

Navigating Precision Medicine Implementation in Qatar and the Region

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Disclosure Slide

I have NO actual or potential conflicts of interest to disclose



Objectives

- To describe at high level the Qatari Biomedical Research Ecosysem
- To introduce Sidra Medicine as an academic medical center advancing Precision Medicine for women and children
- To describe implementation of Precision Medicine in clinical practice
- To describe the regulatory and ethical pathways governing (enabling) clinical implementation
- To invite future collaborations on biomedical and health research in the region



Qatar's Biomedical Research Landscape

Qatar Foundation's Education City



Key Biomedical Stakeholders





















Sidra Medicine

Women and Children's Research Hospital

Sidra Medicine Tripartite Mission

Patient Care

Research

Education



Vision

To be a beacon of learning, discovery and exceptional care, ranked among top **academic medical centers** in the world.

Sidra Medicine in Numbers

- 400 beds
- 250,000 patient visits per year
- >90% pediatric tertiary care
- 37 clinical sub-specialty
- ~3500 staff (~250 in Research)
- >260 active research studies (incl.
 - >80 international collaborations)



Sidra's Research Programs and Platforms

Translational Research Programs (Gold Cohorts)

Rare Metabolic and Mendelian Disorders



Childhood Diabetes Inborn Errors of Met. Newborn Seg Neurological Disorders



Autism Epilepsy Neuromuscular

Cancer



Pediatric Solid Tumors Leukemia Immune Dysregulation



Inborn Erros of Immun. Immune Deficiencies IBD

Congenital Structural Malformations



Craniofacial Disorders Congenital Heart Dis. Genitourinary Malform. Reproductive & Perinatal Health



Infertility
Pre-term birth
Gestational Diabetes

Core Research Platforms

(National Cores)

Diagnostic Cores (Dx)

- National Genome Lab: WGS (short + LR)
- Zebrafish
- Microbiome
- Biorepository
- Cytometry
- Advanced Microscopy

Therapeutic Cores (Tx)

- GMP Facility
- Cell and Gene Therapy Core
- iPS Core
- Clinical Trials Office
- Biostats Core



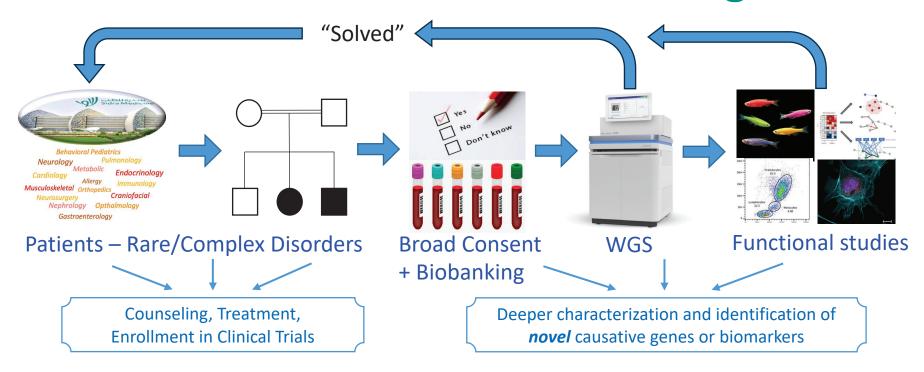
Sidra's Core Genome Lab



- National Reference Lab for Genomics
- Industry-leading sequencers/automation
 - Incl Long- & Ultra-Long Read Seq.
 - Certified service provider for PacBio
- CAP-accredited clinical Whole Genome Sequencing (cWGS)
- High-throughput biobanking/processing of many types of biological samples
- Integrated Bioinformatics & Data Science
- Array-based & qPCR-based panel testing
- Multi-omics discovery platforms
- Single-cell workflows



Sidra's Precision Medicine Program



- "Solved" families are fed back to the healthcare system,
 - Work with pharma/biotech to bring advanced therapies to Qatar
- "Unsolved" families undergo further research
 - Multi-omics, zebrafish, functional cellular assays, etc.

















High impact discoveries over the years...

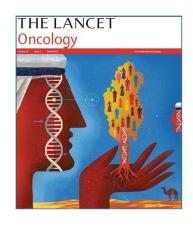
Publish ~100 papers per year on population genomics and rare disease; Avg IF ~8.7





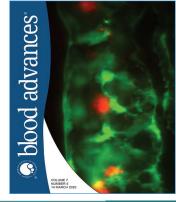
First journal cover (2016)

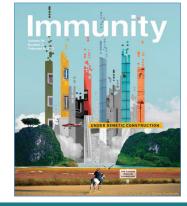


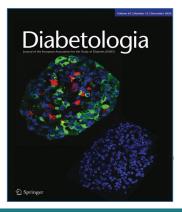










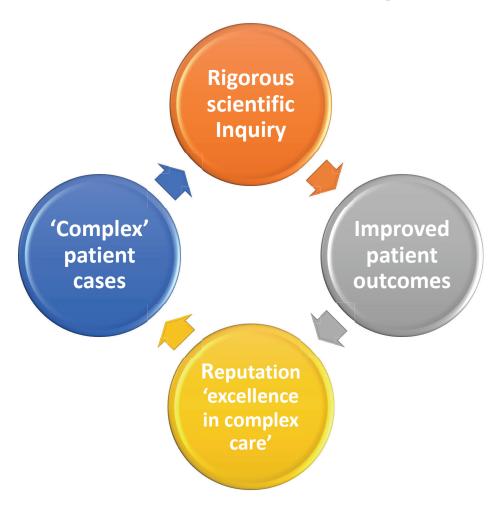


Clinical Implementation of Precision Medicine



The virtuous cycle of Precision Medicine

Precision Medicine = Research-guided care





'Every medical device, biomarker or lab test was once a research study'

Navigating Clinical Implementation

 Clinical Implementation of Precision Medicine is a constant feedback loop between bench and bedside

Bench to Bedside

Research Side:

- Translational Research Programs focused on disease-cohorts
- Diagnostic cores: elucidate disease etiology/mechanism
- Therapeutic cores: identify and develop novel therapies

Hospital Side:

- Multi-disciplinary Teams (MDTs)
- New Lab Developed Tests (LDTs)
- Advanced therapies (Clinical Trials)
- Implementation Science: measure impact of *research-guided* care

Bedside to Bench

Key Enablers:

Infrastrure and People Policies and Regulation

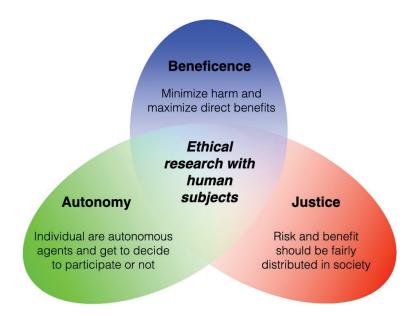


Policy and Regulation governing Clinical Implementation of Precision Medicine



Regulatory Landscape

- Regulation and Ethics are key factors to govern, and <u>enable</u>, research-driven care
- In the Qatar/GCC, the 2 sources of regulatory authority are:
 - Ministry of Health IRB (secular authority)
 - Religious Law Islamic Bioethics (religious authority)

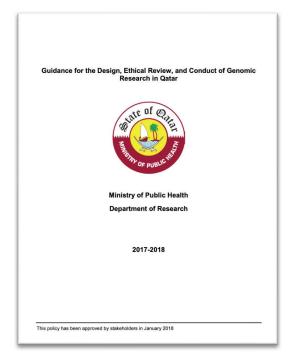




Regulatory Landscape

Regulation, IRB & Ethics

- Qatar's regulatory framework builds on the US 45 CFR 46 (Common Rule), Belmont Report and ICH-GCP
- Launched Genomic Research Guidelines in 2017:
 - Design and conduct of genomic studies
 - Biobanking + Data ownership, storage, and use
 - Patient Privacy, Risks, Ethical considerations
- Informed consent as the cornerstone of all biomedical research
 - Sample & data sharing
 - Commercialization including use by pharma
 - Recontact, future studies
- MoPH requires all IRB professionals to be certified (PRIM&R CIP)
- Flexible regulation (eg. stem cell use, expedited review categories)
- National reliance agreements facilitate multi-site collaborations
- Regular external audits/accreditations, including MoPH, JCI, AAHRPP

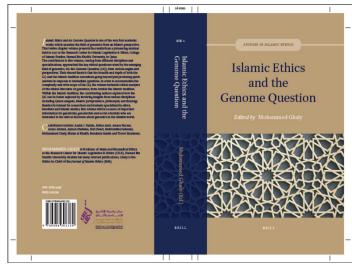




Regulatory Landscape

Islamic Ethics

- Biomedical research is a collective duty (fard kifayah), particularly as it stands to benefit society (ummah)
 - HGP: man's long journey to know oneself
- Guiding Principles: permissibility, beneficence and collective benefit (maslaha) over individual cost
- Higher Objectives of Shari'a:
 - Justice & equity, eg. in access to advanced diagnostics & access to new therapies
- Data sharing and return of actionable results is mandated where direct benefit or avoidance of harm, e.g. BRCA1/2
- Sensitive findings (e.g. non-paternity) disclosure prohibited if harm outweighs benefit







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GENOMICS IN THE GULF REGION AND ISLAMIC ETHICS







Summary

- Prec. Med. thrives where research, care and ethics are seamlessly integrated
- In Qatar, multiple stakeholders work in tandem to **build capacity** and advance precision medicine implementation, from bedside to bench and back
- At Sidra Medicine, key research programs combine with advanced core platforms to turn research insights into improved patient outcomes
- Robust regulatory frameworks (IRB, genomic & clinical trial guidelines) enable responsible sharing, innovation, regular dialogue with regulatory bodies
- Islamic bioethics **encourages** scientific inquiry & participation in research
- Global collaborations and data sharing accelerate discovery & benefit patients not just within the region, but for the Arab diaspora globally



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Sidra's 11th Annual Precision Medicine Conference Save the Date – 2-4 Dec, 2025

