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Published in:
Analytical Chemistry

DOI:
[10.1021/acs.analchem.5b04392](https://doi.org/10.1021/acs.analchem.5b04392)

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Document Version
Publisher's PDF, also known as Version of record

Publication date:
2016

[Link to publication in University of Groningen/UMCG research database](#)

Citation for published version (APA):

Schimmelmann, A., Qi, H., Coplen, T. B., Brand, W. A., Fong, J., Meier-Augenstein, W., Kemp, H. F., Toman, B., Ackermann, A., Assonov, S., Aerts-Bijma, A. T., Brejcha, R., Chikaraishi, Y., Darwish, T., Elsner, M., Gehre, M., Geilmann, H., Groeing, M., Helie, J-F., ... Werner, R. A. (2016). Organic Reference Materials for Hydrogen, Carbon, and Nitrogen Stable Isotope-Ratio Measurements: Caffeines, n-Alkanes, Fatty Acid Methyl Esters, Glycines, L-Valines, Polyethylenes, and Oils. *Analytical Chemistry*, 88(8), 4294-4302. <https://doi.org/10.1021/acs.analchem.5b04392>

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Organic Reference Materials for Hydrogen, Carbon, and Nitrogen Stable Isotope-Ratio Measurements: Caffeines, *n*-Alkanes, Fatty Acid Methyl Esters, Glycines, L-Valines, Polyethylenes, and Oils

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Supporting Information

ABSTRACT: An international project developed, quality-tested, and determined isotope- δ values of 19 new organic reference materials (RMs) for hydrogen, carbon, and nitrogen stable isotope-ratio measurements, in addition to analyzing pre-existing RMs NBS 22 (oil), IAEA-CH-7 (polyethylene foil), and IAEA-600 (caffeine). These new RMs enable users to normalize measurements of samples to isotope- δ scales. The RMs span a range of $\delta^2\text{H}_{\text{VSMOW-SLAP}}$ values from -210.8 to $+397.0$ mUr or ‰, for $\delta^{13}\text{C}_{\text{VPDB-LSVEC}}$ from -40.81 to $+0.49$ mUr and for $\delta^{15}\text{N}_{\text{Air}}$ from -5.21 to $+61.53$ mUr. Many of the new RMs are amenable to gas and liquid chromatography. The RMs include triads of isotopically contrasting caffeines, C_{16} *n*-alkanes, *n*- C_{20} -fatty acid

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Received: November 19, 2015

Accepted: March 14, 2016

Published: March 14, 2016

methyl esters (FAMES), glycines, and L-valines, together with polyethylene powder and string, one *n*-C₁₇-FAME, a vacuum oil (NBS 22a) to replace NBS 22 oil, and a ²H-enriched vacuum oil. A total of 11 laboratories from 7 countries used multiple analytical approaches and instrumentation for 2-point isotopic normalization against international primary measurement standards. The use of reference waters in silver tubes allowed direct normalization of $\delta^2\text{H}$ values of organic materials against isotopic reference waters following the principle of identical treatment. Bayesian statistical analysis yielded the mean values reported here. New RMs are numbered from USGS61 through USGS78, in addition to NBS 22a. Because of exchangeable hydrogen, amino acid RMs currently are recommended only for carbon- and nitrogen-isotope measurements. Some amino acids contain ¹³C and carbon-bound organic ²H-enrichments at different molecular sites to provide RMs for potential site-specific isotopic analysis in future studies.

Semiautomated, economical, and fast stable isotope-ratio analysis (SIRA) in continuous flow mode with online elemental analysis (EA), gas chromatography (GC), or liquid chromatography (LC) interfaced with isotope-ratio mass spectrometry (IRMS) has become a routine and invaluable tool in science and industry, including applications in ecology, (paleo)environmental research, fossil fuel and biofuel research, food authentication, forensics, medical diagnostics, paleoceanography, archeology, and other fields.^{1–4} Accurate determinations of relative stable isotope ratios of hydrogen ($\delta^2\text{H}$), carbon ($\delta^{13}\text{C}$), and nitrogen ($\delta^{15}\text{N}$) require two-point normalization⁵ using at least two isotopic reference materials (RMs, also called standards) with contrasting isotopic compositions to (i) anchor the isotopic scale and (ii) compensate for differences in responses of instruments, which commonly compress isotope- δ scales.^{6–9} The development of suitable organic stable isotope RMs has not kept pace with the rapid development of analytical continuous flow IRMS methods, forcing many practitioners to resort to a single RM and accept the increased analytical uncertainties that accompany ignorance of the relevant isotope- δ scale compression factor. In order to minimize matrix effects, a standard should be chemically as similar as possible to the unknown sample, and therefore organic RMs should be used to normalize δ values of organic samples. The fundamental “principle of identical treatment of sample and RM” in the analytical process¹⁰ does not permit the use of pulses of “standard gases” of CO₂, H₂, or N₂ as primary anchors of isotope- δ scales because such gas pulses have not passed through the same GC, EA, combustion of high-temperature-conversion (HTC), etc. interfaces before being admitted into the mass-spectrometer. The proper role of “standard gas pulses” is that of a mediator between the introduction of RMs within the same sequence, to correct for drift of the IRMS over time, whether during the course of a long GC analysis or between sequential EA analyses. Instead, the RMs (local or international) are interspersed among unknowns, and only through them can the measurements be normalized to isotope- δ scales. These newly developed organic RMs will enable the stable isotope community to adhere more closely to established guidelines and recommended practices when measuring stable isotope ratios of organic samples.

The development of new organic RMs was prompted mainly by the lack of appropriate RMs at a time when novel online IRMS methods were implemented by numerous disciplines. The lack of GC-amenable organic RMs has been particularly problematic.^{11,12} The calibration of in-house standards by individual laboratories was typically hampered by the inability to comparatively measure organic hydrogen and the hydrogen in VSMOW and SLAP while adhering to the principle of identical treatment of standard and sample. Our development of new RMs is based on direct isotopic comparisons between organic hydrogen and standard waters sealed in silver tubes.¹³

Chemical choices of new RMs were guided by the need for chemical stability, low volatility and toxicity, homogeneity, and

ease of handling. In particular, it is preferred that RMs for hydrogen stable isotopes should not be hygroscopic and should contain no exchangeable hydrogen, even though this is not always possible for the wide variety of matrixes sought in this project.

We comply with IUPAC guidelines for the International System of Units (SI) when expressing δ values¹⁴ and we use the unit mUr instead of ‰ when expressing $\delta^2\text{H}$, $\delta^{13}\text{C}$, and $\delta^{15}\text{N}$ values^{1,15} because the use of the ‰ symbol has been deprecated.¹⁶ The term urey (symbol Ur) is suited for diverse isotope scales of all elements. A δ value in the traditional form of -15‰ can be expressed as -15 mUr . The $\delta^2\text{H}$ values are normalized to a scale in which VSMOW = 0 mUr and SLAP = -428 mUr ⁶; the $\delta^{13}\text{C}$ values are normalized to a scale in which NBS 19 is $+1.95\text{ mUr}$ and LSVEC is -46.6 mUr ¹⁷; the $\delta^{15}\text{N}$ values are normalized to a scale in which atmospheric nitrogen = 0 mUr and USGS32 = $+180\text{ mUr}$.^{18,8}

■ MATERIALS AND METHODS

Raw Materials and Preparation of New Organic RMs. *n*-Hexadecanes (USGS67, USGS68, USGS69). Two batches of 125 and 175 mL from Fluka Analytical (CAS No. 544-76-3) with GC-based purities of at least 99.9% and two isotopically enriched *n*-hexadecanes from Aldrich (C₁₆H₃₄, ²H fraction = 98%; *n*-hexadecane-1,2-¹³C₂, ¹³C fraction = 99%) were used in the preparation of 3 *n*-hexadecanes with different $\delta^2\text{H}$ and $\delta^{13}\text{C}$ values (Table 2). Each stirred, homogeneous liquid was subdivided into glass containers and flame-sealed under argon. Users will receive aliquots of 50 μL sealed in glass tubes as RMs for $\delta^2\text{H}$ and $\delta^{13}\text{C}$ normalization.

***n*-Heptadecanoic Acid Methyl Ester (USGS76).** Custom-synthesized C₁₇ fatty acid methyl ester (C₁₈H₃₆O₂, CAS No. 1731-92-6) was obtained from Sigma-Aldrich with a purity of $\geq 99\%$. The supply of 200 g was melted, subdivided as a homogeneous liquid into glass ampules, and flame-sealed under argon. Users will receive an aliquot of this RM of 50 μL sealed in a glass tube for $\delta^2\text{H}$ and $\delta^{13}\text{C}$ normalization.

Icosanoic Acid Methyl Esters (USGS70, USGS71, USGS72). Also known as methyl eicosanoate or arachidic acid methyl ester (C₂₁H₄₂O₂, CAS No. 1120-28-1), a 2-kg batch was custom-synthesized by Spectrum Chemical Mfg. Corp. (New Brunswick, New Jersey) with a purity of 99.9%. A fraction of 800 g of the original material termed icosanoic acid methyl ester (USGS70) was melted to a homogeneous liquid and dripped through a glass capillary into liquid nitrogen, where it shock-solidified to $\sim 2\text{-mg}$ of waxy beads with uniform isotopic composition. The remaining original ester was subdivided into two aliquots for adding medium and high ²H and ¹³C enrichments at the methyl group via transesterification with methan-²H-ol (i.e., ²H¹H₂COH, CAS No. 4206-31-9; ²H fraction = 98%, Aldrich) and methanol-¹³C (¹³C fraction = 99%, Aldrich). Each transesterified product was distilled under vacuum and only the center fractions were used to ultimately produce waxy beads as described above.

Multigram amounts of beads of each type were flame-sealed into glass containers under argon. The three different icosanoic acid methyl esters have increasing $\delta^2\text{H}$ and $\delta^{13}\text{C}$ values in the order from USGS70, USGS71, to USGS72 (Table 2). Users will receive aliquots of 100 mg of beads in glass vials as RMs for $\delta^2\text{H}$ and $\delta^{13}\text{C}$ normalization.

Glycines (USGS64, USGS65, USGS66). A total of 10 kg of glycine were purchased from Acros Organics ($\text{C}_2\text{H}_5\text{NO}_2$, CAS No. 56-40-6, purity $\geq 99\%$). Three kilograms were dissolved in ultrapure water and the homogeneous solution dripped through a glass capillary into liquid nitrogen where drops were flash frozen. Freeze-drying these pellets then produced fine-grained glycine (USGS64) with uniform isotopic composition. The two glycines enriched in ^2H , ^{13}C , and ^{15}N (USGS65 and USGS66) were prepared by small additions of isotopically spiked glycines to the unspiked original glycine from Acros Organics. The following spikes were purchased from ICON: (i) glycine-2,2-d $_2$, ^2H fraction = 98%; (ii) glycine-1- ^{13}C , ^{13}C fraction = 99%; (iii) glycine-2- ^{13}C , ^{13}C fraction = 99%; (iv) glycine- ^{15}N , ^{15}N fraction = 99%. The homogeneous solutions of glycines USGS65 and USGS66 in ultrapure water were dripped into liquid nitrogen and freeze-dried as described above. Aliquots of glycines were flame-sealed under vacuum into multiple round-bottom Pyrex flasks. ^1H NMR analyses at JAMSTEC indicated that $>99.99\%$ of total hydrogen in all three RMs is in glycine. Bulk $\delta^2\text{H}$, $\delta^{13}\text{C}$, and $\delta^{15}\text{N}$ values increase in the order from USGS64, USGS65, to USGS66 (Table 2). Glycines USGS65 and USGS66 feature contrasting ^{13}C -enrichments at molecular sites 1 and 2, namely, in an atom ratio 1:2 in USGS65 and 2:1 in USGS66 for future use in site-specific carbon isotopic measurements. Users will receive aliquots of 0.5 g in glass vials as RMs for $\delta^{13}\text{C}$ and $\delta^{15}\text{N}$ normalization. Note that the preliminary bulk $\delta^2\text{H}$ values are based on partially exchangeable hydrogen in glycine that may not be conservative upon handling in a moist atmosphere. The use of these RMs for normalization of $\delta^2\text{H}$ measurements will have to await the development of analytical methods that can reliably account for the influence of exchangeable hydrogen in individual amino acids.

L-Valines (USGS73, USGS74, USGS75). Four kilograms of L-valine ($\text{C}_5\text{H}_{11}\text{NO}_2$, CAS No. 72-18-4) were donated by the Amino GmbH (Frellstedt, Germany). The L-valine was mixed in an industrial-strength blender without adding any isotopic spikes, sieved through a 40-mesh-metal sieve, and was labeled USGS73. The two ^2H , ^{13}C , and ^{15}N -enriched L-valines USGS74 and USGS75 were prepared from small additions of isotopically spiked L-valines to amounts of 2 and 1.6 kg of unspiked L-valine from Alfa-Aesar, respectively. The following spikes were used: (i) L-valine-3-d $_1$, ^2H fraction = 98%, Cambridge Isotope Laboratories, Inc.; (ii) L-valine-2-d $_1$, ^2H fraction = 99%, CDN Isotopes; (iii) L-valine-2- ^{13}C , ^{13}C fraction $\geq 99\%$, Isotech Laboratories, Inc.; (iv) L-valine-1- ^{13}C , ^{13}C fraction = 99%, Cambridge Isotope Laboratories, Inc.; (v) L-valine- ^{15}N , ^{15}N fraction = 98%, Aldrich. Unlike glycine, the lower solubility of L-valine made it impractical to dissolve kilogram amounts in ultrapure water. Instead, for USGS74 and USGS75, only the small amounts of isotopic spikes were jointly dissolved in ultrapure water and added to powdered, water-wet regular L-valine. Each paste was stirred for 5 h and subsequently extruded as strings into liquid nitrogen. The frozen strings were freeze-dried, mixed in an industrial-strength blender, sieved through a 40-mesh-metal sieve, and crushed in a stainless-steel, refrigerated shatter box. Dry powder was finally tumbled for 12 weeks with

stainless steel balls under nitrogen in rotating metal cans until the medium-enriched L-valine USGS74 was demonstrated by analyses at the USGS in Reston, Virginia to be sufficiently homogeneous at the 0.1 mg level. In spite of tumbling for 6 months, the highly enriched L-valine USGS75 only reached acceptable homogeneity at the 0.2 mg level. In order to approach bulk isotopic mean values, 3 g of the tumbled L-valine no. 3 were dissolved in ultrapure water, dripped into liquid nitrogen, freeze-dried, and aliquots were sent to participating laboratories. The purity of L-valines was evaluated at JAMSTEC using ^1H NMR. The purity is 99.999% for USGS73 from the Amino GmbH, whereas 99.54% for USGS74 and 99.50% for USGS75 may reflect a combination of the different Alfa Aesar raw material batches, the added isotopic spikes, and the extended homogenization efforts. At 99.5% purity, the magnitude of contamination remains limited. Bulk $\delta^2\text{H}$, $\delta^{13}\text{C}$ and $\delta^{15}\text{N}$ values increase in the order from USGS73, USGS74, to USGS75 (Table 2). L-valines USGS74 and USGS75 feature contrasting ^2H - and ^{13}C -enrichments at different molecular sites (i.e., carbon atoms in positions 1, 2, and 3). Future methods for site-specific carbon and hydrogen isotopic characterization may find value in the USGS74 ^2H -spike ratio 2:1 for H-molecular positions 3 and 2 as well as ^{13}C -spike ratio 2:1 for C-molecular positions 2 and 1. In contrast, USGS75 has a ^2H -spike ratio 1:2 for H-molecular positions 3 and 2 and a ^{13}C -spike ratio 1:2 for C-molecular positions 2 and 1. Aliquots of L-valines were flame-sealed under vacuum into round-bottom Pyrex flasks. Users will receive aliquots of 0.5 g in glass vials as RMs for $\delta^{13}\text{C}$ and $\delta^{15}\text{N}$ normalization. As with the glycines, the preliminary $\delta^2\text{H}$ values are based on the total hydrogen, which includes partially exchangeable hydrogen that may not be conservative upon handling in a moist atmosphere. The use of these RMs for normalization of $\delta^2\text{H}$ measurements will have to await the development of analytical methods that can reliably account for the influence of exchangeable hydrogen in individual amino acids. Tumbled L-valine USGS75 has limited homogeneity below a sample size of 0.2 mg.

Caffeines (USGS61, USGS62, USGS63). In total, 2 kg of caffeine ($\text{C}_8\text{H}_{10}\text{N}_4\text{O}_2$, CAS No. 58-08-2) from Alfa Aesar with a purity of 99.5% (chromatographic purity $>99.9\%$) were mixed in an industrial-strength blender, sieved through a 20-mesh-metal sieve, and labeled caffeine USGS61. However, this caffeine was only moderately depleted in ^2H . It was not possible to find caffeine with a combined strong depletion in ^2H and ^{13}C because industrial decaffeination mixes coffees from different areas and yields isotopic ranges far smaller than those observed in natural caffeines.^{19,20} Instead, these three caffeines form a sequence USGS61–USGS62–USGS63 where only the abundances of ^{13}C and ^{15}N are increasing, whereas the sequence for increasing ^2H abundance is USGS62–USGS61–USGS63. Isotopically fractionated caffeines for spiking included (i) caffeine-d $_3$ (1-methyl-d $_3$), ^2H fraction = 99.8%, ICON Isotopes, (ii) caffeine 3-methyl- ^{13}C , ^{13}C fraction = 99%, Cambridge Isotope Laboratories, Inc.; (iii) caffeine 1,3- $^{15}\text{N}_2$, ^{15}N fraction = 99%, Cambridge Isotope Laboratories, Inc. Caffeine USGS62 contains 1.8 kg of ^2H -depleted caffeine from the Coffein Compagnie (Bremen, Germany) with added ^{13}C and ^{15}N -spiked caffeines. Caffeine USGS63 contains 900 g of caffeine from Alfa Aesar with added ^2H , ^{13}C , and ^{15}N -spiked caffeines. Similar to the preparative routes described for L-valines above, for caffeines USGS62 and USGS63 only the small amounts of isotopic spikes were jointly dissolved in ultrapure water and added to powdered, water-wet regular caffeine. Each paste was stirred for 5 h and

subsequently extruded as strings into liquid nitrogen. The frozen strings were freeze-dried, mixed in an industrial-strength blender, sieved through a 20-mesh-metal sieve, and crushed in a stainless-steel, refrigerated shatter box. Homogeneity was demonstrated to be satisfactory at the 0.1-mg level by the USGS in Reston, Virginia. All caffeines were tested to confirm the absence of crystal water, by heating preweighed aliquots in vacuum to 80 °C for at least 3 h and reweighing after cooling. No significant weight loss was detected. Aliquots of caffeines were flame-sealed under vacuum into multiple round-bottom Pyrex flasks. Users will receive aliquots of 0.5 g in glass vials as RMs for $\delta^2\text{H}$, $\delta^{13}\text{C}$, and $\delta^{15}\text{N}$ normalization. Note that caffeine can have a strong physiological effect and express toxicity.

Vacuum Oils (NBS 22a, USGS78). We purchased 19 L of Fisher Scientific Maxima C Plus Vacuum Pump Oil representing a triple-distilled hydrocarbon fraction that had been hydrogenated to reduce the abundance of aromatic components with exchangeable hydrogen and to increase chemical stability. The vapor pressure at 25 °C of 0.133 mPa guarantees the absence of volatile components. A volume of 4.3 L (i.e., 3.35 kg) of vacuum oil was enriched in ^2H by heating it to 60 °C and dissolving 250 mg of perdeuterated *n*-tetracosane ($\text{C}_{24}\text{H}_{50}$, ^2H fraction = 99.1%, MSD Isotopes) to yield a homogeneous solution. In total, 14 L of the original vacuum oil NBS 22a and 4.3 L of the ^2H -enriched vacuum oil USGS78 were flame-sealed under vacuum into multiple round-bottom Pyrex flasks for long-term storage. Users will receive 1 mL aliquots sealed in glass containers as RMs for $\delta^2\text{H}$ and $\delta^{13}\text{C}$ normalization.

Polyethylene Powder (USGS77) and Extruded String NDF-PE77. Custom-synthesized low density polyethylene powder with a grain size of ~ 0.1 mm was purchased from Alfa Aesar (10 kg, $(\text{CH}_2\text{CH}_2)_n$, CAS No. 9002-88-4). Multiple round-bottom Pyrex flasks were filled and flame-sealed under vacuum. Users will receive aliquots of 1 g in glass vials as RMs for $\delta^2\text{H}$ and $\delta^{13}\text{C}$ normalization. In total, 2 kg of the polyethylene powder were extruded to form a string with a diameter of ~ 1 mm in collaboration with the Australian Nuclear Science and Technology Organisation (ANSTO). Isotopic measurements at the USGS and Indiana University indicated that USGS77 powder and the string NDF-PE77 have indistinguishable isotopic compositions. Some laboratories may prefer string NDF-PE77 over powder if electrostatic dispersion of powder is problematic. String NDF-PE77 is available exclusively from ANSTO.²¹

Online and Off-Line Isotopic Analyses. Eleven participating laboratories used a variety of analytical approaches for measuring $\delta^2\text{H}$, $\delta^{13}\text{C}$, and $\delta^{15}\text{N}$ values (Table 1). The isotopic homogeneity of new RMs was tested and confirmed at the 0.1 mg level, except for L-valine no. 3 USGS75 where homogeneity required 0.2 mg-aliquots. Ten laboratories used online methodologies based on a variety of elemental analyzer (EA) and gas chromatographic (GC) interfaces as well as mass spectrometers from different manufacturers. All laboratories used combinations of primary and secondary international measurement standards VSMOW2, SLAP2, NBS 19, NBS 22, LSVEC, USGS32, USGS40, USGS41, IAEA-N-1, IAEA-N-2, IAEA-CH-7, and IAEA-600 for two-point calibrations (normalization) of isotope δ -scales anchored by VSMOW, SLAP, VPDB, LSVEC, atmospheric nitrogen, and USGS32. An additional, new ^2H -enriched reference water IAEA-604 with $\delta^2\text{H} = +799.9$ mUr was used for normalization of RMs having positive $\delta^2\text{H}$ values.²⁸ Waters and oils were crimp-sealed into segments of silver tubing for use in online EA-interfaces.¹³ The use of reference waters in silver tubes made it possible to directly measure $\delta^2\text{H}$ values of

organic materials against known $\delta^2\text{H}$ values of reference waters²⁹ following the principle of identical treatment.¹⁰

For online measurements of $\delta^2\text{H}$ values in nitrogen-containing organic compounds with an EA interface, the conventional high-temperature conversion in a glassy carbon environment (also termed TC/EA) has recently been found to fractionate hydrogen isotopes due to the generation of HCN as a pyrolytic byproduct.^{23,27} The use of chromium for high-temperature conversion of organic hydrogen to gaseous H_2 eliminated the bias both for EA²³ and GC interface applications²⁴ and resulted in online $\delta^2\text{H}$ values that were comparable with off-line measurements at Indiana University. The traditional off-line combustion at Indiana University of milligram amounts of organic material with copper oxide (CuO) at 800 °C in evacuated and sealed “quartz” ampules overnight offers sufficient time at high temperature to yield quantitatively H_2O , CO_2 , and N_2 after cooling to 300 °C. Immediately after combustion, after cooling to 200 °C, individual ampules with break-seals were transferred to the inlet system of a glass vacuum line. Cryogenic separation of combustion products in a vacuum line yielded pure N_2 (collected with a Toepler pump and flame-sealed in a glass tube) and CO_2 (collected and sealed in a glass tube at the temperature of liquid nitrogen). A subsequent step of reducing H_2O to H_2 in the presence of uranium metal at 800 °C is a classical method for preparation of H_2 for stable isotope mass spectrometry³⁰ and was applied uniformly to reference waters and for waters from combustion of organics.³¹ We found no difference in isotopic composition between reference waters that were either “pre-combusted” with CuO in quartz glass or directly admitted to hot uranium in the vacuum line.

Complete oxidative combustion of organic materials was used by all laboratories to generate CO_2 and N_2 analyte gases in online or off-line modes for measurements of $\delta^{13}\text{C}$ and $\delta^{15}\text{N}$ values. At Indiana University, CO_2 reference gases were prepared from NBS 19 and LSVEC carbonates with anhydrous phosphoric acid at 25 °C, followed by additional “combustion” of the CO_2 with CuO at 800 °C in evacuated and sealed “quartz” ampules overnight to match the oxygen stable isotopic composition of CO_2 from combusted organic materials.

■ RESULTS

Only 2-point-normalized data were accepted. Additional data, including averages and standard deviations, and associated graphs are provided in the [Supporting Information](#). All 2-point-normalized $\delta^2\text{H}$ values were accepted except for those that had been measured online with high-temperature-conversion methodologies (e.g., conventional TC/EA) for nitrogen-containing compounds in the absence of chromium to scavenge nitrogen. All $\delta^{13}\text{C}$ values were accepted except for the measurements of four materials from a single laboratory that had used peak jumping when measuring $\delta^{13}\text{C}$ and $\delta^{15}\text{N}$ values in the same runs. We eliminated these four sets of $\delta^{13}\text{C}$ values as outliers (see [Supporting Information](#)). We also eliminated the $\delta^{15}\text{N}$ values of one material from a single laboratory due to outlier status. All accepted data were used for Bayesian Random Effects statistical analysis to yield mean values with standard uncertainties and expanded uncertainties ($\sim 95\%$ uncertainty bounds; see description of statistical details in the [Supporting Information](#)).³² There was one exception to the statistical treatment; because the $\delta^{13}\text{C}$ values of the regular vacuum oil NBS 22a and the ^2H -enriched vacuum oil USGS78 are known to be essentially identical, the results for these two materials were harmonized as described in section 2 of the [Supporting Information](#).

Table 1. Stable Isotope Approaches and Equipment of Participating Laboratories^a

Laboratory	$\delta^2\text{H}$ measurements	$\delta^{13}\text{C}$ and $\delta^{15}\text{N}$ measurements
California Institute of Technology, Pasadena, California, USA	Online, Thermo Scientific Delta ^{plus} XP, Thermo Trace GC-Ultra gas chromatograph connected via Thermo GC/C pyrolysis interface, operated at 1440 °C. ²²	n.a.
ETH Zürich, Switzerland	Online, Costech Zero-Blank autosampler, HEKAtech HTO with reversed flow, SiC outer tube and glassy carbon inner tube with glassy carbon chips without graphite crucible; reactor 1450 °C, GC 40 °C, He flow ~90 mL min ⁻¹ ; trap filled with charcoal, Ascarite and Mg(ClO ₄) ₂ between reactor tube and GC column; connected via Thermo ConFlo III interface to Thermo Scientific Delta ^{plus} XP.	n.a.
Helmholtz Centre Leipzig, Germany	Online, Euro 3000 Elemental Analyzer (EuroVector, Italy), oven filled with chromium, one-oven system, 1050 °C, He flow ~100 mL min ⁻¹ via ConFlo IV connected to Thermo Scientific MAT 253. ^{23,24}	Online, Euro 3000, two-oven system, via ConFlo IV connected to Thermo Scientific MAT 253; peak jump from N ₂ to CO ₂ during each run; combustion at 1020 °C, reduction at 600 °C, GC 60 °C, He flow 100 mL min ⁻¹ .
Helmholtz Centre Munich, Germany	Online, thermal decomposition by pyrolysis (1480 °C/SiC-tube filled with glassy carbon and ~1 g nickel-coated graphite), He flow 90 mL min ⁻¹ . Separation column at 90 °C, Thermo ConFlo II Interface, Thermo Scientific MAT 253.	Online, combustion by EA (EuroVector) via ConFlo II Interface, Thermo Scientific MAT 253; oxidation 1000 °C, reduction 600 °C. Separation column at 85 °C. He flow 70 mL min ⁻¹ . Measurements of $\delta^{13}\text{C}$ and $\delta^{15}\text{N}$ via peak jumping.
Indiana University, Bloomington, USA	Off-line combustion in evacuated and sealed SiO ₂ ampules with CuO, Cu, Ag at 800 °C, reduction of H ₂ O to H ₂ with uranium at 800 °C on vacuum line, purification and collection of analyte gases H ₂ , CO ₂ , and N ₂ on vacuum line, manual dual-inlet measurements with Thermo Scientific Delta XP IRMS.	Online, Thermo Scientific Delta ^{plus} XP IRMS coupled to Flash EA1112 via ConFlo III; oxidation 1050 °C, reduction 750 °C.
JAMSTEC, Japan	n.a.	Online, Thermo Scientific Delta ^{plus} XP IRMS coupled to Flash EA1112 via ConFlo III; oxidation 1050 °C, reduction 750 °C.
Max-Planck-Institute, Jena, Germany	Online, Thermo Scientific Delta ^{plus} XL, HEKAtech HTO with reversed flow connected via Thermo ConFlo III interface. SiC outer tube and glassy carbon inner tube filled with glassy carbon chips without graphite crucible. Reactor at 1450 °C, GC 90 °C, helium flow 60 mL min ⁻¹ . Later, the upper glassy carbon chip layer was mixed with chromium chips. ²⁷	Online, Thermo Scientific Delta ^{plus} XP, CE 1100 EA connected via ConFlo III interface. Oxidation 1050 °C, reduction 620 °C, GC 40 °C. He flow 90 mL min ⁻¹ . ^{25,10,26}
James Hutton Institute, Dundee, United Kingdom	Online, Thermo Scientific Delta ^{plus} XP coupled to TC/EA via ConFlo III; TC/EA reactor: ceramic tube with inner glassy carbon tube filled with sieved glassy carbon chips and graphite crucible; TC/EA 1425 °C; GC 85 °C; He flow 90 mL min ⁻¹ .	Online, Thermo Scientific Delta V Advantage coupled to Flash 1112 EA via ConFlo IV; combustion at 980–1000 °C; reduction 640 °C; GC 42 °C; He flow 100 mL min ⁻¹ .
Université du Québec, Canada	n.a.	Online, continuous flow, no peak jumping, via either (i) Isoprime 100, Elementar Vario MicroCube, or (ii) Isoprime Vision and Elementar Vario PyroCube.
University of Groningen, The Netherlands	Online, GVI Isoprime. <i>Method No. 1</i> : glassy carbon tube with glassy carbon chips, 1450 °C. <i>Method No. 2</i> : quartz tube filled with Patinal Cr powder at 1030 °C; HekaTech HTC with zero-blank autosampler, coupled to GVI Isoprime via open split "ref box"; GC at 90 °C. He flow 90 mL min ⁻¹ , top-feed arrangement.	Online, Elementar Isotope Cube EA in CN arrangement with zero-blank autosampler, connected with open split reference box to Isoprime 100; combustion at 980 °C, reduction at 650 °C, He flow 240 mL min ⁻¹ .
U.S. Geological Survey, Reston, Virginia, USA	Online, Thermo Scientific Delta ^{plus} XP, Thermo Scientific TC/EA connected via Thermo ConFlo IV interface. <i>Method No. 1</i> : Glassy carbon tube filled with glassy carbon chips without graphite crucible. Reactor 1450 °C, GC 90 °C, He flow 120 mL min ⁻¹ . <i>Method No. 2</i> : Removed glassy carbon tube. Filled ceramic tube from bottom to top with glassy carbon chips, quartz wool and 3 cm chromium powder. Reactor 1150 °C, GC 90 °C, He flow 120 mL min ⁻¹ .	Online, Thermo Scientific Delta V Plus, Costech EA connected via ConFlo II interface; oxidation 1020 °C, reduction 650 °C, GC 60 °C, He flow 110 mL min ⁻¹ .

^an.a. = not applicable.

Table 2. Alphabetical Listing of New and Pre-Existing (*) Organic Reference Materials with Bayesian Random Effects Mean Values with Associated Combined Standard Uncertainties^a

Reference ID	Chemical name	Structure or composition of material	Ring-test reference values with combined standard uncertainties (mUr or ‰)		
			$\delta^2\text{H}_{\text{VSMOW-SLAP}}$	$\delta^{13}\text{C}_{\text{VPDB-LSVEC}}$	$\delta^{15}\text{N}_{\text{Air}}$
USGS61	caffeine		+96.9 ± 0.9	−35.05 ± 0.04	−2.87 ± 0.04
USGS62	caffeine		−156.1 ± 2.1	−14.79 ± 0.04	+20.17 ± 0.06
USGS63	caffeine		+174.5 ± 0.9	−1.17 ± 0.04	+37.83 ± 0.06
IAEA-600*	caffeine		−156.1 ± 1.3	−27.73 ± 0.04	+1.02 ± 0.05
USGS64	glycine		no values indicated due to the presence of exchangeable hydrogen	−40.81 ± 0.04	+1.76 ± 0.06
USGS65	glycine			−20.29 ± 0.04	+20.68 ± 0.06
USGS66	glycine			−0.67 ± 0.04	+40.83 ± 0.06
USGS67	<i>n</i> -hexadecane	$\text{C}_{16}\text{H}_{34}$	−166.2 ± 1.0	−34.50 ± 0.05	n.a.
USGS68	<i>n</i> -hexadecane		−10.2 ± 0.9	−10.55 ± 0.04	n.a.
USGS69	<i>n</i> -hexadecane		+381.4 ± 3.5	−0.57 ± 0.04	n.a.
USGS70	icosanoic acid methyl ester (C_{20} FAME)	$\text{C}_{20}\text{H}_{39}\text{OOCH}_3$	−183.9 ± 1.4	−30.53 ± 0.04	n.a.
USGS71	icosanoic acid methyl ester (C_{20} FAME)		−4.9 ± 1.0	−10.50 ± 0.03	n.a.
USGS72	icosanoic acid methyl ester (C_{20} FAME)		+348.3 ± 1.5	−1.54 ± 0.03	n.a.
USGS73	L-valine		no values indicated due to the presence of exchangeable hydrogen	−24.03 ± 0.04	−5.21 ± 0.05
USGS74	L-valine			−9.30 ± 0.04	+30.19 ± 0.07
USGS75	L-valine			+0.49 ± 0.07	+61.53 ± 0.14
USGS76	methylheptadecanoate (C_{17} FAME)	$\text{C}_{17}\text{H}_{33}\text{OOCH}_3$	−210.8 ± 0.9	−31.36 ± 0.04	n.a.
IAEA-CH-7*	polyethylene foil	$(\text{C}_2\text{H}_4)_n$	−99.2 ± 1.2	−32.14 ± 0.05	n.a.
USGS77	polyethylene powder (also extruded string)	$(\text{C}_2\text{H}_4)_n$	−75.9 ± 0.6	−30.71 ± 0.04	n.a.
NBS 22*	oil	n.a.	−117.2 ± 0.6	−30.02 ± 0.04	n.a.
NBS 22a	vacuum oil, regular	n.a.	−120.4 ± 1.0	−29.72 ± 0.04	n.a.
USGS78	vacuum oil, ^2H -enriched	n.a.	+397.0 ± 2.2	−29.72 ± 0.04	n.a.

^aThe unit mUr is synonymous with deprecated ‰; n.a., not applicable; FAME, fatty acid methyl ester. The $\delta^2\text{H}_{\text{VSMOW-SLAP}}$ values are normalized to a scale in which $\delta^2\text{H}$ of SLAP = −428 mUr; the $\delta^{13}\text{C}$ values are normalized to a scale in which NBS 19 is +1.95 mUr and LSVEC is −46.6 mUr; the $\delta^{15}\text{N}$ values are normalized to a scale in which the $\delta^{15}\text{N}$ of atmospheric nitrogen = 0 mUr and that of USGS32 = +180 mUr.

A statistical overview of mean δ values from a Bayesian Random Effects model approach with combined standard uncertainty is presented in Table 2. Multiple graphs in the Microsoft Excel file (see Supporting Information) document the typically excellent agreement of data among participating laboratories and testify to the group's analytical agreement as a consequence of shared primary and secondary RMs, strict adherence to 2-point normalization, and high levels of analytical proficiency. Combined standard uncertainties take into account not only measurement uncertainties from Bayesian statistical analysis of our group's combined results but additionally include published uncertainties of δ values from RMs that were used for normalization, for example, VSMOW2 and SLAP2.⁸ A primary reason that uncertainties are larger is that most laboratories used materials for normalization other than anchor materials, and many of these materials had significant uncertainty associated with them. These are shown in Tables S3–S5 of the Supporting Information Excel file. The inclusion of these sources of uncertainty enlarged our confidence intervals in comparison to earlier ring-tests.^{33,17} Some primary international RMs like NBS 19 and VSMOW define anchors and fixed points along isotope- δ scales and by definition have zero uncertainty.⁷ There is evidence that USGS41 L-glutamic acid is hygroscopic, likely due to the presence of pyroglutamic acid,³⁴ and its measured $\delta^{13}\text{C}$ value can be too low; therefore, the

uncertainty of USGS41 used for normalization has been increased from the value of 0.05 mUr⁸ to 0.25 mUr. There is evidence that LSVEC can increase in $\delta^{13}\text{C}$ value, perhaps by reaction with atmospheric carbon dioxide,³⁴ and the uncertainty of LSVEC used for normalization has been increased from 0 mUr¹⁷ to 0.15 mUr.

The Bayesian Random Effects mean δ values of organic RMs are presented for $\delta^2\text{H}_{\text{VSMOW-SLAP}}$, $\delta^{13}\text{C}_{\text{VPDB-LSVEC}}$, and $\delta^{15}\text{N}_{\text{Air}}$ (Figure 1A–C) in ascending order with assigned combined standard uncertainties (mUr) and range from −210.8 to +397.0 mUr for $\delta^2\text{H}_{\text{VSMOW-SLAP}}$, from −40.81 to +0.49 mUr for $\delta^{13}\text{C}_{\text{VPDB-LSVEC}}$, and from −5.21 to +61.53 mUr for $\delta^{15}\text{N}_{\text{Air}}$. A modified, alternative assessment scheme for the calculation of combined standard uncertainties for these new RMs is discussed in component 6 of the Supporting Information.

DISCUSSION

The principle of identical treatment of sample and RM¹⁰ was followed by each participating laboratory according to their own interpretation. Some laboratories employed identical sample preparation and analysis conditions (e.g., the same microbalances, tin or silver capsules from the same batch, concurrent analyses of RMs and quality control samples within the same EA or GC batch run with unknowns), while other laboratories additionally

Figure 1. (A) Bayesian Random Effects of mean $\delta^2\text{H}_{\text{VSMOW-SLAP}}$ values 22 organic RMs in ascending order with assigned combined standard uncertainties (mUr). The $\delta^2\text{H}_{\text{VSMOW-SLAP}}$ values of L-valines USGS74 and USGS75 are rough estimates without assigned uncertainties due to limited data. The $\delta^2\text{H}_{\text{VSMOW-SLAP}}$ values of amino acids have not been corrected for exchangeable hydrogen. Assigned combined uncertainties of amino acids do not reflect isotopic interference by geographically and seasonably variable air humidity and isotopic composition. (B) Bayesian Random Effects mean $\delta^{13}\text{C}_{\text{VPDB-LSVEC}}$ values of 22 organic RMs in ascending order with assigned combined uncertainties (mUr). (C) Bayesian Random Effects mean $\delta^{15}\text{N}_{\text{Air}}$ values of 10 organic RMs in ascending order with assigned combined standard uncertainties (mUr).

adhered to the notion that RMs, quality control samples, and unknowns should reflect a chemical matrix match with similar chemical compositions. For example, some laboratories used the carbonates NBS 19 and LSVEC for two-point $\delta^{13}\text{C}$ scale normalization of online EA measurements, whereas others used organic RMs such as USGS40, USGS41, and IAEA-CH-7. The remarkable agreement of data from multiple, independent approaches testifies to the practicality and suitability of careful laboratory practices, both online and off-line.

In addition to 19 new organic RMs, our ring-test recalibrated 3 older organic RMs NBS 22 (oil), IAEA-CH-7 (polyethylene foil), and IAEA-600 (caffeine) and arrived at new δ values (with tightened confidence intervals and lower combined standard uncertainties) that supersede earlier recommended values.^{35,36,8} The quality of the results is evidenced by the $\delta^{13}\text{C}_{\text{VPDB-LSVEC}}$ value of NBS 22 of -30.02 ± 0.04 mUr (Table 2) compared to the earlier value of -30.03 ± 0.05 mUr.¹⁷

The hydrogen exchange of organic hydrogen in functional groups can currently not be controlled satisfactorily in monomeric amino acids to warrant the recommendation of bulk hydrogen $\delta^2\text{H}$ values for glycine and L-valine RMs. We refrain from listing any bulk $\delta^2\text{H}$ values of amino acids in Table 2 for RM purposes, but the few preliminary bulk $\delta^2\text{H}$ values on nonequilibrated amino acids graphed on a coarse scale (Figure 1A) demonstrate that the ^2H -spiking of USGS65, USGS66, USGS74, and USGS75 in molecular positions bound to carbon atoms was successful. This bodes well for refined and reproducible hydrogen-isotopic characterization and use as RMs by future analytical techniques capable of measuring only isotopically nonexchangeable hydrogen in monomeric amino acids. We note that the much higher abundance of exchangeable hydrogen in monomeric amino acids poses more difficult challenges than for analytically less demanding polypeptides, such as collagen and keratin, where the formation of peptide bonds has eliminated much of the original exchangeable hydrogen from amino acid building blocks.

Some glycine and L-valine RMs have been isotopically spiked with ^2H (bonded to carbon) and ^{13}C at specific molecular positions resulting in distinct, contrasting abundances of molecular isotopologues. Future analytical techniques capable of discerning the intramolecular isotopic distribution at natural ^2H and ^{13}C abundances (e.g., high-resolution NMR)³⁷ may take advantage of this additional, built-in quality of these new RMs. We note that problems in an earlier laboratory intercomparison of hydrogen isotope abundances measured via IRMS and site-specific natural isotope fractionation measured by nuclear magnetic resonance (SNIF-NMR) have been likely caused by incomplete hydrogen yields during high-temperature conversion without the use of chromium.³⁸

Logistic and administrative constraints precluded the development of RMs with characteristics that may complicate international distribution, such as hydrocarbon gases, mixtures of compounds in a common organic solvent, and highly toxic substances.

Organic RMs are sensitive to UV radiation and heat. Although none of the new RMs expresses significant volatility (except for hexadecane and FAMES under vacuum), and only amino acids have a small propensity for slow oxidation, it is strongly recommended that refrigerated, dark, and dry storage be used for all organic RMs. Nonpolymeric RMs are subject to microbial degradation in the presence of moisture. Glass containers should be warmed to room temperature prior to opening in order to avoid condensation of atmospheric water.

CONCLUSIONS

Nineteen new organic reference materials for hydrogen, carbon, and nitrogen stable isotope-ratio measurements have been developed and analyzed using primary and secondary hydrogen, carbon, and nitrogen isotopic reference materials by an international group of laboratories and will be distributed jointly by the USGS,³⁹ IAEA,⁴⁰ and Indiana University.⁴¹ Polyethylene string NDF-PE77 is available from ANSTO.²¹ Revised δ values are being recommended for pre-existing reference materials NBS 22 (oil), IAEA-CH-7 (polyethylene foil), and IAEA-600 (caffeine). The use of reference waters VSMOW2, SLAP, and IAEA-604 in silver tubes made it possible to directly calibrate $\delta^2\text{H}$ values of organic materials against known $\delta^2\text{H}$ values of reference waters following the principle of identical treatment. The available organic reference materials include various compound classes to facilitate adherence to the principle of identical treatment of organic sample and reference material. It is recommended that efforts be invested in the future to check for heterogeneity of prepared aliquots for routine distribution. It may also be appropriate to revisit the scheme for reference material uncertainty evaluation in light of emerging new conventions and statistical approaches (details provided in component 6 of the [Supporting Information](#)).

Recommended δ values of reference materials are subject to occasional revision, which may not result in prompt updates on Web sites of distributing agencies. Each user of isotopic reference materials is encouraged to consult the Web site of the Commission on Isotopic Abundances and Atomic Weights⁴² (CIAAW) as a dedicated and up-to-date source of information about current δ values of isotopic reference materials.

Future analytical techniques capable of (i) distinguishing between molecular isotopologues with site-specific ^2H and ^{13}C enrichments and (ii) measuring only isotopically nonexchangeable hydrogen in amino acids will take advantage of the fact that some of the new glycine and L-valine reference materials have been spiked at distinct molecular positions with ^2H (bonded to carbon) and ^{13}C . Refrigerated, dark, and dry storage is strongly recommended for organic RMs.

ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge on the [ACS Publications website](#) at DOI: [10.1021/acs.analchem.5b04392](https://doi.org/10.1021/acs.analchem.5b04392).

Table S75 offering details in addition to [Table 1](#) of the main publication; method to incorporate uncertainty of measurement results and uncertainty of each substance used for normalization to calculate a combined uncertainty for each of the 22 materials and 11 laboratories in this study; Bayesian statistical analysis of stable isotope data to arrive at mean values; statistical model for combining $\delta^{13}\text{C}$ data from regular and ^2H -enriched vacuum oils in order to arrive at identical $\delta^{13}\text{C}$ values for both oils; comparison of a user's measurement with the RM δ -value; statistical model to calculate combined standard uncertainties for property values of the new RMs ([PDF](#))

Tables S1–S74 (i.e., sheets) offering a comprehensive listing of all data with detailed graphic overviews (with color) ([XLSX](#))

Tables S1–S74 (i.e., sheets) offering a comprehensive listing of all data with detailed graphic overviews (without color) ([XLSX](#))

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Notes

The authors declare no competing financial interest.

ACKNOWLEDGMENTS

We are indebted to Marilyn Fogel, Libby Stern, and Michael Wieser for assisting with the initiation of this project. Stefanie Poppenhäger (Amino GmbH, Frellstedt, Germany; www.amino.de) donated pure L-valine. Tetyana Gilevska, Henk Jansen, Steffen Kümmel, and Harald Lowag provided analytical assistance. The manuscript benefited from constructive reviews by L. I. Wassenaar, Linda Stalker, and two anonymous reviewers. This project was supported by U.S. National Science Foundation Grant EAR-1052927. The National Deuteration Facility is partly supported by the National Collaborative Research Infrastructure Strategy (NCRIS), an initiative of the Australian Government. The support of the U.S. Geological Survey National Research Program made this report possible. Any use of trade, firm, or product names is for descriptive purposes only and does not imply endorsement by the U.S. Government.

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