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NIST Micronutrients Measurement Quality Assurance Program Summer 2012 Comparability Studies

Results for Round Robin LXXII Fat-Soluble Vitamins and Carotenoids in Human Serum and Round Robin 37 Ascorbic Acid in Human Serum

> David L. Duewer Jeanice B. Thomas

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March, 2013



U.S. Department of Commerce Rebecca Blank, Acting Secretary

National Institute of Standards and Technology Patrick D. Gallagher, Under Secretary of Commerce for Standards and Technology and Director (This page intentionally blank)

Abstract

The National Institute of Standards and Technology coordinates the Micronutrients Measurement Quality Assurance Program (MMQAP) for laboratories that measure fat- and water-soluble vitamins and carotenoids in human serum and plasma. This report describes the design of and results for the Summer 2012 MMQAP measurement comparability improvement studies: 1) Round Robin LXXII Fat-Soluble Vitamins and Carotenoids in Human Serum and 2) Round Robin 37 Total Ascorbic Acid in Human Serum. The materials for both studies were shipped to participants in June 2012; participants were requested to provide their measurement results by September 28, 2012.

Keywords

Human Serum Retinol, α-Tocopherol, γ-Tocopherol, Total and *Trans*-β-Carotene Total Ascorbic Acid

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Introduction

Beginning in 1988, the National Institute of Standards and Technology (NIST) has coordinated the Micronutrients Measurement Quality Assurance Program (MMQAP) for laboratories that measure fat- and water-soluble vitamins and carotenoids in human serum and plasma. The MMQAP provides participants with measurement comparability assessment through use of interlaboratory studies, Standard Reference Materials (SRMs) and control materials, and methods development and validation. Serum-based samples with assigned values for the target analytes (retinol, alpha-tocopherol, gamma/beta-tocopherol, *trans*- and total beta-carotene, and total ascorbic acid) and performance-evaluation standards are distributed by NIST to laboratories for analysis.

Participants use the methodology of their choice to determine analyte content in the control and study materials. Participants provide their data to NIST, where it is compiled and evaluated for trueness relative to the NIST value, within-laboratory precision, and concordance within the participant community. NIST provides the participants with a technical summary report concerning their performance for each exercise and suggestions for methods development and refinement. Participants who have concerns regarding their laboratory's performance are encouraged to consult with the MMQAP coordinators.

All MMQAP interlaboratory studies consist of individual units of batch-prepared samples that are distributed to each participant. For historical reasons these studies are referred to as "Round Robins". The MMQAP program and the nature of its studies are described elsewhere. [1,2]

Round Robin LXXII: Fat-Soluble Vitamins and Carotenoids in Human Serum

Participants in the MMQAP Fat-Soluble Vitamins and Carotenoids in Human Serum Round Robin LXXII comparability study (hereafter referred to as RR72) received one lyophilized and four liquid-frozen human serum test samples for analysis. Unless multiple vials were previously requested, participants received one vial of each serum. These sera were shipped on dry ice to participants in June 2012. The communication materials included in the sample shipment are provided in Appendix A.

Participants are requested to report values for all fat-soluble vitamin-related analytes that are of interest to their organizations. Not all participants report values for the target analytes, and many participants report values for non-target analytes.

The final report delivered to every participant in RR72 consists of three documents:

- A cover letter for the current study, a brief description of the other two documents, and a discussion of our analysis of the overall results that may be of broad interest. This cover letter is reproduced as Appendix B.
- The "All-Lab Report" that lists all of the reported measurement results, a number of consensus statistics for analytes reported by more than one participant, and the mean median and pooled SD from any prior distributions of the serum. This report also provides a numerical "score card" for each participant's measurement comparability for the more commonly reported analytes. This All-Lab Report is reproduced as Appendix C.

• An "Individualized Report" that graphically analyzes each participant's results for all analytes reported by at least five participants. This report also provides a graphical summary of their measurement comparability. The graphical tools used in the Individualized Report are described in detail elsewhere [3]. An example Individualized Report is reproduced as Appendix D.

Round Robin 37: Vitamin C in Human Serum

Participants in the MMQAP Vitamin C in Human Serum Round Robin 37 comparability study (hereafter referred to as RR37) received four frozen serum test samples, one frozen control serum, and a solid ascorbic acid control material for analysis. Unless multiple vials were previously requested, participants received one vial of each material. These sample materials were shipped on dry ice to participants in June 2012. The communication materials included in the sample shipment are provided in Appendix E.

The test and control serum materials were prepared by adding equal volumes of 10 % metaphosphoric acid (MPA) to human serum that had been spiked with ascorbic acid. While these samples contain some dehydroascorbic acid, its content is variable. Therefore, the participants report only total ascorbic acid (TAA, ascorbic acid plus dehydroascorbic acid). Participants are also encouraged to prepare calibration solutions from the supplied solid control to enable calibrating their serum measurements to the same reference standard.

The final report delivered to every participant in RR37 consists of three documents:

- A cover letter for the current study, a brief description of the other two documents, and a discussion of our analysis of overall results that may be of broad interest. This cover letter is reproduced as Appendix F.
- The "All-Lab Report" that summarizes all of the reported measurement results and provides several consensus statistics. This All-Lab Report is reproduced as Appendix G.
- An "Individualized Report" that graphically analyzes each participant's results for TAA, including a graphical summary of their measurement comparability. The graphical tools used in the Individualized Report are described in detail elsewhere [3]. An example Individualized Report is reproduced as Appendix H.

References

- Duewer DL, Brown Thomas J, Kline MC, MacCrehan WA, Schaffer R, Sharpless KE, May WE, Crowell JA. NIST/NCI Micronutrients Measurement Quality Assurance Program: Measurement Repeatabilities and Reproducibilities for Fat-Soluble Vitamin-Related Compounds in Human Sera. Anal Chem 1997;69(7):1406-1413.
- 2 Margolis SA, Duewer DL. Measurement Of Ascorbic Acid in Human Plasma and Serum: Stability, Intralaboratory Repeatability, and Interlaboratory Reproducibility. Clin Chem 1996;42(8):1257-1262.
- 3 Duewer DL, Kline MC, Sharpless KE, Brown Thomas J, Gary KT, Sowell AL. Micronutrients Measurement Quality Assurance Program: Helping Participants Use Interlaboratory Comparison Exercise Results to Improve Their Long-Term Measurement Performance. Anal Chem 1999;71(9):1870-1878.

Appendix A. Shipping Package Inserts for RR72

The following three items were included in each package shipped to an RR72 participant:

- Cover letter
- Datasheet
- Packing List and Shipment Receipt Confirmation Form

The cover letter and datasheet were enclosed in a sealed waterproof bag along with the samples themselves. The packing list was placed at the top of the shipping box, between the cardboard covering and the foam insulation.



UNITED STATES DEPARTMENT OF COMMERCE National Institute of Standards and Technology Gaithersburg, Maryland 20899-

June 25, 2012

Dear Colleague:

Enclosed are samples for the second fat-soluble vitamins and carotenoids in serum study Round Robin LXXII (RR72) for the 2012 NIST Micronutrients Measurement Quality Assurance Program. The set of samples (Sera 387- 391) consists of one vial of lyophilized serum and one vial each of four liquid-frozen serum samples for analysis along with a form for reporting your results. These samples should be stored in the dark at or below –20 °C upon receipt.

When reporting your results, please submit one value for each analyte for a given serum sample. If a value obtained is below your limit of quantification, please indicate this result on the form by using NQ (*Not Quantified*). Results are due to NIST by **September 28, 2012**. Results received more than <u>two weeks</u> after the due date may not be included in the summary report for RR72. The feedback report concerning the study will be distributed in November 2012.

Samples should be allowed to stand at room temperature under subdued light until thawed. We recommend that sample mixing be facilitated with 3 to 5 min agitation in an ultrasonic bath or at least 15 min at room temperature with intermittent swirling. (CAUTION: Vigorous shaking will cause foaming and possibly interfere with accurate measurement. The rubber stopper contains phthalate esters that may leach into the sample upon intermittent contact of the liquid sample with the stopper. These esters absorb strongly in the UV region and elute near retinol in most LC systems creating analytical problems.) Water should not be added to the liquid-frozen samples.

For consistency, we request that laboratories use the following absorptivities (dL/g • cm): retinol, 1843 at 325 nm (ethanol); retinyl palmitate, 975 at 325 nm (ethanol); α -tocopherol, 75.8 at 292 nm (ethanol); γ -tocopherol, 91.4 at 298 nm (ethanol); α -carotene, 2800 at 444 nm (hexane); β -carotene, 2560 at 450 nm (ethanol), 2592 at 452 nm (hexane); and lycopene, 3450 at 472 nm (hexane).

Please report your results for RR72 by e-mail to david.duewer@nist.gov or fax to 301-977-0685. If you have questions or comments regarding this study, please call me at (301) 975-3120 or e-mail me at jbthomas@nist.gov.

Sincerely.

Jeanice Brown Thomas Program Coordinator/Research Chemist Analytical Chemistry Division Material Measurement Laboratory



Date:

Round Robin LXXII: Human Sera

NIST Micronutrients Measurement Quality Assurance Program

Analyte	387	388	389	390	391	Units*
total retinol						
trans-retinol						
retinyl palmitate						
α -tocopherol						
γ/β-tocopherol						
δ-tocopherol						
total β-carotene						
trans-β-carotene						
total cis-β-carotene						
total α -carotene						
total lycopene						
trans-lycopene						
total β -cryptoxanthin						
total α -cryptoxanthin						
total lutein						
total zeaxanthin						
total lutein&zeaxanthin						
total coenzyme Q10						
ubiquinol (QH ₂)						
ubiquinone (Qox)						
phylloquinone (K1)						
25-hydroxyvitamin D						
Phytoene						
Phytofluene						
,						

* we prefer $\mu g/mL$

Were the samples frozen when received? Yes | No

Comments:

Fat-Soluble Vitamins Round Robin LXXII NIST Micronutrients Measurement Quality Assurance Program

Packing List and Shipment Receipt Confirmation Form

This box contains: one vial each of the following five FSV M²QAP sera

Serum	Form	Reconstitute?	Vial/Cap
#387	Lyophilized	Yes	2 mL amber, red cap
#388	Liquid frozen	No	2 mL amber, green cap
#389	Liquid frozen	No	2 mL amber, red cap
#390	Liquid frozen	No	2 mL amber, gold cap
#391	Liquid frozen	No	2 mL amber, green cap

Please 1) Open the pack immediately

- 2) Check that it contains all of the above samples
- 3) Check if the vials are intact
- 4) Store the sera at -20 °C or below until analysis
- 5) Complete the following information
- Fax the completed form to us at 301-977-0685 (or email requested information to david.duewer@nist.gov)

1) Date this shipment arrived: _____

2) Are all five sera vials intact? Yes | No If "No", which one(s) were damaged?

- 3) Was there any dry-ice left in cooler? Yes | No
- 4) Did the samples arrive frozen? Yes | No
- 5) At what temperature are you storing the serum samples? _____ °C
- 6) When do you anticipate analyzing these samples? _____

Your prompt return of this information is appreciated.

The M²QAP Gang

Appendix B. Final Report for RR72

The following two pages are the final report as provided to all participants:

- Cover letter.
- An information sheet that:
 - o describes the contents of the "All-Lab" report,
 - o describes the content of the "Individualized" report,
 - o describes the nature of the test samples and details their previous distributions, if any, and
 - o summarizes aspects of the study that we believe may be of interest to the participants.



November 30, 2012

Dear Colleague:

Enclosed is the summary report of the results for round robin LXXII (RR72) of the 2012 NIST Micronutrients Measurement Quality Assurance Program (M²QAP) for the fat-soluble vitamins and carotenoids in human serum. Included in this report are: 1) a summary of data and measurement comparability scores for all laboratories, 2) a detailed graphical analysis of your results; and 3) a graphical summary of your measurement comparability. RR72 (Sera 387 - 391) consisted of one vial of lyophilized serum and one vial each of four liquid-frozen serum samples. Details regarding the samples can be found in the enclosed report.

Your overall measurement comparability is summarized in the "Score Card" summary, page 6 of the All Lab Report. Combined results rated 1 to 3 are within 1 to 3 standard deviations of the assigned value, respectively; those rated 4 are >3 standard deviations from the assigned value. Similar information is presented graphically in the "target plots" that are the last page of your Individualized Report. If you have concerns regarding your laboratory's performance, please contact us for consultation.

Samples for the first set of fat-soluble vitamins and carotenoids in serum interlaboratory exercise (RR73) of the 2013 M²QAP will be shipped **starting January 22, 2013**. Please contact us immediately if this schedule is problematic for your laboratory.

If you have questions or concerns regarding this report, please contact David Duewer at 301-975-3935; e-mail: david.duewer@nist.gov or me at 301-975-3120; e-mail: jbthomas@nist.gov; or fax: 301-977-0685.

Sincerely,

Jeanice Brown Thomas, M.B.A. Research Chemist Chemical Sciences Division Material Measurement Laboratory

Enclosures

Cc: L.C. Sander

David L. Duewer, Ph.D. Research Chemometrician Chemical Sciences Division Material Measurement Laboratory



The NIST MMQAP Round Robin LXXII (RR72) report consists of:

Page	All-Lab Report
1-4	A listing of all results and statistics for analytes reported by more than one participant.
5	A listing of the analytes reported by only one participant and a legend for the list of results and statistics.
6	The text Comparability Summary ("Score Card") of measurement performance.
Page	Individualized Report
1	Your values, the number of labs reporting values, and our assigned values.
2 to n	"Four Plot" summaries of your current and past measurement performance, one page for each analyte you report that is also reported by at least eight other participants.

n+1 The graphical Comparability Summary (target plot) of measurement performance.

Samples. Five samples were distributed in RR72.

Serum	Description	Prior Distributions
387	Lyophilized, native serum prepared in 1999. Serum #380 is the liquid-frozen partner of this sample.	#270:RR49-3/01, #276:RR50-9/01, #367:RR68-9/10, #377:RR70-9/11
	Liquid-frozen, native serum prepared in 1999. Serum #377 is the lyophilized partner of this sample.	#267:RR49-9/00, #274:RR50-9/01, #368:RR68-9/10, #380:RR70-9/11
389	Liquid-frozen, native, multi-donor, prepared in 2009. This is Level I of SRM 968e.	#357:RR66-9/09, #365:RR67-3/10, #375:RR69-3/11
390	Liquid-frozen, native, single-donor serum prepared in 2011. This material was designed to have low contents of α - and β -carotene and lycopene but a high content of β -cryptoxanthin.	#383:RR71
391	Liquid-frozen, native, multi-donor, prepared in 2008	#356:RR65-3/09, #360:RR66-9/09

Results

- 1) <u>SRM 968e Stability</u>. There was no significant change in the median level or measurement variability of any measurand in the SRM 968e Level I material.
- 2) <u>Lyophilized vs Liquid-frozen Stability</u>. After 13 years, there is no change in the median level or the measurement variability in any analyte in the lyophilized #387 material or the #388 liquid-frozen material prepared from the same serum pool. The analyte levels in the lyophilized material are slightly lower than in the liquid-frozen material due to reconstituting *with* 1.0 ml water rather than reconstituting *to* a total volume of 1.0 ml.

Appendix C. All-Lab Report for RR72

The following six pages are the "All-Lab Report" as provided to all participants, with two exceptions:

- the participant identifiers (Lab) have been altered.
- the order in which the participant results are listed has been altered.

The data summary in the "All-Lab Report" has been altered to ensure confidentiality of identification codes assigned to laboratories.

ol, μg/mL γ/β-Tocopherol	5 2 3 0 2 4 1 9 0 1 8 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3 3	7.34 10.50 2.21 2.30 1.79 1.72		6.50 10.00	6.40	6.90 10.80	6.45 10.42 2.32 2.45 1.92	6.49 10.06 2.21 2.29 1.85 1.72	6.59	6.90	7.00 7.40 10.60		6.16	8.00 11	7.13 6.70 9.90		8.49 12.59 2.23 2.27 1.77 1.85	6.96 9.11 2.30 2.19 1.66 1.75	7.25 10.53 3.94 2.08 1.69	6.85		6.96 10.89 2.60 2.78 2.16 1.96	6.66 9.92 2.05 2.11 1.69 1.67	7.13 7.28 11.49 2.05 2.17 1.90 1.78 2.18		5.50 7.20 8.60		8.10 12.10	7.86 11.53 2.68 2.76 2.19 2.04 2.	25 25 13 13 13 13 13	6.16 8.60 2.04 2.08 1.66 1.55	6.90 10.42 2.23 2.29 1.79 1.75	2.78 2.19	0.20 0.17 0.16	7 9 8	33 29 32 21 22 19 16 19	10.56 2.21 2.33 1.81 1.	0.67 0.80 0.21 0.22 0.16 0.23	_	0.71 0.59 0.81 0.23 0.23 0.19 0.19 0.24
α-Tocop	300 6 10	6.58		6.70		6.70	6.20	6.01	6.30	6.50	6.90	6.02		8.00	6.40 6.70 7		7.29	6.00	6.60	6.60	5.58	6.47	61 5.81	6.38 6.69 7		5.10 6.30 5		7.50	6.87	25	5.58	6.50	7.30 8.00 8	0.62 0.46 (10 7	38 39	5.97 6.35 (51 0.62	6.10 6.50 6	0.62 0.56 (
Retinyl Palmitate, µg/mL	200 309 309 390 2 0 0 20 0 0 16 0 0 16 C	0.020 0.009 0.008					0.017 0.022 0.014 0.015 0.044												0.020 0.019 0.005 0.013 0.053				0.020 0.022 0.012 0.013 0.048							5 5 5	0.019 0.005 0.008	0.020 0.012 0.013	0.016	0.002 0.002 0.005 0.003 0.006	9 41 23	10 10 8 5 9	0.021 0.024 0.010 0.019 0.045	0.011 0.005 0.012		0.011 0.011 0.011 0.011 0.015
g/mL																0.374 0.629				0.444 0.737													0.444 0.737			0	0.692	0.097		
trans-Retinol, µg/ml																0.281				0.358 0.											0.281	0.319	0.358			7	0.34			
trans-R	200															4 0.542				2 0.676										~		~	2 0.676			2 12	\sim	0 0.048		
200	4	1 9	22	9	0	0	80	4	55	0	0	0	00	0		9 0.53	9	2		13 0.632	6	0	9	2	0	6		90	9			15 0.583		9	5	30	0.61		5	69
100	_ د د				0 0.630									0 0.790		74		8 0.722				8 0.790				2 0.826			0.7					4 0.040	7		0 0.740			17 0.059
, µg/mL	1 0 487												0.370			м									0	2 0.542			0.4				§ 0.542		10	2 31	7 0.460		1 0.464	0.037
Total Retinol, µg/mL	209															74				0.358					0	0.392			0				3 0.406	0.019		32	0.357		0.361	0:030
Total	0 711	0.664	0.672	0.687	0.580	0.760	0.700	0.681	0.636	0.630	0.620	0.690	0.610	0.740		/4		0.731	0.650	0.678	0.670	0.699	0.672	0.720	0.690	0.783		0.626	0.657	27	0.580			0.049	7	37	0.665			0.054
200	0 664	0.651	0.621	0.668	0.550	0.600	0.658	0.650	0.660	0.570	0.620	0.660	0.530		0.660	/4		0.732	0.630	0.638	0.531	0.691	0.654	0.665	0.660	0.668		0.607	0.644	27	0.530	0.654	0.732	0.024	4	35	0.636	0.048	0.654	0.052
	Lau FSV-RA	FSV-BB	FSV-BC	FSV-BD	FSV-BE	FSV-BF	FSV-BG	FSV-BJ	FSV-BK	FSV-BL	FSV-BM	FSV-BN	FSV-BO	FSV-BQ	FSV-BR	FSV-BS	FSV-BU	FSV-BV	FSV-BW	FSV-CC	FSV-CE	FSV-CG	FSV-CI	FSV-CZ	FSV-DD	FSV-DV	FSV-EE	FSV-FK	FSV-FZ	z	Min	Median	Max	SD	S	Npast	Medianpast	SDpast	NAV	NAU

All Lab Report

Page 1/6

Results
Laboratory
LXXII
Robin
Round

ug/mL		0.010 0.046 0.010 0.046		0.012 0.053	nq 0.036			0.019 0.065 0.010 0.042			0.017 0.041			0.015 0.063						11 12	0.010 0.036			0.0			0.012 0.046 0.004 0.007	0.014 0.049
Total α-Carotene, μg/mL				0.010 0.	ı bu			0.016 0.			0.007 0.			0.009 0.	0.013 0.					11	0.005 0.				47	16	0.009 0. 0.004 0.	0.007 0.
al α-Ca	388	0.020 0.006 0.018 0.005		0.019 0	bu			0.037 0 0.017 0			0.016 0			0.023 0	0.026 0					11	0.016 0				67 7		0.021 0 0.006 0	0.020 0
Tot	387	0.016 (0.019)))))))))))))))))))))))))))))))))))		0.017 (bu			0.039 (0.036)			0.022 (0.025 (11					୫ ୪		0.019 0.006	0.022 (
,mL	391	0.016 0.015												0.017						n	0.015		0.017		c	٥	0.014 0.002	0.016
Total cis-β-Carotene, μg/mL	390	0.001 0.003												ра						2	0.001	0.002	0.003			C		
-Carote	389	0.007 0.006												na						2	0.00		0.007		L	ŋ	0.005	
al cis-β	388	0.019 0.022 0.019 0.019												0.026 0.025						e	0.019	0.022	0.025			x	0.024 0.007	0.022
Tot																				۳ ر			0.026				0.025 0.007	0.019
٦L	391	2 0.262 0.230		_	_	_	_	_	_	0.111	_	_	_	0.416 0.444 0.082 0.040 0.252	_	_	_	_	_	4	0.111			0.0	_		0.247 0.012	0.241
trans-β-Carotene, μg/mL	390	3 0.042) 0.041								0.607 0.435 0.106 0.029 0.111				0.040						4	0.029	3 0.041		0.00			3 0.044 5 0.004	0.083 0.041
-Carote		5 0.083 5 0.080								5 0.106				4 0.082						4		0 0.083		0.00			1 0.083 8 0.006	
trans-β	7 388	.0 0.475 :4 0.455								7 0.43				6 0.44						4	6 0.435			8 0.015			6 0.471 7 0.038	2 0.450
	387	8 0.440 0 0 0.424			4			م ۲			<u> </u>	+ 00				0				20				0 0.018			4 0.436 4 0.037	4 0.432
	391	0.278 0.290	0.340		0.254			0.259		ΛI	0.228			0.336		0.220				15	0.2			0.0			0.274 0.034	0.274
e, µg/mL	390	0.043 0.056	0.050	0.042	0.054			0.047		≥0.029	0.040	0.039		0.063	0.050	0.068				15	0.037	0.050	0.068	0.010		18	0.047 0.009	0.050
arotene	389	060.0 0.096	0.110 0.075	0.089	0.094			0.086		≥0.106	0.081	0.082		0.087	0.099	0.072				15	0.072	0.088	0.114	0.010		77	0.091 0.014	0.088
Total β-Caroten	388	0.496 0.496	0.580 0.593	0.541	0.495			0.536			0.403	0.509		0.465 0.469	0.559	0.407				15	0.403	0.496	0.593	0.059	2 6	17	0.512 0.067	0.496
F	387	0.459 0.462	0.620	0.514	0.457			0.514 0.473			0.406	0.505		0.808 0.442	0.564	0.382				15	0.382	0.482	0.808	0.047	D 0	202	0.484 0.063	0.482
	391	0.271 0.265												0.179						ო	0.17	0.265	0.271		L	Ω	0.261 0.026	0.265
hg/mL	390	0.101 0.110 0.094 0.075 0.271 0.109 0.117 0.100 0.059 0.265												0.188 0.189 0.164 0.255 0.179						n	0.059	0.075	0.255		•	4		
ō-Tocopherol, µg/mL	389	0.094 0.100												0.164						n	0.094	0.100	0.164		L	Ω	0.110 0.025	0.109 0.117 0.100 0.075
ð-Toco	388	0.110 0.117												0.189						e		0.117	0.189			٥	0.109 0.101	0.117
	387		-																	с С	0.101	0.109	0.188	_		0	0.101	
	Lab	FSV-BA FSV-BB FSV-BD	FSV-BD FSV-BE FSV-BE	FSV-BG	FSV-BJ FSV-RK	FSV-BL	FSV-BM	FSV-BN FSV-BO	FSV-BQ FSV-BR	FSV-BS	FSV-BU	FSV-BW	FSV-CC	FSV-CG	FSV-CI	FSV-CZ	FSV-DU FSV-DV	FSV-EE	FSV-FK		Min	Median	Max	02.5	کر :	Npast	Medianpast 0.101 0.109 0.110 0.063 SDpast 0.040 0.101 0.025 0.043	NAV

٦L	391	0.029 0.016																												2	0.016	0.023	0.029			œ	0.020	0.009		
g/	390	0.027 0 0.019 0																												2	0.019 0		0.027 0			4	025 0	0.013 0		
toxanth	389	0.022 0																												2	0.016 0	0.019 0	0.022 0			4	018 0	003		
a-Crypt	388	0.033 0																												2	0.023 0.016	0.028 0	0.033 0			LC.	027 0	005 0		
Total		0.023 0																													0.022 0	0.023 C	0.023 0			LC.	0.023 0.027 0.018 0.025	0.006 0.005 0.003 0.013		
L	391	0.076 0				.068	0.046				0.055	000.			0.048	0.070	0.055				0.085	0										0.058 (0.085 (0.015	26	17			0.058	0.015
in, µg/r		182 C				0.186 0	0.107 0				0.164 0	5			0.082 0	0.189 0	0.188 0															0.173 0	0.201 0		13	14	163 0	0.048		0.036 0
oxanth	389	0.057 C				0.056 0	nq C				0.052 0	010			0.073 0	0.071 0	0.044 0				0.68.0								,	თ	0.044 0.082	0.056 0	0.073 0	0.015 0	26	17	050 0	0.011 0		0.015 0
Q	388	0.060 0.057 0.182 0.048 0.046 0.153				0.056 0	bu				0.052 0				0.064 0	0.064 0	0.044 0				0.67 0									თ	0.044 0	0.056 0	0.067 0	0.012 0		22	0.050 0.050 0.163	0.010 0		0.013 0
Total	387	0.050 0.060 0.057 0.182 0.045 0.048 0.046 0.153				0.053 0.056	bu				0.049 0	0+0-0			0.080 0	0.060 0	0.044 0				0.063 0.067 0.068 0.201	2000										0.050 0	0.080 0	0.006 0	12	2		0.011 0		0.012 0
	391	0.183 0.156 0				0.194 (_				0	0															0.183 (0.205 (œ			L	0.033
		0.080 0				0.086 0									0.043 0						086 0	, , , , , , , , , , , , , , , , , , ,							1		0.043	0.080 0	0.086 0	0.009 0	£	y		0.019	0.080 0	0.014 0
ωĴ	389					0.116 0									0.088 0						121 0								1		0.088 0	0.113 0	0.121 0	0.011 0	10	œ	116 0	0.018		0.020 0
ns-Lyco	388	135 C													.065 0						144 0	-								ß	0.065 0	0.135 C	0.152 0	0.014 0	10	10	138 0	0.018	135 0	0.024 0
tra	387	0.123 0.135 0.113 0.114 0.125 0.099				0.138 0.152									0.082 0.065 0.088 0.043 0.061						0 134 0 144 0 121 0 086 0 205										0.082 0	0.123 0	0.138 0	0.016 0	13	10	0.124 0.138 0.116	0.023 0		0.021 0
						_	0.415				0.385	007.				0.349	0.351	0.413			0 390											0.351 (0.415 (0.058 (16				0.078 (
	390	0.151 0.318 0.164 0.303				0.190 0	0.200 0				0.195 0					0.177 0					0 185 0	2								Ξ	2	2	8	2	5	ст С			0.177 0	0.045 0
cene,						.222 0					0.230 0	5 +			27	194	216	0.216 0			1236 0									-1	0.174 C	0.216 C	4 0.270 0.236 0.20	0.021 0	10	17	0.226 0.171	0.032 0	216	052
5	388).205 C				0.245 0).270 C				0.267 0				0.186 0	0.214 0.7	0.214 0	0.257 0			0 240 0	2								;	0.175 C	0.218 C	0.270 0	0.040 0	18	22				0.053 0.
	387	0.188 0.205 0.202 0.199 0.218 0.199				0.237 C	0.238 0.270 0.225				0.254 0				0.252 0	0.205 0	0.222 0	0.274 0			0 230 0 240 0 236	,								;	Min 0.171 0.175).230 C	0.274 0	0.036 0	15	21	224 0	0.032 0		0.055 0
L			FSV-BC		FSV-BE			F0V-BK	FSV-BL	FSV-BM	FSV-BN 0		FSV-BQ		FSV-BS 0	FSV-BU 0	FSV-BV 0	FSV-BW 0		ESV-CE			FSV-CZ			י ר י ר			Z-7-72	z	Min	Median 0.230 0.218	Max 0	SD 0	S	Nnast	Mediannast 0.224 0.233	SDpast 0		NAU 0
	Lab	ES.	ίς Γ	οú	ΣΩ	FS'	S C	D D	S I	-S-	ίς Π	5	FS	N.	ĘS	FS	FS	FSV	FS	с Ц	о (С - Ц	р Ц -	S ST) č	ρί	δİ	ה ר ו	Ϋ́	Σ Γ			ž					Media	S		

Phylloquinone (K1), ng/mL	389 390 391			3.305 0.051 0.202 2.021												0.263 0.091 0.776						2 2 2	0.263 0.091 0.776	0.447 0.177 1.399	0.631 0.262 2.021			0 0 0		
Phylloquinc	387 388			2.980 3.302												1.433 1.429 0.263						2	1.433 1.429	2.210 2.397	2.986 3.365 (0		
Coenzyme Q10, µg/mL	387 388 389 390 391		č	0.330 0.31 0.300 1.16 0.86		0.760 0.86 0.910 1.04 0.95				0.553 0.47 0.768 0.94 0.58		0.720 0.67 0.840 0.92 0.91		0.780 0.97 0.910 1.37 0.81		0.83 0.840	0.881 1.14			0.823 0.68 1.436 0.88 0.67		8 8 8 8 8	0.768	0.85 0.891 1.07	1.207 1.28 1.436 1.37 1.08	0.22 0.052 0.16 0	10 26 6 15 17	10 8 9	0.814 0.80 0.857 1.08 0.92	
Total Lutein&Zeaxanthin, µg/mL	387 388 389 390 391	0.121 0.121 0.097 0.113 0.100 0.170 0.175 0.147 0.173 0.160			0.123 0.131 0.104 0.110 0.100				0.128 0.159 0.185	0.120 0.129 0.110 0.120 0.103	0.191 0.170 0.206 0.115 0.104	0.139 0.113	0.124 0.127 0.109 0.146 0.076		0146 0156 0149 0172 0136	0.098 0.092 0.104						10 10 10 10 10	0.098 0.092 0.104	0.127 0.130 0.125 0.118 0.100	0.191 0.175 0.206 0.185 0.160	0.018 0.035 0.016 0.0	8 14 28 13 10	21 18 14	0.122 0.127 0.107 0.131 0.097	140.0 010.0 470.0
Total Zeaxanthin, μg/mL	387 388 389 390 391	0.054 0.055 0.050 0.066 0.053								0.027 0.029 0.015 0.015 0.019	0.065 0.060 0.053 0.057 0.053					0.026 0.032 0.026 0.031 0.018						4 4 4 4	0.026 0.029 0.015 0.015 0.018	0.041 0.044 0.038 0.044 0.036	0.065 0.060 0.053 0.066 0.053	0.020 0.020 0.026 0.0	51 45 53 59 71	0	0.034 0.036 0.034 0.029	
Total Lutein, µg/mL	Lab 387 388 389 390 391	FSV-BA FSV-BB 0.115 0.120 0.098 0.107 0.106	FSV-BC FSV-BD	FSV-BE FSV-BF		FSV-BJ 0.078 0.086 0.080 0.089 0.081	SV-BN ESV-BI	ESV-BM		0.093 0.100 0.095 0.105 0.084	0.126 0.110 0.153 0.058 0.051	FSV-BU	FSV-BV	FSV-BW		0.069 0.066 0.066 0.073 0.054	FSV-CZ	-SV-DD	-SV-DV	FSV-EE	FSV-FK	N 5 5 5 5 5	0.066 0.058 0.051	Median 0.093 0.100 0.095 0.089 0.081 0	Max 0.126 0.120 0.153 0.107 0.106	0.033 0.021 0.022 0.024 0.038	CV 36 21 23 27 46		Medianpast 0.086 0.088 0.072 0.097 0.060 0	0.010 0.040 0.010

Analytes Reported By One Laboratory

Analyte	Lab	387	388	389	390	391
25-hydroxyvitamin D	FSV-BN	0.0021	0.0040	0.0054	0.0063	0.0047
Phytoene	FSV-BS	nd	nd	nd	0.089	0.081
Phytofluene	FSV-BS	0.123	0.119	0.132	0.110	0.116

Term	Legend
N	Number of (non-NIST) quantitative values reported for this analyte
Min	Minimum (non-NIST) quantitative value reported
Median	Median (non-NIST) quantitative value reported
Max	Maximum (non-NIST) quantitative value reported
SD	Adjusted median absolute deviation from the median of the non-NIST results
CV	Coefficient of Variation for (non-NIST) results: 100*SD/Median
Npast	Mean of N(s) from past RR(s)
Medianpast	
	Pooled SD from past RR(s)
ODpast	
NAV	NIST Assigned Value
	= (Median + NIST)/2 for analytes reported by NIST
	= Median for analytes reported by \geq 5 labs but not NIST
NAU	
14/10	S is the maximum of (0.05*NAV, SD, SD _{past} , eSD) and S _{btw} is the standard
	deviation between Median and NIST. The expected long-term SD, eSD,
	is defined in: Duewer et al., Anal Chem 1997;69(7):1406-1413.
nd	Not detected (i.e., no detectable peak for analyte)
ng	Detected but not quantitatively determined
×	Concentration greater than or equal to x
italics	Not explicitly reported but calculated by NIST from reported values

Lab	TR	аT	g/bT	bC	aC	TLy	TbX	TLu	L&Z
FSV-BA	1	1	1	1	1	1	1		1
FSV-BB	1	1	1	1	1	1	1	1	2
FSV-BC	1								
FSV-BD	1	1							
FSV-BE	2	1	1	2					
FSV-BF	2	1		2					
FSV-BG	1	1	1	1	1	1	1		1
FSV-BJ	1	1	1	1		1	2	1	
FSV-BK	1	1							
FSV-BL	2	1							
FSV-BM	1	1							
FSV-BN	1	1		1	2	1	1		2
FSV-BO	3	1	1	2	1	1	1	1	1
FSV-BQ	1	2							
FSV-BR	1	1							
FSV-BS	3			3	4	2	2	2	2
FSV-BU	2	3	1	2	1	1	1		1
FSV-BV	2	2	1	1	1	1	1		1
FSV-BW	1	1	4	1	1	1			
FSV-CC	1	1		-					
FSV-CE	2	2		3			-		
FSV-CG	1	1	2	1	1	1	2	-	2
FSV-CI	1	1	1	1	2			2	2
FSV-CZ	1	1	1	2					
FSV-DD	1	_							
FSV-DV	2	2							
FSV-FK	1	2	•						
FSV-FZ	1	2	3	10			4.0		10
n	28	25	13	16	11	11	10	5	10
	TR	аT	g/bT	bC	aC	TLy	TbX	TLu	L&Z
% 1	68	72	77	56	73	91	70	60	50
% 2	25	24	8	31	18	9	30	40	50
%3	7	4	8	13	0	0	0	0	0
% 4	0	0	8	0	9	0	0	0	0
/ U 1	•	•	~	•	•	•	•	~	v

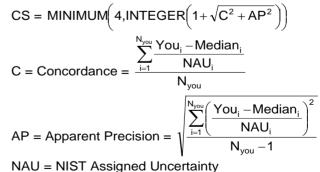
Comparability Summary

Label	Definition
Lab	Participant code
TR	Total Retinol
aT	α-Tocopherol
g/bT	γ/β-Tocopherol
bC	Total β-Carotene
tbC	trans-β-Carotene
aC	Total α-Carotene
TLy	Total Lycopene
TbX	Total β-Cryptoxanthin
TLu	Total Lutein
ΤZ	Total Zeaxanthin
L&Z	Total Lutein & Zeaxanthin
n	number of participants providing quantitative data
% 1	Percent of $CS = 1$ (within 1 SD of medians)
% 2	Percent of CS = 2 (within 2 SD of medians)
% 3	Percent of $CS = 3$ (within 3 SD of medians)
% 4	Percent of CS = 4 (3 or more SD from medians)
	· · · · · · · · · · · · · · · · · · ·

"Comparability Score"

The Comparability Score (CS) summarizes your measurement performance for a given analyte relative to the consensus medians in this study. CS is the average distance (in units of standard deviation) of your measurement performance characteristics from the consensus performance. CS is calculated when the number of quantitative values you reported, N_{you} , is at least two and at least six participants reported quantitative values for the analyte.

We define CS as follows:



For further details, please see

Duewer DL, Kline MC, Sharpless KE, Brown Thomas J, Gary KT. Micronutrients Measurement Quality Assurance Program: Helping participants use interlaboratory comparison exercise results to improve their long-term measurement performance. Anal Chem 1999;71(9):1870-8.

Appendix D. Representative Individualized Report for RR72

Each participant in RR72 received an "Individualized Report" reflecting their reported results. Each report included a detailed analysis for analytes that were assayed by at least five participants. The following analytes met this criterion in RR72:

- Total Retinol
- Retinyl Palmitate
- α-Tocopherol
- γ/β -Tocopherol
- Total β-Carotene
- Total α-Carotene
- Total Lycopene
- *trans*-Lycopene
- Total β-Cryptoxanthin
- Total Lutein
- Total Lutein & Zeaxanthin
- Coenzyme Q10

The following twelve pages are the "Individualized Report" for the analytes evaluated by participant FSV-BA.

Please check our records against your records. Send corrections and/or updates to...

Micronutrients Measurement Quality Assurance Program

National Institute of Standards and Technology

100 Bureau Drive Stop 8392 Gaithersburg, MD 20899-8392 USA

Individualized Report

Individualized Round Robin LXXII Report: FSV-BA

Summary

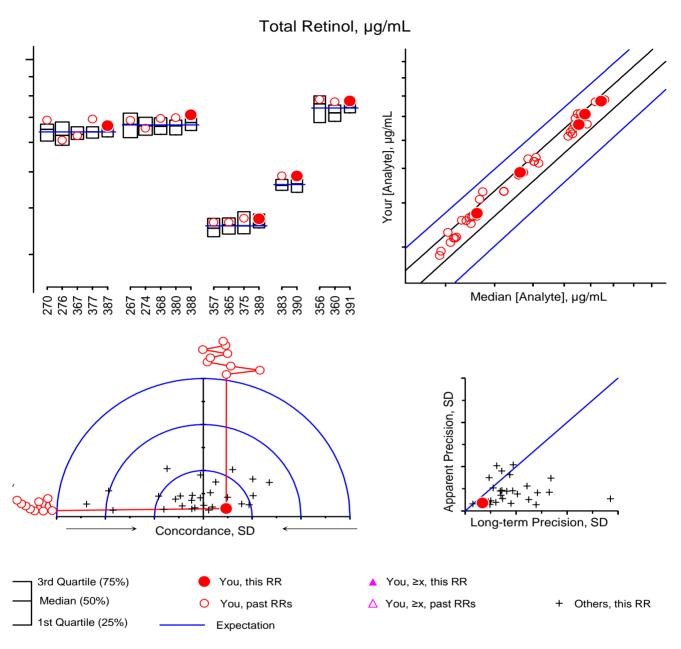
				J		5									
	Ser	Serum 387	~	Ser	Serum 388		Ser	Serum 389		Ser	Serum 390	_	Seru	Serum 391	
Analyte	NoV	NAV n	c	You	NAV n	c	You	NAV n	c	You	NAV n	c	You	NAV	c
Total Retinol	0.664	0.654	27	0.711		27	0.374	0.361	27	0.361 27 0.487		27		0.745	26
Retinyl Palmitate	0.02	0.02	5	0.0	0.0	S	0.0	0.0	S	5 0.02 0		S	0.04	0.04	S
a-Tocopherol	5.89	6.10	25	6.19	6.50 25 (25	6.66	6.90	25	6.72	6.90	25		10.42	25
γ/β-Tocopherol	2.300	2.235	13	2.438	2.290	13	1.901	1.794	13	1.831	1.750	13	2.561	2.350	13
ō-Tocopherol	0.101	0.109	ო	0.110	0.117	ო	0.094	0.100	ო	0.075	0.075	ო	0.271	0.265	ო
Total β-Carotene	0.459	0.482	15	0.496	0.496	15	60.	0.088	15	0.043	050.050	15	0.278	0.274	15
trans-β-Carotene	0.440	0.432	4	0.475	0.450 4 0	4	.08	0.083	4	0.042	0.041	4	0.262	0.241	4
Total cis-β-Carotene	0.019	0.019	ო	0.022	0.022	ო	00.		2	0.001		2	0.016	0.016	ო
Total α-Carotene	0.016	0.022	5	0.020	0.020	1	0.006	0.007	1	0.010	0.014	;	0.046	0.049	12
Total Lycopene	0.188	0.230	5	0.205	0.218 11	;-		0.216	7	0.151	0.177	7	0.318	0.351	1
trans-Lycopene	0.123	0.123	5	0.135	0.135 5	2	0.113	0.113	ß	0.080	0.080	S	0.183	0.183	S
Total β-Cryptoxanthin	0.050	0.050	ი	0.060	0.056	б	0.057	0.056	თ	0.182	0.173	10	0.076	0.058	10
Total α-Cryptoxanthin	0.023		2	0.033		2	0.022		2	0.027		2	0.029		2
Total Lutein&Zeaxanthin	0.121	0.127	10	0.121	0.130	10	0.097	0.125	10	0.113	0.118	10	0.100	0.100	10

You : Your reported values for the listed analytes (micrograms/milliliter)

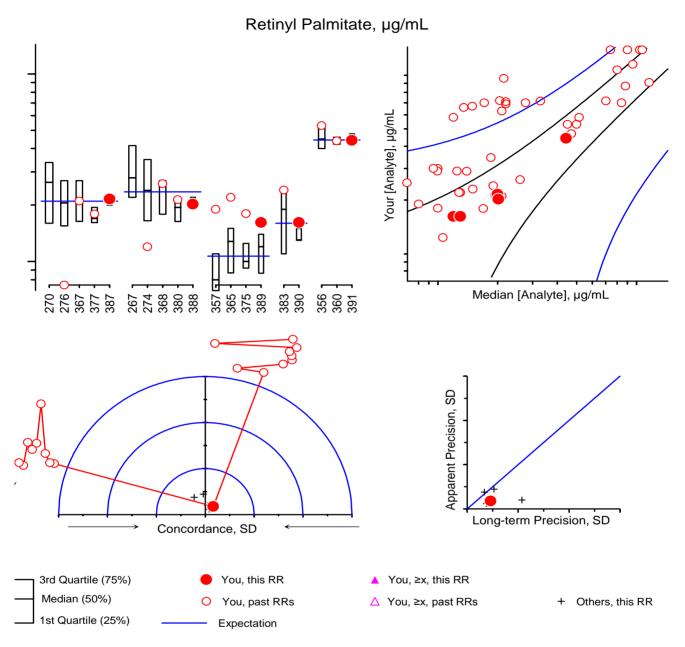
NAV : NIST Assigned Values, here equal to this RR's median

n: Number of non-NIST laboratories reporting quantitative values for this analyte in this serum

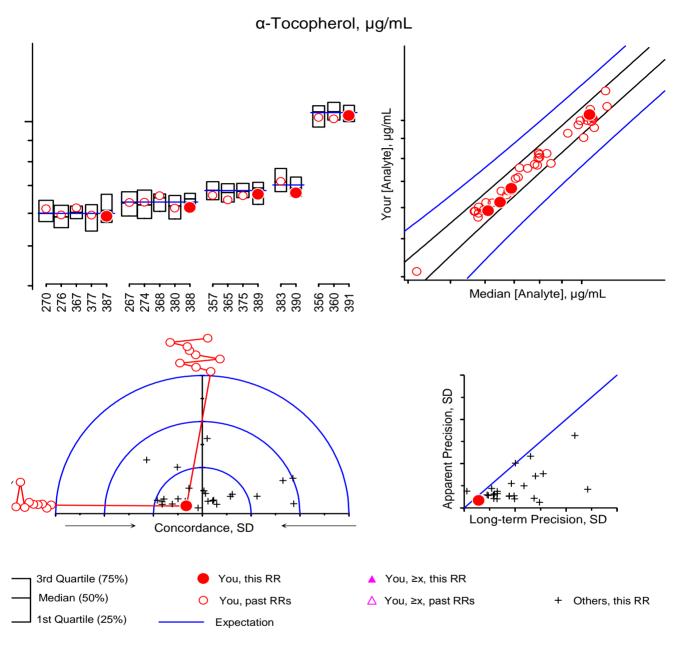
Set 1 of 33



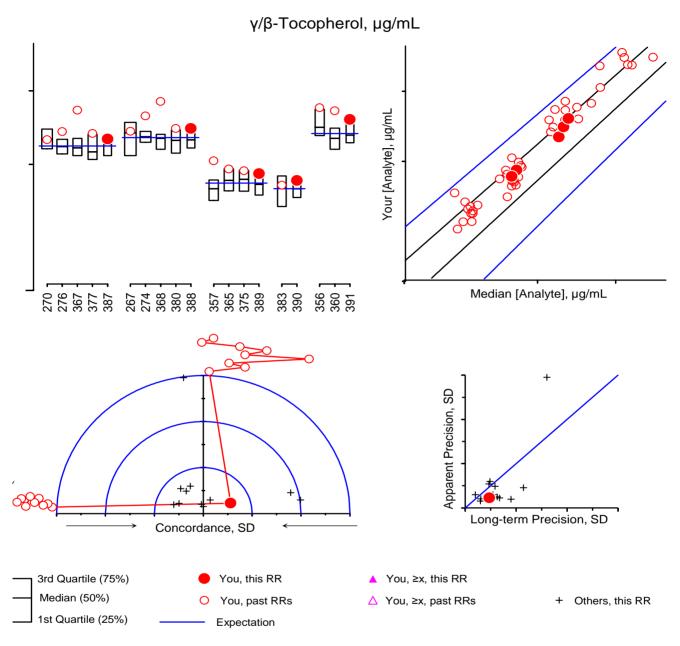
<u>erum</u>	Comments	<u>History</u>
387	Lyophilized, same native pool as #388	49#270, 50#276, 68#367, 70#377
388	Fresh-frozen, same native pool as #387	49#267, 50#274, 68#368, 70#380
389	Fresh-frozen, native, multi-donor: SRM 968e I	66#357, 67#365,69#375
390	Fresh-frozen, native, single-donor	71#383
391	Fresh-frozen, native, multi-donor	65#356, 66#360



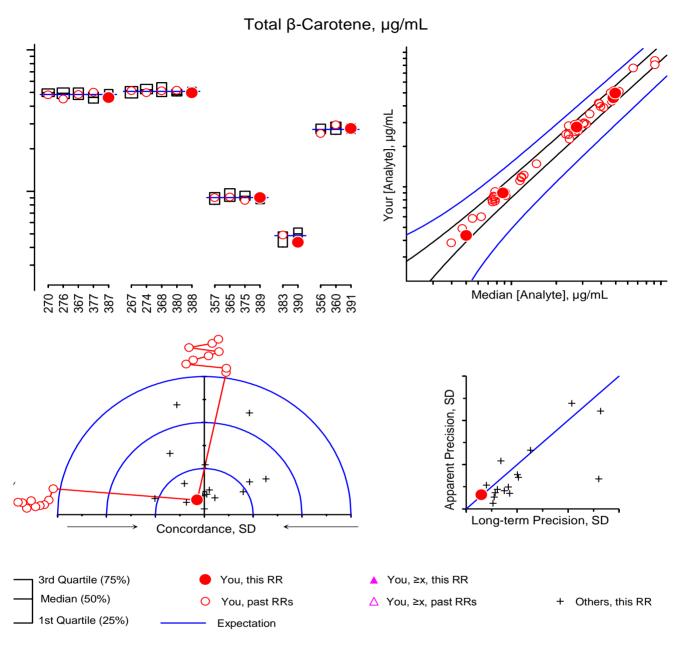
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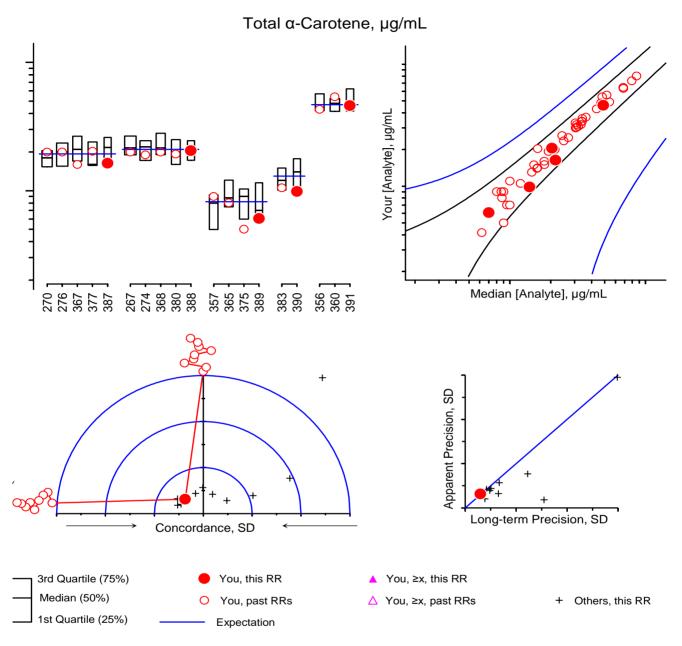
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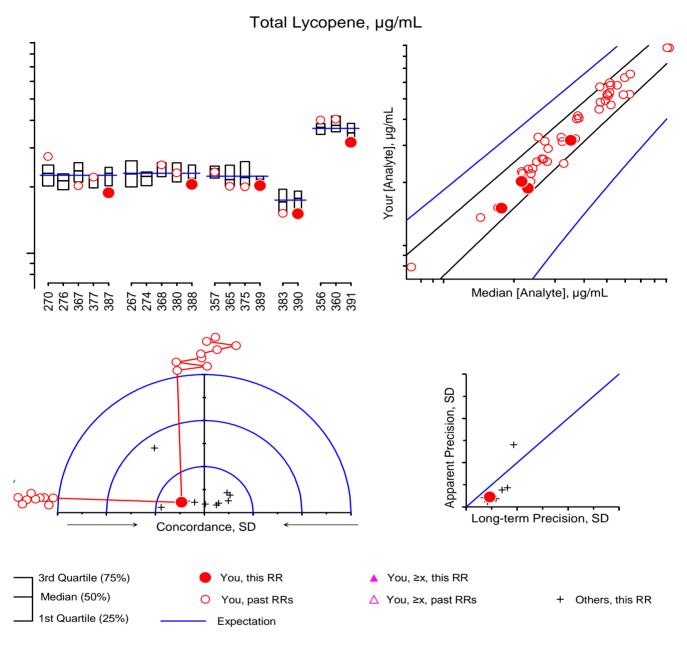
<u>erum</u>	Comments	<u>History</u>
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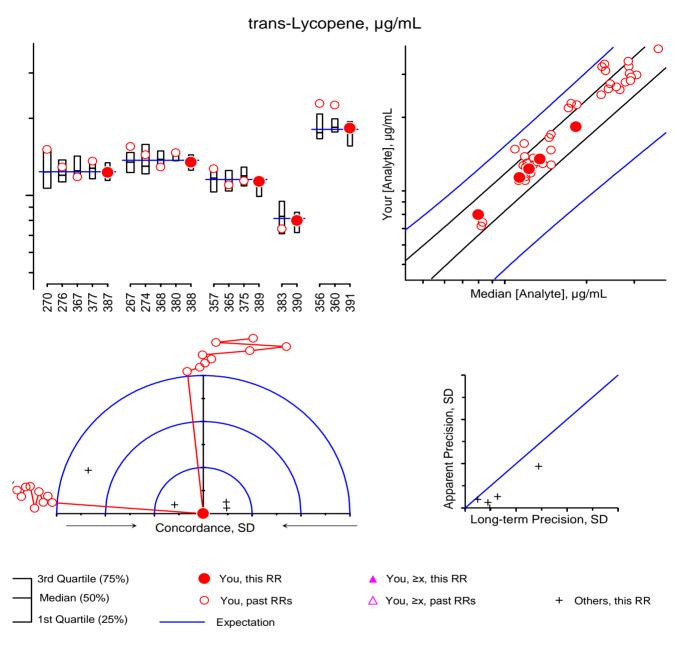
erum	Comments	<u>History</u>
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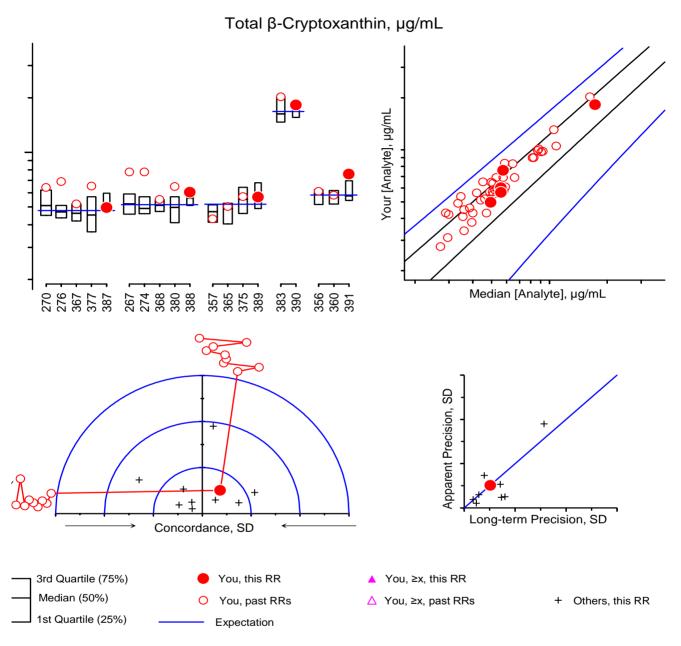
erum	Comments	<u>History</u>
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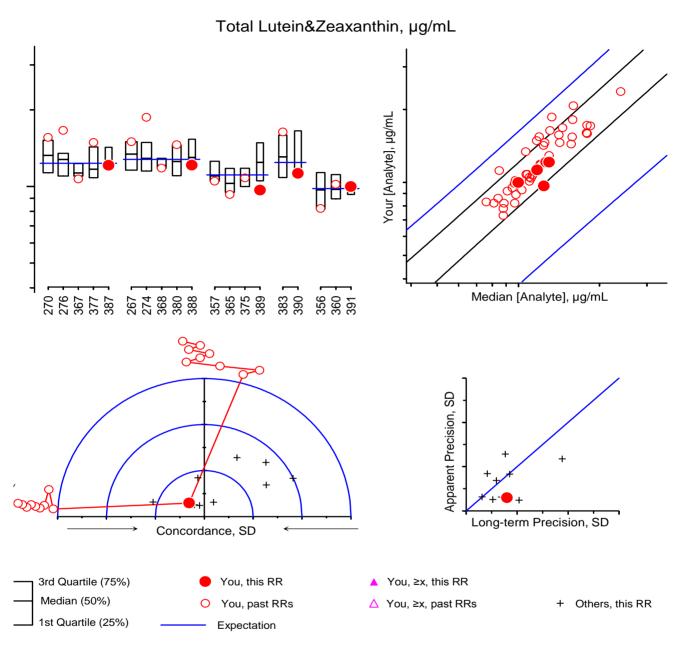
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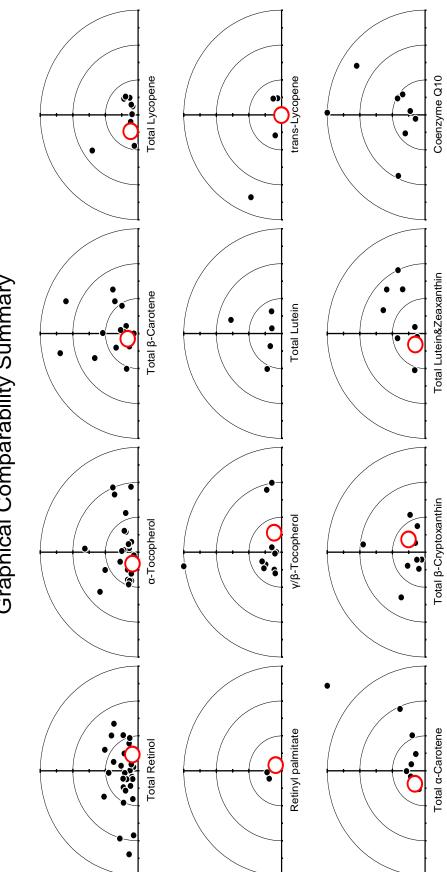
<u>erum</u>	Comments	<u>History</u>
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388	Fresh-frozen, same native pool as #387	49#267, 50#274, 68#368, 70#380
389	Fresh-frozen, native, multi-donor: SRM 968e I	66#357, 67#365,69#375
390	Fresh-frozen, native, single-donor	71#383
391	Fresh-frozen, native, multi-donor	65#356, 66#360

Set 1 of 33





Appendix E. Shipping Package Inserts for RR37

The following five items were included in each package shipped to an RR37 participant:

- Cover letter
- Protocol for Preparation and Analysis of the Ascorbic Acid Solid Control Material
- Preparation and Validation of Ascorbic Acid Solid Control Material Datasheet
- Analysis of Control Materials and Test Samples Datasheet
- Packing List and Shipment Receipt Confirmation Form

The cover letter, preparation protocol, and the two datasheets were enclosed in a sealed waterproof bag along with the samples themselves. The packing list was placed at the top of the shipping box, between the cardboard covering and the foam insulation.



UNITED STATES DEPARTMENT OF COMMERCE National Institute of Standards and Technology Gaithersburg, Maryland 20899-

June 25, 2012

Dear Colleague:

The samples within this package constitute Vitamin C Round Robin 37 (RR37) of the 2012 Micronutrients Measurement Quality Assurance Program. RR37 consists of four vials of frozen serum *test samples* (#371, #372 #373, and #374), one vial each of two frozen *control serum samples* (CS #3 and CS #4), and one vial of ascorbic acid *solid control material* (Control). Please follow the attached protocols when you prepare and analyze these samples. If you cannot prepare the *solid control* solutions gravimetrically, please prepare equivalent solutions volumetrically and report the exact volumes used. (Routine 0.5 g gravimetric measurements are generally 10-fold more accurate than routine 0.5 mL volumetric measurements.)

Please use the *control sera* to validate the performance of your measurement system <u>before</u> you analyze the *test samples*. The target value for CS #3 is $(15.5 \pm 1.6; 13.9 \text{ to } 17.1) \mu \text{mol/L}$ and the target for CS #4 is $(46.1 \pm 4.6; 41.5 \text{ to } 50.7) \mu \text{mol/L}$. We expect your results for both of these controls to be within this $\pm 10\%$ target range. If either of your result is significantly outside this range, your analysis system may not be suited to the analysis of MPA-preserved samples. In this case, please do **not** proceed to the analysis of the *test samples* but contact us at 301-975-3120 or <u>jbthomas@nist.gov</u>.

Please be aware that sample contact with any oxidant-contaminated surface (vials, glassware, etc.) may degrade your measurement system's performance (SA Margolis and E Park, "Stability of Ascorbic Acid in Solutions in Autosampler Vials", *Clinical Chemistry* 2001, 47(8), 1463-1464). You should suspect such degradation if you observe unusually large variation in replicate analyses.

Please report your results (using the attached form) for RR37 by e-mail to <u>david.duewer@nist.gov</u> or fax to 301-977-0685 by **September 28, 2012**. If you have questions or comments regarding this study, please call me at (301) 975-3120 or e-mail me at <u>jbthomas@nist.gov</u>.

Sincerely,

Jeanice Brown Thomas Program Coordinator/Research Chemist Analytical Chemistry Division Chemical Science and Technology Laboratory

Enclosures: Protocols, Preparation and Analysis of Control Materials and Analysis of Test Samples RR37 Report Form for Ascorbic Acid Solid Control Material Preparation RR37 Report Form for Control Material and Test Sample Analyses



Micronutrient Measurement Quality Assurance Program for Vitamin C

Please Read Through Completely BEFORE Analyzing Samples

Protocol for Preparation and Analysis of the Ascorbic Acid Solid Control Material

The *ascorbic acid solid control material* (in the amber vial) should be prepared and used in the following manner:

- 1) Prepare at least 500 mL of 5% mass fraction metaphosphoric acid (MPA) in distilled water. This solution will be referred to as the "Diluent" below.
- Weigh 0.20 to 0.22 g of the ascorbic acid solid control material to 0.0001 g (if possible), dissolve it in the Diluent in a 100 mL volumetric flask, and dilute with the Diluent to the 100 mL mark. Weigh the amount of Diluent added to 0.1 g. Record the weights. The resulting material will be referred to as the "Stock Solution" below.
- 3) Prepare three dilute solutions of the Stock Solution as follows:

<u>Dilute Solution 1:</u> Weigh 0.500 mL of the Stock Solution to 0.0001 g into a 100 mL volumetric flask; dilute with Diluent to the 100 mL mark. Record the weight.

<u>Dilute Solution 2:</u> Weigh 0.250 mL of the Stock Solution to 0.0001 g into a 100 mL volumetric flask; dilute with Diluent to the 100 mL mark. Record the weight.

<u>Dilute Solution 3:</u> Weigh 0.125 mL of the Stock Solution to 0.0001 g into a 100 mL volumetric flask; dilute with Diluent to the 100 mL mark. Record the weight.

4) Calculate and record the total ascorbic acid concentrations, [TAA], in these Dilute Solutions. If you follow the above gravimetric preparation directions, the [TAA] in μmol/L is calculated:

 $[\mathsf{TAA}]_{\mathsf{DS}} = \frac{(g \operatorname{Stock} \operatorname{Solution} \operatorname{in} \operatorname{Dilute} \operatorname{Solution}) \cdot (g \operatorname{AA} \operatorname{in} \operatorname{Stock} \operatorname{Solution}) \cdot (56785 \ \mu \mathsf{mol/g} \cdot \mathsf{L})}{(g \operatorname{AA} \operatorname{in} \operatorname{Stock} \operatorname{Solution}) + (g \operatorname{Diluent} \operatorname{in} \operatorname{Stock} \operatorname{Solution})}$

For example, if you prepared the Stock Solution with 0.2000 g of solid ascorbic acid and 103.0 g of Diluent, then 0.5 mL of the Stock Solution should weigh (0.2+103)/200 = 0.52 g and $[TAA]_{DS1} = (0.52 \text{ g})(0.2 \text{ g})\cdot(56785 \mu \text{mol/g}\cdot\text{L})/(0.2 + 103 \text{ g}) = 57.2 \mu \text{mol/L}$. Likewise, 0.25 mL of the Stock Solution should weigh 0.26 g and $[TAA]_{DS2} = 29.4 \mu \text{mol/L}$ and 0.125 mL should weigh 0.13 g and $[TAA]_{DS3} = 14.2 \mu \text{mol/L}$.

5) Measure the ultraviolet absorbance spectrum of Dilute Solution 1 against the Diluent as the blank using paired 1 cm path length cuvettes. Record the absorbance at 242, 243, 244, and 245 nm. Record the maximum absorbance (A_{max}) within this region. Record the wavelength (λ_{max}) at which this maximum occurs.

The extinction coefficient ($E^{1\%}$) of ascorbic acid at λ_{max} (using a cell with a 1 cm path length) of Dilute Solution #1 can be calculated:

 $E^{1\%}(\frac{dL}{g \cdot cm}) = \frac{(A_{max}) \cdot ((g \text{ AA in Stock Solution}) + (g \text{ Diluent in Stock Solution}))}{(g \text{ Stock Solution in Dilute Solution 1}) \cdot (g \text{ AA in Stock Solution})}$

If your spectrophotometer is properly calibrated, λ_{max} should be between 243 and 244 nm and $E^{1\%}$ should be 550 ± 30 dL/g·cm. If they are not, you should recalibrate the wavelength and/or absorbance axes of your spectrophotometer and repeat the measurements.

- 6) Measure and record the concentration of total ascorbic acid in all three dilute solutions and in the 5% MPA Diluent in duplicate using *exactly* the same method that you will use for the serum control materials and test samples, including any enzymatic treatment. We recommend that you analyze these solutions in the following order: Diluent, Dilute Solution 1, Dilute Solution 2, Dilute Solution 3, Dilute Solution 3, Dilute Solution 2, Dilute Solution 1, Diluent.
 - a) Compare the values of the duplicate measurements. *Are you satisfied that your measurement precision is adequate?*
 - b) Compare the measured with the calculated [TAA] values. This is most conveniently done by plotting the measured values on the y-axis of a scatterplot against the calculated values on the x-axis. The line through the four {calculated, measured} data pairs should go through the origin with a slope of 1.0. *Are you satisfied with the agreement between the measured and calculated values?*

Do <u>**not**</u> analyze the serum control materials or test samples until you are satisfied that your system is performing properly!

Once you have confirmed that your system is properly calibrated, analyze the serum controls CS #3 and CS #4 (see protocol below). The target range for CS #3 is (15.5 \pm 1.6; 13.9 to 17.1) µmol/L and the target range for CS #4 is (46.1 \pm 4.6; 41.5 to 50.7) µmol/L.

If either of your measured values is not within its target range, please review your sample preparation procedure and whether you followed *exactly* the same measurement protocol the solutions prepared from the solid control material as you used for these serum controls. If the protocols differ, please repeat from Step 6 using the proper protocol. If the proper protocol was used, your measurement system may not be suitable for MPA-preserved samples; please contact us at 301-975-3120 or jbthomas@nist.gov.

Do <u>not</u> analyze the test samples until you are satisfied that your system is performing properly and is suitable for the analysis of MPA-preserved serum!

Protocol for Analysis of the Serum Control Materials and Test Samples

The *serum control material* and *test samples* are in sealed ampoules. They were prepared by adding equal volumes of 10% MPA to spiked human serum. We have checked the samples for stability and homogeneity. Only the total ascorbic acid is stable. While these samples contain some dehydroascorbic acid, its content is variable. Therefore, only <u>total ascorbic acid</u> should be reported. The *serum control material* and *test samples* should be defrosted by warming at 20 °C for not more than 10 min otherwise some irreversible degradation may occur.

Each *serum test sample* contains between 0.0 and 80.0 μ mol of total ascorbic acid/L of solution. The total ascorbic acid in each ampoule should be measured in duplicate. Please report your results in μ mol/(L of the sample solution) rather than μ mol/(L of serum NIST used to prepare the sample).

Participant #: _____

Date:

Vitamin C Round Robin 37

NIST Micronutrient Measurement Quality Assurance Program

Preparation and Validation of Ascorbic Acid Solid Control Material

STOCK SOLUTION

Mass of ascorbic acid in the Stock Solution	g
Mass of 5% MPA Diluent added to the 100 mL volumetric flask	g

DILUTE SOLUTION 1

Mass of added stock solution (0.5 mL)	g
Mass of 5% MPA Diluent added to the 100 mL volumetric flask	g
Absorbance of Dilute Solution 1 at 242 nm	AU
Absorbance of Dilute Solution 1 at 243 nm	AU
Absorbance of Dilute Solution 1 at 244 nm	AU
Absorbance of Dilute Solution 1 at 245 nm	AU
Absorbance of Dilute Solution absorbance maximum	AU
Wavelength of maximum absorbance	nm
Calculated E ^{1%}	dL/g∙cm
Calculated [TAA] _{DS1}	µmol/L

DILUTE SOLUTION 2

Mass of added stock solution (0.25 mL)	g
Mass of 5% MPA Diluent added to the 100 mL volumetric flask	g
Calculated [TAA] _{DS2}	_μmol/L

DILUTE SOLUTION 3

Mass of added stock solution (0.125 mL)	g
Mass of 5% MPA Diluent added to the 100 mL volumetric flask	g
Calculated [TAA] _{DS3}	µmol/L

Please return by September 28, 2012

MMQAP
100 Bureau Drive, Stop 8392
Gaithersburg, MD 20899-8392

Fax: 301-977-0685 Email: david.duewer@nist.gov Participant #: _____

Date:

Vitamin C Round Robin 37

NIST Micronutrient Measurement Quality Assurance Program

Analysis of Control Materials and Test Samples

Sample	Replicate 1	Replicate 2	Units
Dilute Solution 1			µmol/L of Dilute Solution
Dilute Solution 2			µmol/L of Dilute Solution
Dilute Solution 3			μmol/L of Dilute Solution
5% MPA Diluent			μmol/L of Diluent
CS # 3			μ mol/L of Sample <i>Target: (15.5 ±1.6) µmol/L</i>
CS # 4			μmol/L of Sample <i>Target: (46.1 ±4.6) μmol/L</i>
Serum Test Sample #371			µmol/L of Sample
Serum Test Sample #372			µmol/L of Sample
Serum Test Sample #373			μmol/L of Sample
Serum Test Sample #374			μmol/L of Sample

Were samples frozen upon receipt? Yes | No

Analysis method: HPLC-EC | HPLC-Fluor DAB | HPLC-OPD | HPLC-UV | AO-OPD | Other If "Other", please describe:

Nature of samples you typically analyze: native | MPA-preserved | DTT-preserved | Other If "Other", please describe:

COMMENTS:

Please return by September 28, 2012

Fax: 301-977-0685 Email: david.duewer@nist.gov Vitamin C Round Robin 37 NIST Micronutrients Measurement Quality Assurance Program

Packing List and Shipment Receipt Confirmation Form

This box contains one vial each of the following **seven** VitC M²QAP samples:

Label	Form
VitC #371	Liquid frozen (1:1 serum:10% MPA)
VitC #372	Liquid frozen (1:1 serum:10% MPA)
VitC #373	Liquid frozen (1:1 serum:10% MPA)
VitC #374	Liquid frozen (1:1 serum:10% MPA)
CS #3	Liquid frozen (1:1 serum:10% MPA)
CS #4	Liquid frozen (1:1 serum:10% MPA)
Control	Solid AA

Please 1) Open the pack immediately

- 2) Check that it contains one vial each of the above samples
- 3) Check if the samples arrived frozen
- 4) Store the samples at -20 °C or below until analysis
- 5) Complete the following information
- 6) Fax the completed form to us at 301-977-0685 (or email requested information to david.duewer@nist.gov)

1) Date this shipment arrived: _____

2) Are all of the vials intact? Yes | No

If "No", which one(s) were damaged?

- 3) Was there any dry-ice left in cooler? Yes | No
- 4) Did the samples arrive frozen? Yes | No
- 5) At what temperature are you storing the samples? _____°C
- 6) When do you anticipate analyzing these samples? _____

Your prompt return of this information is appreciated.

The M²QAP Gang

Appendix F. Final Report for RR37

The following five pages are the final report as provided to all participants:

- Cover letter.
- An information sheet that:
 - o describes the contents of the "All-Lab" report,
 - o describes the content of the "Individualized" report,
 - o describes the nature of the test samples and details their previous distributions, if any, and
 - o summarizes aspects of the study that we believe may be of interest to the participants.



December 7, 2012

Dear Colleague:

Enclosed is the summary report of the results for Round Robin 37 (RR37) for the measurement of total ascorbic acid (TAA, ascorbic acid plus dehydroascorbic acid) in human serum. Included in this report are a summary of data for all laboratories and an individualized summary of your laboratory's measurement performance. The robust median is used to estimate the consensus value for all samples, the "adjusted median absolute deviation from the median" (MADe) is used to estimate the expected standard deviation, and we estimate the coefficient of variation (CV) as 100×MADe/median.

RR37 consisted of four test samples (#371, #372,, #373, and #374), one vial each of two frozen control serum control samples (CS #3 and CS #4), and one vial of solid control material (Control) for preparation of TAA control solutions. Details regarding the samples can be found in the enclosed report.

If you have concerns regarding your laboratory's performance, we suggest that you obtain and analyze a unit of Standard Reference Material (SRM) 970 Vitamin C in Frozen Human Serum. SRM 970 can be purchased from the NIST SRM Program at <u>www.nist.gov/srm</u>; phone: 301-975-6776; fax: 301-948-3730. If your measured values do not agree with the certified values, we suggest that you contact us for consultation.

Samples for the first vitamin C round robin (RR38) of the 2013 MMQAP will be shipped starting January 22, 2013. Please contact us immediately if this schedule is problematic for your laboratory.

If you have questions or concerns regarding this report, please contact David Duewer at david.duewer@nist.gov or me at jbthomas@nist.gov, 301-975-3120, or fax: 301-977-0685.

Sincerely,

Jeanice Brown Thomas, M.B.A . Research Chemist Analytical Chemistry Division Material Measurement Laboratory

Enclosures

cc: L. C. Sander

David L. Duewer, Ph.D. Research Chemometrician Analytical Chemistry Division Material Measurement Laboratory



The NIST MMQAP Vitamin C Round Robin 37 (RR37) report consists of:

Page	"Individualized" Report
1	Summary of your reported values for the nominal 55 mmol/L solution you prepared from the ascorbic acid solid control sample, the serum control sample, and the four serum test samples.
2	Graphical summary of your RR37 sample measurements.
Page	"All-Lab" Report
1	A tabulation of results and summary statistics for total ascorbic acid [TAA] in the RR37 samples and control/calibration solutions.
Serum-B	ased Samples. Two serum controls and four test samples were distributed in RR37.
CS#3	a (13.9 to 17.1) µmol/L material ampouled in late 2009
CS#4	a (41.5 to 50.7) µmol/L material ampouled in late 2009
S37:1	SRM 970 level 1, ampouled in mid-1998, previously distributed as an "Unknown" in RRs 11 to 16, 19, 20 23, 25, 29, 31, and 34
S37:2	Ampouled in late 2001, previously distributed in RRs 17, 18, 20, 22, 23, 27, and 31
S37:3	SRM 970 level 2, ampouled in mid-1998, previously distributed as an "Unknown" in RRs 11 to 15, 18, 20 22, 25, 29, and 36
S37:4	Ampouled in late 2001, previously distributed in RRs 16, 17, 20, 21, 23, 27, and 30

Results.

- All participants who prepared the four 5% metaphosphoric acid (MPA) control/calibration solutions (the three "Dilute Solutions" and the "Diluent") did so correctly. The criteria used to evaluate this success are: the density of the 5% MPA solution (≈1.03 gm/mL), the observed wavelength maximum of "Dilute Solution #1" (≈244 nm), the observed absorbance at that maximum (≈0.58), and the calculated E^{1%}#1" (≈560 dL/g·cm).
- 2) The Measured = a+b*Gravimetric calibration parameters for the control/calibration solutions (columns 10 to 13 of the All-Lab Report) indicate that the measurement systems for all participants are linear (R^2 close to 1 and the root-mean-square (RMS) residual close to 0.0) and well calibrated (intercepts close to 0 and slopes close to 1).
- 3) The Measured = p+q*Median regression parameters for samples S37:1 to S37:4 (columns 23 to 26 of the All-Lab Report) confirm the linearity of all measurement systems (R² close to 1 and RMS close to 0.0). However, the intercepts and slopes continue to indicate that there are systematic differences in the response of the various measurement procedures to the TAA in the 5% aqueous metaphosphoric acid (MPA) of the test samples.
- 4) The variability of the measured values after calibration to the control solutions (the final 6 columns of the All-Lab Report) continues to be as great or greater than that of the measured values themselves. The primary source of the systematic differences in response of the measurement procedures appears to be the presence of serum components in the sample matrix, not the MPA preservative. On this basis, we intend to drop the standard solution arm of the MMQAP Vitamin C Round Robin studies beginning with RR38.

Appendix G. "All-Lab Report" for RR37

The following single page is the "All-Lab Report" as provided to all participants, with two exceptions:

- the participant identifiers (Lab) have been altered.
- the order in which the participant results are listed has been altered.

The data summary in the "All-Lab Report" has been altered to ensure confidentiality of identification codes assigned to laboratories.

am for Total Ascorbic Acid	N
licronutrients Measurement Quality Assurance Program for Total Ascorbic Acid	"Round Robin" 37 - Fall 2012
Micronutrients Measure	

	I	+	N	8	6	7	2	2	<u>م</u>	_	4	6	2	2	4	2	2	2	20	0	0
		S37:4	42.2	44.8	47.9	46.7	45.2	47.2	50.5	42.	40.4		45.2	З.	40.4	42.2	45.2	47.2	50.5	4.0	
	L	S37:3	26.7	26.9	27.6	27.4	25.8	29.5	32.2	25.9	23.9	6	27.3	2.4	23.9	25.9	26.9	27.6	32.2	1.4	5
	, µmol/	S37:2	19.7	21.7	22.1	21.8	20.0	21.6	26.4	21.0	17.1	6	21.3	2.5	17.1	20.0	21.6	21.8	26.4	0.8	4
	Calibrated, µmol/I	S37:1	7.7	8.7	8.3	8.3	7.6	10.1	11.5	8.5	7.1	6	8.7	1.4	7.1	7.7	8.3	8.7	11.5	0.9	11
	Са	CS#4 S	47.2	46.6	47.5	46.0	46.2	45.8	53.3	41.9	40.4	6	46.1	3.6	40.4	45.8	46.2	47.2	53.3	1.6	e
		CS#3 C	15.9	12.6	15.6	14.9	14.8	16.5	22.7	14.9	13.1	6	15.7	2.9	12.6	14.8	14.9	15.9	22.7	1.4	10
	an	RMS	0.5	0.7	1.1	0.8	1.1	1.0	0.6	0.5	1.1										
	q*Medi:	R ² R	0.999	0.999	0.997	0.998	J.996	7.997	0.999	0.999	0.996										
	Measured = p+q*Median	Slope	1.00 0.1	0.99 0.	.11	.10	.03 0.	.00	.08	0.96 0.	0.96 0.										
	Measur	Inter SI	-0.48 1	0.20 0	-1.36 1	-1.50 1	-2.13 1	-0.07 1	2.09 1	0.44 0	-1.75 0										
Samples		S37:4 Ir	44.0 -0	44.7 0	48.9 -1		44.7 -2	44.9 -0	50.1 2	43.3 0	41.8 -1	6	45.6	2.8	41.8	44.0	44.7	48.2	50.1	2.0	5
Sam		S37:3 S:	27.8	26.8	28.2	28.1	25.4	27.8	31.7	26.6	24.8	6	27.5	2.0	24.8	26.6	27.8	28.1	31.7	1.5	5
	Imol/L	S37:2 S3	20.5	21.7	22.6	22.2	19.7	20.2	25.9	21.5	17.7	6	21.3	2.3	17.7	20.2	21.5	22.2	25.9 :	1.7	8
	Measured, µmol/l	S37:1 S3	8.0 2	8.8	8.6	8.2	7.4	9.1	10.9	8.5	7.3	6	8.5	1.1	7.3 1	8.0	8.5	8.8	0.9	0.7	8
	Mea	CS#4 S3	49.2	46.5	48.4	47.5	45.7	43.6	53.0 1	43.2	41.8	6	46.5	3.5	41.8	43.6	46.5	48.4	53.0 1	4.1	6
		CS#3 CS	16.5 4	12.6 4	16.0 4	15.1 4	14.6 4	15.2 4	22.2 5	15.1 4	13.5 4	6	15.7 4	2.7	12.6 4	14.6 4	15.1 4		22.2 5		6
					<u>`</u>			-				7		8.7	_		·			2.6	0.5
ution 1	tometry	E ¹ %	0 541.6	0 560.3	2 558.0	0 569.4	3 560.8		^a 349.9 ^a	4 559.7	0 551.4	7	0 557.3	1 8.	0 541.6	2 554.7	3 559.7	1 560.5	0 569.4		
Dilute Solution	Spectrophotometry	A_{max}	0.5510	0.5990	0.5952	0.6140	0.5933		0.340^{a}	0.5694	0.5450		0.5810	0.026	0.5450	0.5602	0.5933	0.5971	0.6140	0.0307	5.2
ē	Spe	λ_{\max}	243.	243.5	244.	243.6	243.3		255 ^a	243.9	242.	2	243.3	0.7	242.0	243.2	243.5	243.8	244.0	0.6	0.26
MPA	Density	g/mL	1.036	1.030	1.024	1.026	1.031	1.031	1.015	1.032	1.028	6	1.028	0.006	1.015	1.026	1.030	1.031	1.036	0.0	0.33
ĺ		RMS	0.4	0.8	0.2	0.6	0.4	2.2	1.5	0.5	0.2	z	Average	ß	Min	%25	Median	%75	Max	eSD	S
	Measured = a + b*Grav	R ² RI	1.000	0.999	1.000	1.000	1.000	0.994	0.997	1.000	000.		Ave				Me				
	ed = a	Slope	1.04 1.	0.99 0.	1.02 1.	1.04 1.	0.99 1.	0.97 0.	1.00 0.	1.04 1.	1.04 1.										
les	Measui	Inter SI	-0.03	.11	0.15	-0.43	-0.19	-0.69	-0.61	-0.35	-0.07										
ר Samp		MPA II	0.0	0.0	0.0	0.0	0.0	0.0	0.7	0.0	0.0	6	0.1	0.2	0.0	0.0	0.0	0.0	0.7	0.0	
ibratio	μmol/L	Dil:3 N	15.0	14.5	15.8	16.8	14.7	13.5	13.4	13.5	14.3	6	14.6	1.2	13.4	13.5	14.5	15.0	16.8	1.4	10
Control / Calibration Samples	Measured, µmol/L	Dil:2	29.9	31.2	31.3	32.1	29.4	23.4	26.8 `	30.2	30.2	6	29.4	2.7	23.40	29.35	30.18	31.23	32.15	1.6	5
Contr	Mea	Dil:1	60.0	60.1	61.7	63.7	59.7	53.9	55.5	59.7	58.2	6	59.2	3.0	53.9 2	58.2 2	59.7 3	60.1 3	63.7 3	2.2	4
	Ĺ	Dil:3	14.8	14.9	15.2	16.8	14.9	13.8	15.1	13.9	13.8	6	14.8	0.9	13.8	13.9	14.9	15.1	16.8	0.4	с
	Grav, µmol/L	Dil:2	28.3	30.3	30.6	31.8	30.2	27.7	28.1	29.2	29.4	6	29.5	1.3	27.67	28.33	29.39	30.31	31.83	1.6	5
	Gra∿	Dil:1	57.8	60.7	60.6	61.2	60.1	55.3	55.1	57.8	56.1	6	58.3	2.4	55.1 2	56.1 2	57.8 2	60.6 3	61.2 3	3.7	9
	1	Date	25/09/12	06/08/12	20/08/12	15/08/12	12/09/12	27/09/12	29/08/12	12/09/12	10/08/12	z	Average 58.3	SD	Min	%25	Median	%75	Max	MADe	S
				-									A				2				
		Lab	VC-MA	VC-MB	VC-MC	VC-MG	VC-MH	VC-MI	VC-MJ	VC-MN	VC-NM										
																		(20	,	

a) 5% Trichloroacetic acid solution

Appendix H. Representative "Individualized Report" for RR37

Each participant in RR37 received an "Individualized Report" reflecting their reported results. The following two pages are the "Individualized Report" for participant "VC-MA".

Vitamin C "Round Robin" 37 Report: Participant VC-MA

					MPA	Dilu	te Solutio	Control/Calibration Solutions							
					Density Spectrophotometry					Y _{meas} = Inter + Slope* X _{grav}					
Date	RR		Method		g/mL		λ_{max}	A _{max}	E ^{1%}	Inter	Slope	R^2	SEE		
02/24/10	32	HPLC-EC			1.035		242.0	0.566	545.1	0.3	1.03	1.000	0.46		
09/27/10	33	HPLC-EC			1.037		244.0	0.560	540.5	0.4	1.08	1.000	0.43		
02/28/11	34	HPLC-EC			1.039		244.0	0.575	555.2	0.6	1.14	1.000	0.78		
08/24/11	35	HPLC-EC			1.039		242.0	0.568	547.7	0.0	1.03	1.000	0.24		
02/02/12	36	HPLC-EC			1.035		244.0	0.561	550.5	-0.4	1.05	1.000	0.55		
09/25/12	37	HPLC-EC			1.035		243.0	0.551	541.6	0.0	1.04	1.000	0.41		
				Mean	1.037		243.2	0.56	546.8		Po	oled SEE	0.50		
				SD	0.002		1.0	0.01	5.6						
							nmol/Lsa	- ·		1					
Date	RR		Sample		Rep ₁	Rep_2	F_{adj}	Mean	SD_{dup}	N	Mean	SD _{repeat}	SD _{reprod}		
09/23/98	11		S11:1		15.5	13.9	0.5	7.4	0.6	14	8.2	0.3	0.5		
04/02/99	12		S12:1		14.5	15.8	0.5	7.6	0.5						
09/17/01	13		S13:1		8.4	8.5	1.0	8.5	0.1						
09/27/01	14		S14:3		8.0	7.7	1.0	7.8	0.2						
09/18/01	15		S15:1		8.9	8.7	1.0	8.8	0.1						
11/18/02	16		S16:1		8.8	8.8	1.0	8.8	0.0						
11/13/03	19		S19:4		7.8	8.6	1.0	8.2	0.5						
02/23/04	20		S20:3		8.3	8.1	1.0	8.2	0.1						
10/17/05	23		S23:4		8.6	8.8	1.0	8.7	0.1						
08/28/06	25		S25:1		8.7	8.5	1.0	8.6	0.2						
08/11/08	29		S29:2		8.3	8.4	1.0	8.3	0.1						
09/10/09	31		S31:3		7.3	8.1	1.0	7.7	0.5						
02/28/11	34		S34:1		8.5	8.3	1.0	8.4	0.1						
09/25/12	37		S37:1		8.0	8.1	1.0	8.0	0.0						
09/23/98	11		S11:2		50.7	47.7	0.5	24.6	1.1	12	26.7	0.5	1.8		
04/02/99	12		S12:2		49.5	45.9	0.5	23.9	1.3						
09/17/01	13		S13:2		27.6	27.7	1.0	27.7	0.1						
09/27/01	14		S14:4		25.7	26.4	1.0	26.0	0.5						
09/18/01	15		S15:2		25.4	25.6	1.0	25.5	0.2 0.3						
03/20/03 02/23/04	18 20		S18:3 S20:4		28.8 25.9	29.2 25.2	1.0 1.0	29.0 25.5	0.5						
02/23/04 03/08/05	20 22		S20.4 S22:4		23.9 29.4	29.2 29.4	1.0	25.5 29.4	0.0						
08/28/06	25		S25:2		23.4	27.4	1.0	27.5	0.0						
08/11/08	29		S29:3		27.0	25.7	1.0	25.6	0.1						
02/02/12	36		S36:3		28.3	28.5	1.0	28.4	0.1						
09/25/12	37		S37:3		27.9	27.8		27.8	0.1						
12/12/02	17		S17:2		23.3	23.4	1.0	23.4	0.1	8	23.0	0.5	1.4		
03/20/03	18		S18:1		22.7	23.7	1.0	23.2	0.7						
02/23/04	20		S20:1		25.1	24.1	1.0	24.6	0.7						
03/08/05	22		S22:2		22.7	22.7	1.0	22.7	0.0						
10/17/05	23		S23:2		25.5	24.4	1.0	24.9	0.8						
10/05/07	27		S27:1		22.9	22.4	1.0	22.6	0.4						
09/10/09	31		S31:1		21.5	22.6	1.0	22.1	0.7						
09/25/12	37		S37:2		20.4	20.7	1.0	20.5	0.2						
11/18/02	16		S16:3		49.9	44.9	1.0	47.4	3.5	8	48.3	1.3	2.2		
12/12/02	17		S17:3		49.7	49.1	1.0	49.4	0.4						
02/23/04	20		S20:2		50.6	50.0	1.0	50.3	0.4						
09/13/04	21		S21:4		47.1	47.0	1.0	47.0	0.0						
10/17/05	23		S23:3		49.8	48.8	1.0	49.3	0.7						
10/05/07	27		S27:2		48.6	47.6	1.0	48.1	0.8						
03/03/09	30		S30:2		51.2	50.4	1.0	50.8	0.6						
09/25/12	37		S37:4		43.9	44.1	1.0	44.0	0.1						

Please check our records against your records. Send corrections and/or updates to...

Micronutrients Measurement Quality Assurance Program National Institute of Standards and Technology

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Total Ascorbic Acid, µmol/mL 000000 Your [Total Ascorbic Acid], µmol/L Total Ascorbic Acid], µmol/L \$17:2 \$18:1 \$20:1 \$20:1 \$22:2 \$23:2 \$23:2 \$23:1 \$31:1 \$37:2 S16:3 S17:3 S27:3 S21:4 S21:4 S23:3 S27:2 S37:4 S37:4 RR25 RR27 RR27 RR25 RR25 RR31 RR34 RR37 RR37 RR37 RR37 RR1 Median [Total Ascorbic Acid], µmol/L Apparent Precision, SD Concordance, SD 3rd Quartile (75%) You, this RR Median (50%) You, past RRs Others, this RR O 1st Quartile (25%)

For details of the construction and interpretation of these plots, see: Duewer, Kline, Sharpless, Brown Thomas, Gary, Sowell. Anal Chem 1999;71(9):1870-8.

Comments

Sample

Set 1 of 9

S37:1 SRM970 Lv I - distributed in RRs 11, 12, 13, 14, 15, 16, 19, 20, 23, 25, 29, 31, and 34

S37:2 Distributed in RRs 17, 18, 20, 22, 23, 27, and 31

S37:3 SRM970 Lv II - distributed in RRs 11, 12, 13, 14, 15, 18, 20, 22, 25, 29, and 36 S37:4 Distributed in RRs 16, 17, 20, 21, 23, 27, and 30