Documentation, Codebook, and Frequencies

RBC Folate and Serum Folate

Laboratory

Survey Years: 2005 to 2006

SAS Transport File: FOLATE_D.XPT



NHANES 2005-2006 Data Documentation

Laboratory Assessment: RBC Folate and Serum Folate (FOLATE_D)

First Published: January 2008 Last Revised: N/A

Component Description

The objectives of this component are: 1) to provide data for monitoring secular trends in measures of nutritional status in the U.S. population; 2) to evaluate the effect of people's habits and behaviors such as physical activity and the use of alcohol, tobacco, and dietary supplements on people's nutritional status; and 3) to evaluate the effect of changes in nutrition and public health policies including welfare reform legislation, food fortification policy, and child nutrition programs on the nutritional status of the U.S. population. These data will be used to estimate deficiencies and toxicities of specific nutrients in the population and subgroups, to provide population reference data, and to estimate the contribution of diet, supplements, and other factors to serum levels of nutrients. Data will be used for research to further define nutrient requirements as well as optimal levels for disease prevention and health promotion.

Eligible Sample

Participants aged 1 year and older who do not meet any of the exclusion criteria are eligible.

Description of Laboratory Methodology

Serum folate is measured by using the Bio-Rad Laboratories "Quantaphase II Folate" radioassay kit. The assay is performed by combining serum or a whole blood hemolysate sample with ¹²⁵I-folate and cyanide. The mixture is boiled to inactivate endogenous folate-binding proteins. The reduced folate and its analogs are stabilized by DTT during the heating. The mixture is cooled and then combined with immobilized affinity-purified porcine intrinsic factor and folate-binding proteins. The addition of these substances adjusts and buffers the pH of the reaction mixture to 9.2. The reaction mixture is then incubated for 1 hour at room temperature.

During incubation, the endogenous and labeled folate competes for the limited number of binding sites on the basis of their relative concentrations. The reaction mixtures are then centrifuged and decanted. Labeled and unlabeled folate, binding to immobilized binding proteins, is concentrated in the bottom of the tube in the form of a pellet. The unbound folate in the supernatant are discarded, and the

radioactivity associated with the pellet is counted. Standard curves are prepared by using the pre-calibrated folate standards in a human serum albumin base. The concentration of the folate in the participant's serum or folate in a participant's whole blood is calculated from the standard curve.

In the erythrocyte folate procedure, the sample is first diluted 1:11 with a solution of 1 g/dL ascorbic acid in water and either incubated for 90 min prior to assay or frozen immediately for later assay. The 90-min incubation or the freeze-thaw is necessary for hemolysis of the red blood cells; either allows the endogenous folate conjugates to hydrolyze the conjugated pterylpolyglutamates prior to assay. The sample is further diluted 1:2 with a protein diluent (human serum albumin), resulting in a matrix similar to that of the standards and serum samples.

There were no changes to the equipment, lab method or lab site from the previous 2 years.

A detailed description of the laboratory method used can be found on the NHANES website.

Laboratory Quality Control and Monitoring

The NHANES quality assurance and quality control (QA/QC) protocols meet the 1988 Clinical Laboratory Improvement Act mandates. Detailed QA/QC instructions are discussed in the NHANES Laboratory/Medical Technologists Procedures Manual (LPM). Read the LABDOC file for detailed QA/QC protocols.

A detailed description of the quality assurance and quality control procedures can be found on the NHANES website.

Data Processing and Editing

Serum specimens are processed, stored, and shipped to the Division of Environmental Health Laboratory Sciences, National Center for Environmental Health, and Centers for Disease Control and Prevention for analysis.

Detailed specimen collection and processing instructions are discussed in the NHANES LPM. Vials are stored under appropriate frozen (–20°C) conditions until they are shipped to National Center for Environmental Health for testing.

Two derived variables were created in this data file. The formula for their derivation is as follows:

The serum folate in ng/mL was converted to nmol/L by multiplying by 2.265.

The RBC folate in ng/mL RBC was converted to nmol/L RBC by multiplying by 2.265.

Detailed instructions on specimen collection and processing can be found on the NHANES website.

Analytic Notes

The analysis of NHANES 2005–2006 laboratory data must be conducted with the key survey design and basic demographic variables. The NHANES 2005–2006 Household Questionnaire Data Files contain demographic data, health indicators, and other related information collected during household interviews. The Household Questionnaire Data Files also contain all survey design variables and sample weights required to analyze these data. The Phlebotomy Examination file includes auxiliary information on duration of fasting, the time of day of the venipuncture, and the conditions precluding venipuncture. The Household Questionnaire and Phlebotomy Exam files may be linked to the laboratory data file using the unique survey participant identifier SEQN.

References

N/A

Locator Fields

Title: RBC Folate and Serum Folate **Contact Number:** 1-866-441-NCHS

Years of Content: 2005–2006 First Published: January 2008

Last Revised: N/A

Access Constraints: None
Use Constraints: None

Geographic Coverage: National

Subject: RBC Folate and Serum Folate **Record Source:** NHANES 2005–2006

Survey Methodology: NHANES 2005–2006 is a stratified multistage probability sample of the civilian

non-institutionalized population of the U.S.

Medium: NHANES Web site; SAS transport files

National Health and Nutrition Examination Survey Codebook for Data Production (2005-2006)

RBC Folate and Serum Folate (FOLATE_D)

January 2008



SEQN	Target				
BEQIV	B(1 Yrs. to 150 Yrs.)				
Hard Edits	SAS Label				
	Respondent sequence number				
English Text: Respondent sequence number.					
English Instructions:					

LBXRBF		Target			
		B(1 Yrs. to 150 Yrs.)			
Hard Edits SAS Label					
		Folate, RBC (ng/mL RBC)			
English Text: Folate,	RBC (ng/m	ng/mL RBC)			
English Instructions:	English Instructions:				
Code or Value	I	Description	Count	Cumulative	Skip to Item
52 to 1840	Ra	nge of Values	8374	8374	
		Missing	1066	9440	

LBDRBFSI		Target			
		B(1 Yrs. to 150 Yrs.)			
Hard Edits	1	SAS Label			
		Folate, RBC(nmol/L RBC)			
English Text: Folate,	RBC(nmol	ol/L RBC)			
English Instructions:	English Instructions:				
Code or Value	I	Description	Count	Cumulative	Skip to Item
117.8 to 4167.6	Ra	nge of Values	8374	8374	
		Missing	1066	9440	

LBXFOL	Target				
LDAI OL	B(1 Yrs. to 150 Yrs.)				
Hard Edits	SAS Label				
	Folate, serum (ng/mL)				
English Text: Folate, serum (ng/mL)					
English Instructions:					

Code or Value	Description	Count	Cumulative	Skip to Item
0.7 to 171	Range of Values	8143	8143	
	Missing	1297	9440	

LBDFOLSI		Target				
EDDI GEGI		B(1 Yrs. to 150 Yrs.)				
Hard Edits		SAS Label				
		Folate, serum (nmol/L)				
English Text: Folate, serum (nmol/L)						
English Instructions:						
Code or Value	Description	Count	Cumulative	Skip to Item		

Code or Value	Description	Count	Cumulative	Skip to Item
1.6 to 387.3	Range of Values	8143	8143	
	Missing	1297	9440	