2017 National Health Interview Survey (NHIS) Public Use Data Release

Family Disability Questions (FDB) File Description

Family Disability Questions File (FDB)

Core Content

The Family Disability Questions file contains a set of six core disability questions first developed and used on the American Community Survey (ACS) and subsequently adopted by several federal data collection systems. Originally included on the NHIS as part of a larger effort to develop and test a standard set of disability questions, HHS adopted the question set as a minimum standard for survey questions on disability in 2011.

The six questions in the 2017 Family Disability Questions section (FDB) asked about difficulties hearing, seeing, concentrating, walking, dressing/bathing, and doing errands. These disability questions were initially asked from October 2008 through December 2009 on the Family Disability questionnaire at both the person level (FDB) and the family level (FDA), but the FDA was dropped in 2010. Disability questions were included in the 2011 and 2012 Sample Adult (ADB) and Sample Child (CDB) questionnaires, in addition to the Family Disability questionnaire (FDB), but the ADB and CDB were not included in 2017.

Background

The U. S. Census has a history of including questions about disability to satisfy a variety of stakeholder needs. The 2000 Census of Population and the 2000–2007 questionnaires of the Census Bureau's ACS included six disability questions: a combined sensory (vision and hearing) question; separate mobility, self-care, and cognition questions; and two independent living questions (based on daily activities and working). Before each decennial census, other agencies and researchers join with the Census Bureau to develop survey questions most appropriate for their needs and to determine if existing questions need modification. The user community had expressed some dissatisfaction with the disability measures on the 2000 Census and 2000–2007 ACS.

In response, the Census Bureau and other stakeholder agencies refined the disability questions to bring them into line with recent changes in the definition of disability and the conceptualization of the components of that definition. A work group was formed under the auspices of the Office of Management and Budget and led by NCHS. That work group researched the theoretical approach to the definitions based on the latest national and international ideas about disability, analyzed available data to test their conceptualizations, identified and examined agency mandates for collecting disability data, discussed question content and wording, and sponsored cognitive testing of a new question set. The modified disability questions were subsequently tested in the ACS Content Test of 2006, along with other question additions and modifications (Brault et al., 2007).

The underlying concept behind the choice of questions was to identify the subpopulation that is at a greater risk than the general population of experiencing restrictions in social participation, for example, restrictions in employment, education, or civic life. The objective was thus to measure equalization of opportunities. Four basic domains of functioning (vision, hearing, mobility, and cognitive functioning) were identified that would define the largest component of the population of people with disabilities. These four domains could be used individually or combined in order to assess equalization of opportunities for people with disabilities. Also, two more domains were identified that could be used for monitoring independent living and the need for services: the ability to take care of oneself (self-care, in particular, the ability to bathe and dress oneself), and the ability

to move around the community (independent living, in particular, the ability to visit a doctor's office or go shopping). The new set included separate questions for vision and hearing, refined the mobility question, expanded the cognitive functioning question, continued the inclusion of a self-care question, and improved the question on independent living. The question about work, which had been included in earlier censuses, was not retained.

Technical Notes

A separate, stand-alone public use data file was created for the FDB variables, rather than appending these variables to the 2017 Person File. This file is called the Family Disability Questions File. Families were randomly selected to either receive the FDB questions or for the sample adult to receive the Adult Functioning and Disability (AFD) questions. The FDB questions were asked of approximately all family members. Approximately one-half of the respondents from the 2017 Person File was selected for the FDB questions. Therefore, a different weight (WTFA_FDB) was generated for persons in the FDB File. This weight was designed to produce annual-level estimates calculated based on data included in the files.

In 2017, a question was added to the FDB section to capture prevalence estimates on cognitive disability. This question (PDFCAUSE) asks about causes of difficulty concentrating, remembering, or making decisions. An other-specify option is also available to respondents who give answers that do not fit into the existing categories, and these responses are captured in the variable PDFSPEC.

Analysts wishing to perform analyses of the NHIS disability questions data (e.g., cross-survey comparisons of disability prevalence rates to gain insight into possible survey context effects on estimates of disability; or produce estimates and perform comparisons within key subgroups such as age, sex, and race/ethnicity) will need to merge the FDB file with one or more health data files. Information on merging data files and combining multiple years of data can be found in Appendix IV of the 2017 NHIS Survey Description Document.

Reference

Brault M, Stern S, Raglin D. Evaluation Report Covering Disability. 2006 American Community Survey Content Test Report P.4. U.S. Census Bureau. Available at https://www.census.gov/library/working-papers/2007/acs/2007 Brault 01.html.