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.
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.

Residence
- 38% parent or relative’s home
- 32% Community-based Group Residential Settings
- 18% own home or apartment
- 7% foster care or host home
- 4% ICF/ID or other institutional setting
- 2% other or don’t know

89% Have ID diagnosis

39% mild
29% moderate
12% severe
8% profound
12% unspecified or unknown

Mental Health Diagnoses (not mutually exclusive)
- 32% mood disorder
- 27% anxiety disorder
- 11% psychotic disorder
- 11% other mental health diagnosis

58% male
43 average age

53% under guardianship
2018-19 In-Person Survey

- **96%** reported having taken part in last service planning meeting
- **81%** understood what was being talked about at last planning meeting
- **94%** reported that they chose or had input in services getting 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