

National Core Indicators® 2018-19 In-Person Survey (IPS) Report

Executive Summary



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The National Core Indicators (NCI) are standard measures used across states to assess the outcomes of services provided to individuals with intellectual/developmental disabilities and their families. Indicators address key areas of concern such as employment, respect/rights, service planning, community inclusion, choice, and health and safety. The data that result from NCI surveys are often used to inform strategic planning, produce legislative reports, and prioritize quality improvement initiatives. Many states also share the data with stakeholder groups such as Quality Councils and use the stakeholder feedback to help set priorities and establish policy direction.

The NCI In-Person Survey is administered to individuals with a developmental disability who receive at least one service other than case management. Not all states that participate in NCI administer the In-Person Survey on an annual basis. Of the 46 states, District of Columbia and 22 sub-state entities who participated in NCI during the 2018-19 data collection cycle, 37 states submitted a valid sample of In-Person Survey data: Alabama (AL), Arizona (AZ), Arkansas (AR), Arizona (AZ), Colorado (CO), Connecticut (CT), Delaware (DE), Florida (FL), Georgia (GA), Hawaii (HI), Indiana (IN), Kansas (KS), Kentucky (KY), Maine (ME), Michigan (MI), Minnesota (MN), Missouri (MO), North Carolina (NC), Nebraska (NE), New Hampshire (NH), New Jersey (NJ), New York (NY), Ohio (OH), Oklahoma (OK), Oregon (OR), Pennsylvania (PA), Rhode Island (RI), South Carolina (SC), South Dakota (SD), Tennessee (TN), Texas (TX), Utah (UT), Virginia (VA), Vermont (VT), Washington (WA), Wisconsin (WI), and Wyoming (WY). This Final Report provides a summary of results based on data submitted by June 30, 2018.

The following are weighted NCI national averages for a selection of survey items. Complete breakouts by state of items in the In-Person Survey can be found in their respective chapters of this report.

2018-19 In-Person Survey

22,009 adults
with IDD
participated
across
37 states*

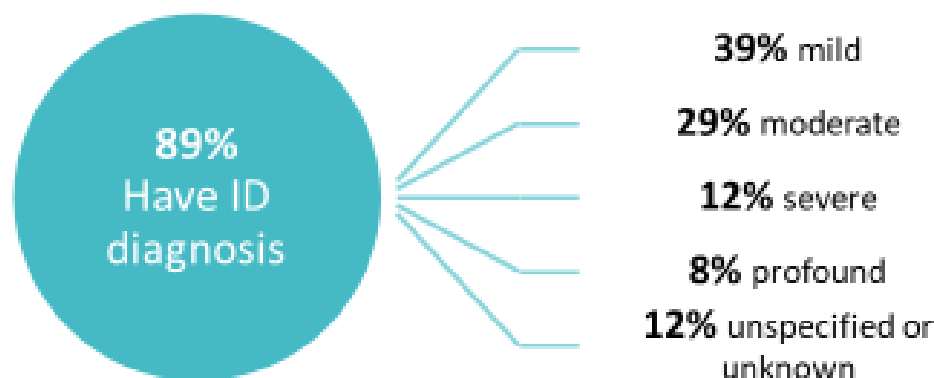
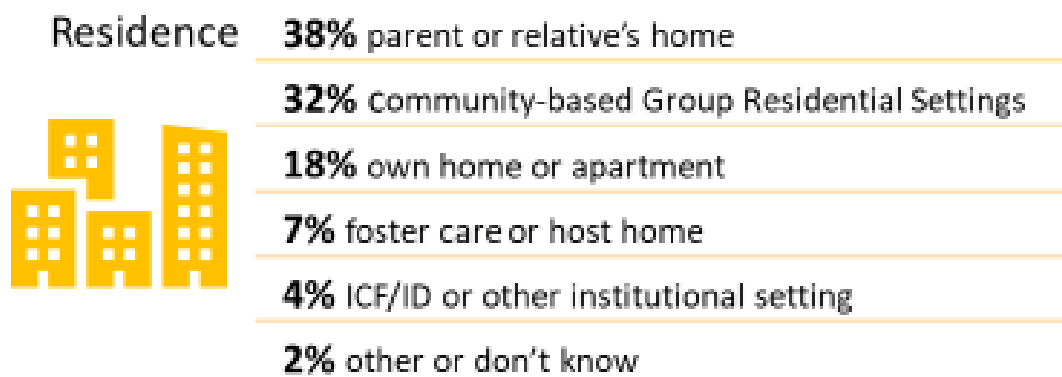
NCI Averages:

- Include all participating states
- Data are weighted

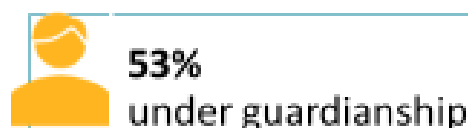
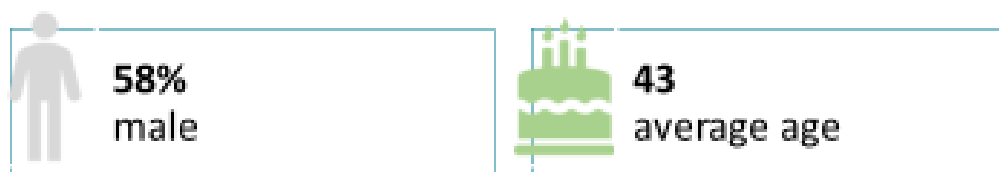
*AL, AR, AZ, CO, CT, DE, FL, GA, HI, IN, KS, KY, ME, MI, MN, MO, NC, NE, NH, NJ, NV, NY, OH, OK, OR, PA, RI, SC, SD, TN, TX, UT, VA, VT, WA, WI, WY

Respondents

Primarily adults with IDD age 18 and older receiving at least one service (in addition to case management) from the state DD agency. Some information may come from proxy respondents as well as administrative records.



Mental Health Diagnoses (not mutually exclusive)



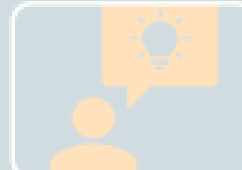
2018-19 In-Person Survey



96%
reported
having taken
part in last
service
planning
meeting



81%
understood
what was
being talk
about at last
planning
meeting



94%
reported
that they
chose or had
input in
services gets
as part of



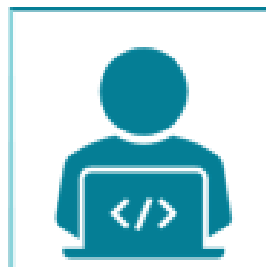
83%
know who to
ask to make
changes to
service plan

79%
report having
friends who are
not staff or
family

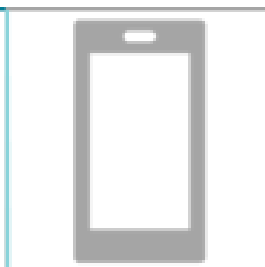
79%
report that they
can see friends
when they want

81%
report having
other ways of
talking or
chatting with
friends when
cannot see them

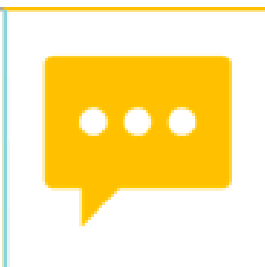
48%
want more help
to make or keep
in contact with
friends



89%
report being able
to use phone or
internet when
wanted



56%
report having a
cell phone or
smartphone



44%
of those who do
not have a cell
phone or
smartphone want
a cell phone or
smartphone