



National Institute of Standards & Technology

Certificate of Analysis

Standard Reference Material[®] 972a

Vitamin D Metabolites in Frozen Human Serum

This Standard Reference Material (SRM) is intended for use as an accuracy control in the critical evaluation of methods for determining the amount-of-substance concentration of vitamin D metabolites in human serum. This SRM can also be used as a quality assurance tool for assigning values to in-house control materials for these constituents. A unit of SRM 972a consists of four vials (Levels 1 through 4) of frozen serum with different concentration levels of 25-hydroxyvitamin D [25(OH)D] and 24R,25-dihydroxyvitamin D₃ [24R,25(OH)₂D₃]. Measurement of total 25(OH)D concentration in serum, the sum of 25-hydroxyvitamin D₂ [25(OH)D₂] and 25-hydroxyvitamin D₃ [25(OH)D₃], is generally considered a reliable indicator of vitamin D status. The concentration of 3-epi-25-hydroxyvitamin D₃ [3-epi-25(OH)D₃] is generally not included in total 25(OH)D, but this metabolite poses a potential measurement interference for some vitamin D metabolite assays. Measurement of 24R,25(OH)₂D₃ in serum is considered as a catabolism marker and an indicator of kidney disease. Each vial of SRM 972a contains approximately 1 mL of serum.

Each of the four levels of SRM 972a was prepared with a specific target level of 25(OH)D. While some measurement methods might be applicable to each of the four levels of SRM 972a, it is recognized that some methods may not be applicable to some levels. Individual users will need to assess which level or levels best suit their particular needs. Levels 1, 2, and 3 of SRM 972a were prepared from pools of human serum with endogenous concentrations of 25(OH)D. Level 4 was prepared from a pool of human serum that was fortified with 3-epi-25-hydroxyvitamin D₃.

Certified Values: The certified values for 25(OH)D₃, 25(OH)D₂, 3-epi-25(OH)D₃, and 24R,25(OH)₂D₃ are provided in Table 1. A NIST certified value is a value for which NIST has the highest confidence in its accuracy in that all known or suspected sources of bias have been investigated or taken into account [1]. The certified values for 25(OH)D₃, 25(OH)D₂, and 3-epi-25(OH)D₃ are based on the consensus of results from isotope dilution liquid chromatography mass spectrometry (ID-LC-MS) [2] and isotope dilution liquid chromatography tandem mass spectrometry (ID-LC-MS/MS) [3] procedures performed at NIST, and from two different ID-LC-MS/MS procedures performed at the Centers for Disease Control and Prevention (CDC), Atlanta, GA [4,5]. The certified values for 24R,25(OH)₂D₃ are based on the results from a reference measurement procedure using ID-LC-MS/MS performed at NIST [6]. The NIST and CDC ID-LC-MS/MS methods are recognized as higher-order reference measurement procedures by the Joint Committee for Traceability in Laboratory Medicine (JCTLM) [7,8].

Reference Values: Reference values for 25(OH)D₂ and 3-epi-25(OH)D₃ are provided in Table 2. Reference values are noncertified values that are the best estimate of the true values based on available data; however, the values do not meet the NIST criteria for certification and are provided with associated uncertainties that may reflect only measurement precision, may not include all sources of uncertainty, or may reflect a lack of sufficient statistical agreement among multiple analytical methods [1]. The reference values for 25(OH)D₂ and 3-epi-25(OH)D₃ are based on the agreement of results from ID-LC-MS and ID-LC-MS/MS procedures performed at NIST and from ID-LC-MS/MS results provided by the CDC.

Certified and Reference Values for total 25(OH)D are provided in Tables 3 and 4 primarily for assays that are suitable for use with SRM 972a but do not measure the vitamin D metabolites separately.

Expiration of Certification: The certification of **SRM 972a** is valid, within the measurement uncertainty specified, until **31 January 2023**, provided the SRM is handled and stored in accordance with the instructions given in this certificate (see "Instructions for Storage and Use"). The certification is nullified if the SRM is damaged, contaminated, or otherwise modified.

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Certificate Issue Date: 31 March 2021
Certificate Revision History on Last Page

Steven J. Choquette, Director
Office of Reference Materials

Maintenance of SRM Certificate: NIST will monitor this SRM over the period of its certification. If substantive technical changes occur that affect the certification before the expiration of this certificate, NIST will notify the purchaser. Registration (see attached sheet or register online) will facilitate notification.

Support for the development of SRM 972a was provided in part by the National Institutes of Health Office of Dietary Supplements (NIH-ODS). Technical consultation was provided by C.T. Sempos, J.M. Betz and P.M. Coates of NIH-ODS.

Overall direction and coordination of the analytical measurements leading to the certification of this SRM were performed by J. Camara of the NIST Chemical Sciences Division, and S.S.-C. Tai, formerly of NIST.

Acquisition of the material was performed by K.W. Phinney of the NIST Biomolecular Division and K.E. Sharpless of the NIST Office of Special Programs. Certification measurements were performed by M. Bedner and G. Hahn, of the NIST Chemical Sciences Division, S.S.-C. Tai, and R.S.C. Chia, a guest scientist at NIST from the Health Sciences Authority of Singapore. Certification measurements were also performed by S. Momin, E.M. Mineva, K. Maw, S. Encisco, M. Chaudhary-Webb, and R.L. Schleicher at the CDC. Additional measurements in support of the development of SRM 972a were performed by M.A. Nelson, B.E. Lang, and L.C. Sander of the NIST Chemical Sciences Division, and M.M. Schantz, L.T. Sniegowski, formerly of NIST.

Statistical analysis was provided by B. Toman and J.H. Yen of the NIST Statistical Engineering Division.

Support aspects involved in the issuance of this SRM were coordinated through the NIST Office of Reference Materials.

NOTICE AND WARNINGS TO USERS

Warning: SRM 972a IS INTENDED FOR LABORATORY RESEARCH USE ONLY. THIS IS A HUMAN SOURCE MATERIAL. HANDLE PRODUCT AS A BIOHAZARDOUS MATERIAL CAPABLE OF TRANSMITTING INFECTIOUS DISEASE. The supplier of the serum has reported that each donor unit of serum used in the preparation of this product has been tested by a FDA-approved method and found non-reactive/negative for hepatitis B surface antigen (HbsAg), human immunodeficiency (HIV) 1 and 2 antibodies, and hepatitis C virus (HCV). However, no known test method can offer complete assurance that hepatitis B virus, hepatitis C virus, HIV, or other infectious agents are absent from this material. Accordingly, this human blood-based product should be handled at the Biosafety Level 2 or higher as recommended for any POTENTIALLY INFECTIOUS HUMAN SERUM OR BLOOD SPECIMEN in the Centers for Disease Control and Prevention/National Institutes of Health Manual [9].

This SRM was developed after an appropriate human subjects research determination by NIST.

INSTRUCTIONS FOR STORAGE AND USE

Storage: Until required for use, SRM 972a should be stored in the dark at a temperature between $-20\text{ }^{\circ}\text{C}$ and $-80\text{ }^{\circ}\text{C}$.

Use: SRM 972a is provided as a set of four vials of frozen serum. The vial (or vials) to be used should be allowed to thaw at room temperature for at least 30 min under subdued light. The contents of the vial should then be gently mixed prior to removal of a test portion for analysis. Precautions should be taken to avoid exposure to strong UV light and direct sunlight.

SOURCE, PREPARATION, AND ANALYSIS⁽¹⁾

Source and Preparation: SRM 972a was prepared by Solomon Park Research Laboratories (Kirkland, WA). Four serum pools were prepared. The naturally occurring concentrations of vitamin D metabolites in the human serum pools used to prepare Levels 1, 2, and 3 have not been modified. Level 4 is a human serum pool enriched with 3-epi-25(OH)D₃.

Analysis: Value assignment of the concentrations of 25(OH)D₃, 25(OH)D₂, and 3-epi-25(OH)D₃ in SRM 972a are based on the combination of results provided from two analytical methods at NIST (ID-LC-MS and ID-LC-MS/MS)

⁽¹⁾ Certain commercial equipment, instruments or materials are identified in this certificate to adequately specify the experimental procedure. Such identification does not imply recommendation or endorsement by the National Institute of Standards and Technology, nor does it imply that the materials or equipment identified are necessarily the best available for the purpose.

and from two ID-LC-MS/MS analytical procedures at CDC. Value assignment of the concentrations of 24R,25(OH)₂D₃ are based on the results from ID-LC-MS/MS at NIST.

Measurement of 25(OH)D₃, 25(OH)D₂, and 3-epi-25(OH)D₃ by ID-LC-MS (NIST): Serum (450 mg) and an internal standard solution containing ²H₆-25(OH)D₃, ²H₃-25(OH)D₂, and ²H₃-3-epi-25(OH)D₃ were combined in glass tubes, proteins were precipitated, and the metabolites were extracted into hexane twice. The hexane phases were combined and evaporated to dryness at 40 °C under nitrogen. The residues were reconstituted with methanol and vortex-mixed. Extracts were analyzed by using LC-MS with (1) an Ascentis Express F5 pentafluorophenylpropyl column (Supelco, Bellefonte, PA) and (2) a Zorbax SB-CN cyanopropyl stationary phase column (Agilent Technologies, Palo Alto, CA). Analyses on the pentafluorophenylpropyl column were performed under isocratic conditions with a mobile phase of 26 % water and 74 % methanol at a flow rate of 0.8 mL/min. All solvent compositions represent volume fractions in percent. The column temperature was maintained at 15 °C. A step gradient to 100 % methanol was incorporated into the method at the end of the run to elute retained matrix components. Analyses on the cyanopropyl column were performed under isocratic conditions with a mobile phase of 33 % water and 67 % methanol (Levels 1, 2, and 4 of SRM 972a) or 34 % water and 66 % methanol (Level 3) at a flow rate of 1.0 mL/min. A step gradient to 100 % methanol was incorporated into the method at the end of the run to elute retained matrix constituents. The column temperature was maintained at 45 °C.

Atmospheric pressure chemical ionization (APCI) with positive polarity was used for both chromatographic methods. The [M – H₂O + H]⁺ ions were monitored and used for quantification of all species. The ions monitored included *m/z* 383 for 25(OH)D₃ and for 3-epi-25(OH)D₃; *m/z* 386 for ²H₃-3-epi-25(OH)D₃; *m/z* 389 for ²H₆-25(OH)D₃; *m/z* 395 for 25(OH)D₂; and *m/z* 398 for ²H₃-25(OH)D₂.

Measurement of 25(OH)D₃, 25(OH)D₂, and 3-epi-25(OH)D₃ by ID-LC-MS/MS (NIST): Serum (1.0 g to 2.0 g) was spiked with an appropriate internal standard solution [²H₆-25(OH)D₃, ²H₃-25(OH)D₂, or ²H₃-3-epi-25(OH)D₃]. After equilibration at room temperature for 1 h, the pH of each sample was adjusted to pH 9.8 ± 0.2 with carbonate buffer. Analytes were extracted twice from the serum matrix with a mixture of hexane and ethyl acetate. The combined extracts were dried under nitrogen at 45 °C, and the residues were reconstituted with methanol for LC-MS/MS analysis. Extracts were analyzed using either an Ascentis Express F5 or a Zorbax SB-CN column under isocratic conditions with water:methanol mobile phases. APCI in the positive-ion mode and multiple reaction monitoring (MRM) mode were used. The following transitions were monitored: *m/z* 401 → *m/z* 383 for 25(OH)D₃ and 3-epi-25(OH)D₃; *m/z* 407 → *m/z* 389 for ²H₆-25(OH)D₃; *m/z* 404 → *m/z* 386 for ²H₃-3-epi-25(OH)D₃; *m/z* 413 → *m/z* 395 for 25(OH)D₂; and *m/z* 416 → *m/z* 398 for ²H₃-25(OH)D₂.

Measurement of 25(OH)D₃, 25(OH)D₂, and 3-epi-25(OH)D₃ by ID-LC-MS/MS (CDC): Samples of SRM 972a (100 µL) were spiked with the following isotopically-labeled internal standards: ²H₆-25(OH)D₃, ²H₃-25(OH)D₂, and ²H₃-epi-25(OH)D₃. Each serum sample was extracted using 1.5 mL hexane, and the supernatant was collected, dried under nitrogen at 25 °C, and reconstituted in 69 % methanol in water (volume fractions). Analytes were eluted from the extract (isocratic mobile phase same composition as the diluent) at a flow rate of 0.4 mL/min on a Hypersil GOLD pentafluorophenyl column (Thermo Fisher Scientific, Waltham, MA) at 28 °C and detected using APCI in positive-ion mode. Two transitions per vitamin D metabolite along with one transition per internal standard were monitored: 25(OH)D₃: *m/z* 383 → *m/z* 365 and *m/z* 383 → *m/z* 105; ²H₆-25(OH)D₃: *m/z* 389 → *m/z* 371; epi-25(OH)D₃: *m/z* 383 → *m/z* 365 and *m/z* 383 → *m/z* 105; ²H₃-epi-25(OH)D₃: *m/z* 386 → *m/z* 368; 25(OH)D₂: *m/z* 395 → *m/z* 377 and *m/z* 395 → *m/z* 209; ²H₃-25(OH)D₂: *m/z* 398 → *m/z* 380. Analytes were quantitated using six-point linear calibration curves traceable to SRM 2972 or SRM 2972a 25-Hydroxyvitamin D2 and D3 Calibration Solutions, and internal standards were used to correct for recovery.

Measurement of 25(OH)D₃ and 25(OH)D₂, by ID-LC-MS/MS (CDC Reference Measurement Procedure): Serum SRM 972a (500 µL) was accurately measured and spiked (gravimetrically) with pre-determined amounts of isotopically-labeled internal standards (ISTD), ²H₆-25(OH)D₃ and ²H₃-25(OH)D₂, to get an approximate 1:1 mass ratio of analyte to ISTD in glass tubes. The pH of each sample was adjusted to approximately pH 9.8 with aqueous sodium carbonate (0.1 g/mL), and the metabolites were extracted with hexanes twice. The combined extracts were dried under vacuum at 45 °C, reconstituted in 75 % methanol in water, vortex mixed, and filtered (0.45 µm PVDF, 96-well plate). Analytes were analyzed on LC-MS/MS with gradient elution (the ratio of methanol to water was gradually increased from 65 % to 80 % over 20 minutes) at a flow rate of 0.45 mL/min on a Acquity HSS PFP column (1.8 µm x 150 mm x 3.0 mm) (Waters Corp, Milford, MA) at 22 °C. APCI in the positive-ion mode and selected reaction monitoring (SRM) mode were used for detection with the following transitions: *m/z* 383 → *m/z* 365 (quantitation) and *m/z* 383 → *m/z* 105 (confirmation) for 25(OH)D₃; *m/z* 389 → *m/z* 371 for ²H₆-25(OH)D₃; *m/z* 395 → *m/z* 377 (quantitation) and *m/z* 395 → *m/z* 209 (confirmation) for 25(OH)D₂; and *m/z* 398 → *m/z* 380 for ²H₃-25(OH)D₂.

Measurement of 24R,25(OH)₂D₃ by ID-LC-MS/MS (NIST): Serum (1.5 g to 2.0 g) was spiked with an internal standard solution containing ²H₆-24R,25(OH)₂D₃. After equilibration at room temperature for 1 h, the pH of each sample was adjusted to pH 9.8 ± 0.2 with carbonate buffer. The 24R,25(OH)₂D₃ was extracted twice from the serum matrix with a mixture of hexane and ethyl acetate. The combined extracts were dried under nitrogen at 45 °C, and the residues were reconstituted with methanol for LC-MS/MS analysis. Extracts were analyzed using an Ascentis Express C₁₈ column under isocratic conditions with a water:methanol mobile phase. APCI in the positive-ion mode and multiple reaction monitoring (MRM) mode were used. The following transitions were monitored: *m/z* 417 → *m/z* 381 for 24R,25(OH)₂D₃ and *m/z* 423 → *m/z* 387 for ²H₆-24R,25(OH)₂D₃.

Homogeneity Analysis: The homogeneity assessment was made at the time the certification analyses were performed. A stratified sampling plan was devised to test for homogeneity across the lot of vials. There was no apparent trend in the data when plotted against the sequence in which the vials were prepared, with the exception of 3-epi-25(OH)D₃ in Level 4. An additional component of uncertainty related to possible inhomogeneity has been included in the expanded uncertainty for this analyte in Level 4.

Certified Values for 25(OH)D₂, 25(OH)D₃, 3-epi-25(OH)D₃, and 24R,25(OH)₂D₃: The 25(OH)D₃ and 3-epi-25(OH)D₃ values are weighted means of the results from analyses at NIST using ID-LC-MS and ID-LC-MS/MS and from CDC using one ID-LC-MS/MS procedure. The 25(OH)D₂ values are consensus means obtained using the Bayesian hierarchical model in the NIST Consensus Builder [10] on the results from analyses at NIST using ID-LC-MS and ID-LC-MS/MS and from CDC using two ID-LC-MS/MS procedures. The 24R,25(OH)₂D₃ values are the results from analyses at NIST using ID-LC-MS/MS. The uncertainty provided with each certified value is an expanded uncertainty about the consensus mean to cover the measurand with approximately 95 % confidence; it incorporates Type B uncertainty components related to the analyses and expresses both the observed difference between the results from the methods and their respective uncertainties, consistent with the Guide to the Expression of Uncertainty in Measurement and with its Supplement 1 [11–14]. The expanded uncertainties are calculated as $U = ku_c$, where u_c is the combined uncertainty and k is a coverage factor corresponding to approximately 95 % confidence for each analyte [11]. For the certified values shown in Table 1, $k = 2$. The measurands are the total concentrations of the analytes listed in Table 1. Metrological traceability is to the International System of Units (SI) derived units of mass fraction, expressed as nanograms per gram; mass concentration, expressed as nanograms per milliliter; and amount-of-substance concentration, expressed as nanomoles per liter.

Table 1. Certified Values for 25(OH)D₂, 25(OH)D₃, 3-epi-25(OH)D₃, and 24R,25(OH)₂D₃ in SRM 972a

	ng/g		ng/mL ^(a)		nmol/L ^(b)	
Level 1						
25-hydroxyvitamin D ₃	28.1	± 1.1	28.8	± 1.1	71.8	± 2.7
3-epi-25-hydroxyvitamin D ₃	1.77	± 0.10	1.81	± 0.10	4.5	± 0.2
24R,25-dihydroxyvitamin D ₃	2.60	± 0.10	2.66	± 0.10	6.38	± 0.23
Level 2						
25-hydroxyvitamin D ₂	0.79	± 0.08	0.81	± 0.06	2.0	± 0.2
25-hydroxyvitamin D ₃	17.7	± 0.4	18.1	± 0.4	45.1	± 1.0
3-epi-25-hydroxyvitamin D ₃	1.25	± 0.09	1.28	± 0.09	3.2	± 0.2
24R,25-dihydroxyvitamin D ₃	1.38	± 0.05	1.41	± 0.05	3.39	± 0.12
Level 3						
25-hydroxyvitamin D ₂	12.9	± 0.3	13.2	± 0.3	32.0	± 0.8
25-hydroxyvitamin D ₃	19.4	± 0.4	19.8	± 0.4	49.5	± 1.1
24R,25-dihydroxyvitamin D ₃	1.58	± 0.06	1.62	± 0.06	3.88	± 0.13
Level 4						
25-hydroxyvitamin D ₃	28.8	± 0.9	29.4	± 0.9	73.4	± 2.3
3-epi-25-hydroxyvitamin D ₃	25.4	± 2.1	26.0	± 2.2	64.8	± 5.4
24R,25-dihydroxyvitamin D ₃	2.58	± 0.09	2.64	± 0.09	6.32	± 0.22

^(a) Mass concentration levels were calculated from mass fractions using measured serum densities: Level 1, 1.02326 g/mL; Level 2, 1.02196 g/mL; Level 3, 1.02294 g/mL; and Level 4, 1.02295 g/mL.

^(b) Molar concentration levels were calculated from mass concentration levels using the relative molecular masses. The relative molecular masses are 412.65 g/mol for 25(OH)D₂, 400.64 g/mol for 25(OH)D₃ and 3-epi-25(OH)D₃ and 416.64 g/mol for 24R,25(OH)₂D₃. The equivalent conversion factors are 2.4234 for 25(OH)D₂, 2.4960 for 25(OH)D₃ and 3-epi-25(OH)D₃ and 2.4002 for 24R,25(OH)₂D₃.

Reference Values for 25(OH)D₂ and 3-epi-25(OH)D₃: Values are weighted means of the results from analyses at NIST using ID-LC-MS and ID-LC-MS/MS and at CDC using ID-LC-MS/MS. The measurands are the concentrations of analytes listed in Table 2 as determined by the indicated methods. Metrological traceability is to the ID-LC-MS/MS method, with values expressed in SI derived units of mass fraction, expressed as nanograms per gram; mass concentration, expressed as nanograms per milliliter; and amount-of-substance concentration, expressed as nanomoles per liter. The uncertainty provided with each reference value is an expanded uncertainty about the weighted mean to cover the measurand with approximately 95 % confidence; it incorporates Type B uncertainty components related to the analyses and expresses both the observed difference between the results from the methods and their respective uncertainties, consistent with the ISO/JCGM Guide and with its Supplement 1 [11–14]. The expanded uncertainty is calculated as $U = ku_c$, where u_c is the combined uncertainty and k is a coverage factor corresponding to approximately 95 % confidence for each analyte [11]. For the reference values shown in Table 2, $k = 2$.

Table 2. Reference Values for 25(OH)D₂ and 3-epi-25(OH)D₃ in SRM 972a

	ng/g	ng/mL ^(a)	nmol/L ^(b)
Level 1			
25-hydroxyvitamin D ₂	0.52 ± 0.06	0.54 ± 0.06	1.3 ± 0.2
Level 3			
3-epi-25-hydroxyvitamin D ₃	1.14 ± 0.14	1.17 ± 0.14	2.9 ± 0.4
Level 4			
25-hydroxyvitamin D ₂	0.54 ± 0.10	0.55 ± 0.10	1.3 ± 0.2

^(a) Mass concentration levels were calculated from mass fractions using measured serum densities: Level 1, 1.02326 g/mL; Level 3, 1.02294 g/mL; and Level 4, 1.02295 g/mL.

^(b) Molar concentration levels were calculated from mass concentration levels using the relative molecular masses. The relative molecular masses are 412.65 g/mol for 25(OH)D₂ and 400.64 g/mol for 25(OH)D₃ and 3-epi-25(OH)D₃. The equivalent conversion factors are 2.4234 for 25(OH)D₂ and 2.4960 for 25(OH)D₃ and 3-epi-25(OH)D₃.

Certified and Reference Values for Total 25(OH)D: Vitamin D levels in serum are typically reported as the total of 25(OH)D₃ and 25(OH)D₂. The values for total 25(OH)D, as the sum of the individual values for 25(OH)D₃ and 25(OH)D₂, are shown in Table 3 for certified values and Table 4 for reference values. The uncertainty provided with each value is an expanded uncertainty about the total 25(OH)D that covers the measurands with approximately 95 % confidence; it incorporates Type B uncertainty components related to the analyses and their respective uncertainties of the two analytes, consistent with the ISO/JCGM Guide and its Supplement 1 [11–14]. The expanded uncertainty is calculated as $U = ku_c$, where u_c is the combined uncertainty and k is a coverage factor corresponding to approximately 95 % confidence for the analytes [11]. For the values shown in Table 3 and Table 4, $k = 2$.

Table 3. Certified Values for Total 25(OH)D in SRM 972a^(a)

	ng/g	ng/mL ^(b)
Level 2		
Total 25(OH)D	18.5 ± 0.4	18.9 ± 0.4
Level 3		
Total 25(OH)D	32.3 ± 0.5	33.0 ± 0.5

^(a) Certified values for total 25(OH)D are based on the combination of certified values for both 25(OH)D₂ and 25(OH)D₃. The measurands are the total concentrations of the analytes listed. Metrological traceability is to the SI derived units of mass fraction, expressed as nanograms per gram; and mass concentration, expressed as nanograms per milliliter.

^(b) Mass concentration levels were calculated from mass fractions using measured serum densities: Level 2, 1.02196 g/mL; and Level 3, 1.02294 g/mL.

Table 4. Reference Values for Total 25(OH)D in SRM 972a^(a)

	ng/g		ng/mL ^(b)	
Level 1				
Total 25(OH)D	28.7	± 1.1	29.3	± 1.1
Level 4				
Total 25(OH)D	29.3	± 0.9	30.0	± 0.9

^(a) Reference values for total 25(OH)D are based on the combination of a reference value for 25(OH)D₂ and a certified value for 25(OH)D₃. The measurands are the amounts of substance listed, as determined by the indicated method. Metrological traceability is to the ID-LC-MS/MS method, with values expressed in SI derived units of mass fraction expressed as nanograms per gram; and mass concentration, expressed as nanograms per milliliter.

^(b) Mass concentration levels were calculated from mass fractions using measured serum densities: Level 1, 1.02326 g/mL; and Level 4, 1.02295 g/mL.

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