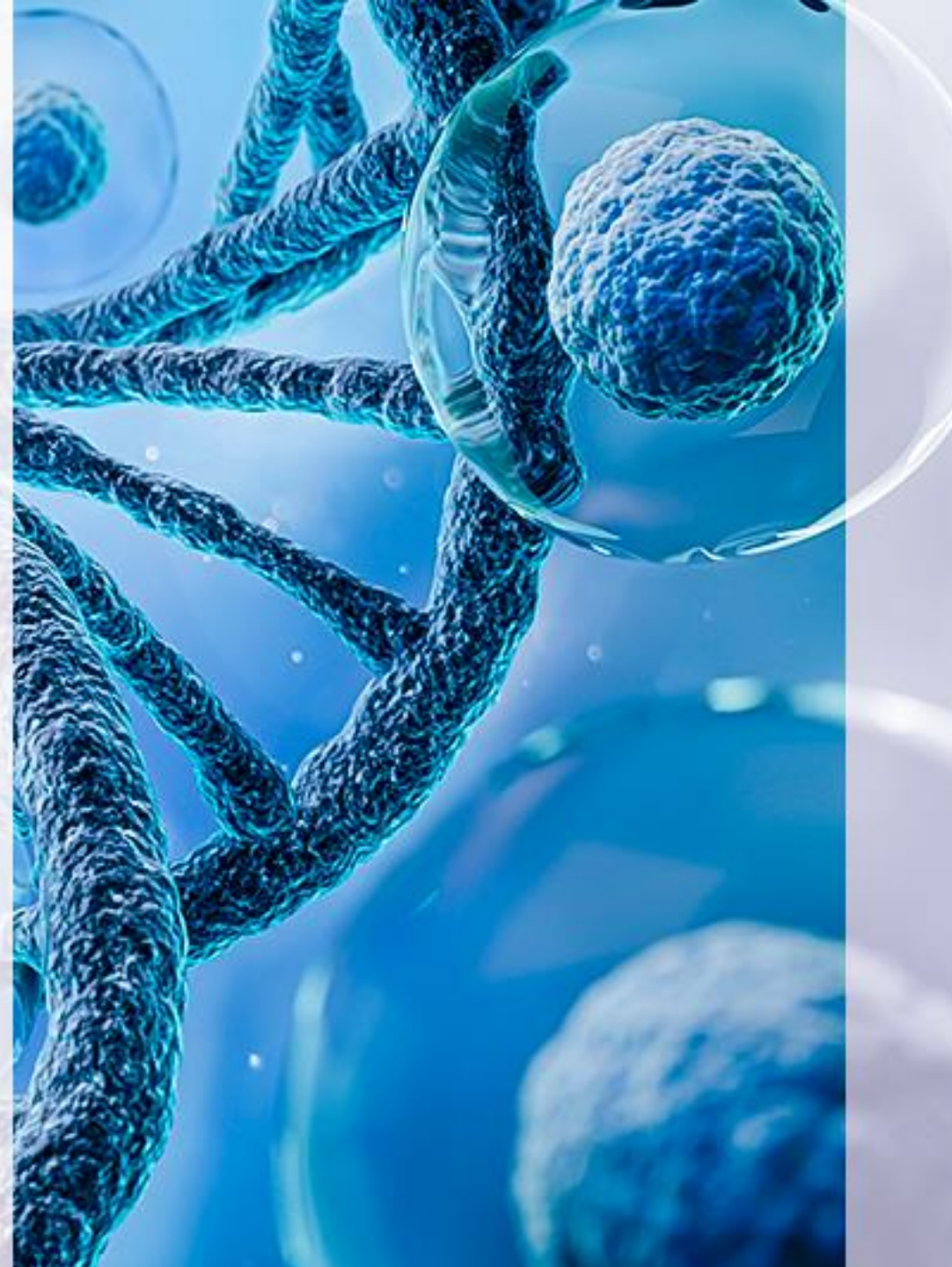


**BAYLOR**  
GENETICS

# INTRODUCTION TO BAYLOR GENETICS





## EMPOWERING YOU WITH ANSWERS THAT MATTER

### Mission

We empower patients, healthcare providers, and partners with trusted insights, translating scientific innovations into accessible clinical solutions.

### Vision

As a pioneer in precision medicine, we unlock the power of genetics to transform healthcare and improve lives.

# // PROFILE AND OVERVIEW

## BAYLOR GENETICS

### BUSINESS

Genetic Testing Services

### JOINT VENTURE

H.U. Group Holdings  
Baylor College of Medicine

### PRODUCTS & SERVICES

Diagnostic Genomics  
Partner Lab Services

### HEADQUARTERS

Texas Medical Center - Houston

### CERTIFIED LAB



45+ years  
of Innovation

2024 Fast Company  
Innovators Award



400+  
Employees



3000+  
Customizable  
Test Menu



350+  
Scientific  
Publications



Research  
Commercialization

Baylor College of Medicine  
#1 NIH funded in genetics



4 Million+  
Clinical Tests

Performed to date



80+  
Partners

Life Sciences, Pharma  
Precision Medicine

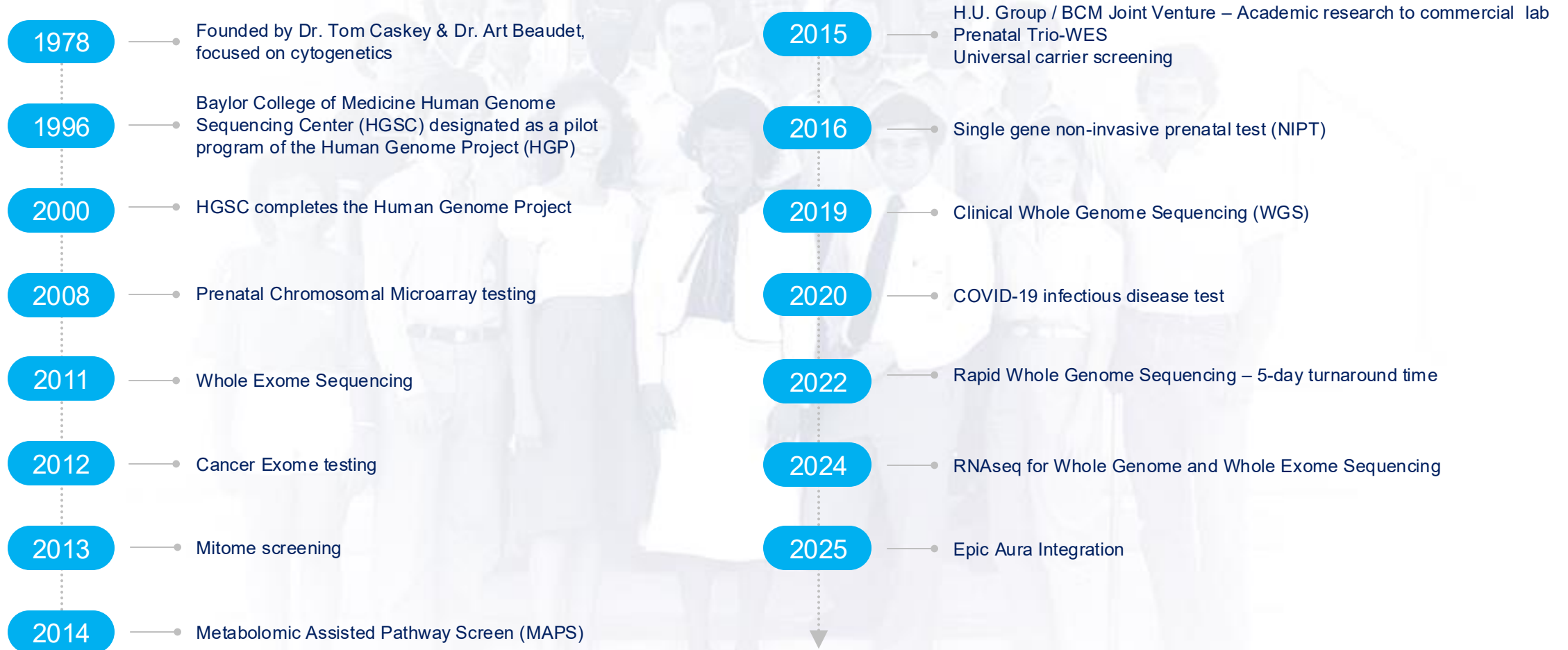


National &  
Global Reach

50 states – 16 Countries

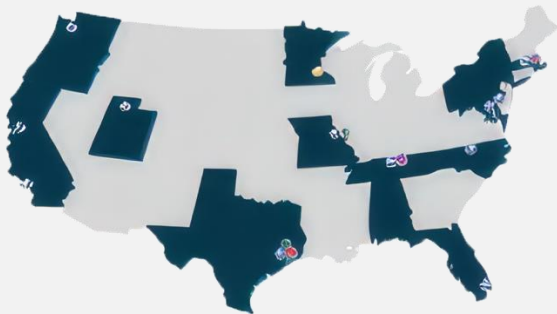
# // TRACK RECORD OF INNOVATION

## SCIENTIFIC AND CLINICAL IMPACT



# // DEEP EXPERTISE IN RARE DISEASE

Baylor Genetics  
is the sole sequencing  
core partner for the  
Undiagnosed Diseases  
Network (UDN), funded by  
National Institutes of Health



2,396

participants  
evaluated or ~1  
every other day

2,229

participants with  
exome and/or  
genome  
sequencing

718

30% of  
participants  
diagnosed

- UDN leverages Baylor Genetics expertise with Whole Genome and Exome Sequencing, and unique capabilities to deliver answers for patients with ultra-rare diseases
- Providing data-driven information and insights to improve health outcomes



## COLLABORATING ON COMPLEX CASES

- 10-year partnership
- 14 sites at top U.S. children's hospitals and 10 diagnostic centers of excellence
- Working with clinical and research experts to solve the most challenging cases



# // PROVEN LEADERSHIP TEAM

## DECADES OF EXPERTISE IN HEALTH AND TECH

### MANAGEMENT TEAM



**Kengo Takishima**  
Chairman & Chief Executive Officer



**Christine Eng, MD**  
Chief Medical Officer  
& Chief Quality Officer



**Chris Sands**  
SVP, Sales & GM,  
Diagnostic Genomics



**Shannon Kieran**  
VP, Product



**Jerry Wang**  
SVP, Business Development,  
Emerging Business, & Strategic Alliances



**Fan Xia, PhD**  
Chief Genomics Officer



**Ji He, PhD**  
Chief Technology Officer



**Li Shen**  
Chief Accounting Officer



**Kourtney Walsh**  
Director, Client Services



**David Berger**  
Chief Legal Officer



**Linda Ballard**  
Chief Human Resources Officer

### SCIENTIFIC ADVISORY BOARD



**Brendan Lee, MD, PhD**  
Robert and Janice McNair Endowed  
Chair in Molecular and Human  
Genetics, Professor & Chairman,  
Molecular and Human Genetics,  
Baylor College of Medicine



**Sharon E. Plon, MD, PhD**  
Dan L. Duncan Comprehensive  
Cancer Center Professorship,  
Baylor College of Medicine



**Ignatia Barbara  
Van den Veyver, MD**  
Professor, Departments of Obstetrics and  
Gynecology and Molecular and Human  
Genetics, Baylor College of Medicine

### SCIENCE AND MEDICINE

- 15+ PhDs and MDs
- 40+ ABMCG-certified lab directors, researchers, genetic counselors

### TECHNOLOGY AND BUSINESS

- Broad experience at enterprises including GE, GeneDx, Invitae, MD Anderson, PerkinElmer, Pfizer, and Roche
- Recognized on association boards, fellowships, 2024 top healthcare leader lists and merit awards

### DOMAIN EXPERTISE

- Clinical and commercial
- Healthcare, biotech, life sciences, pharma, technology, strategy

# // SERVING PATIENT AND PROVIDER NEEDS



430,000+

Individuals  
planning a family



60,000+

Pregnant women  
seeking diagnoses



57,000+

Newborn babies



200,000+

Children



250,000+

People at risk for  
cancer

Patients  
~ to date

Healthcare  
Providers

Genetic Counselors

OB/GYNs

Critical care

Pediatricians

Oncologists

Medical  
Conditions

Reproductive Health

Prenatal

Rare Diseases

Developmental  
Delays

Inherited  
Mutations

Tests

Carrier screening

Non-invasive  
prenatal testing

Whole Genome / Exome Sequencing

Hereditary cancer

# // COMPREHENSIVE MULTI-OMIC TESTING



## Whole Genome / Exome

- Comprehensive tests for complex cases
- High diagnostic yield in one test
- Fastest **rapid turnaround** – as soon as 5 days
- Re-analysis



## Chromosomal Microarray

- Product range and flexibility
- Exon coverage of 5000+ genes
- 100k+ microarrays
- Complementary test options



## Reproductive Health

- Prenatal multigene sequencing and carrier screening
- **PreSeek™** non-invasive prenatal screen
- **GeneAware™** reproductive carrier screen



## Oncology

- Somatic and germline
- Hereditary Cancer with RNA seq reflex
- CMA exon coverage
- Cytogenetics and FISH



## Biochemical

- Enzymes, proteins and metabolites
- **Global MAPS™** one-of-a-kind single test for inborn metabolic disorders



## Molecular Diagnostics

- Customizable testing
- Disease-specific panels with WGS reflex
- Nuclear and Mitochondrial
- RNA sequencing



## Advantages

Multidisciplinary assessment, industry-leading rapid turnaround time, flexible sample types, personalized treatment guidance, clinical support



# // SCIENTIFIC RESEARCH AND KNOWLEDGE SHARING

## Collaborative Research



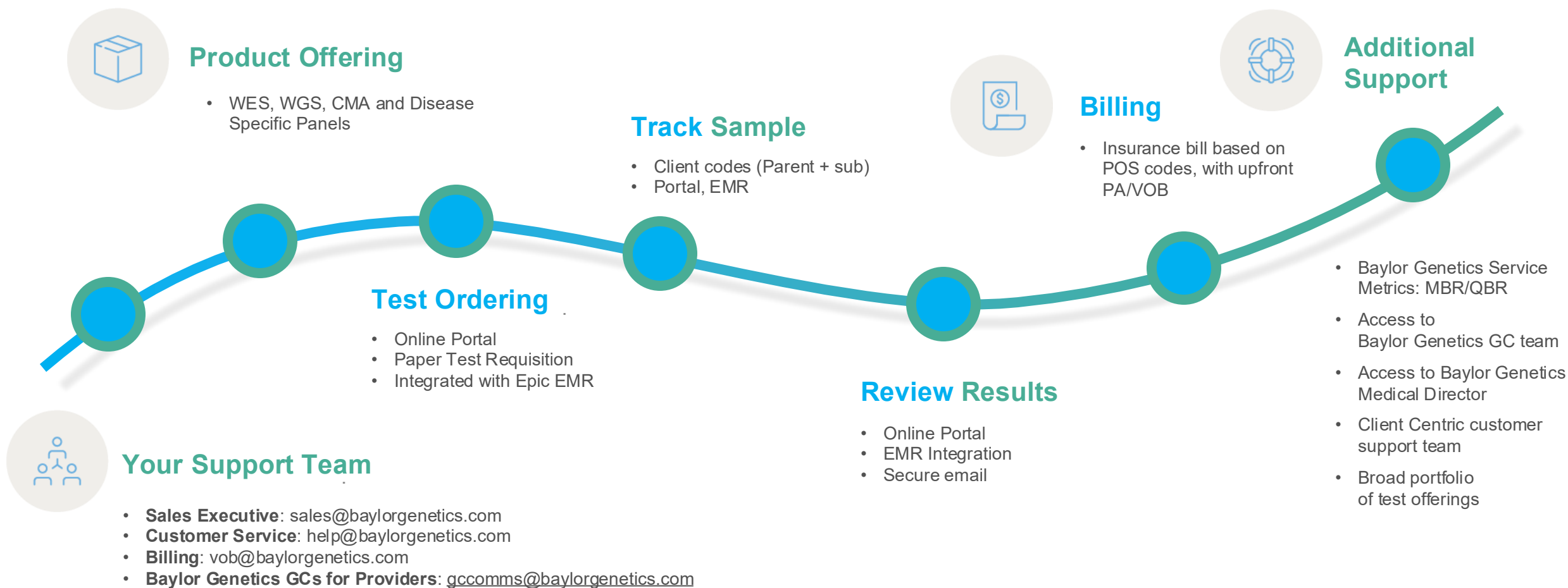
PreNatalSeq

## Peer Reviewed Publications and Presentations

- 350+ research studies and presentations
- 8 scientific studies<sup>1</sup> in major medical journals
- 15+ abstracts at top 2024 industry conferences
- 100+ new disease mechanisms discovered
- Data sharing with ClinVar



# // CUSTOMER-CENTRIC SERVICE MODEL



# // SUPPORTING PROVIDERS WITH OUR GENETIC EXPERTS

## Genetic Counselor Support Team



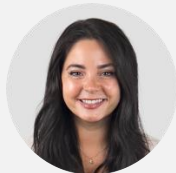
Laura Andolina  
MS, CGC



Allie Heritage  
MS, CGC



Thomas Felton  
MS, CGC



Romy Fawaz  
MS, CGC



Grace Hollingsworth  
MS, CGC



Lexi Isaacs  
MS, CGC

## Medical Affairs Team



Jason Chibuk  
MS, CGC



Rob Rigobello  
MS, CGC



Sydney Lau  
MS, CGC



Leah Campbell  
MS, CGC



Eileen Barr  
MS, CGC



Lisa Salz  
MS, CGC

## SCIENTIFIC LEADERSHIP

- 15+ PhDs and MDs
- 40+ ABMCG-certified lab directors, researchers, genetic counselors

## PROVIDER SUPPORT

Our experienced client support team is available to handle your day-to-day needs.

## DIRECT ACCESS TO BAYLOR'S GENETIC COUNSELORS

The genetic counseling team is available to provide clinical support to healthcare providers.

# // STATE-OF-THE-ART TESTING LABORATORY



## AUTOMATED WORKFLOWS

- Robotics, predictive analytics
- AI, machine learning
- 300,000 samples / yr



## MULTI-OMICS PLATFORM

- Flexible, scalable testing capabilities
- Custom solutions and assays
- 2 petabytes of sequencing data / yr



## ADVANCED TECHNOLOGY

- NovaSeq X sequencing high throughput clinical apps - 1st in Texas
- 73,000 sf testing hub



## CURATED DATASETS

- 3.6 million+ validated clinical patient datasets
- Clinical trial, biomarker, drug discovery support



## PARTNER SERVICES

- Strategic partnerships - life sciences, biotech & pharma
- Assay development, testing, validation and collaborative research



## ANALYTICAL EXPERTISE

- 19 board-certified clinical Lab Directors
- Faculty of Baylor College of Medicine



## CERTIFIED & ACCREDITED



CA, FL, MD, PA, RI



# // OUR PARTNERS

## NATIONAL RESEARCH PARTNERS



## TOP-RANKED HEALTH SYSTEMS



// CONTACT US

# BAYLOR GENETICS

[www.baylorgenetics.com](http://www.baylorgenetics.com) // 1-800-411-4363 // [help@baylorgenetics.com](mailto:help@baylorgenetics.com)