

**Health Resources and Services Administration (HRSA)
 Required Data Reporting for Regional Genetics Networks
 from [Funding Opportunity Announcement No. 93.110](#)
 6/1/2017 – 5/31/2020**

HRSA Program Objectives	HRSA Required Data Reporting
<ol style="list-style-type: none"> 1. By May 2018 each RGN will provide services to at least 250 individuals or families within the geographic area served by the RGN. 2. By May 2020 each RGN will provide services to at least 1,500 individuals or families within the geographic area served by the RGN. (also see earlier May 2018 Objective) 3. By May 2020, increase by 20 percent the number of medically underserved patients served by each RGN. 4. By May 2020, increase by 20 percent the percentage of clinical sites that use telehealth/telemedicine to provide genetic services. 5. By May 2020, increase by 20 percent the number of medically underserved patients receiving genetic services through telemedicine visits. 6. By May 2020, increase by 20 percent the number of primary care providers using RGN resources. 	<ol style="list-style-type: none"> 1. Number of individuals or families receiving services within each RGN* 2. Number of medically underserved patients receiving services within each RGN 3. Number of primary care, genetic service providers, and public health officials participating in the regional network infrastructure 4. Number of technical assistance activities/services provided to genetic services providers 5. RGN participation in QI activities to connect the medically underserved to genetic services 6. Number of patients/families receiving education and resource materials 7. Number of health care providers receiving health education/resource materials 8. Number of hits on RGN website and social media by stakeholders, providers and consumers 9. Number of clinical sites that use telehealth/telemedicine to provide genetic services 10. Number of medically underserved patients receiving genetic services through telemedicine visits <p>*An individual or family directly “served” by the RGN is one for whom a genetic appointment has been facilitated through RGN resources, for example through a telehealth program, or indirectly through outreach and education to primary care or public health.</p>

Note: As of November 15, 2017, the [National Coordinating Center for Regional Genetic Networks \(NCC\) Evaluation Work Group](#) is working to develop performance measures that address the above reporting requirements.