

National Survey of Children with Special Health Care Needs (NS-CSHCN)

National Chartbook Indicators

Summary of key differences in measurement: 2001 to 2005-2006

Indicator comparable across survey years

 \bigcirc = Indicator not comparable across survey years

Indicators	Percent of CSHCN meeting indicator		Key differences to consider when comparing results across survey years*
	2001 NS-CSHCN	2005/06 NS-CSHCN	
Child Health			
CSHCN whose conditions affect their activities usually, always or a great deal	23.2% 95% CI: 22.4-24.0	24.0% 95% CI: 23.2-24.7	No changes; same as 2001.
CSHCN with 11 or more days of school absences due to illness	15.8% 95% CI: 15.0-16.6	14.3% 95% CI: 13.7-15.0	Vo changes; same as 2001.
Health Insurance Coverage			
CSHCN without insurance at some point during past year	11.6% 95% CI: 11.0-12.3	8.8% 95% CI: 8.4-9.3	No changes; same as 2001.
CSHCN without insurance at time of survey	5.2% 95% CI: 4.8-5.7	3.5% 95% CI: 3.2-3.8	✓ No changes; same as 2001.
Currently insured CSHCN whose insurance is inadequate	33.8% 95% CI: 32.9-34.7	33.1% 95% CI: 32.3-33.9	Minor changes to the response options for one of the questions; other questions same as 2001.
Access to Care	17.7%	16.1%	Preventive dental care added in 2005/06; other questions same as 2001.
CSHCN with any unmet need for specific health care services	95% CI: 16.9-18.5	95% CI: 15.4-16.7	Wording changes modified the definition of respite
	5.1% 95% CI: 4.7-5.5	4.9% 95% CI: 4.5-5.2	care in 2005/06; other questions same as 2001.
CSHCN with any unmet need for family support services CSHCN needing a referral who have difficulty getting it	21.9% 95% CI: 20.8-23.1	21.1% 95% CI: 19.9-22.4	 New question added in 2005/06 to identify the group of children who qualify as needing referrals.
CSHCN without a usual source of care when sick (or who rely on the emergency room)	9.3% 95% CI: 8.8-9.8	5.7% 95% CI: 5.3-6.1	No changes; same as 2001.
CSHCH without any personal doctor or nurse	11.0% 95% CI: 10.3-11.6	6.5% 95% CI: 6.1-6.9	Question modified in 2005/06 to permit reporting of more than one personal doctor or nurse.

Data Resource Center (DRC) is a project of the Child and Adolescent Health Measurement Initiative at Oregon Health & Science University. DRC is sponsored by the Maternal and Child Health Bureau, Health Resources and Services Administration.

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	2001 NS-CSHCN	2005/06 NS-CSHCN	
Family-Centered Care			
CSHCN without family-centered care	33.2% 95% CI: 32.2-34.1	34.5% 95% CI: 33.6-35.3	New question about availability of interpreter during health care visits added in 2005/06. Nationally, 1.5% of CSHCN needed interpreter services during health care visits.
Impact on Family			
CSHCN whose families pay \$1,000 or more out of pocket in medical expenses per year for child	11.2% 95% CI: 10.6-11.8	20.0% 95% CI: 19.4-20.6	"Co-payments, dental or vision care" newly added to list of expenses for parents to consider as out- of- pocket costs in 2005/06.
CSHCN whose conditions cause financial problems for family	20.9% 95% CI: 20.1-21.7	18.1% 95% CI: 17.5-18.7	✓ No changes; same as 2001.
CSHCN whose families spend 11 or more hours per week providing or coordinated the child's health care	13.5% 95% CI: 12.8-14.2	9.7% 95% CI: 9.2-10.3	S For 2005/06, the wording "providing transportation to appointments" was dropped from list of activities parents are directed to consider as providing health care at home for child.
CSHCN whose conditions cause family members to cut back or stop working	29.8% 95% CI: 29.0-30.7	23.8% 95% CI: 23.1-24.5	Significant differences to question wording and ordering in 2005/06.