



HRSA Overview

Mountain States Regional Genetics Network Winter Meeting

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Maternal and Child Health Bureau (MCHB)

Vision: Healthy Communities, Healthy People



Health Resources and Services Administration

HRSA's Mission:

To improve health outcomes and address health disparities through access to quality services, a skilled health workforce, and innovative, high-value programs.

Department of Health and Human Services (DHHS)

Administration for Children and Families (ACF)

Food and Drug Administration (FDA)

Administration for Community Living (ACL)

Health Resources and Services
Administration (HRSA)

Agency for Healthcare Research and Quality (AHRQ)

Indian Health Service (IHS)

Agency for Toxic Substances and Disease Registry (ASTDR)

National Institutes of Health (NIH)

Centers for Disease Control and Prevention (CDC) Substance Abuse and Mental Health Services Administration (SAMHSA)

Centers for Medicare and Medicaid Services (CMS)





Maternal and Child Health Bureau

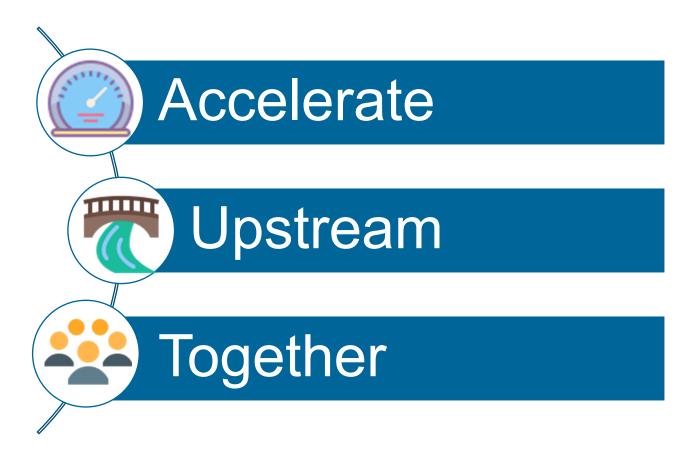


Mission: Improve the health of America's mothers, children, and families.





Paradigm for Improving Maternal and Child Health







Division of Services for Children with Special Health Needs

DSCSHN Mission

To provide national leadership to expand and strengthen access to a system of services for CYSHCN and their families as they grow and transition through adulthood.



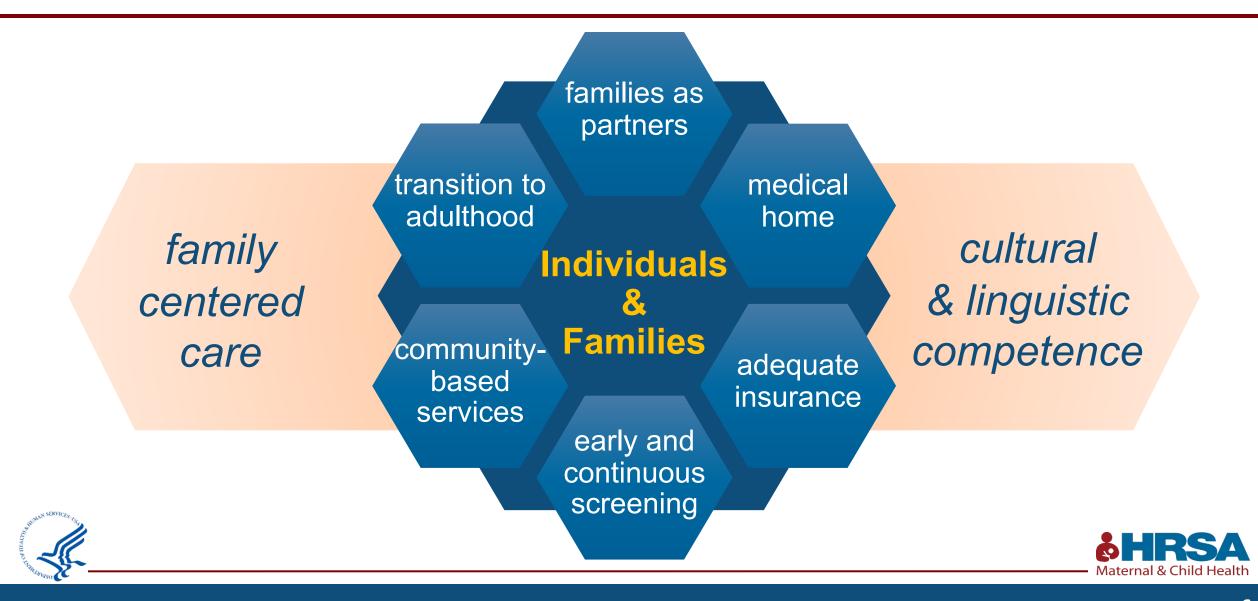
Who do we serve?

Children or youth who have or are at increased risk for a chronic physical, developmental, behavioral, or emotional condition and who also require health and related services or a type or amount beyond that required for children generally.





Six Indicators of a Well-Functioning System



DSCSHN: Program Snapshot

Systems of Care

Medical home

Transition

Financing

Family/Professional Partnership

CSHCN Research Network

CMC CollN

Screening

Newborn screening

Genetics Programs

Vision

Early Hearing Detection and Intervention

Environmental Health

Conditionspecific

Epilepsy

Autism

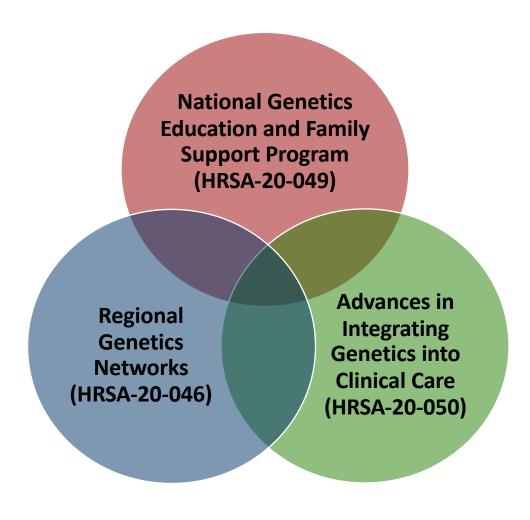
Hemoglobinopathies

Hemophilia





Genetics Programs



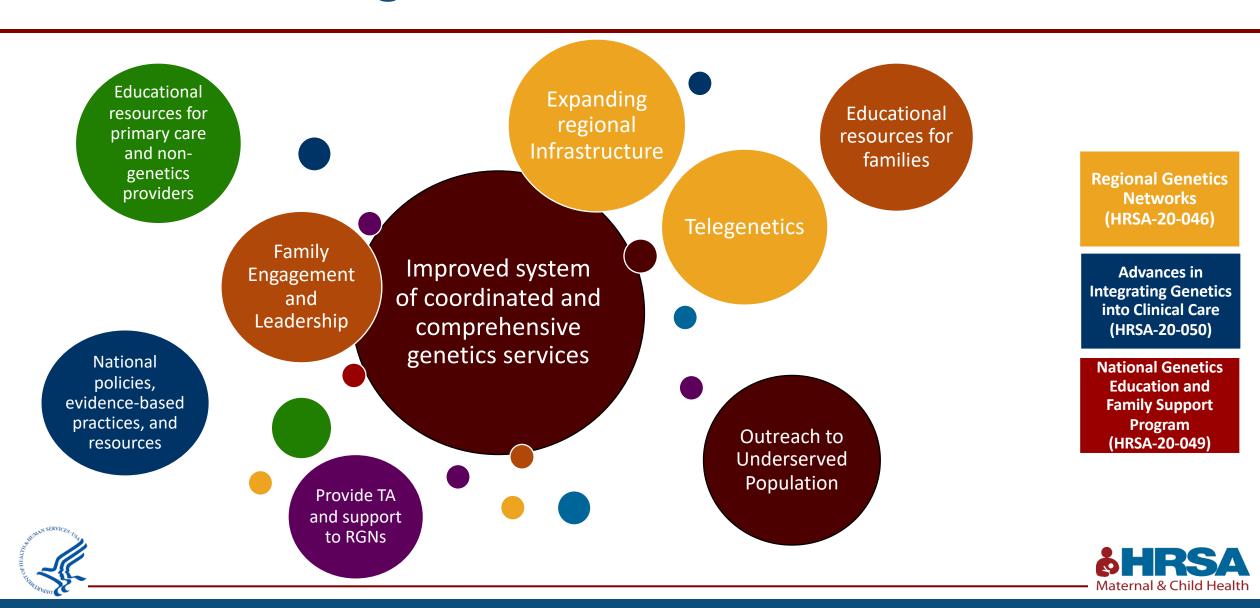
Overarching Goal

To improve access to genetic services, especially for underserved, disadvantaged, geographically isolated, and special needs populations

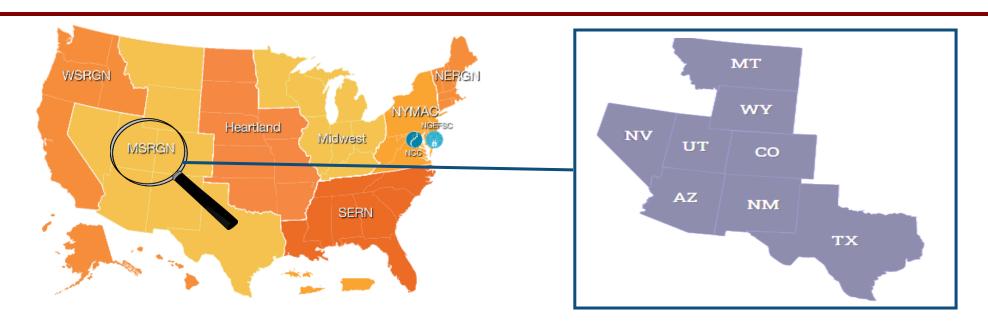




Genetics Programs Priorities



Regional Genetics Networks Priorities 2020-2024



Support a regional infrastructure for the genetics health care delivery system

- Strengthen, build, and expand partnerships, relationship, and infrastructure throughout each state of the region
- Develop and implement mechanisms to identify individuals and their families with or at risk for genetic disorders, especially underserved, and connect them to genetic services



Regional Genetics Networks Priorities 2020-2024

Provide education-related activities for providers, families, individuals, and other stakeholders.

- Develop and expand educational resources for health professionals
- Develop educational resources for families and support to family leaders to participate in MSRGN and genetics care delivery system

Facilitate the use of telehealth and telemedicine in the genetics health care delivery system.

- Provide technical assistance, training, and support to providers on telehealth
- Facilitate implementation and expansion of telehealth in genetics service provision







What and why do we measure?

- RGN Program Objectives 2020-2024
 - Each RGN will facilitate connections to genetic services for at least 2,000 individuals or families with or at risk for genetic conditions, within the geographic area served by the RGN.
 - Increase to at least 33% of the individuals who are medically underserved with or at risk for genetic conditions, served by each RGN.
 - Increase by 20% the number of health care providers receiving education or training through the RGN.
 - Increase by 20% the number of individuals with or at risk for genetic conditions,
 receiving genetic services through telemedicine visits.
 - Increase by 20% the number of RGN resources accessed.
- Program Measures
- Progress Reports Both Quantitative and Qualitative

Manage the work

Transparency

Feedback

Answer stakeholder questions





New Websites

- Newbornscreening.hrsa.gov
 - Provides clear and up-to-date information, materials, and resources about NBS in the United States. These resources help increase awareness, knowledge, and understanding of NBS and genetic conditions.



 Provides information for patients and providers about the latest federal efforts to support and promote telehealth services



EXPLORE NEWBORN SCREENING

About Newborn Screening	Newborn Screening in Your State
Learn how newborn screening can prevent serious health	Find out about the newborn screening programs in each U.S.





Mark your calendar



Next meeting scheduled for February 11-12, 2021 Register Here





Contact Information

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