

NHANES III Data Documentation

Laboratory Assessment: NHANES III Surplus Sera – Antibody to Pertussis Toxin

Years of Coverage: 1991–1994 (Phase 2)

First Published: July 2005

Last Revised: N/A

Component Description	Pertussis antitoxin testing of stored sera specimens from Phase 2 (1991–1994) of NHANES III was conducted to examine the distribution of immunoglobulin g (IgG) levels against three Bordatella pertussis antigens (pertussis toxin [PT], filamentous hemagglutinin [FHA], and fimbria types 2 and 3 [FIM]) to determine population-based antibody levels for the purpose of establishing diagnostic cut points for a proposed serologic test to pertussis.
Eligible Sample	Participants aged 6–49 years from Phase 2 of NHANES III with stored sera (N = 6,040).
Description of Laboratory Methodology	IgG against PT, FHA, and FIM in serum were measured by ELISA methods (1) at Vanderbilt University Medical Center. Antibody levels were quantitated with respect to a reference serum sample that was calibrated against the U.S. Reference Pertussis Serum, Human Lot 3 (HRP3). The lower limit of quantitation (LLQ) was determined to be 20 EU (ELISA units/ml) (2). The diagnostic cut-off of anti-PT IgG was ≥ 94 EU (2). The cut-off has a diagnostic sensitivity of 80% and a specificity of 93%.
Laboratory Quality Control and Monitoring	See Ref. 2 for details on quality control for these data.
Data Processing and Editing	<p>Data was received after all the antitoxin testing was complete. The data on these results were not edited.</p> <p>Data Access: All data are publicly available.</p> <p>See Ref. 2 to see complete description of the antigens included in the data file.</p>

Analytic Notes

There are four variables in the pertussis antitoxin file:

SEQN: Survey participant identifier

PTPTX: Pertussis toxin

PTPFHA: Pertussis filamentous hemagglutinin

PTPFIM: Pertussis fimbria types 2 and 3

Missing results are due to quality control issues during testing. All antibody levels are reported in EU. These variables are available on Phase 2 participants. Please use “wtpfhx2” examination weights in all data analyses.

References

1. Meade BD, Deforest D, Edwards KM, *et al.* Description and evaluation of serologic assays used in a multicenter trial of acellular pertussis vaccines. *Pediatrics*. 1995;96:570–575.
2. Baughman AL, Bisgard KM, Edwards KM, *et al.* Establishment of diagnostic cutoff points for levels of serum antibodies to pertussis toxin, filamentous hemagglutinin, and fimbriae in adolescents and adults in the United States. *Clin Diagn Lab Immunol*. 2004;11:1045–1053.

Locator Fields

Title: Pertussis antitoxin

Contact Number: 1-866-441-NCHS

Years of Content: 1991–1994

First Published: July 2005

Revised: N/A

Access Constraints: None

Use Constraints: None

Geographic Coverage: National

Subject: Pertussis antitoxin

Record Source: NHANES 2003–2004

Survey Methodology: NHANES 2003–2004 is a stratified multistage probability sample of the civilian non-institutionalized population of the U.S.

Medium: NHANES Web site; SAS transport files