

Roles and Responsibilities of a Quality Council: Using Data to Improve System Performance

Valerie J. Bradley RQC Summit August 22, 2019

What To Expect

- Overview of rationale for quality councils
- Mission of quality councils
- Responsibilities of members
- Ways to review and understand data
- Turning data into quality improvement
- Highlights from NCI In-Person Survey
- Examples of how data has been used around the country





Parable of the Blind Men and the Elephant: Each Man Brought a Different Interpretation

To Fully Understand Quality Multiple Perspectives Should be Present

Program participants

Family members

Providers

Direct support staff

Public managers

Advocates



Why Should We Care About Quality?

We have created a movement and made promises to people with disabilities and their families

Ideology alone does not create a stable and reliable system of supports

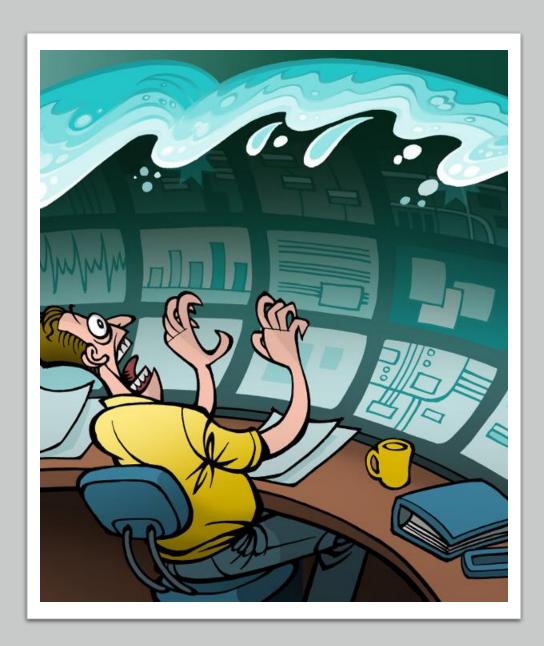
The greater the investment the greater the expectations

Unless we build quality in at the beginning, it is very hard to retrofit a program later





What is the Context You're Working in?



Multiple Data Sources

- Incident management systems
- Certification and licensing
- Payment and claims data
- National Core Indicators
- Individual and provider reviews
- HCBS rule compliance
- Waiver assurances

Complexity



- I/DD systems have multiple layers and players
- Data comes from different parts of the system
- One data point does not necessarily tell you all you need to know
- Collecting and trending data is one way to cut through the complexity

Multiple Demands for Data



- Compliance with HCBS Rule
- Compliance with Court Orders
- Accountability to the legislature
- HCBS waiver assurances
- Fiscal accountability
- Transparent data for people with disabilities and families



Mission of Regional and State Quality Councils

Quality Council Responsibilities

To bring your particular perspective to bear when reviewing performance data

To identify priority issues reflected in the data

To develop recommendations and quality projects aimed at issues that arise in the data and that can reasonably be expected to result in system improvement

To find ways to use data to improve system and participant outcomes

To follow up and track progress on any recommendations or projects recommended by the Council

To explore whether quality processes generate data relevant for quality improvement



What Does it Mean to be a Participant on a Quality Council?



- Each person's input is needed to ensure that multiple perspectives are represented
- Active participation is important given the need for continuity. This means attending meetings
- This is not a forum for individual issues or grievances. This is a forum for discussing system level issues identified in the data
- A quality council relies on people bringing an open and sometimes skeptical mind to the discussion not specific agendas

Priorities: You Can't Do Everything!

Participant Outcomes?

- Employment?
- Inclusion?
- Health and Well-Being?
- Other?

Health and Safety?

- Abuse, Neglect, Exploitation?
- Mortality?

Provider capabilities

• Compliance issues?

System Change Goals

- Implementation milestones?
- Improvements in quality assurance?

Staffing/Training Issues?



Review Data -- Cautions



Validity and Reliability of Data

Sample size (big enough?) and characteristics (random?)

Date of data collection? Has any change happened in the interim?

Beware of averages – pay attention to the ranges

Is the data biased? (e.g., self report)

ALWAYS make sure you:



1

Analyze the analysis.

2

Identify BIG issues that may compromise the data.

3

Do NOT generalize the findings beyond their limits.

4

BALANCE your review.— take into consideration other sources of information.

NEVER:



1

Make assumptions about the data – ask questions.

2

Expand the findings to whole population — unless it is appropriate.

3

Treat the data as "significant" unless it says it is.

4

Jump to conclusions without checking other sources.

Spotting Potential Misinterpretations

The numbers of reported incidents of restraints goes up

 This could indicate a serious problem. However, it may also be positive indicator if reforms have been put into place to incentivize providers to report restraints, this may be a positive indicator. Therefore important to understand the whole context.

The numbers of admissions to hospitals increases

• This trend could mean that there are serious issues at the provider level with respect to protection of health. However, another explanation might be that providers in the state have been trained to use the Health Risk Screening Tool to identify red flags that suggest the need for medical treatment and close surveillance. Such surveillance may have resulted in more necessary hospitalizations



Criteria for Identifying Issues



- Has an impact on a large number of individuals or a profound impact on a smaller group of individuals
- Important for compliance with federal or state requirements
- Issue has high cost(s) to the system: \$, timing, etc.
- Problem is growing/worsening
- Likely to be amenable to improvement
- Improvement is likely in a reasonable amount of time
- Improvement is likely with reasonable resource expenditure
- Aligns with other agency priorities



Quality Improvement Plans

- State problem & how improvement will be measured.
- Explore any additional steps to examine issue
- Name QI initiative and identify who is responsible (e.g., state I/DD agency? Council workgroup? DD Council?)
- Brainstorm possible action steps, potential barriers & methods to minimize the barriers.
- Select action steps and how the results of the improvement plan will be measured
- Implement the plan
- Measure progress

Cycle of Quality Improvement

Embed the reform, or refine if aims weren't met



What is the goal? What are the steps

Did the intervention work?

Assign tasks and implement

Selected
Findings
from InPerson NCI
Survey in
Virginia

Selected finding from National Core Indicators Data: 2017-2018





WHAT IS NATIONAL CORE INDICATORS?

Multi-state collaboration of state DD agencies

Measures performance of public systems for people with intellectual and developmental disabilities

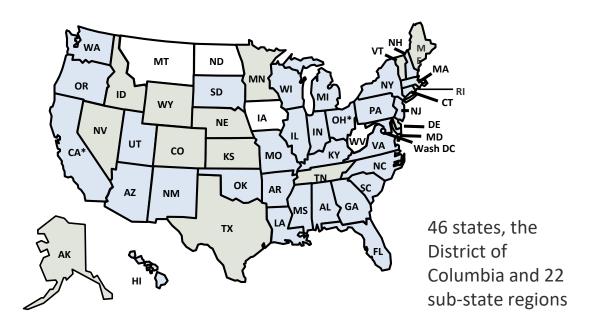
Assessment of performance in several areas, including employment, community inclusion, choice, rights, and health and safety

Launched in 1997 in 13 participating states

Supported by participating states

NASDDDS - HSRI Collaboration

National Core Indicators State Participation 2016-2017



Please note: not all NCI participating states participate in all NCI surveys each year.

NCI is a Person-Centered Tool that Provides Information on:

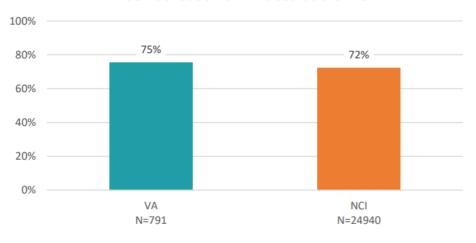


- Individual characteristics of people receiving services
- The locations where people live
- The activities they engage in during the day including whether they are working
- The nature of their experiences with the supports that they receive (e.g., with case managers, ability to make choices, self-direction)
- The context of their lives friends, community involvement, safety
- Health and well-being, access to healthcare

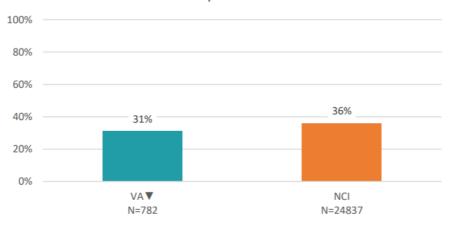
Wellness



Exercises or does physical activity at least once a week at least 10 minutes at a time



Exercises for the muscles in arms, legs, back, and/or chest

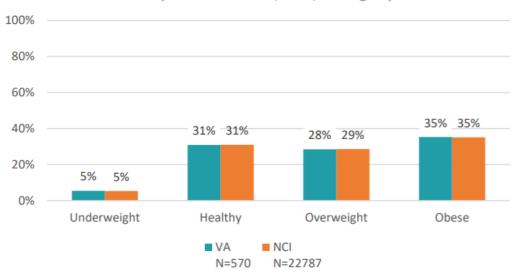


Exercises or does physical activity at least 3 times per week that makes the muscles in the arms, back, and/or chest work hard

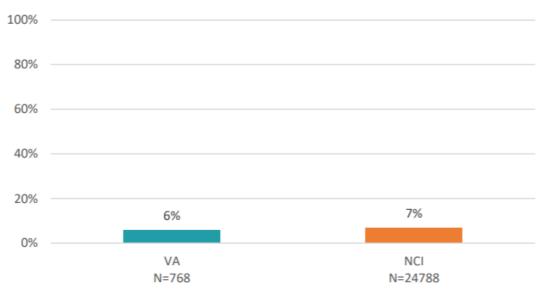
Wellness



Body Mass Index (BMI) category



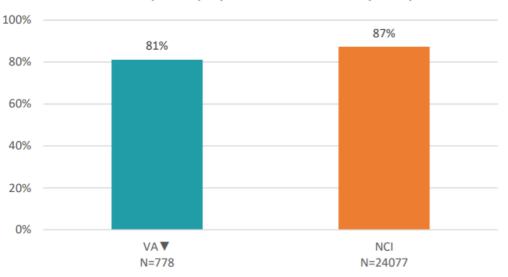
Uses tobacco products

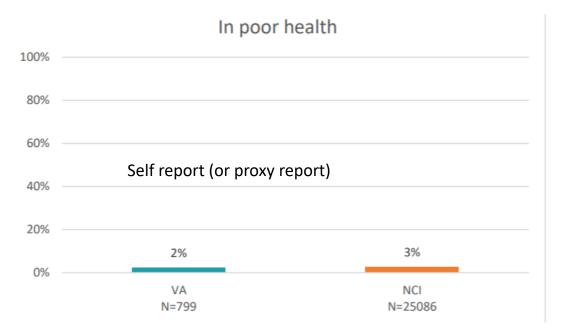


Health



Had a complete physical exam in the past year

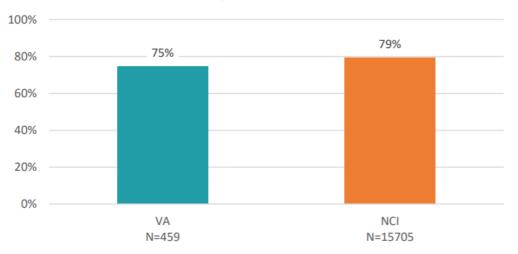




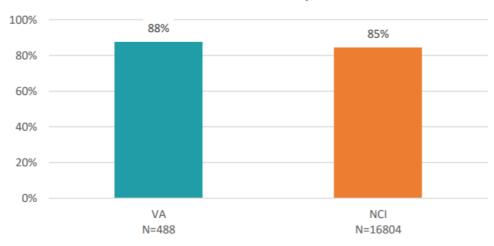
Community Inclusion



Able to go out and do the things s/he like to do in the community as often as s/he wants



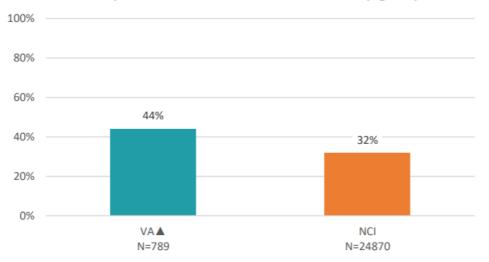
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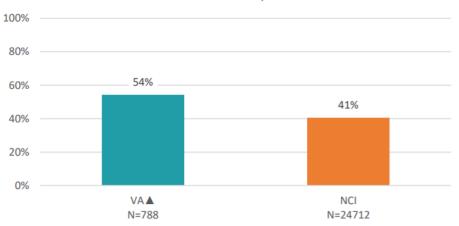
Community Inclusion



Participated as a member in community group

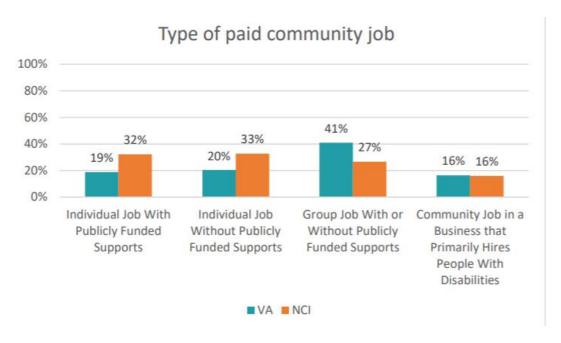


Went out to religious service or spiritual practice at least once in the past month

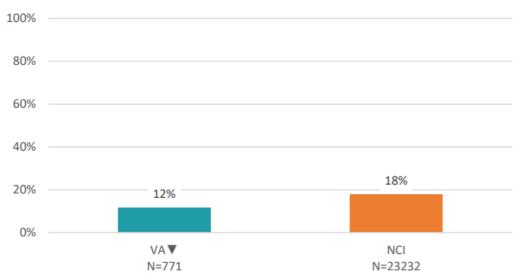


Employment





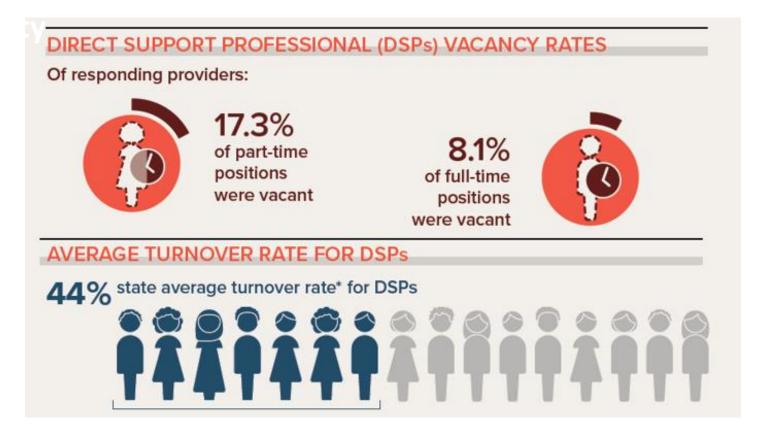
Has a paid job in the community





Data on the Direct Support Workforce NCI Staff Stability Survey

2017 NCI Staff Stability Report: Direct Support Professional Workforce in NCI States



All data are from a sample of 3,334 provider agencies from 19 states (Arizona, Connecticut, Georgia, Illinois, Indiana, Kentucky, Maryland, Missouri, Nebraska, New York, Ohio, Oklahoma, Oregon, South Carolina, South Dakota, Tennessee, Utah, Vermont), and the District of Columbia. Nationally aggregated data are weighted.

Direct Support Professional Workforce

HEALTH INSURANCE

Of responding providers:



Of responding providers who offer health insurance:



69% offer health insurance to only full-time DSPs



38% require DSPs be employed at the agency for a certain length of time to be eligible for health insurance



Of DSPs who left positions* in calendar year 2017:

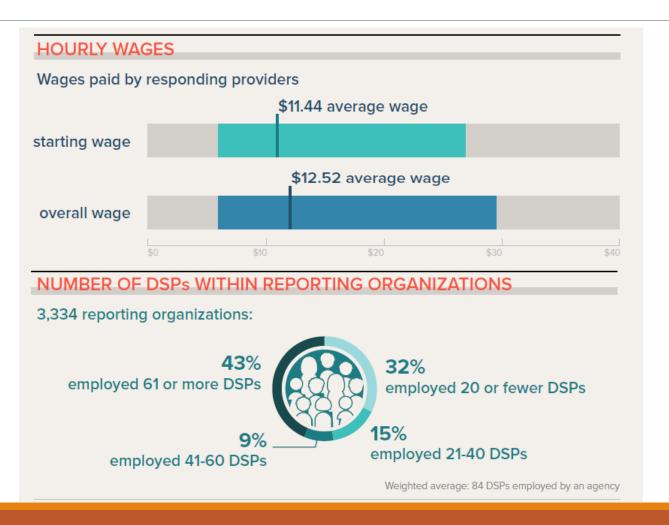
40%	21%	39%
left in fewer than	left between	left after
6 months	6 & 12 months	12 months or more

AVERAGE TENURE OF DSPs

Of DSPs employed within reporting organizations, as of December 31, 2017:

20%	16%	65%
have been	have been there	have been there
there fewer than 6 months	6-12 months	12 or more months

Direct Support Professional Workforce



Issues that NCI Data Have Shed Light On

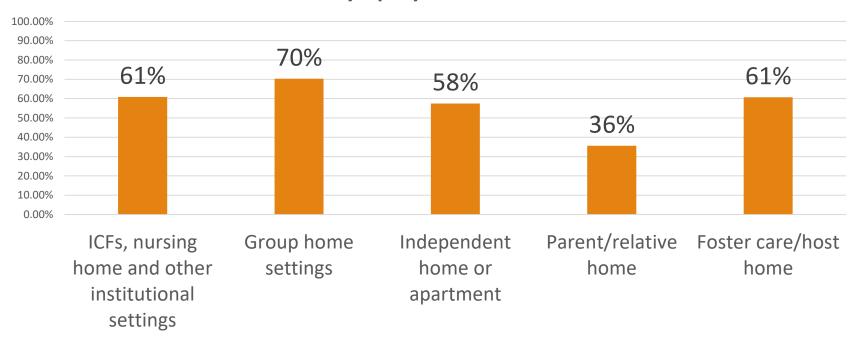






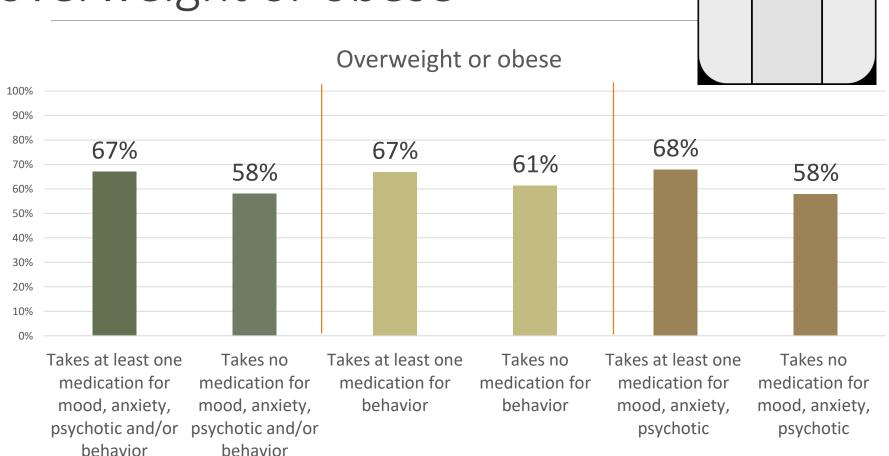


Percentage taking at least one medication for mood, anxiety, psychotic and/or behavior



Medication by residence type

Those taking medication were significantly more likely to be overweight or obese



Data Brief: Friendship and Life Outcomes

Expanded Friendships: Has friends who are not staff/family

Limited friendships: No friends, or friends are only staff and/or family

Employment:

Individuals who had expanded friendships were more likely to be employed in a paid community job

Rights and Privacy:

Those who had expanded friendships reported **having more privacy**; they also reported **having their rights respected** at greater rates.

Community Inclusion

Those with expanded friendships were more likely to have **gone into the community to take part in specific activities**; they also reported participating in these activities with greater frequency.

People with ASD were less likely to be independent of guardianship

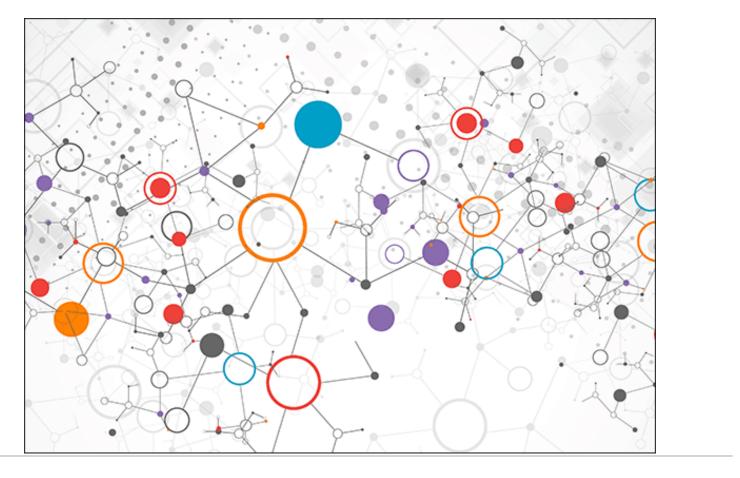
People with ASD were less likely than those without ASD to say they had friends who weren't family or staff

People with ASD were less likely to report having had at least some input in critical life choices such as choosing staff, daily schedule, free-time, etc.

People with ASD were significantly more likely to live in the family home, and significantly less likely to live in their own home and apartment

Article: Outcomes for those with ASD

Hiersteiner, D., Bradley, V., Ne'eman, A., Bershadsky, J. & Bonardi, A. (2017) Putting the research in context: The life experience and outcomes of adults on the autism spectrum receiving services in 29 states. *Inclusion* 5(1) 45-59



HOW STATES USE NCI DATA

Michigan



- 1. Supported Decision Making-because of the high rates of guardianship in Michigan, the DD Council developed a Supported Decision Making project to train people around the state to use SDM in the daily lives of people with IDD
- 2. Home and Community Based Services-developing a crosswalk between NCI and HCBS Settings Rules variables to validate data
- 3. Community of Practice-analyzing NCI data to inform Michigan's CoP on cultural and linguistic competence in service delivery

Missouri



- Identified risk factors for abuse neglect based on NCI case study that included loneliness, lack of friends, feeling unsafe
- Developed indicators based on NCI data
- 3. Developed strategies to identify interventions to prevent social isolation

Other State Initiatives Using NCI Data



New York

Publishes comparison data against other states

Targeted campaigns to decrease obesity rates

Ohio Council of Governments

Tracks person centered practices and changed the terminology of their planning process

Kentucky

Issues formal report on service quality and community participation; data was basis for system redesign

Massachusetts

Tracks and acts on health and wellness and safety data

