





Sera Leadership Team



Zhenya Lindgardt President & CEO

- Former CEO, The Commons Project Foundation
- Former VP of Platform and Customer Engagement, Uber Technologies
- Former Senior Partner and Managing Director, The Boston Consulting Group
- MBA, Harvard University



Austin Aerts

Chief Financial Officer

- Former VP, Finance and Corporate Controller, Sera
- · Former finance team member, Myriad
- Former auditor, Ernst & Young LLP
- · Master of Accounting University of Utah; CPA



Jay Boniface, Ph.D.

Chief Scientific Officer



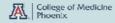


Myriad genetics



Michael Foley, M.D.

Medical Advisor







Paul Kearney, Ph.D,

Chief Data Officer









Doug RoachVP of Commercial



SEARLE



Benjamin Jackson

General Counsel





Robert G. Harrison
Chief Information Officer

Nyriad genetics



