
M110 BHRN Data Collection

Presented to Oversight and Accountability Council
May 4, 2022



SB 755, Section 22, 5e

(e) Outcomes for individuals receiving treatment and other social services under section 2, chapter 2, Oregon Laws 2021 (Ballot Measure 110 (2020)), including, but not limited to, the following:

(A) Whether access to care increased since December 3, 2020, and, if data is available, whether, since December 3, 2020:

(i) The number of drug and alcohol treatment service providers increased.

(ii) The number of culturally specific providers increased.

(iii) Overdose rates have decreased.

(iv) Access to harm reduction services has increased.

(v) More individuals are accessing treatment than they were before December 3, 2020.

(vi) Access to housing for individuals with substance use disorder has increased.

(B) Data on Behavioral Health Resource Networks and recipients of grants and funding under section 2, chapter 2, Oregon Laws 2021 (Ballot Measure 110 (2020)), including:

(i) The outcomes of each network or recipient, including but not limited to the number of clients with substance use disorder receiving services from each network or recipient, the average duration of client participation and client outcomes.

(ii) The number of individuals seeking assistance from the network or recipients who are denied or not connected to substance use disorder treatment and other services, and the reasons for the denials.

(iii) The average time it takes for clients to access services and fulfill their individual intervention plan and the reason for any delays, such as waiting lists at referred services.

(iv) Whether average times to access services to which clients are referred, such as housing or medically assisted treatment, have decreased over time since December 3, 2020.

(v) Demographic data on clients served by Behavioral Health Resource Networks, including self-reported demographic data on race, ethnicity, gender and age.

Required data specific to BHRNs

- Who is/is not receiving services
 - Number of clients served/denied
 - Reasons for denials
 - Self-reported demographic data
- Service measures
 - Average wait times to access services
 - Average duration of service/time to fulfill intervention plan
 - Reasons for any delays
 - **Outcomes**

Phased approach to data collection

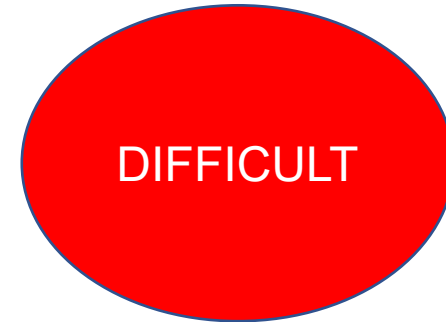
The required data to be collect ranges from



Phased approach to data collection

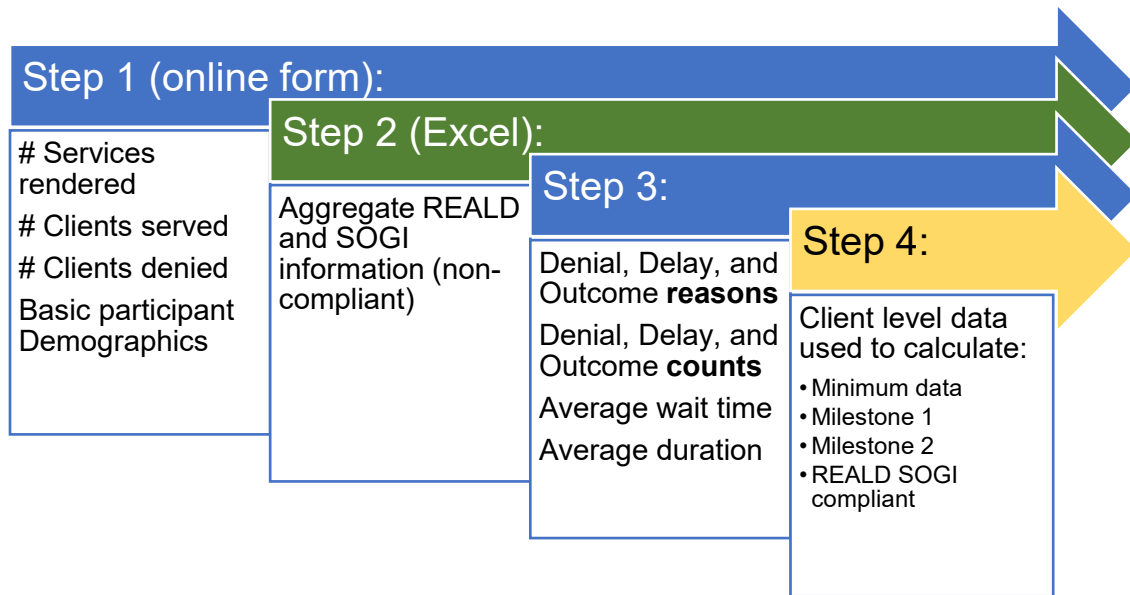


- One entity for a county
- 7 service elements to report



- Multiple entities by county
 - 7 service elements to report
 - Collecting details that allow for reporting at aggregate
- Multiple entities & BHRNs by county
 - 7 service elements to report
 - Collecting details that allow for reporting at person level
 - Allowing for 360-degree view of individual

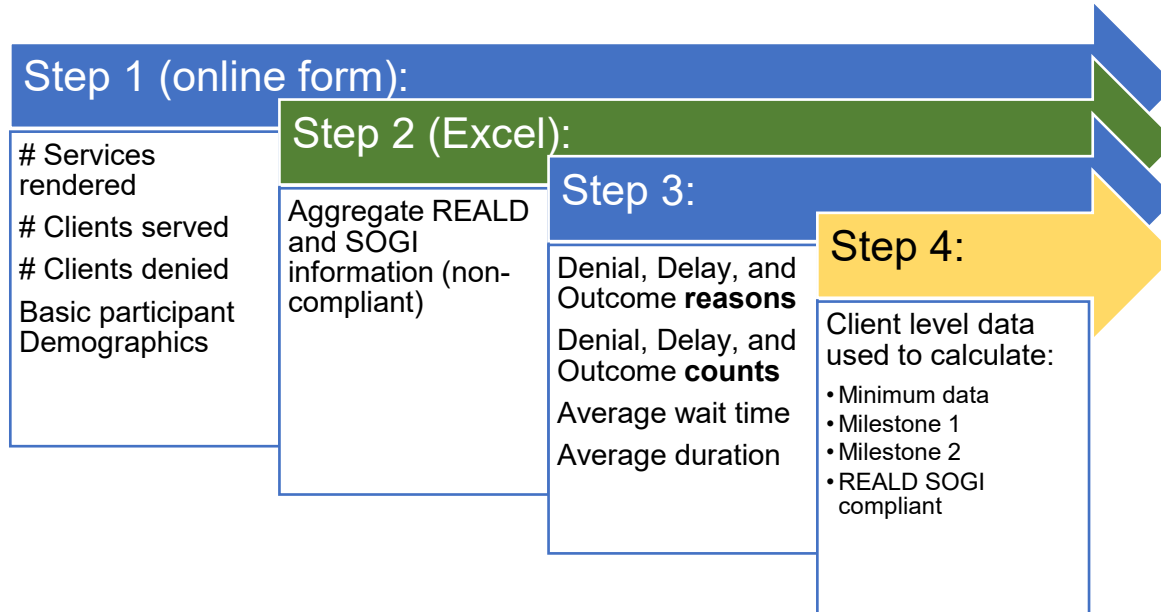
Phased approach to data collection



Pros and Cons

- Steps can be tied to dates to drive adoption
- Allows all awardees to express work completed
- Clearly describes direction for future data collection efforts
- Reflects the many varied capabilities of BHRN entities
- Mutually beneficial for BHRNs and OHA; Technical Assistance can be targeted
- **Reporting across BHRNs may be limited initially**

Phased approach to data collection



STEP 1:
Allows us to view things at high level and establish the amount of people and services being delivered by element

STEP 2:
Add details related to demographics and pushes providers to REALD and SOGI compliance

STEP 3:
Incorporates details associated with outcomes and provides insight into what barriers to services may exist

STEP 4:
Allows us to look at this for an individual point of view and understand the relationship between services and outcomes

Additive

Thank You

Health
Oregon
Authority

Grant language about BHRN data reporting

Added May 19, 2022

Data reporting language in BHRN grant

- To reduce the initial burden on recipients, data reporting will be implemented in a phased approach over the first year of the award. Throughout the first year, recipients shall participate in brief surveys to assess data collection and reporting capabilities, which will inform benchmark requirements set by the OHA. All recipients, regardless of the initial funding date, shall meet program-wide reporting deadlines.
- Recipients shall meet first year benchmark reporting requirements as described below.

Data reporting language, continued

- Benchmark 1: By the end of the first two quarters, recipients shall report aggregated data for:
 - The number of clients served and services rendered
 - The number of clients denied or not connected to services
 - Demographic data on clients served
- Benchmark 2: (Date TBD) Recipients shall report aggregated data for:
 - Benchmark 1 requirements
 - Demographic data (REALD/SOGI) collected in accordance with OAR 943 Division 070.

Data reporting language, continued

- Benchmark 3: (Date TBD) Recipients shall report aggregated data for:
 - Benchmarks 1 and 2 requirements
 - The average wait time for clients to access services
 - The average duration of client participation
 - Client outcomes*
 - The reasons for service denials*
 - The reasons for service delays*

*Specific categories to be defined by the OAC

Data reporting language, continued

- Recipients will be provided with an online reporting tool to submit aggregated data elements.
- After the first year of the award (June 2023), recipients will shift from reporting aggregated data to client-level data on the measures listed above.
- Throughout the period of the award, each recipient shall complete brief surveys on topics including, but not limited to: organization characteristics, outreach activities, accessibility of services, and interactions between BHRN entities.