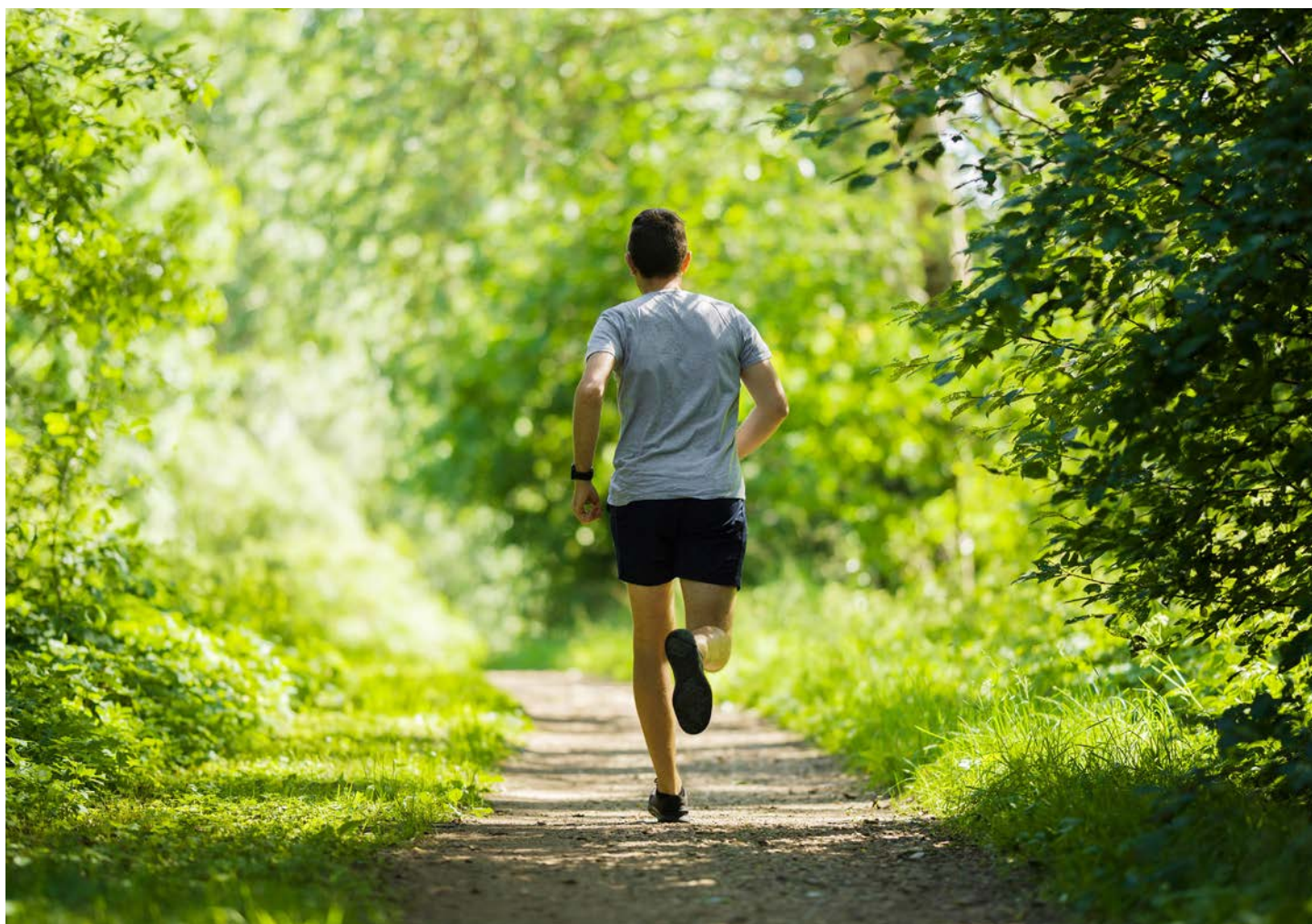




Cambridge Isotope Laboratories, Inc.
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RESEARCH PRODUCTS

Stable Isotope-Labeled Mixtures, Sets, and Kits For Mass Spectrometric Applications



Cambridge Isotope Laboratories, Inc.

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Table of Contents

Mixtures and Sets

Class-Specific Mixes

Amino Acid Mixes and Sets	3
Bile Acid Mixes and Sets	10
Carnitine/Acylcarnitine Mixes and Sets	13
Fatty Acid Mixes and Extracts	16
Nucleic Acid Sets	17
Organic Acid Mixes	19
Other Mixes and Sets	21
Steroid Mixes and Sets	25

Pathway-Specific Mixes

TCA Mixes and Sets	27
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Kits

Metabolomic QC Kits	29
<i>E. coli</i> and Yeast Extract Kits	35
PeptiQuant™ Plus Assay Kits	43
INLIGHT® Glycan Tagging Kit	46

Q&As	47
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The global and targeted measurement of biomolecules continues to be two areas of growing focus in analytical chemistry and biomedicine. Drivers for this research include efforts to better understand the underlying mechanisms of disease pathogenesis and to improve precision medicine through the qualitative/quantitative screening of candidate biomarkers. To address these objectives, mass spectrometry (MS)-based approaches are increasingly utilized and have been aided by advancements in experimental methodologies, instrumentation technologies, and bioinformatic tools.

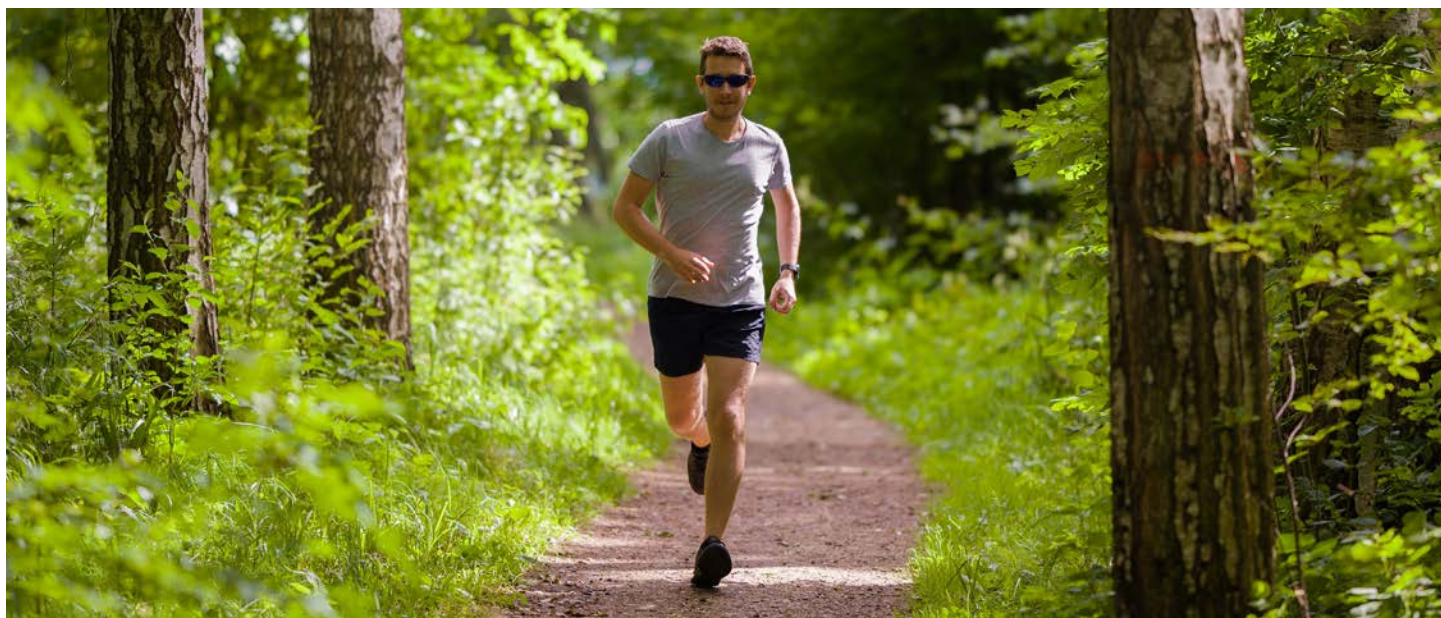
To help advance research in MS omics and MS/MS screening, Cambridge Isotope Laboratories, Inc. (CIL) is pleased to offer the largest variety of stable isotope-labeled mixtures, sets, and kits. The mixes are formulated neat and/or as solutions and are readily available for immediate use. In addition to the packaged mix(es), the kits include a user manual and shipping documents (e.g., certificate of analysis, safety data sheet). The manual outlines general procedures and processing examples for user reference, as well as troubleshooting notes and analysis results. Please see isotope.com for product details, pricing, and to inquire about customized mixes.

Benefits

- Easy to implement
- Offers end user flexibility
- Reduces development time and cost
- Enhances data quality
- Improves reproducibility
- Renders confidence in analytical results

Features Overview

- Mixes supplied in neat and/or solution form
- Preferentially ¹³C- and/or ¹⁵N-enriched
- Site-specific or uniform labeling
- Established specification guidelines
- Procedural guides in kit manuals
- Broad-spanning applications (from QC to quantification)



Mixtures and Sets

The mixtures available off-the-shelf are class- or pathway-specific for omics and MS/MS screening applications. These mixes are amenable for use in quality control and qualification/quantification exercises using targeted, semi-targeted, or untargeted MS-based methodologies. Outlined below is an overview of our current mix offerings, as well as details into their compositions and usage specifications (i.e., reconstitution guidelines, storage, and stability). For reference purposes, example results and published manuscripts are also provided.

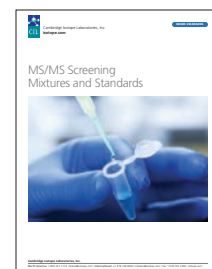
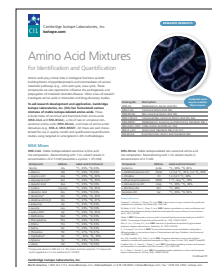
Class-Specific Mixes

Amino Acid Mixes and Sets

Amino acids (AAs) play critical roles in biological functions as both building blocks of peptides/proteins and intermediates of various metabolic pathways (e.g., citric acid cycle, urea cycle). These compounds are also reported to influence the pathogenesis and propagation of metabolic disorders/disease. To aid continued development and application, CIL has formulated a number of stable isotope-labeled (and unlabeled) AA mixtures. These include mixes of the canonical amino acids (MSK-A2 and MSK-CAA), the rare or unnatural non-canonical amino acids (MSK-NCAA), and a series of reference standard AA mixes (e.g., NSK-A).

Overview

Catalog No.	Description	No. of Metabolites	Unit Size
MSK-A2	Metabolomics Amino Acid Mix	17	1.2 mL
MSK-CAA	Canonical Amino Acid Mix	20	1 vial
MSK-NCAA	Non-canonical Amino Acid Mix	7	1 vial
MSK-CNCAA	Canonical/Non-canonical Amino Acid Mix Set	20 (CAA), 7 (NCAA)	2 × 1 vial
MSK-MSM1	Metabolomics Screening and Monitoring Standards Mix 1	14	1 vial
NSK-A	Amino Acid Standards Mix A	12	1 vial, 10 vials
NSK-A1	Amino Acid Standards Mix A1	12	1 vial, 10 vials
NSK-AB	Standards Mix Sets A and B	12 (A), 8 (B)	2 × 10 vials
NSK-A1B	Standards Mix Sets A1 and B	12 (A1), 8 (B)	2 × 10 vials
NSK-ABBG1	Standards Mix Sets A, B, and B-G1	12 (A1), 8 (B), 5 (B-G1)	3 × 10 vials
NSK-AA3	3-Plex Amino Acid Standards Mix	3	1 vial, 10 vials
NSK-AA3-10X	3-Plex Amino Acid Standards Mix (10x)	3	1 vial, 10 vials
NSK-BCAA	Branched-chain Amino Acid Standards Mix	4	1 vial



Click on the thumbnails or visit isotope.com for more information.

MSK-A2 and MSK-CAA Mixes

Compositions

The A2 mix (in 0.1 M HCl) consists of 17 amino acids (in solution), and the CAA mix (dried down) comprises 20. Reconstituting the CAA mix in 1 mL solvent (e.g., water) results in 2.5 mM concentrations (exception: L-cystine at 1.25 mM). **Note:** This CAA mix should not be reconstituted in 0.1 M HCl as Asn and Gln are unstable in acid.

Compound	Abbrev.	Label and Enrichment	Conc. (mM)
L-Alanine	Ala	$^{13}\text{C}_3$, 99%; ^{15}N , 99%	2.5
L-Arginine-HCl	Arg	$^{13}\text{C}_6$, 99%; $^{15}\text{N}_4$, 99%	2.5
L-Asparagine*	Asn	$^{13}\text{C}_4$, 99%; $^{15}\text{N}_2$, 99%	2.5
L-Aspartic acid	Asp	$^{13}\text{C}_4$, 99%; ^{15}N , 99%	2.5
L-Cystine	Cys-Cys	$^{13}\text{C}_6$, 99%; $^{15}\text{N}_2$, 99%	1.25
L-Glutamic acid	Glu	$^{13}\text{C}_5$, 99%; ^{15}N , 99%	2.5
L-Glutamine*	Gln	$^{13}\text{C}_5$, 99%; $^{15}\text{N}_2$, 99%	2.5
Glycine	Gly	$^{13}\text{C}_2$, 99%; ^{15}N , 99%	2.5
L-Histidine-HCl	His	$^{13}\text{C}_6$, 97%; $^{15}\text{N}_3$, 97%	2.5
L-Isoleucine	Ile	$^{13}\text{C}_6$, 99%; ^{15}N , 99%	2.5
L-Leucine	Leu	$^{13}\text{C}_6$, 99%; ^{15}N , 99%	2.5
L-Lysine-2HCl	Lys	$^{13}\text{C}_6$, 99%; $^{15}\text{N}_2$, 99%	2.5
L-Methionine	Met	$^{13}\text{C}_5$, 99%; ^{15}N , 99%	2.5
L-Phenylalanine	Phe	$^{13}\text{C}_9$, 99%; ^{15}N , 99%	2.5

Continued ►

Compound	Abbrev.	Label and Enrichment	Conc. (mM)
L-Proline	Pro	$^{13}\text{C}_5$, 99%; ^{15}N , 99%	2.5
L-Serine	Ser	$^{13}\text{C}_3$, 99%; ^{15}N , 99%	2.5
L-Threonine	Thr	$^{13}\text{C}_4$, 97%; ^{15}N , 97%	2.5
L-Tryptophan*	Trp	$^{13}\text{C}_{11}$, 99%; $^{15}\text{N}_2$, 99%	2.5
L-Tyrosine	Tyr	$^{13}\text{C}_9$, 99%; ^{15}N , 99%	2.5
L-Valine	Val	$^{13}\text{C}_5$, 99%; ^{15}N , 99%	2.5

*Compounds absent in MSK-A2.

Example Results

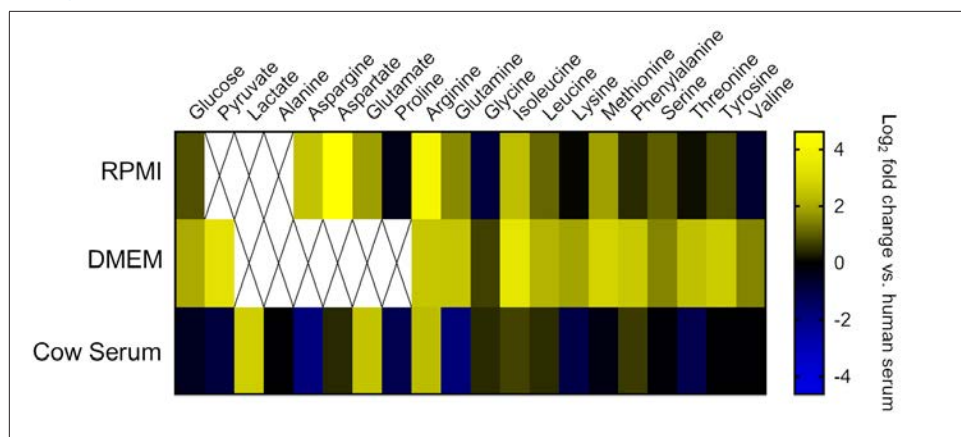


Figure. Application of the MSK-A2 mix to compare the quantitative profiles of cow serum and standard media (see PMID: 28826492 for additional information).

Example References

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- Muir, A.; Danai, L.V.; Gui, D.Y.; et al. **2017**. Environmental cystine drives glutamine anaplerosis and sensitizes cancer cells to glutaminase inhibition. *Elife*, 6, e27713.

Companion unlabeled standard mixes and kits may be available; please inquire.

Chemical purity (CP) is 98% or greater, unless otherwise indicated.

For research use only. Not for use in diagnostic procedures.

MSK-NCAA Mix

Composition

Reconstituting the dried-down NCAA mix with 1 mL solvent (e.g., 1:1 v/v methanol:water) results in 2.5 mM for each compound.

Compound	Abbrev.	Label and Enrichment
β-Alanine	β-Ala	¹³ C ₃ , 98%; ¹⁵ N, 96%
L-Azidohomoalanine-HCl	hAHA	1,2,3,4- ¹³ C ₄ , 99%; 2,4- ¹⁵ N ₂ , 98%
L-Citrulline	Cit	1,2,3,4,5- ¹³ C ₅ , 98%
L-Dihydroxyphenylalanine	DOPA	1- ¹³ C, ring- ¹³ C ₆ , 99%
L-Homoarginine-HCl	Harg	¹³ C ₇ , 98%; ¹⁵ N ₄ , 98%
L-Ornithine-HCl	Orn	¹³ C ₅ , 98%
Sarcosine-HCl	Sar	¹³ C ₃ , 99%; ¹⁵ N, 98%

“I use several of CIL’s metabolomics mixes in my LC-MS analysis. They make quantitative metabolomic work convenient with a single internal standard mix spike and provide the corresponding unlabeled mix for absolute quantitation. The ready mixes save time from the tedious task of making up individual solutions of each analyte for quantitation. I find the different mixes applicable to a variety of analyses in complicated matrices that provide metabolic insight to my studies.”

— Andrew Downey

Senior Scientist, Axcella Health Inc. (USA)

Usage Specifications

Mix Type	MSK-A2	MSK-CAA	MSK-NCAA
Form	1.2 mL solution	dried down	
Before reconstitution:			
Storage	2-8°C; protect from light		
Recommended retest	5 years from date of manufacture	2 years from date of manufacture	
Upon reconstitution:			
Storage	N/A	4°C; protect from light	
Recommended retest	N/A	4 weeks	

MSK-MSM1 Mix

Composition

Reconstituting a labeled MSM1 mix in 10 mL of solvent (e.g., water) yields an equimolar concentration of 500 μM. **Note:** Corresponding unlabeled mix has identical compound composition but yields a 10 mM equimolar concentration upon a 1 mL solvent addition.

Compound	Abbrev.	Label and Enrichment
L-allo-Isoleucine	allo-Ile	¹³ C ₆ , 97%
DL-2-Aminoadipic acid	Aad	2,5,5-D ₃ , 98%
DL-2-Aminobutyric acid	AABA	D ₆ , 98%
4-Aminobutyric acid	GABA	¹³ C ₄ , 97%
L-Argininosuccinic acid	ASA	arginine- ¹³ C ₆ , 99%; ¹⁵ N ₄ , 99%
DL-Cystathionine	Cyst	3,3,4,4-D ₄ , 98%
L-Cysteine, N-acetyl	NAC	cysteine- ¹³ C ₃ , 97%; ¹⁵ N, 97%
Ethanolamine	MEA	1,1,2,2-D ₄ , 98%
π-Methyl-L-histidine	1-MH	methyl-D ₃ , 98%
τ-Methyl-L-histidine	3-MH	methyl-D ₃ , 98%
DL-Homocystine	Hcys	3,3,3',3',4,4,4',4'-D ₈ , 98%
o-Phospho-L-serine	SEP	¹³ C ₃ , 99%; ¹⁵ N, 98%
o-Phosphorylethanolamine	PE	1,1,2,2-D ₄ , 98%
Taurine	T	¹³ C ₂ , 99%; ¹⁵ N, 98%

Usage Specifications

Criteria	Recommendation
Before reconstitution:	
Storage	2-8°C; protect from light
Recommended retest	2 years from date of manufacture
Upon reconstitution:	
Storage	-80°C; protect from light
Recommended retest	10 months*

*Metabolite stability demonstrated through 10 months with MSM1 mix solubilized in water stored at -80°C.

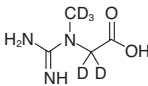
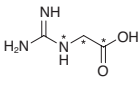
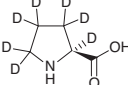
Companion unlabeled standard mixes and kits may be available; please inquire.

Chemical purity (CP) is 98% or greater, unless otherwise indicated.

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NSK-AA3 Mix**Composition**

Reconstituting the AA3 mix in 1 mL of solvent (e.g., 50% acetonitrile) will produce the tabulated concentrations below. **Note:** NSK-AA3-10X has 10x the specified concentrations.

Standard (Abbreviation)	Label and Enrichment	MW (Da)	Conc. (μM)	Structure
Creatine (Cre)	N-Methyl-D ₃ ; glycine-2,2-D ₂ , 99%	154.18	500	
Guanidinoacetic acid (GAA)	1,2- ¹³ C ₂ , 97%; 3- ¹⁵ N, 97% (CP 97%)	120.09	50	
L-Proline (Pro)	D ₇ , 97%	122.17	500	

Usage Specifications

Criteria	Recommendation
Before reconstitution:	
Storage	2-8°C; protect from light
Recommended retest	3 years from date of manufacture
After reconstitution:*	
Storage	-20°C
Recommended retest	3 months

*Represents minimum stability period when AA3 mix is reconstituted with 1:1 purified water:acetonitrile.

Companion unlabeled standard mixes and kits may be available; please inquire.

Chemical purity (CP) is 98% or greater, unless otherwise indicated.

For research use only. Not for use in diagnostic procedures.

NSK-A Mix

Composition

Reconstituting a given NSK-A vial's contents in 1 mL of high-purity solvent (1:1 water:methanol recommended) will produce the tabulated concentrations below. To facilitate complete dissolution, it is recommended to vortex manually for 1 minute then auto-vortex for a minimum of 30 seconds. **Note:** A combined set of NSK-A with the carnitine/acylcarnitine mixes NSK-B (NSK-AB) or NSK-B-G1 (NSK-ABBG1) is also available.

Compound	Abbrev.	Label and Enrichment	Conc. (μM)
L-Alanine	Ala	2,3,3,3-D ₄ , 98%	500
L-Arginine-HCl	Arg	5- ¹³ C, 99%; 4,4,5,5-D ₄ , 95%	500
L-Aspartic acid	Asp	2,3,3-D ₃ , 98%	500
L-Citrulline	Cit	5,5-D ₂ , 98%	500
DL-Glutamic acid	Glu	2,4,4-D ₃ , 98%	500
Glycine	Gly	2- ¹³ C, 99%; ¹⁵ N, 98%	2500
L-Leucine	Leu	5,5,5-D ₃ , 99%	500
L-Methionine	Met	methyl-D ₃ , 98%	500
L-Ornithine-HCl*	Orn	5,5-D ₂ , 98%	500
L-Phenylalanine	Phe	ring- ¹³ C ₆ , 99%	500
L-Tyrosine	Tyr	ring- ¹³ C ₆ , 99%	500
L-Valine	Val	D ₈ , 98%	500

*NSK-A1 contains Orn 3,3,4,4,5,5,-D₆, 98% instead of 5,5-D₂, 98%. The remaining components and concentrations are equivalent.

Usage Specifications

Criteria	Recommendation
No. of uses	960 samples/vial
Before reconstitution:	
Storage	≤25°C; protect from light
Recommended retest	4 years from date of manufacture
Upon reconstitution:	
Storage	2-8°C in a tightly sealed vial Note: Storing the sealed vial in a second sealed container helps maintain the integrity of the solution.
Recommended retest	4 weeks

Companion unlabeled standard mixes and kits may be available; please inquire.

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Example Results

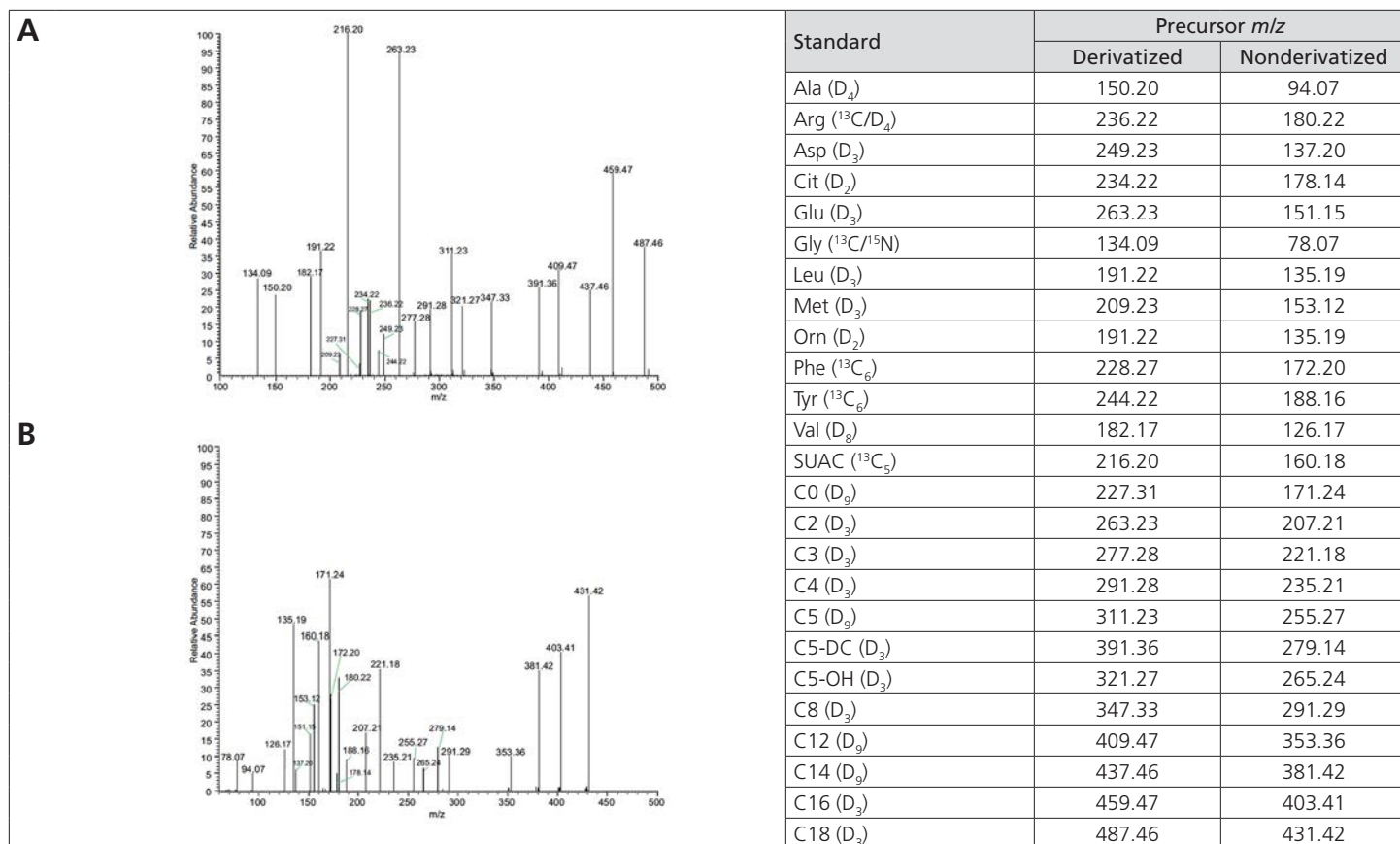


Figure. Full scan MS1 spectra of the DBS internal standards, with the derivatized standards shown in **A** and the nonderivatized in **B**. The standard mixture consists of stable isotope-labeled amino acids (NSK-A and -T) and carnitine/acylcarnitines (NSK-B and NSK-B-G). Please refer to Thermo Scientific technical note #73398 for method and application details.

Example References

- Li, S.; Huang, B.; Liu, M.; et al. **2022**. The association between leucine and diabetic retinopathy in different genders: a cross-sectional study in Chinese patients with type 2 diabetes. *Front Endocrinol*, 13, 806807-806815.
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Technical Note

Xie, X.; Kozak, M. **2020**. Simultaneous analysis of amino acids, acylcarnitines, and succinylacetone in dried blood spots for research using nonderivatized and derivatized methods. (Thermo Scientific technical note #73398)

“We have been using CIL’s NSK-A and NSK-B reference standards in our LC-MS/MS method for newborn screening tests since 2001. From that time more than 4,000,000 newborns have been screened. The products maintain good stability after dissolution and the high analysis repeatability enabled us to stabilize their use as internal standards. We appreciate the high-quality products, timely deliveries, and excellent customer relations.”

– Mariusz Oltarzewski

Head of Department of Screening Tests and Metabolic Diagnostics, Institute of Mother and Child (Poland)

Companion unlabeled standard mixes and kits may be available; please inquire.

Chemical purity (CP) is 98% or greater, unless otherwise indicated.

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NSK-BCAA Mix

Composition

Reconstituting the BCAA mix in 1 mL of solvent (e.g., 0.1 M HCl) will produce the tabulated concentrations below.

Standard (Abbreviation)	Label and Enrichment	Conc. (μM)	Q1 m/z	Q3 m/z
L-Allo-isoleucine (Alle)	$^{13}\text{C}_6$, 97%; ^{15}N , 97%	400	139.2	92.2
L-Isoleucine (Ile)	D_{10} , 98%	400	142.2	96.2
L-Leucine (Leu)	5,5,5- D_3 , 99%	400	135.2	89.2
L-Valine (Val)	$^{13}\text{C}_5$, 99%; ^{15}N , 99%	400	124.1	77.1

Usage Specifications

Criteria	Recommendation
Before reconstitution:	
Storage	$\leq 25^\circ\text{C}$; protect from light and moisture
Recommended retest	5 years from date of manufacture
After reconstitution:*	
Storage	4°C
Recommended retest	5 weeks

*Represents minimum stability period when the BCAA mix is reconstituted with 100% water.

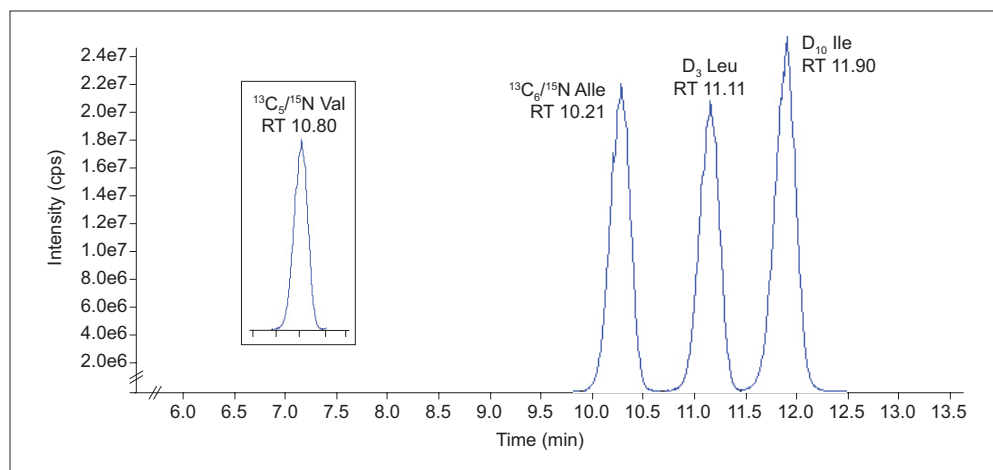


Figure. Representative XICs of the BCAA mix metabolites measured by RPLC-MRM/MS (ESI+, 5500+ QqQ). Separation of the isobaric metabolites is shown in the main plot, with Val in the inset. Displayed are the $[\text{M}+\text{H}]^+$ ions (see Table for precursor and product ion values).

Companion unlabeled standard mixes and kits may be available; please inquire.

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Bile Acid Mixes and Sets

Bile acids (BAs) are steroid-like compounds that act as a detergent in the breakdown of fats. This family of compounds comprises primary BAs (synthesized in the liver) and secondary BAs (produced in the gut by bacteria). These are essential regulatory compounds that are involved in various metabolic processes (e.g., cholesterol and lipid metabolism) and signaling interactions (e.g., in glucose and energy homeostasis). Investigations into their synthesis/ metabolism, disease linkage, and biomarker potential are examples of the type of research studies being undertaken. To aid further research and development efforts in this space, CIL has formulated stable isotope-labeled (and unlabeled) BA mixes. These dried-down mixes are constructed with the unconjugated and conjugated BAs in separate vials.

Overview

Catalog No.	Description	No. of Metabolites	Unit Size
MSK-BA1	Bile Acid Standards Mix 1 – Unconjugated	6	1 vial
MSK-BA1-25X	Bile Acid Standards Mix 1 (25x) – Unconjugated	6	1 vial
MSK-BA2	Bile Acid Standards Mix 2 – Conjugated	10	1 vial
MSK-BA2-20X	Bile Acid Standards Mix 2 (20x) – Conjugated	10	1 vial
MSK-BA-A	Bile Acid Standards Mix Sets 1 and 2	6 (BA1), 10 (BA2)	2 × 1 vial
MSK-BA3	Bile Acid Standards Mix 3 – Unconjugated Sulfates	6	1 vial
MSK-BA4	Bile Acid Standards Mix 4 – Conjugated Sulfates	10	1 vial

Note: Companion unlabeled mixes may be available; please inquire.

Compositions

Dissolving the standard BA-A mixes in 1 mL of solvent (e.g., 1:1 v/v methanol:water) yields equimolar concentrations of 100 µM, whereas reconstitution of the concentrated mixes (i.e., BA1-25X and BA2-20X) yields 1 mg/mL per analyte.

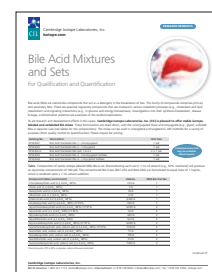
Unconjugated BA Mix (MSK-BA1)			
Compound	Abbrev.	Type of BA	Label and Enrichment
Chenodeoxycholic acid	CDCA	Primary	2,2,4,4-D ₄ , 98%
Cholic acid	CA	Primary	2,2,4,4-D ₄ , 98%
Deoxycholic acid	DCA	Secondary	2,2,4,4-D ₄ , 98%
Lithocholic acid	LCA	Secondary	2,2,4,4-D ₄ , 98%
β-Muricholic acid	β-MCA	Primary	2,2,3,4,4-D ₅ , 99%
Ursodeoxycholic acid	UDCA	Secondary	2,2,4,4-D ₄ , 98% (CP 95%)

Conjugated BA Mix (MSK-BA2)			
Compound	Abbrev.	Type of BA	Label and Enrichment
Glychenodeoxycholic acid	GCDCA	Primary	2,2,4,4-D ₄ , 98% (CP 97%)
Glycocholic acid	GCA	Primary	2,2,4,4-D ₄ , 98% (CP 96%)
Glycodeoxycholic acid	GDCA	Secondary	2,2,4,4-D ₄ , 98%
Glycolithocholic acid	GLCA	Secondary	2,2,4,4-D ₄ , 98%
Glycoursodeoxycholic acid	GUDCA	Secondary	2,2,4,4-D ₄ , 98% (CP 97%)
Taurochenodeoxycholic acid, sodium salt	TCDCA	Primary	2,2,4,4-D ₄ , 98% (CP 97%)
Taurocholic acid, sodium salt	TCA	Primary	2,2,4,4-D ₄ , 98%
Taurodeoxycholic acid, sodium salt	TDCA	Secondary	2,2,4,4-D ₄ , 98%
Tauroolithocholic acid, sodium salt	TLCA	Secondary	2,2,4,4-D ₄ , 98%
Tauroursodeoxycholic acid, sodium salt	TUDCA	Secondary	2,2,4,4-D ₄ , 98%

Companion unlabeled standard mixes and kits may be available; please inquire.

Chemical purity (CP) is 98% or greater, unless otherwise indicated.

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Compositions (continued)

Reconstituting each sulfated BA mix in 1 mL of solvent (e.g., 50% methanol) will produce an equimolar concentration of 100 µM.

Unconjugated Sulfated BA Mix (MSK-BA3)			
Compound	Abbrev.	Type of BA	Label and Enrichment
Chenodeoxycholic acid 3-sulfate, disodium salt	CDCA-3S	Primary	2,2,4,4-D ₄ , 98%
Cholic acid 3-sulfate, disodium salt	CA-3S	Primary	2,2,4,4-D ₄ , 98%
Deoxycholic acid 3-sulfate, disodium salt	DCA-3S	Secondary	2,2,4,4-D ₄ , 98%
Lithocholic acid 3-sulfate, disodium salt	LCA-3S	Secondary	2,2,4,4-D ₄ , 98%
β-Muricholic acid 3-sulfate, disodium salt	β-MCA-3S	Primary	2,2,3,4,4-D ₅ , 99%
Ursodeoxycholic acid 3-sulfate, disodium salt	UDCA-3S	Secondary	2,2,4,4-D ₄ , 98%

Conjugated Sulfated BA Mix (MSK-BA4)			
Compound	Abbrev.	Type of BA	Label and Enrichment
Glychenodeoxycholic acid 3-sulfate, disodium salt	GCDCA-3S	Primary	2,2,4,4-D ₄ , 98%
Glycocholic acid 3-sulfate, disodium salt	GCA-3S	Primary	2,2,4,4-D ₄ , 98%
Glycodeoxycholic acid 3-sulfate, disodium salt	GDCA-3S	Secondary	2,2,4,4-D ₄ , 98%
Glycolithocholic acid 3-sulfate, disodium salt	GLCA-3S	Secondary	2,2,4,4-D ₄ , 98%
Glycoursodeoxycholic acid 3-sulfate, disodium salt	GUDCA-3S	Secondary	2,2,4,4-D ₄ , 98%
Taurochenodeoxycholic acid 3-sulfate, disodium salt	TCDCA-3S	Primary	2,2,4,4-D ₄ , 98%
Taurocholic acid 3-sulfate, disodium salt	TCA-3S	Primary	2,2,4,4-D ₄ , 98%
Taurodeoxycholic acid 3-sulfate, disodium salt	TDCA-3S	Secondary	2,2,4,4-D ₄ , 98%
Taurolithocholic acid 3-sulfate, disodium salt	TLCA-3S	Secondary	2,2,4,4-D ₄ , 98%
Tauroursodeoxycholic acid 3-sulfate, disodium salt	TUDCA-3S	Secondary	2,2,4,4-D ₄ , 98%

Usage Specifications

Criteria	MSK-BA1 and MSK-BA2	MSK-BA3 and MSK-BA4
Before reconstitution:		
Storage	-20°C; protect from light	-20°C; protect from light
Recommended retest	2.5 years from date of manufacture	2 years from date of manufacture
After reconstitution:		
Storage	4°C	4°C
Recommended retest	4 weeks	

“Having a concentrated mix of isotopically labeled bile acid standards greatly increases the quality control within an assay. It eliminates having to weigh out each individual standard and increases precision/accuracy of batches over time while reducing human error during prep. It saves time with the ability to reconstitute one vial instead of having to individually prepare each labeled bile acid before creating an in-house mix. Alike CIL's standard labeled mixes, their concentrated mixes MSK-BA1-25X and MSK-BA2-20X are reproducibly prepared allowing for increased confidence in the detection/quantification of bile acids by LC-MS analysis.”

– Mary McMillin

Research Specialist, University of Chicago, Host-Microbe Metabolomics Facility (USA)

“The materials that Cambridge Isotope Laboratories (CIL) supply are always of the highest quality in terms of chemical and isotopic purity. Recently, we have embarked on two new major research areas involving the analyses of bile acids (BAs) and per- and polyfluorinated alkylsubstances (PFAS). The standards/mixes from CIL that enable these, and other, projects are precisely what we have come to expect – perfection!”

– David C. Muddiman, PhD

Jacob and Betty Belin Distinguished Professor,

Department of Chemistry, North Carolina State University

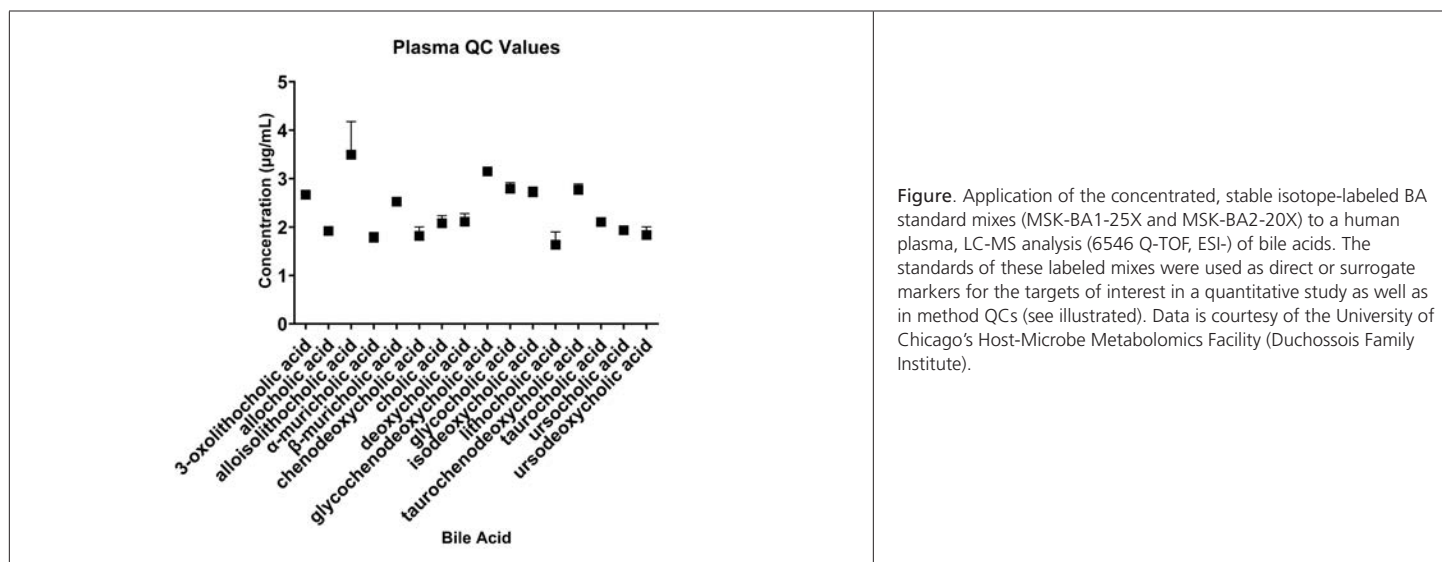
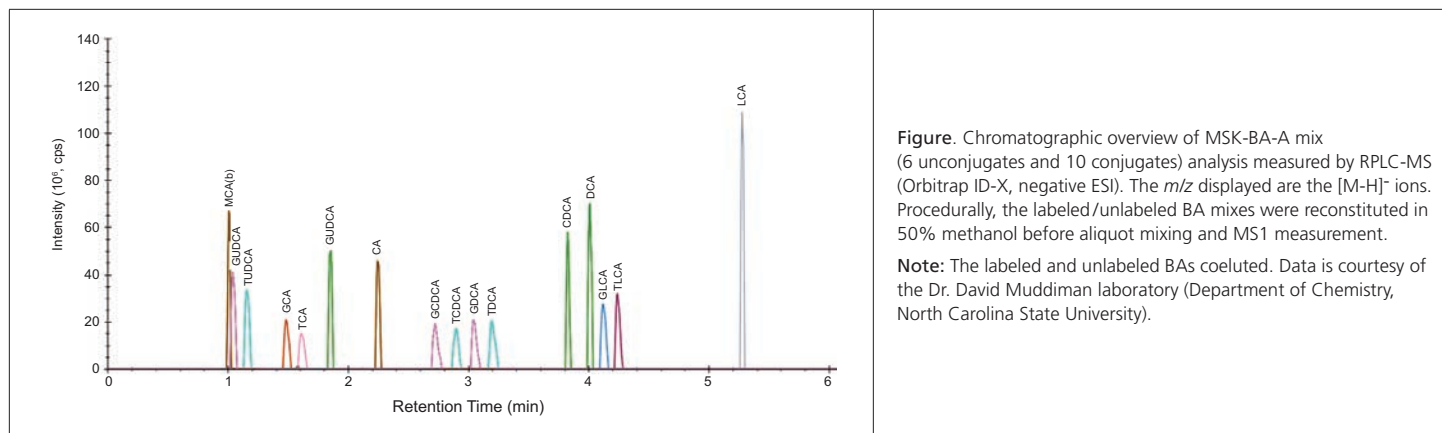
Director of Molecular Education, Technology, and Research Innovation Center (METRIC)

Companion unlabeled standard mixes and kits may be available; please inquire.

Chemical purity (CP) is 98% or greater, unless otherwise indicated.

For research use only. Not for use in diagnostic procedures.

Example Results



Example References for MSK-BA-A Mixes

- McMillan, A.S.; Zhang, G.; Dougherty, M.K.; et al. **2024**. Metagenomic, metabolomic, and lipidomic shifts associated with fecal microbiota transplantation for recurrent *Clostridioides difficile* infection. *mSphere*, 9(10), e0070624-e0070650.
- Engevik, M.E.; Thapa, S.; Lillie, I.; et al. **2024**. Repurposing dried blood spot (DBS) device technology to examine bile acid profiles in human dried fecal spot (DFS) samples. *Am J Physiol Gastrointest Liver Physiol*, 326(2), G95-G106.
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- Foley, M.H.; Walker, M.E.; Stewart, A.K.; et al. **2023**. Bile salt hydrolases shape the bile acid landscape and restrict *Clostridioides difficile* growth in the murine gut. *Nat Microbiol*, 8(4), 611-628.

Application Notes for MSK-BA-A Mixes

- Baker, P.R.S.; van Soest, R.; Kapil, S.; et al. **2025**. Quantitative and qualitative bile acid analysis on the ZenoTOF 8600 System using EAD. (SCIEX application note MKT-35189-A)
- Baker, P.R.S.; Proos, R.; Seferovic, M.D.; Horvath, T.D. **2023**. Quantitative analysis and structural characterization of bile acids using the ZenoTOF 7600 System. (SCIEX application note MKT-30747-A)
- Horvath, T.D.; Engevik, A.C.; Percy, A.J. **2022**. Bile acid analysis in mouse samples to study liver cholestasis etiology. (CIL application note #52)

Companion unlabeled standard mixes and kits may be available; please inquire.

Chemical purity (CP) is 98% or greater, unless otherwise indicated.

For research use only. Not for use in diagnostic procedures.

Carnitine/Acylcarnitine Mixes and Sets

Carnitine and acylcarnitines play an essential role in fatty acid metabolism. Metabolism disorders of fatty acid oxidation and several organic acidurias impose major clinical manifestations (e.g., hypoketotic hypoglycemia, skeletal myopathy, liver disease, and/or failure). These are largely attributed to enzymatic deficiencies and can be monitored through carnitine/acylcarnitine measurement. To aid MS/MS screening studies, CIL has formulated stable isotope-labeled (and unlabeled) standard carnitine/acylcarnitine mixes.

Overview

Catalog No.	Description	No. of Metabolites	Unit Size
NSK-B	Carnitine/Acylcarnitine Standards Mix B	8	1 vial, 10 vials
NSK-B-G1	Acylcarnitine Standards Mix 1 Supplement to NSK-B	5	1 vial, 10 vials
NSK-B-G2	Acylcarnitine Standards Mix 2 Supplement to NSK-B	5	1 vial, 10 vials
NSK-BBG1	Carnitine/Acylcarnitine Standards Mix Sets B and B-G1	8 (B), 5 (B-G1)	2 × 1 vial
NSK-AB	Standards Mix Sets A and B	12 (A), 8 (B)	2 × 10 vials
NSK-A1B	Standards Mix Sets A1 and B	12 (A1), 8 (B)	2 × 10 vials
NSK-ABBG1	Standards Mix Sets A, B, and B-G1	12 (A), 8 (B), 5 (B-G1)	3 × 10 vials

Compositions

Reconstituting a given NSK-B or NSK-B-G1 vial's contents in 1 mL of highly pure methanol will produce the tabulated concentrations below. To facilitate complete dissolution, it is recommended to vortex manually for 1 minute then auto-vortex for a minimum of 30 seconds. **Note:** A combined set of NSK-B and NSK-A (NSK-AB) or NSK-B-G1 (NSK-BBG1) is also available.

NSK-B			
Compound	Abbrev.	Label and Enrichment	Conc. (μM)
L-Carnitine	C0	trimethyl-D ₉ , 98%	152
O-Acetyl-L-carnitine·HCl	C2	N-methyl-D ₃ , 98%	38
O-Propionyl-L-carnitine·HCl	C3	N-methyl-D ₃ , 98%	7.6
O-Butyryl-L-carnitine·HCl	C4	N-methyl-D ₃ , 98%	7.6
O-Isovaleryl-L-carnitine·HCl	C5	N,N,N-trimethyl-D ₉ , 98%	7.6
O-Octanoyl-L-carnitine·HCl	C8	N-methyl-D ₃ , 98%	7.6
O-Myristoyl-L-carnitine·HCl	C14	N,N,N-trimethyl-D ₉ , 98%	7.6
O-Palmitoyl-L-carnitine·HCl	C16	N-methyl-D ₃ , 98%	15.2

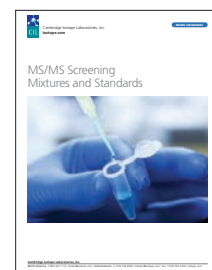
NSK-B-G1			
Compound	Abbrev.	Label and Enrichment	Conc. (μM)
O-Glutaryl-L-carnitine·ClO ₄	C5-DC	N-methyl-D ₃ , 98% (CP 97%)	15.2
3-Hydroxyisovaleryl-L-carnitine·ClO ₄	C5-OH	N-methyl-D ₃ , 98%	7.6
O-Dodecanoyl-L-carnitine·HCl	C12	N,N,N-trimethyl-D ₉ , 98%	7.6
O-3-DL-Hydroxypalmitoyl-L-carnitine·ClO ₄	C16-OH	N-methyl-D ₃ , 98%	15.2
O-Octadecanoyl-L-carnitine·HCl	C18	N-methyl-D ₃ , 98%	15.2

NSK-B-G2			
Compound	Abbrev.	Label and Enrichment	Conc. (μM)
O-Malonyl-L-carnitine·ClO ₄	C3-DC	N-methyl-D ₃ , 98% (CP 95%)	7.6
Tiglyl-L-carnitine·ClO ₄	C5:1	N,N,N-trimethyl-D ₉ , 98% (CP 90%)	7.6
O-Hexanoyl-L-carnitine·HCl	C6	N-methyl-D ₃ , 98%	7.6
O-Decanoyl-L-carnitine·HCl	C10	N-methyl-D ₃ , 98%	7.6
Tetradec-5- <i>cis</i> -enoyl-L-carnitine·ClO ₄	C14:1	N,N,N-trimethyl-D ₉ , 98% (CP 90%)	7.6

Companion unlabeled standard mixes and kits may be available; please inquire.

Chemical purity (CP) is 98% or greater, unless otherwise indicated.

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Usage Specifications

Mix Type	NSK-B	NSK-B-G1	NSK-B-G2
No. of uses	960 samples/vial		
Before reconstitution:			
Storage	-20°C; protect from light	2-8°C; protect from light	-20°C; protect from light
Recommended retest	1.5 years from date of manufacture	2 years from date of manufacture	2 years from date of manufacture
After reconstitution:			
Storage	2-8°C in a tightly sealed vial	2-8°C in a tightly sealed vial	2-8°C in a tightly sealed vial
	Note: Storing the sealed vial in a second sealed container helps maintain the integrity of the solution		
Recommended retest	4 weeks		

Example Results

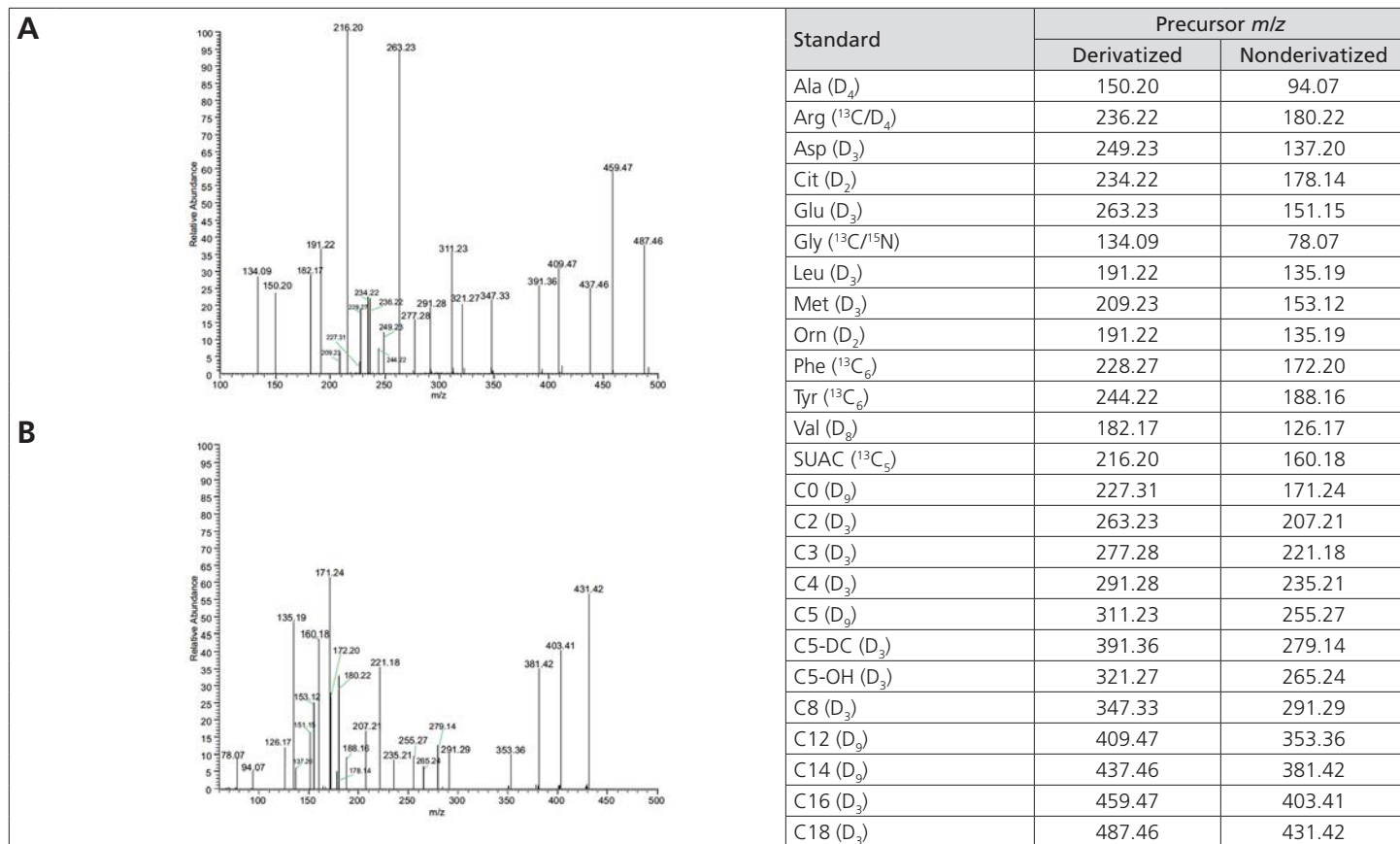


Figure. Full scan MS1 spectra of the DBS internal standards, with the derivatized standards shown in **A** and the nonderivatized in **B**. The standard mixture consists of stable isotope-labeled amino acids (NSK-A and -T) and carnitine/acylcarnitines (NSK-B and NSK-B-G). Please refer to Thermo Scientific technical note #73398 for method and application details.

Companion unlabeled standard mixes and kits may be available; please inquire.

Chemical purity (CP) is 98% or greater, unless otherwise indicated.

For research use only. Not for use in diagnostic procedures.

Example References

- Kilroe, S.P.; Von Ruff, Z.D.; Arentson-Lantz, E.J.; et al. **2025**. Human skeletal muscle disuse atrophy has profound and negative effects on the muscle metabolome and lipidome. *Am J Physiol Endocrinol Metab*, 328(6), E962-E978.
- Habib, A.; Nazri, M.I.B.A.; Rahman, S.A.; et al. **2023**. Dataset from dried blood spot acylcarnitine for detection of Carnitine-Acylcarnitine Translocase (CACT) deficiency and Carnitine Palmitoyl Transferase 2 (CPT2) deficiency. *Data Brief*, 48, 109090-109096.
- Mak, J.; Peng, G.; Le, A.; et al. **2023**. Validation of a targeted metabolomics panel for improved second-tier newborn screening. *J Inherit Metab Dis*, 46(2), 194-205.
- Daas, S.; Salah, N.A.; Anikster, Y.; et al. **2023**. Addition of galactose-1-phosphate measurement enhances newborn screening for classical galactosemia. *J Inherit Metab Dis*, 46(2), 232-242.
- Habib, A.; Azize, N.A.A.; Rahman, S.A.; et al. **2021**. Novel mutations associated with carnitine-acylcarnitine translocase and carnitine palmitoyl transferase 2 deficiencies in Malaysia. *Clin Biochem*, 98, 48-53.
- Brailova, M.; Clerfond, G.; Trésorier, R.; et al. **2020**. Inherited metabolic diseases and cardiac pathology in adults: diagnosis and prevalence in a cardiometabo study. *J Clin Med*, 9(3), 694-707.
- Varkey, A.; Devi, S.; Mukhopadhyay, A.; et al. **2020**. Metabolome and microbiome alterations related to short-term feeding of a micronutrient-fortified, high-quality legume protein-based food product to stunted school age children: a randomized controlled pilot trial. *Clin Nutr*, 39(11), 3251-3261.
- Brennenstuhl, H.; Kohlmüller, D.; Gramer, G.; et al. **2020**. High throughput newborn screening for aromatic L-amino-acid decarboxylase deficiency by analysis of concentrations of 3-O-methyldopa from dried blood spots. *J Inherit Metab Dis*, 43(3), 602-610.
- Cao, B.; Wang, D.; Pan, Z.; et al. **2019**. Characterizing acylcarnitine biosignatures for schizophrenia: a longitudinal pre- and post-treatment study. *Transl Psychiatry*, 9(1), 19.
- Puskarich, M.A.; Evans, C.R.; Karnovsky, A.; et al. **2018**. Septic shock nonsurvivors have persistently elevated acylcarnitines following carnitine supplementation. *Shock*, 49(4), 412-419.

Technical Note

Xie, X.; Kozak, M. **2020**. Simultaneous analysis of amino acids, acylcarnitines, and succinylacetone in dried blood spots for research using nonderivatized and derivatized methods. (Thermo Scientific technical note #73398)

“In clinical chemistry-based applications of mass spectrometry, the first lesson the laboratory learns is the requisite nature of stable isotope-enriched standards for quantification of metabolites in biological fluids. In newborn screening of amino acids and acylcarnitines, Cambridge Isotope Laboratories, Inc., set the standard for quantification of these metabolites in dried blood spots. As research and development of the newborn screening analysis by mass spectrometry progressed, it was clear that a half dozen isotope-labeled internal standards would not be adequate for the analysis of an amino acid and acylcarnitine profile, together comprising a range of 500 separate mass units and more than 30 important metabolites, most of which require accurate quantification. When screening began to expand beyond research, it was clear that weighing out small quantities of individual standards would reduce accuracy and introduce unnecessary error. Therefore, together, we set out to develop sets of standards for amino acids and acylcarnitine analysis that would enable quantification. We started this development more than 20 years ago adding, changing and improving these standards. CIL, together with the early developers of tandem mass spectrometry-based newborn screening, set the standard by which all other laboratories follow. CIL's commitment to supporting the metabolic and newborn screening community is exceptional. It is our good fortune in the clinical chemistry and mass spectrometry community to have CIL as part of our laboratory solutions.”

— Donald H. Chace, PhD MSFS FACB
Medolac Laboratories (USA)

Fatty Acid Mixes and Extracts

Fatty acids and lipids are important biological compounds that are essential to the regulation and control of cellular functions and metabolic pathways. These biomolecules are also tied to the energetic balance of an organism. Their qualitative/quantitative analysis has emerged to better understand the underlying pathophysiology, as well as to identify new biomarkers or diagnose existing ones. To aid such initiatives, CIL offers an array of fatty acid mixes and extracts. These dried-down products comprise uniform or partially labeled fatty acids in different forms (i.e., free acid, methyl ester) as research-grade material.

Overview

Catalog No.	Description	No. of Metabolites	Unit Size
MSK-SBCFA	Short-/Branched-chain Fatty Acid Mix	8	1 vial
CLM-8455	Fatty Acid Algal Extract (U- ¹³ C, 98%)	5-10	0.25 g, 1 g
DLM-8572	Fatty Acid Algal Extract (U-D, 96%)	5-10	Please inquire
CDLM-8376	Fatty Acid Algal Extract (U- ¹³ C, 98%; U-D, 97%)	5-10	0.25 g, 1 g
CLM-8381	Fatty Acid, Methyl Ester Algal Extract (U- ¹³ C, 98%) (unlabeled terminal ester) CP 95%	5-10	0.25 g, 1 g
DLM-2497	Fatty Acid, Methyl Ester Algal Extract (U-D, 96%)	5-10	Please inquire

Note: Companion unlabeled mixes and extracts may be available; please inquire.

SBCFA Composition

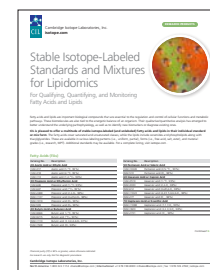
Composition of the stable isotope-labeled SBCFA mix. Reconstituting in 1 mL of solvent will produce the specified equimolar concentrations.

Note: The unlabeled SBCFA mix (MSK-SBCFA-US) has an equivalent composition.

Description (label and enrichment)	Abbrev.	Chemical Formula	Conc. (μM)
Sodium acetate (1,2- ¹³ C ₂ , 99%)	2:0	¹³ C ₂ H ₃ O ₂ Na	150
Sodium propionate (¹³ C ₃ , 99%)	3:0	¹³ C ₃ H ₅ O ₂ Na	150
Sodium butyrate (¹³ C ₄ , 99%)	4:0	¹³ C ₄ H ₇ O ₂ Na	150
Sodium isobutyrate (¹³ C ₄ , 99%)	iso4:0	¹³ C ₄ H ₇ O ₂ Na	150
2-Methylbutyric acid (methyl- ¹³ C, 99%)	2-MB 4:0	¹³ CC ₄ H ₁₀ O ₂	150
Sodium pentanoate (¹³ C ₅ , 99%)	5:0	¹³ C ₅ H ₉ O ₂ Na	150
Isovaleric acid (2,3,4- ¹³ C ₃ , 3-methyl- ¹³ C, 99%)	iso5:0	¹³ C ₄ CH ₁₀ O ₂	150
Hexanoic acid (1- ¹³ C, 99%)	6:0	¹³ CC ₅ H ₁₂ O ₂	150

Usage Specifications

Criteria	Recommendation
Before reconstitution:	
Storage	2-8°C; protect from light
Recommended retest	2 years from date of manufacture



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Algal Extracts Compositions

The fatty acid extracts are derived from *Agmenelum quadriplicatum* algae. Please inquire for details as the compositions can vary by lot.

Usage Specifications

Mix Type	Catalog No.	Before Reconstitution	
		Storage	Recommended Retest
Fatty acids	CLM-8455 DLM-8572 CDLM-8376	2-8°C; protect from light	2 years
Fatty acid, methyl esters	CLM-8381 DLM-2497	2-8°C; protect from light	5 years

Example References

Song, W.; Beigneux, A.P.; Weston, T.A.; et al. **2023**. The lipoprotein lipase that is shuttled into capillaries by GPIHBP1 enters the glycocalyx where it mediates lipoprotein processing. *Proc Natl Acad Sci U S A*, 120(44), e2313825120-2313825131.

Rahim, M.; Ragavan, M.; Deja, S.; et al. **2022**. INCA 2.0: A tool for integrated, dynamic modeling of NMR- and MS-based isotopomer measurements and rigorous metabolic flux analysis. *Metal Eng*, 69, 275-285.

Chen, J.; Zou, L.; Lu, G.; et al. **2022**. PFKP alleviates glucose starvation-induced metabolic stress in lung cancer cells via AMPK-ACC2 dependent fatty acid oxidation. *Cell Discov*, 8(1), 52-67.

Hernandez-Saavedra, D.; Sanders, L.; Freeman, S.; et al. **2020**. Stable isotope metabolomics of pulmonary artery smooth muscle and endothelial cells in pulmonary hypertension and with TGF-beta treatment. *Sci Rep*, 10(1), 413-426.

He, C.; Weston, T.A.; Jung, R.S.; et al. **2018**. NanoSIMS analysis of intravascular lipolysis and lipid movement across capillaries and into cardiomyocytes. *Cell Metab*, 27(5), 1055-1066.

Nucleic Acid Sets

Nucleic acids are necessary biomolecules of living systems, being fundamentally important to a multitude of cellular processes. Its basic building blocks are nucleobases (e.g., adenine, cytosine), nucleosides (e.g., adenosine, guanosine), and nucleotides (e.g., ATP, CDP). The qualification/quantification of these compounds is conducted for a number of purposes. This includes the screening of metabolic errors and evaluating the efficacy of drug treatments (be it anticancer, antiviral, or immunosuppressive), among other target areas. CIL offers a variety of stable isotope-labeled nucleic acid sets to help advance MS- and/or NMR-based research. The sets consist of deoxyribonucleoside monophosphates (dNMPs), deoxyribonucleoside triphosphates (dNTPs), and ribonucleoside triphosphate (rNTPs) in different unit configurations.

Overview

Catalog No.	Description	No. of Metabolites	Unit Size
CNLM-7871-SL	Set of 4 2'-deoxyribonucleoside 5'-monophosphates, lithium salt (¹³ C, 98%; ¹⁵ N, 98%) CP 95% (in solution)	4	10 mg
NLM-7512-SL	Set of 4 2'-deoxyribonucleoside 5'-triphosphates, lithium salt (¹⁵ N, 98%) CP 95% (in solution)	4	10 mg, 50 mg
DLM-7511-SL	Set of 4 2'-deoxyribonucleoside 5'-triphosphates, lithium salt (D, 98%) CP 95% (in solution)	4	10 mg, 50 mg
CNLM-7513-SL	Set of 4 2'-deoxyribonucleoside 5'-triphosphates, lithium salt (¹³ C, 98%; ¹⁵ N, 98%) CP 95% (in solution)	4	10 mg, 50 mg
NLM-7519-SL	Set of 4 ribonucleoside 5'-triphosphates, lithium salt (¹⁵ N, 98%) CP 95% (in solution)	4	10 mg, 50 mg
DLM-7518-SL	Set of 4 ribonucleoside 5'-triphosphates, lithium salt (D, 98%) CP 95% (in solution)	4	10 mg, 50 mg
CNLM-7503-SL	Set of 4 ribonucleoside 5'-triphosphates, lithium salt (¹³ C, 98%; ¹⁵ N, 98%) (in solution) CP 95%	4	10 mg, 50 mg
NLM-7519-CA	Set of 4 ribonucleoside 5'-triphosphates, ammonium salt (¹⁵ N, 98%) CP 90% (in solution)	4	4 × 20 µmol 4 × 100 µmol
DLM-7518-CA	Set of 4 ribonucleoside 5'-triphosphates, ammonium salt (D, 98%) CP 95% (in solution)	4	4 × 20 µmol 4 × 50 µmol 4 × 100 µmol
CNLM-7503-CA	Set of 4 ribonucleoside 5'-triphosphates, ammonium salt (¹³ C, 98%; ¹⁵ N, 98%) CP 95% (in solution)	4	4 × 20 µmol 4 × 50 µmol 4 × 100 µmol

dNMP*Composition and Specification*

Catalog No.	dNMP	Conc. (mM)
CNLM-7871-SL	AMP, CMP, GMP, TMP	80

- Supplied on dry ice as a solution of 5 mM Tris-HCl (pH 7.5) in water.
- Store below -20°C; protect from light.

dNTPs*Compositions and Specifications*

Catalog No.	dNTP	Conc. (mM)
NLM-7512-SL	dATP, dCTP, dGTP, dTTP	50-100
DLM-7511-SL		
CNLM-7513-SL		

- Supplied on dry ice as a solution of 5 mM Tris-HCl (pH 7.5) in water (for NLM-7512-SL and CNLM-7513-SL) or D₂O (for DLM-7511-SL)
- Store below -20°C; protect from light.

Example References

Liu, B.; Winkler, F.; Herde, M.; et al. **2019**. A link between deoxyribonucleotide metabolites and embryonic cell-cycle control. *Curr Biol*, 29(7), 1187-1192.

Song, Y.; Marmion, R.A.; Park, J.O.; et al. **2017**. Dynamic control of dNTP synthesis in early embryos. *Dev Cell*, 42(3), 301-308.

rNTPs*Compositions and Specifications*

Catalog No.	rNTP	Conc. (mM)
NLM-7519-SL	rATP, rCTP, rGTP, rTTP	50-100
DLM-7518-SL		
CNLM-7503-SL		

- Supplied on dry ice as a solution of 5 mM Tris-HCl (pH 7.5) in water (for NLM-7519-SL and CNLM-7503-SL) or D₂O (for DLM-7518-SL)
- Store below -20°C; protect from light.

Catalog No.	rNTP	Conc. (mM)
NLM-7519-CA	rATP, rCTP, rGTP, rUTP	20, 50, and/or 100
DLM-7518-CA		
CNLM-7503-CA		

- Supplied on dry ice as a solution (e.g., 100 mM – contains 100 µmol of each rNTP in 1 mL water, pH ~7.5).
- Store at -20°C; protect from light. Recommended retest is 5 years from date of manufacture.

Companion unlabeled standard mixes and kits may be available; please inquire.

Chemical purity (CP) is 98% or greater, unless otherwise indicated.

For research use only. Not for use in diagnostic procedures.

Organic Acid Mixes

Organic acids are byproducts of amino acids and are intermediates in various biochemical pathways. To aid MS-based metabolomic research endeavors, CIL has formulated an isotope-enriched organic acid mix. This single-vial mixture comprises 33 organic acids (MSK-OA) and is also available as an unlabeled mix (MSK-OA-US).

Overview

Catalog No.	Description	No. of Metabolites	Unit Size
MSK-OA	Organic Acid Mix	33	1 vial

Compositions

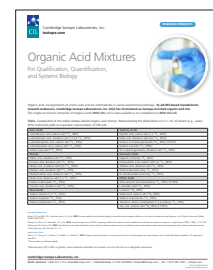
Composition of the stable isotope-labeled organic acid mixture. Reconstituting this dried-down mix in 1 mL of solvent (e.g., water, 1:1 v/v methanol:water) yields an equimolar concentration of 250 µM.

Keto Acids	Hydroxy Acids
α-Ketobutyric acid, sodium salt ($^{13}\text{C}_4$, 98%)	Glycolic acid, sodium salt (1,2- $^{13}\text{C}_2$, 99%)
α-Ketoglutaric acid, disodium salt (1,2,3,4- $^{13}\text{C}_4$, 99%)	Malic acid, disodium salt-H ₂ O ($^{13}\text{C}_4$, 99%)
α-Ketoisocaproic acid, sodium salt ($^{13}\text{C}_6$, 99%)	Sodium D-3-hydroxybutyrate ($^{13}\text{C}_4$, 99%) CP 97%
α-Ketoisovaleric acid, sodium salt ($^{13}\text{C}_5$, 98%)	Sodium L-lactate ($^{13}\text{C}_3$, 98%)
Sodium pyruvate ($^{13}\text{C}_3$, 99%)	DL 2-Hydroxyglutarate, disodium salt ($^{13}\text{C}_5$, 99%)
Diacids	Aromatic Acids
Adipic acid, disodium salt ($^{13}\text{C}_6$, 99%)	Hippuric acid (ring- $^{13}\text{C}_6$, 99%)
Fumaric acid, disodium salt ($^{13}\text{C}_4$, 99%)	Homovanillic acid, sodium salt (1,2- $^{13}\text{C}_2$, 98%)
Maleic acid, disodium salt-H ₂ O ($^{13}\text{C}_4$, 99%)	Phthalic acid, disodium salt ($^{13}\text{C}_4$, 99%)
Malonic acid, disodium salt ($^{13}\text{C}_3$, 99%)	Sodium benzoate (ring- $^{13}\text{C}_6$, 99%)
Methylmalonic acid, disodium salt ($^{13}\text{C}_4$, 99%)	DL-Vanilmandelic acid (ring- $^{13}\text{C}_6$, 99%)
Oxalic acid, disodium salt (1,2- $^{13}\text{C}_2$, 99%)	Other Acids
Sodium isobutyrate ($^{13}\text{C}_4$, 99%)	trans-Aconitic acid (tricarboxyl- $^{13}\text{C}_3$, 99%) CP 95%
Succinic acid, disodium salt ($^{13}\text{C}_4$, 99%)	L-Ascorbic acid ($^{13}\text{C}_6$, 98%)
Monoacids	Creatine ($^{15}\text{N}_3$, 98%)
Sodium acetate (1,2- $^{13}\text{C}_2$, 99%)	Orotic acid, sodium salt ($^{15}\text{N}_2$, 98%)
Sodium butyrate ($^{13}\text{C}_4$, 99%)	Sodium D-gluconate ($^{13}\text{C}_6$, 99%)
Sodium propionate ($^{13}\text{C}_3$, 99%)	Trisodium citrate, hemihydrate (1,5,6-carboxy- $^{13}\text{C}_3$, 99%)
	Uric acid, sodium salt ($^{15}\text{N}_2$, 98%) CP 95%

Usage Specifications

Criteria	Recommendation
Before reconstitution:	
Storage	2-8°C; protect from light
Recommended retest	2 years from date of manufacture
After reconstitution:	
Storage	4°C
Recommended retest	4 weeks

Note: Extended mix stabilities have been demonstrated when solubilized in 50% methanol and stored as specified above.



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Companion unlabeled standard mixes and kits may be available; please inquire.

Chemical purity (CP) is 98% or greater, unless otherwise indicated.

For research use only. Not for use in diagnostic procedures.

Example Results

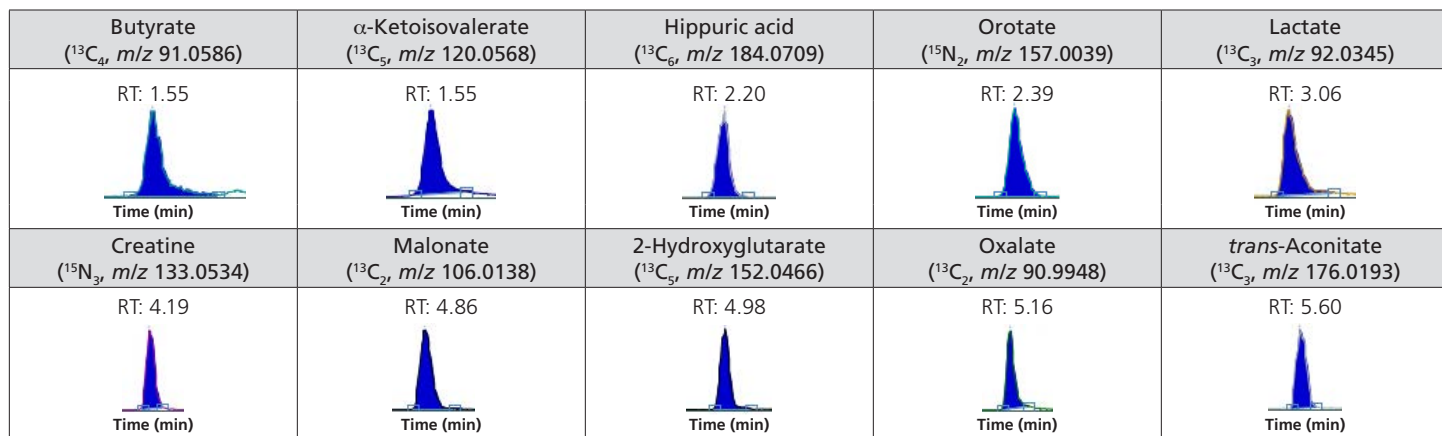


Figure. XICs of example metabolites measured in the labeled OA master mix by HILIC-MS (Orbitrap ID-X, negative ESI). Shown are a collection of OAs from a variety of subset classes. The m/z displayed are the $[\text{M}-\text{H}]^-$ ions. Procedurally, the dried-down master mix was reconstituted in 1 mL of 50% methanol then an aliquot 10-fold diluted before MS1 analysis.

Example Reference

Kilroe, S.P.; Von Ruff, Z.D.; Arentson-Lantz, E.J.; et al. **2025**. Human skeletal muscle disuse atrophy has profound and negative effects on the muscle metabolome and lipidome. *Am J Physiol Endocrinol Metab*, 328(6), E962-E978.

Kalenta, H.; Kilroe, S.P.; Romsdahl, T.B.; et al. **2025**. Constitutively active mTORC1 signaling modifies the skeletal muscle metabolome and lipidome response to exercise. *J Appl Physiol* (1985), 138(5), 1173-1186.

DeArmond, P.D.; Bunch, D.R. **2022**. Quantitation of non-derivatized free amino acids for detecting inborn errors of metabolism by incorporating mixed-mode chromatography with tandem mass spectrometry. *J Mass Spectrom Adv Clin Lab*, 25, 1-11.

Application Note

Petucci, C.*; Percy, A.J.*; Zelenin, A.; Gardell, S.J.; Backiel, K. **2017**. Organic acid quantitation in mouse muscle by ion chromatography-mass spectrometry with isotopically labeled standards. (CIL application note #47)

*These authors contributed equally.

Companion unlabeled standard mixes and kits may be available; please inquire.

Chemical purity (CP) is 98% or greater, unless otherwise indicated.

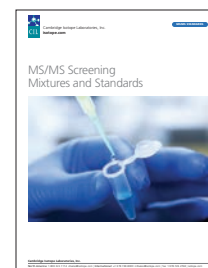
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Other Mixes and Sets

A number of additional stable isotope-labeled mixes are also available at CIL (see overview). Included in this collection is a lysophosphatidylcholine (LPC or lysoPC) mix (NSK-LPC), a metabolomics standard mix (MSK-MET1), and MS/MS standard screening mixes (e.g., NSK-MP, NSK-FA) for basic and translational MS research. The details of these offerings are outlined below.

Overview

Catalog No.	Description	No. of Metabolites	Unit Size
NSK-LPC	Lysophosphatidylcholine Mix	4	1 vial
NSK-NI	Acid Sphingomyelinase Substrate and Internal Standard Mix	S, IS	1 vial
NSK-KR	Galactocerebrosidase Substrate and Internal Standard Mix	S, IS	1 vial
NSK-FA	α -Galactosidase Substrate and Internal Standard Mix	S, IS	1 vial
NSK-GA	Glucocerebrosidase Substrate and Internal Standard Mix	S, IS	1 vial
NSK-MP	α -L-Iduronidase Substrate and Internal Standard Mix	S, IS	1 vial
NSK-PO	Acid α -Glucosidase Substrate and Internal Standard Mix	S, IS	1 vial
MSK-MET1	Metabolomics Standards Mix 1	11	1 vial, 5 vials, 10 vials



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NSK-LPC Mix

Composition

Reconstituting a given LysoPC mix (labeled, NSK-LPC; unlabeled, NSK-LPC-US) in 1 mL of high-purity solvent (e.g., 95:5 v/v methanol:water) will produce the concentrations specified below. To facilitate complete dissolution, it is recommended to sonicate the vial for 3 minutes then auto-vortex for a minimum of 10 seconds.

Standard (Abbreviation)	Label and Enrichment	Conc. (μ M)	Structure
Lysophosphatidylcholine 20:0 (LysoPC 20:0)	eicosanoyl-12,12,13,13-D ₄ , 98%	5.5	
Lysophosphatidylcholine 22:0 (LysoPC 22:0)	docosanoyl-1,2,3,4,5,6- ¹³ C ₆ , 99%	5.5	
Lysophosphatidylcholine 24:0 (LysoPC 24:0)	tetracosanoyl-1,2,3,4,5,6- ¹³ C ₆ , 99%	5.5	
Lysophosphatidylcholine 26:0 (LysoPC 26:0)	hexacosanoyl-1,2,3,4,5,6- ¹³ C ₆ , 99%	5.5	

Usage Specifications

Criteria	Recommendation
No. of uses	~765 samples/vial
Before reconstitution:	
Storage	-20°C; protect from light
Recommended retest	3 years from date of manufacture
After reconstitution:	
Storage	5±3°C or -20±5°C
Recommended retest	6 weeks

“Testing 1,200 to 2,000 newborn specimens every day is a challenging job. In order to ensure high data quality and proper efficacy of an MS-based screening assay (1st and 2nd tier), we utilize CIL's lyophilized LysoPC mix. This 4-plex cocktail helps reduce significantly our laboratory day-to-day variation and preparation time. We also use the LysoPC mixes (both unlabeled and isotope-labeled) in the tuning of our mass spectrometers and in determining response factors. This ensures that the concentrations rendered for each LysoPC is accurately obtained and can be cross-checked from platform-to-platform.”

— Kuldeep Dhillon, Research Scientist Supervisor I
California Department of Public Health (USA)

Companion unlabeled standard mixes and kits may be available; please inquire.

Chemical purity (CP) is 98% or greater, unless otherwise indicated.

For research use only. Not for use in diagnostic procedures.

Example Results

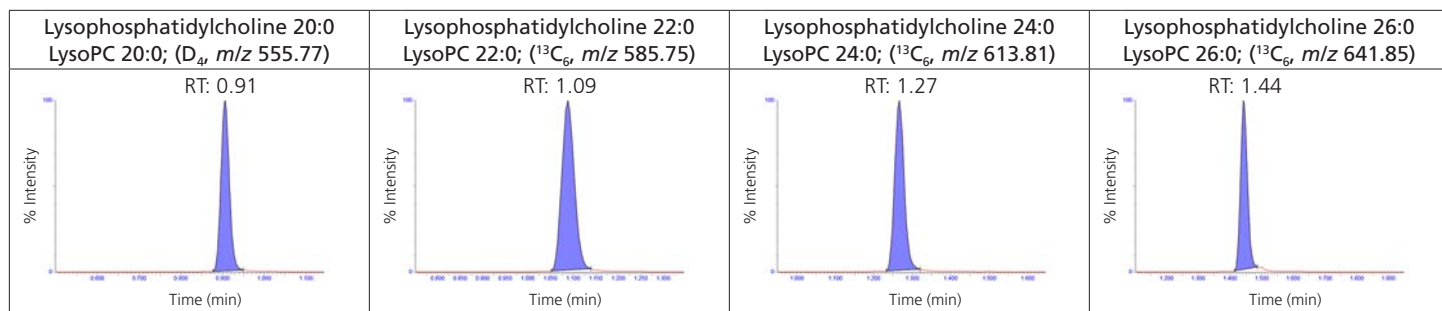


Figure. XICs of four labeled LysoPCs from the NSK-LPC mix measured by LC-MS/MS (positive ESI, Xevo-TQS). These serve as individual calibrators for their unlabeled counterparts in sample screening experiments. Data courtesy of CA Department of Public Health.

Substrate and Internal Standard Mixes

α -Galactosidase Substrate and Internal Standard Mix (NSK-FA)

Each vial contains the following compounds at a substrate:internal standard (S:IS) molar ratio of 500:1.

Substrate	Internal Standard
6-Benzoylamino-hexyl-[2-[4-(3,4,5-trihydroxy-6-hydroxymethyl-tetrahydropyran-2-yloxy)-phenylcarbamoyl]-ethyl]-carbamic acid <i>tert</i> -butyl ester	6-D ₅ -Benzoylamino-hexyl-[2-(4-hydroxy-phenyl-carbamoyl)-ethyl]-carbamic acid <i>tert</i> -butyl ester
C ₃₃ H ₄₇ N ₃ O ₁₀ MW: 645.7 Da	C ₂₇ H ₃₂ N ₃ O ₅ D ₅ MW: 488.5 Da

Glucocerebrosidase Substrate and Internal Standard Mix (NSK-GA)

Each vial contains the following compounds at a substrate:internal standard (S:IS) molar ratio of 50:1.

Substrate	Internal Standard
D-Glucosyl- β 1'-1'- <i>N</i> -dodecanoyl-D- <i>erythro</i> -sphingosine (C12-glucocerebroside)	<i>N</i> -Myristoyl-D- <i>erythro</i> -sphingosine (C14-ceramide)
C ₃₆ H ₆₉ NO ₈ MW: 643.9 Da	C ₃₂ H ₆₃ NO ₃ MW: 509.8 Da

Galactocerebrosidase Substrate and Internal Standard Mix (NSK-KR)

Each vial contains the following compounds at a substrate:internal standard (S:IS) molar ratio of 150:1.

Substrate	Internal Standard
D-Galactosyl- β 1'-1'-octanoyl-D- <i>erythro</i> -sphingosine (C8-galactosylceramide)	<i>N</i> -Decanoyl-D- <i>erythro</i> -sphingosine (C10-ceramide)
C ₃₂ H ₆₁ NO ₈ MW: 587.8 Da	C ₂₈ H ₅₅ NO ₃ MW: 453.7 Da

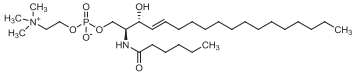
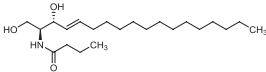
α -L-Iduronidase Substrate and Internal Standard Mix (NSK-MP)

Each vial contains the following compounds at a substrate:internal standard (S:IS) molar ratio of 150:1.

Substrate	Internal Standard
7-(1-Iduronic acid)-oxycoumarin-4-methylamine-(5'- <i>N</i> -boc-aminopentanoyl)-amide	7-Hydroxycoumarin-4-methylamine-(4'- <i>N</i> -boc-aminobutanoyl)-amide
C ₂₆ H ₃₄ N ₂ O ₁₂ MW: 566.6 Da	C ₁₉ H ₂₄ N ₂ O ₆ MW: 376.4 Da

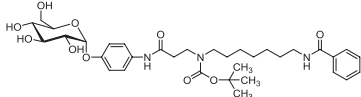
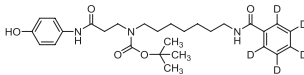
Acid Sphingomyelinase Substrate and Internal Standard Mix (NSK-NI)

Each vial contains the following compounds at a substrate:internal standard (S:IS) molar ratio of 50:1.

Substrate	Internal Standard
<i>N</i> -Hexanoyl-D- <i>erythro</i> -sphingosylphosphorylcholine (C6-sphingomyelin)	<i>N</i> -Butyroyl-D- <i>erythro</i> -sphingosine (C4-ceramide)
C ₂₉ H ₅₉ N ₂ O ₆ P MW: 562.8 Da	C ₂₂ H ₄₃ NO ₃ MW: 369.6 Da
	

Acid α-Glucosidase Substrate and Internal Standard Mix (NSK-PO)

Each vial contains the following compounds at a substrate:internal standard (S:IS) molar ratio of 100:1.

Substrate	Internal Standard
7-Benzoylamino-heptyl-[2-[4-(3,4,5-trihydroxy-6-hydroxymethyl-tetrahydro-pyran-2-yloxy)-phenylcarbamoyl]-ethyl]-carbamic acid <i>tert</i> -butyl ester	7-D ₅ -Benzoylamino-heptyl-[2-(4-hydroxy-phenyl-carbamoyl)-ethyl]-carbamic acid <i>tert</i> -butyl ester
C ₃₄ H ₄₉ N ₃ O ₁₀ MW: 659.8 Da	C ₂₈ H ₃₄ N ₃ O ₅ D ₅ MW: 502.7 Da
	

Usage Specifications

Criteria	Recommendation
Use	~600 samples/vial
Before reconstitution:	
Storage	-20°; protect from light
Recommended retest	2 years from date of manufacture
After reconstitution:	
Storage	5±3°C or -20±5°C
Recommended retest	4 weeks

Example References

Ribas, G.; De Mari, J.F.; Civaliero, G.; et al. **2017**. Validation of a multiplex tandem mass spectrometry method for the detection of selected lysosomal storage diseases in dried blood spots. *JIMES*, 5, 1-7.

Tortorelli, S.; Turgeon, C.T.; Gavrilov, D.K.; et al. **2016**. Simultaneous testing for 6 lysosomal storage disorders and x-adrenoleukodystrophy in dried blood spots by tandem mass spectrometry. *Clin Chem*, 62(9), 1248-1254.

Cho, S.E.; Kwak, J.R.; Lee, H.; et al. **2016**. Triplex tandem mass spectrometry assays for the screening of 3 lysosomal storage disorders in a Korean population. *Clin Chim Acta*, 454, 20-27.

MSK-MET1 Mix**Composition**

Reconstituting a given vial in 1 mL of solvent (e.g., water) will produce the tabulated concentrations below. **Note:** The unlabeled 2DG was included to examine its metabolic uptake in tracer studies.

Compound	Abbrev.	Label and Enrichment	Conc. (mM)
L-Alanine	Ala	2,3,3,3-D ₄ , 98%	20
L-Carnitine	C0	trimethyl-D ₉ , 98%	0.4
Creatinine	Crn	N-methyl-D ₃ , 98%	0.7
2-Deoxy-D-glucose	2DG	unlabeled	50
D-Glucose	Glc	U- ¹³ C ₆ , 99%	400
L-Glutamine	Gln	2,3,3,4,4-D ₅ , 97%	8
L-Leucine	Leu	5,5,5-D ₃ , 99%	5
L-Phenylalanine	Phe	ring-D ₅ , 98%	0.5
Sodium butyrate	BTA	D ₇ , 98%	2
Sodium propionate	PA	D ₅ , 98%	2
Urea	UR	¹⁵ N ₂ , 98%	20

Usage Specifications

Criteria	Recommendation
Before reconstitution:	
Storage	2-8°C; protect from light
Recommended retest	5.5 years from date of manufacture

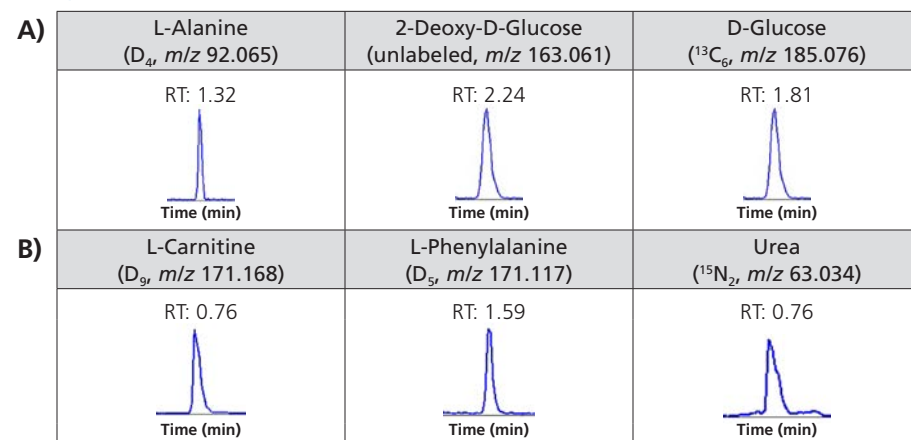
Example Results

Figure. Representative XICs of example metabolites in the MET1 mix measured in mouse plasma by LC-MS analysis. A TripleTOF 6600+ was fronted by a Hypercarb™ Porous Graphitic Carbon HPLC column (100 × 4.6 mm, 3 μm) in **A)** and a Kinetex F5 HPLC column (100 × 2.1 mm, 2.6 μm) in **B)** operated under negative and positive ESI, respectively. Data was courtesy of the Kibbey lab at Yale School of Medicine.

Example Reference

Richiardone, E.; Giolito, M.V.; Al Roumi, R.; et al. **2025**. Acidosis overrides molecular heterogeneity to shape therapeutically targetable metabolic phenotypes in colon cancers. *Cancer Lett*, 613, 217512-217525.

Richiardone, E.; Al Roumi, R.; Lardinois, F.; et al. **2024**. MCT1-dependent lactate recycling is a metabolic vulnerability in colorectal cancer cells upon acquired resistance to anti-EGFR targeted therapy. *Cancer Lett*, 598, 217091-217103.

Tan, H.C.; Hsu, J.W.; Tai, E.S.; et al. **2024**. The impact of obesity-associated glycine deficiency on the elimination of endogenous and exogenous metabolites via the glycine conjugation pathway. *Front Endocrinol*, 15, 1343738-1343749.

Steroid Mixes and Sets

Steroids play vital roles in the regulation of a diverse array of cellular functions and physiological processes. These pertain to development, reproduction, homeostasis, and metabolism, among others. Accurate quantification of this compound class is essential for basic and translation research. To aid MS-based research endeavors in this space, CIL is pleased to offer a few different types of stable isotope-labeled steroid mixes.

Overview

Catalog No.	Description	No. of Metabolites	Unit Size
NSK-S	Steroid Standards Mix S	5	1 vial, 10 vials
NSK-S-40X	Steroid Standards Mix S (40x)	5	1 vial
NSK-S-EXP	Expanded Steroid Standards Mix S	9	1 vial, 10 vials

NSK-S Mix

Composition

Reconstituting a given vial's contents in 1 mL of highly pure methanol will produce the concentrations listed in the table below. To facilitate complete dissolution, it is recommended to vortex manually for 1 minute then auto-vortex for a minimum of 30 minutes. **Note:** NSK-S-40X has 40x the specified concentrations.

Compound	Abbrev.	Label and Enrichment	Conc. (nM)
4-Androstene-3,17-dione	A4	2,2,4,6,6,16,16-D ₇ , 97%	20
Cortisol	F	9,11,12,12-D ₄ , 98%	100
11-Deoxycortisol	11-S	2,2,4,6,6-D ₅ , 98%	20
21-Deoxycortisol	21-S	2,2,4,6,6,21,21,21-D ₈ , 97%	20
17 α -Hydroxyprogesterone	17-OHP	2,2,4,6,6,21,21,21-D ₈ , 98%	20

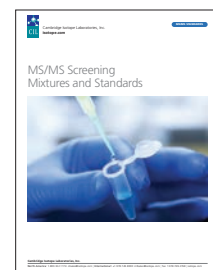
Usage Specifications

Criteria	Recommendation
No. of uses	48 samples/vial
Before reconstitution:	
Storage	2-8°C; protect from light
Recommended retest	5 years from date of manufacture
Upon reconstitution:	
Storage	Store in a tightly sealed vial at 2-8°C. Note: Storing the sealed vial in a second sealed container helps maintain the integrity of the solution.
Recommended retest	4 weeks

Example References

Gervasoni, J.; Schiattarella, A.; Primiano, A.; et al. **2016**. Simultaneous quantification of 17-hydroxyprogesterone, androstenedione, testosterone and cortisol in human serum by LC-MS/MS using TurboFlow online sample extraction. *Clin Biochem*, 49(13-14), 998-1003.

Hicks, R.A.; Yee, J.K.; Mao, C.S.; et al. **2014**. Precursor-to-product ratios reflect biochemical phenotype in congenital adrenal hyperplasia. *Metabolomics*, 10(1), 123-131.



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NSK-S-EXP Mix**Composition**

Reconstituting a given vial's contents in 1 mL of highly pure methanol will produce the concentrations listed in the table below.

To facilitate complete dissolution, it is recommended to vortex manually for 1 minute then auto-vortex for a minimum of 30 minutes.

Compound	Abbrev.	Label and Enrichment	Conc. (μM)
Aldosterone	A	D ₇ , 98%	0.52
4-Androstene-3,17-dione	A4	2,2,4,6,6-D ₅ , 98%	0.12
Corticosterone	B	2,2,4,6,6,17α,21,21-D ₈ , 97%	1.58
Cortisol	F	9,12,12-D ₃ , 98%	2.57
Dehydroepiandrosterone sulfate-sodium salt·2H ₂ O	DHEAS	2,2,3,4,4,6-D ₆ , 95%	21.69
11-Deoxycortisol	11-S	2,2,4,6,6-D ₅ , 98% (CP 97%)	0.54
17-α-Hydroxyprogesterone	17-OHP	2,2,4,6,6,21,21,21-D ₈ , 98%	0.27
Progesterone	P	2,2,4,6,6,17α,21,21,21-D ₉ , 98%	0.14
Testosterone	T	2,2,4,6,6-D ₅ , 98%	0.12

Usage Specifications

Criteria	Recommendation
Before reconstitution:	
Storage	2-8°C; protect from light
Recommended retest	1 year from date of manufacture

Companion unlabeled standard mixes and kits may be available; please inquire.

Chemical purity (CP) is 98% or greater, unless otherwise indicated.

For research use only. Not for use in diagnostic procedures.

Pathway-Specific Mixes

TCA Mixes and Sets

The tricarboxylic acid (TCA) cycle plays an essential role in central carbon and energy metabolism. The study of TCA cycle intermediates (e.g., citrate, α -ketoglutarate, succinate) and its offshoot metabolites (e.g., itaconate and 2-hydroxyglutarate), have proven pivotal in not only understanding their impact on metabolism, but also in profiling their cellular function/fate in oncogenesis, inflammation, and other pathologies (e.g., necrosis, cirrhosis). To aid the analysis of TCA cycle-associated compounds in metabolomic studies, we offer stable isotope-labeled and unlabeled TCA cycle mixes. These mixes are dried down and comprise a collection of TCA cycle and offshoot metabolites.

Overview

Catalog No.	Description	No. of Metabolites	Unit Size
MSK-TCA1	TCA Cycle Standards Mix 1	8	1 vial
MSK-TCA2-A	TCA Cycle Standards Mix 2A	6	1 vial
MSK-TCA-A	TCA Cycle Standards Mix Sets 1 and 2A	8 (TCA1), 6 (TCA2-A)	2 × 1 vials

Compositions

Composition of the stable isotope-labeled TCA mixtures. Reconstituting a given vial in 1 mL of solvent (e.g., water) will produce an equimolar concentration of 100 μ M.

Compound	Abbrev.	Label and Enrichment	Vial No.
Fumaric acid, disodium salt	Fum	$^{13}\text{C}_4$, 99%	1
DL-2-Hydroxyglutaric acid, disodium salt	HG	$^{13}\text{C}_5$, 99%	1
α -Ketoglutaric acid, disodium salt	α -KG	1,2,3,4- $^{13}\text{C}_4$, 99%	1
Malic acid, disodium salt	Mal	$^{13}\text{C}_4$, 99%	1
Sodium L-lactate	Lac	$^{13}\text{C}_3$, 98%	1
Sodium pyruvate	Pyr	$^{13}\text{C}_3$, 99%	1
Succinic acid, disodium salt	SA	$^{13}\text{C}_4$, 99%	1
Trisodium citrate, hemihydrate	CA	1,5,6-carboxy- $^{13}\text{C}_3$, 99%	1
cis-Aconitic acid	CAA	$^{13}\text{C}_4$, 99%	2
L-Aspartic acid	Asp	$^{13}\text{C}_4$, 99%	2
L-Glutamic acid	Glu	$^{13}\text{C}_5$, 99%	2
Isocitric acid, trisodium salt hydrate	Iso	3,4,5,6- $^{13}\text{C}_4$, 98% (mixture of diastereomers)	2
Itaconic acid	IA	$^{13}\text{C}_5$, 99%	2
Potassium phosphoenol pyruvate	PEP	2,3- $^{13}\text{C}_2$, 99%	2

Usage Specifications

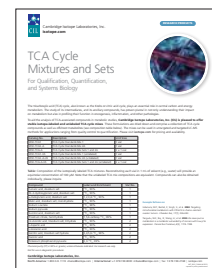
Criteria	Recommendation
Before reconstitution:	
Storage	ambient temperature; protect from light and moisture
Recommended retest	2 years from date of manufacture
After reconstitution:	
Storage	4°C
Recommended retest	4 weeks

Note: Extended mix stabilities have been demonstrated when solubilized in water and stored as specified above.

Companion unlabeled standard mixes and kits may be available; please inquire.

Chemical purity (CP) is 98% or greater, unless otherwise indicated.

For research use only. Not for use in diagnostic procedures.



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Example Results

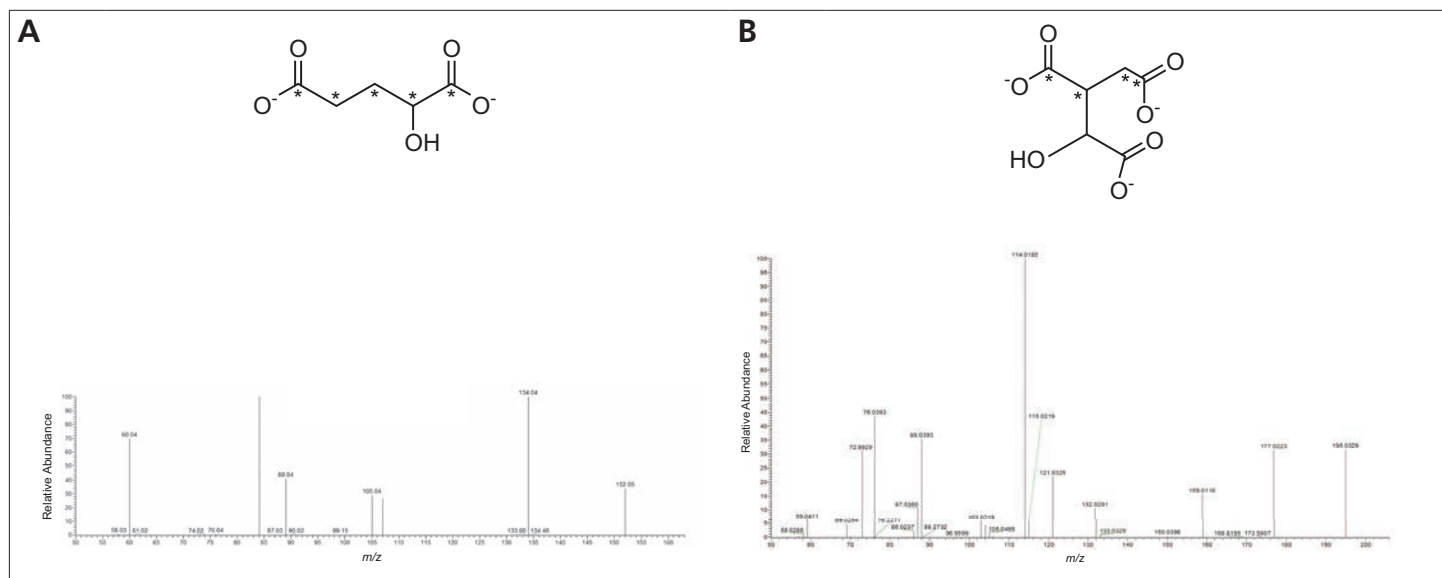


Figure. Example MS/MS spectra of the TCA mixes. These were obtained from the IC-MS analysis (negative ESI, Orbitrap Fusion Tribrid) of 10 μM working solutions. The compounds are $^{13}\text{C}_5$ 2-hydroxyglutarate (from mix 1) in **A**) and $^{13}\text{C}_4$ isocitrate (from mix 2) in **B**). Their complete, unfragmented structures ($^{13}\text{C}_5$ hydroxyglutarate, precursor m/z 152.05; $^{13}\text{C}_4$ isocitrate, precursor m/z 195.03) are illustrated for reference. Data was courtesy of the Metabolomics Facility at MD Anderson Cancer Center.

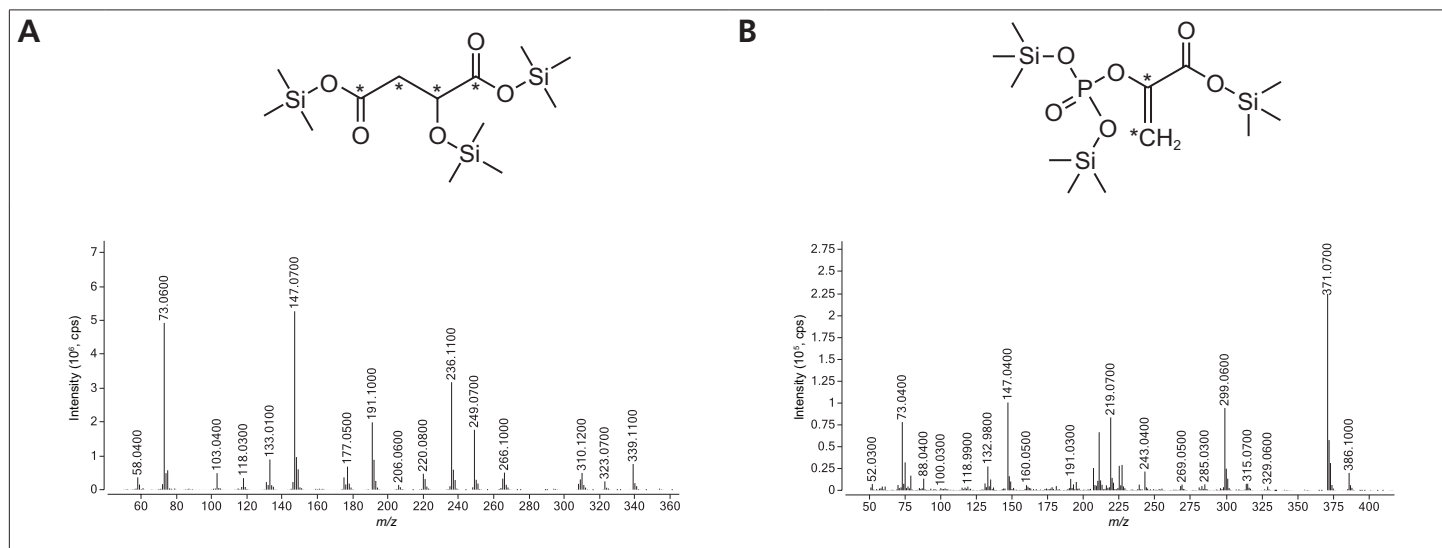


Figure. Example spectra of the labeled TCA mixes. These were obtained from the analysis of 50 μM working solutions by GC-MS (BSTFA with TMCS, EI ionization, scan mode on single quadrupole). The compounds are $^{13}\text{C}_4$ malate (from mix 1) in **A**) and $^{13}\text{C}_2$ phosphoenolpyruvate (from mix 2) in **B**). Their complete, unfragmented structures ($^{13}\text{C}_4$ malate, exact mass 354.15; $^{13}\text{C}_2$ phosphoenolpyruvate, exact mass 386.11) are illustrated for reference. Data was courtesy of the University of Chicago Metabolomics Facility.

“The metabolomic compound mixes by Cambridge Isotope Laboratories have expedited several processes in our GC-MS analysis pipeline. We have used these products, including their novel TCA mixes, to verify metabolite retention times, test derivatization reaction conditions, and to confirm fragmentation patterns detected in biological samples. CIL’s metabolite mixes are ready to go, easy to use, and consistently of high quality!”

— Ashley M. Sidebottom, PhD

Metabolomics Platform Director, University of Chicago (USA)
Host-Microbe Metabolomics Facility, Duchossois Family Institute

Kits

To help facilitate product application, a number of the CIL mix offerings are accompanied with a user manual. The kit manual provides a guide to the preparation and processing possibilities for product implementation, as well as supplying further resources for additional user reference. The majority of the kits supply the supporting documents through a QR code (via product label on kit materials box); the exception is the PeptiQuant™ Plus kits (documents on USB). The accompanying documents minimally include a user manual, certificate of analysis (CoA), safety data sheet (SDS), and product flyer. Overall, the mixes are amenable for use in quality control and qualification/quantification exercises in the MS omics space using targeted, semitargeted, or untargeted LC-MS methodologies. Note that the individual mixes may be obtained separately without the manual. Please inquire.

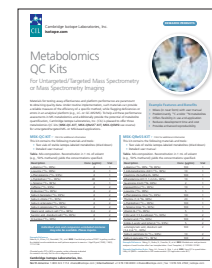
Metabolomic QC Kits

To ensure high-quality metabolomics results, the method and instrument platform must be qualified as being fit-for-purpose. This involves testing for losses or errors in the analytical workflow. To aid such performance assessments in MS metabolomics and potentially other applications (e.g., qualification, quantification), CIL offers three types of QC kits: MSK-QC-KIT, MSK-QReSS-KIT, and MSK-QReSS-EXP-KIT (see details below). These kits comprise vials of dried-down metabolite mixes (see composition tables below) and have been qualified on both low- and high-resolution mass spectrometers operated under an untargeted/targeted metabolomics regimen (see example results below).

Overview

Catalog No.	Description	No. of Metabolites	Kit Contents
MSK-QC-KIT	Metabolomics QC Kit	5 (mix 1), 9 (mix 2)	• 2 vials of ^{13}C-labeled metabolites • document package (user manual, CoA, SDS, product flyer)
MSK-QReSS-KIT*	Metabolomics QReSS Kit	12 (mix 1), 6 (mix 2)	• 2 vials of isotope-labeled metabolites • document package (user manual, CoA, SDS, product flyer)
MSK-QCMSI-KIT	Metabolomics QC Kit for MSI	3 (mix 1), 3 (mix 2)	• Two vials of metabolite mixes (SIL and NAT, lyophilized) • document package (user manual, CoA, SDS, product flyer)

*QReSS™ stands for Quantification, Retention, and System Suitability. This kit was developed and matrix tested in collaboration with SCIEX.



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MSK-QC Mixes/Kits

Mix Compositions

Reconstituting in 1 mL of solvent (e.g., 0.1% FA in 5% ACN) will yield the specified compound concentrations.

Compound	Abbrev.	Label and Enrichment	Conc. (µg/mL)	Mix No.
L-Alanine	Ala	$^{13}\text{C}_3$, 99%	4	1
L-Leucine	Leu	$^{13}\text{C}_6$, 99%	4	1
L-Phenylalanine	Phe	$^{13}\text{C}_6$, 99%	4	1
L-Tryptophan	Trp	$^{13}\text{C}_{11}$, 99%	40	1
L-Tyrosine	Tyr	$^{13}\text{C}_6$, 99%	4	1
D-Glucose	Glc	$^{13}\text{C}_6$, 99%	4	2
D-Sucrose	Suc	$^{13}\text{C}_6$, 98%	4	2
Caffeine	CAF	$^{13}\text{C}_3$, 99%	4	2
Stearic acid, sodium salt	18:0	$^{13}\text{C}_{18}$, 98%	0.4	2
Sodium octanoate	8:0	$^{13}\text{C}_8$, 99%	4	2
Sodium propionate	PA	$^{13}\text{C}_3$, 99%	4	2
Sodium benzoate	BZA	$^{13}\text{C}_6$, 99%	4	2
Succinic acid, disodium salt	SA	$^{13}\text{C}_4$, 99%	4	2
Trisodium citrate	CA	$^{13}\text{C}_3$, 99%	4	2

Usage Specifications

Criteria	Recommendation
Before reconstitution:	
Storage	-20°C (protect from light)
Recommended retest	3 years from date of manufacture

Companion unlabeled standard mixes and kits may be available; please inquire.

Chemical purity (CP) is 98% or greater, unless otherwise indicated.

For research use only. Not for use in diagnostic procedures.

Example Results

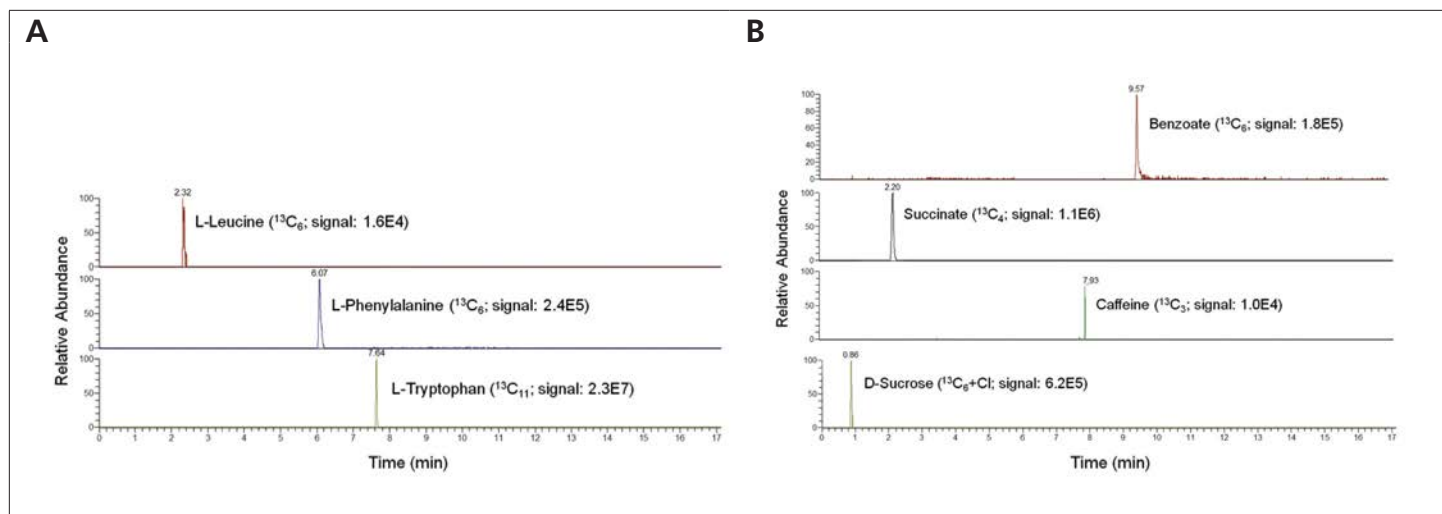


Figure. Representative XICs of a subset of kit metabolites – vial 1 in A and vial 2 in B – measured in human plasma by RPLC-MS (negative ESI, Q Exactive). Note that all isotopically labeled metabolites in the mixes co-eluted with their endogenous analytes in the plasma analyses and their m/z were predominantly $[M-H]^-$.

“I have used products from Cambridge Isotope Laboratories (CIL) for well over 15 years because of the high quality provided. I have collaborated with CIL to develop a new product that would enable improved quality control in MS metabolomics. This process was exciting and engaging. Through collaboration, I found CIL to be a cutting-edge vendor that seeks opportunities to enable scientific discovery and data quality. They seek advice from clients to learn how to better serve them. I always look forward to talking with CIL at conferences and often seek out their booth to visit, not just to talk science, but to also see how their team is doing.”

– Timothy J. Garrett, PhD

Associate Professor in Department of Pathology,
Immunology, and Laboratory Medicine, University of Florida (USA)

MSK-QReSS Mixes/Kits

Mix Compositions

Reconstituting a given QReSS mix in 1 mL of solvent (e.g., 1:1 v/v methanol:water) will yield the specified concentrations.

Compound	Abbrev. or Alt. Name	Label and Enrichment	Conc. (µg/mL)	Mix No.
L-Alanine	Ala	$^{13}\text{C}_3$, 99%; ^{15}N , 99%	100	1
1,4-Butanediamine·2HCl	putrescine	$^{13}\text{C}_4$, 99%	10	1
Creatinine	Crn	<i>N</i> -methyl- D_3 , 98%	100	1
Ethanolamine-HCl	ETA	1,1,2,2- D_4 , 98%	10	1
Guanosine·2H ₂ O	Guo	$^{15}\text{N}_5$, 96%	2	1
Hypoxanthine	HPX	$^{13}\text{C}_5$, 99%	10	1
L-Leucine	Leu	$^{13}\text{C}_6$, 99%	5	1
L-Phenylalanine	Phe	ring- $^{13}\text{C}_6$, 99%	100	1
Thymine	T	1,3- $^{15}\text{N}_2$, 98%	20	1
L-Tryptophan	Trp	$^{13}\text{C}_{11}$, 99%	100	1
L-Tyrosine	Tyr	ring- $^{13}\text{C}_6$, 99%	100	1
Vitamin B ₃	nicotinamide	$^{13}\text{C}_6$, 99%	5	1
Citric acid	CA	1,5,6-carboxyl- $^{13}\text{C}_3$, 99%	10	2
Fumaric acid	FA	$^{13}\text{C}_4$, 99%	100	2
Indole-3-acetic acid	IAA	phenyl- $^{13}\text{C}_6$, 99%	5	2
α-Ketoglutaric acid, disodium salt	α-KG	1,2,3,4- $^{13}\text{C}_4$, 99% (CP 97%)	100	2
Sodium palmitate	16:0	U- $^{13}\text{C}_{16}$, 98%	10	2
Sodium pyruvate	Pyr	$^{13}\text{C}_3$, 99%	100	2

Usage Specifications

Criteria	Recommendation
Before reconstitution:	
Storage	-20°C; protect from light
Recommended retest	2 years from date of manufacture
Upon reconstitution:	
Storage	4°C
Recommended retest	4 weeks

Example Results

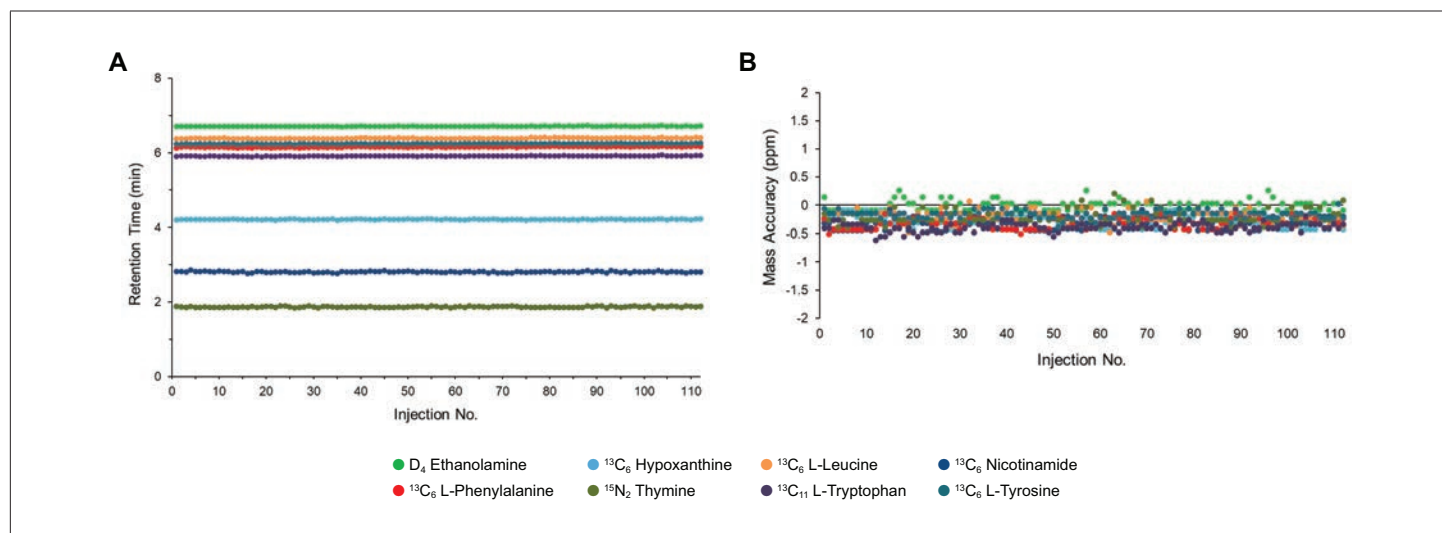


Figure. QC results for a subset of stable isotope-labeled QReSS metabolites measured across calibrants, QCs, and FBS samples. Acquisitions were by HILIC-MS (ESI+, Orbitrap Exploris™ 120). Variability of retention times are shown in **A** and mass accuracy in **B**. For additional information on the application, see CIL application note #51.

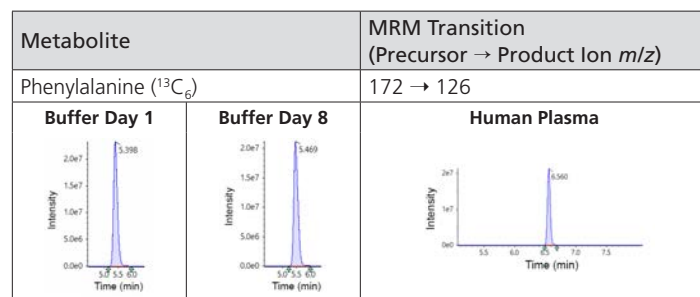
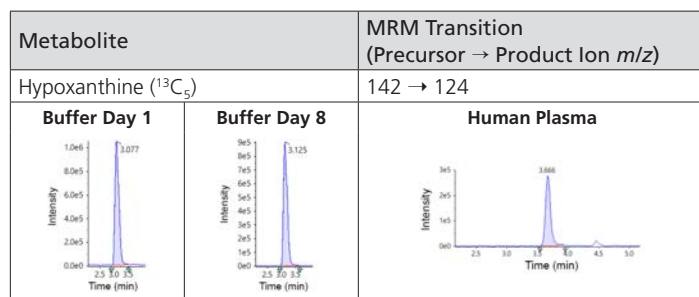


Figure. Example XICs for a subset of labeled QReSS metabolites measured in buffer and human plasma by RPLC-MRM/MS (positive ESI, QTRAP 6500+). In the buffer analysis, a vial of a working stock mix was stored in on autosampler rack (maintained at 4°C) and processed eight days apart. Such measurements, when performed and metric tracked routinely, would constitute a system suitability test. In the human plasma analysis, an aliquot of the working stock mix was matrix spiked and subjected to a metabolomics workflow. The relative signals to its corresponding endogenous analyte (not shown for simplicity) is within an order of magnitude and can be used for relative quantitative applications, with absolute quantitation likely involving calibration curves. For additional information and application demonstrations see CIL application note #49.

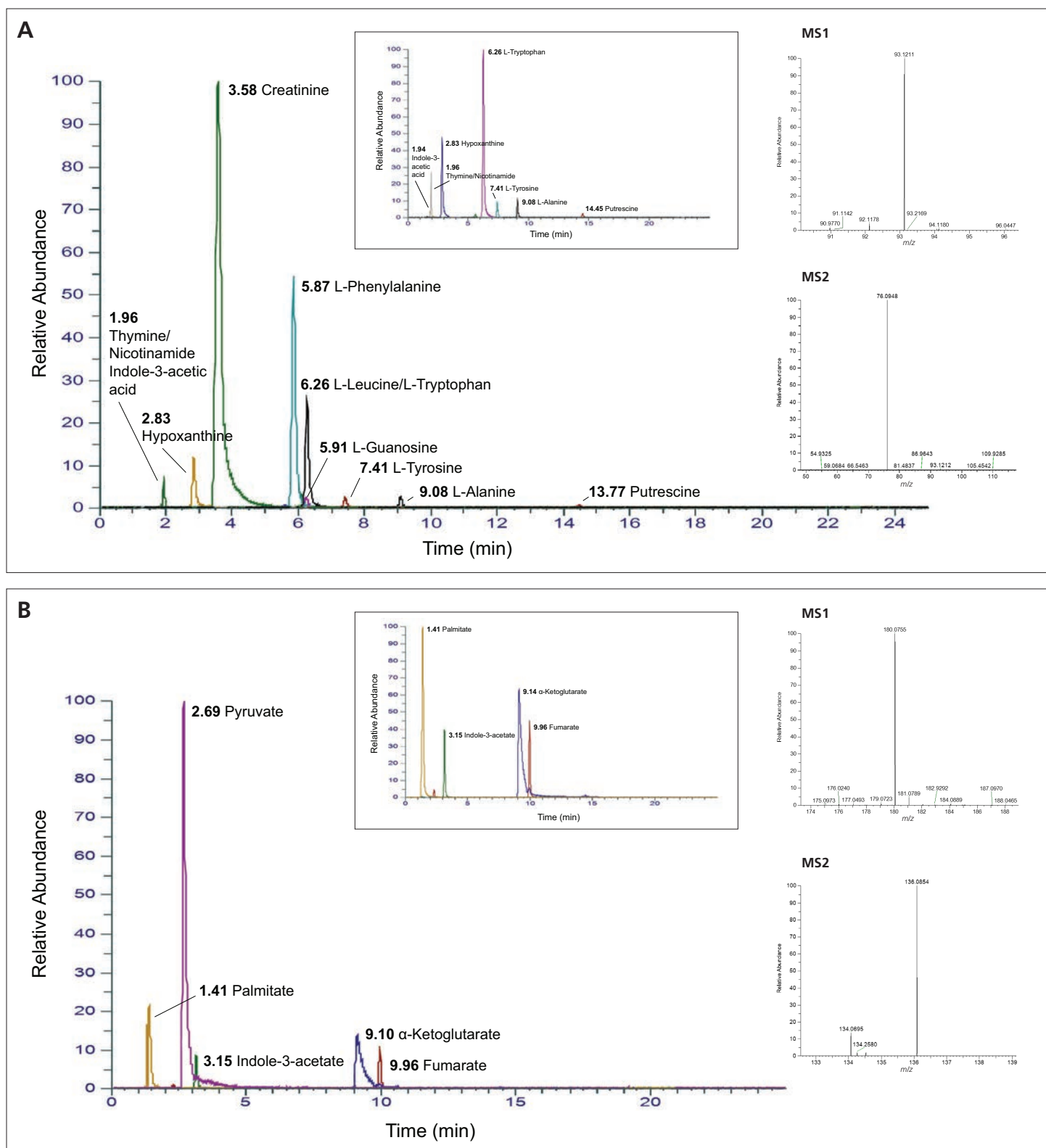


Figure. XIC overview of stable isotope-labeled QReSS metabolites measured matrix-free by HILIC-MS (Q Exactive), with ESI+ in **A** and ESI- in **B**. Chromatographic insets reflect zoomed-in snapshots of lower-abundance metabolites, while the MS1/MS2 spectra reflect putrescine in **A** and indole-3-acetate in **B**. For clarity, the following higher-abundance metabolites were removed in the insets – creatinine (RT 3.58 min), phenylalanine (RT 5.87 min), and leucine (RT 6.26 min) in **A** and pyruvate (RT 2.69 min) in **B**. Data is courtesy of the Jackson Laboratory.

Example References

MSK-QC-KIT

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MSK-QReSS-KIT

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MSK-QCMSI

- Mills, Q.; Kibbe, R.R.; Sohn, A.L.; Percy, A.J.; et al. **2025**. Part A: Implementing an analyte panel and sampling protocol for quality control in mass spectrometry imaging. *Rapid Commun Mass Spectrom*, 39(8), e999.
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Application Notes

MSK-QReSS Mixes/Kits

- Percy, A.J.; Souza, A.; Ntai, I.; et al. **2022**. From QC to quantitation: Utility of QReSS™ metabolites in FBS measurements. (CIL application note #51)
- Percy, A.J.; Proos, R.; Demianova, Z.; et al. **2021**. Standardizing quantitative metabolomics analyses through the QReSS kit. (CIL application note #49)

“The unique feature of our media analysis workflow is the use of the QReSS standard mix. We spike this into media samples before extraction, which helps normalize variabilities in metabolite extraction efficiencies and combat matrix effects. The QReSS mix consists of 18 isotope-labeled metabolites, which spans a multitude of metabolic classes and mimics the largely diverse chemical composition of cell culture media. This serves as an excellent choice for an internal standard mix and helps to improve the accuracy, precision, and robustness of our cell culture method.”

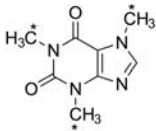
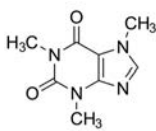
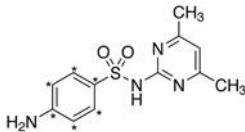
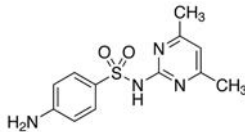
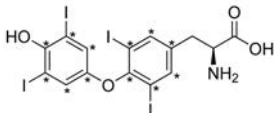
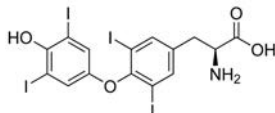
– Hari Kosanam, PhD
Associate Principal Scientist, Vaccine Process Development and Commercialization
Merck & Co. (USA)

MSK-QCMSI Kit

This kit contains two vials of 3-plex mixtures (dried down), where the stable isotope-labeled (SIL) and unlabeled (natural, NAT) metabolites are contained separately. Separation of the NAT and SIL mixes enables flexibility to independently monitor different features of an ionization source, such as ionization efficiency and electrospray stability. The suggested reconstitution of the mixtures produces 1 mL of 1 mM stock solutions, supporting 50 QC experiments or more, depending on a user's experimental strategy. The manual protocol was established by the Muddiman lab on an IR-MALDESI platform,¹⁻³ but can be adapted to alternate imaging platforms (e.g., MALDI, DESI, etc.) and may be suitable for cross-platform applications.

Composition

Table. Composition of the SIL and NAT mixes. Reconstituting each vial in 1 mL of solvent (e.g., 50% ACN) produces 1 mM equimolar stock solutions. The isotopic enrichments of the labeled standards are 99%. The precursor ions m/z are $[M+H]^+$.

SIL Vial (Catalog No. MSK-QCMSI)		NAT Vial (Catalog No. MSK-QCMSI-US)	
¹³ C ₃ Caffeine (m/z 198.0977)		Unlabeled caffeine (m/z 195.0877)	
¹³ C ₆ Sulfamethazine (m/z 285.1111)		Unlabeled sulfamethazine (m/z 279.0910)	
¹³ C ₁₂ L-Thyroxine (m/z 789.7342)		Unlabeled L-thyroxine (m/z 777.6940)	

Usage Specifications

Criteria	Recommendation
Before reconstitution:	
Storage	-20°C; protect from light
Recommended retest	2 years from date of manufacture

References

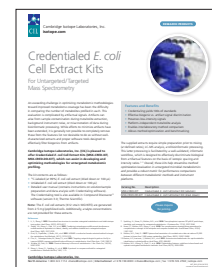
- Kibbe, R.R.; Mills, Q.; Sohn, A.L.; Muddiman, D.C. **2025.** Part B: SLICE-MSI – a machine-learning interface for system suitability testing of mass spectrometry imaging platforms. *Rapid Commun Mass Spectrom*, 39(8), e9990.
- Kibbe, R.R.; Sohn, A.L.; Muddiman, D.C. **2024.** Leveraging supervised machine learning algorithms for system suitability testing of mass spectrometry imaging platforms. *J Proteome Res*, 23(10), 4384-4391.
- Mills, Q.; Kibbe, R.R.; Sohn, A.L.; Percy, A.J.; et al. **2025.** Part A: Implementing an analyte panel and sampling protocol for quality control in mass spectrometry imaging. *Rapid Commun Mass Spectrom*, 39(8), e999.

E. coli and Yeast Extract Kits

Expanded qualification remains a necessity in MS omics today. Validated identifications are desired that can be used to springboard in-depth profiling studies and selective quantifications in preclinical screening studies. To aid method benchmarking (e.g., through a credentialing approach) and improved qualification/quantification studies, CIL offers two types of cell extracts: yeast (*Pichia pastoris*, strain CBS 7435) and *E. coli* (K12 strain MG1655), as outlined in the overview table below. Upon careful and precise solubilization (see reconstitution guidelines below), 100s of metabolites are potentially observable in their U-¹³C and unlabeled form. The metabolites span a broad class range (see example identification tables), having linkage to various biochemical pathways (e.g., citrate and glyoxylate cycle, nucleotide and lipid metabolism) and cellular/molecular processes (e.g., intracellular signaling, immune system, blood coagulation, lipolysis). Please refer to the sample results and references below for examples of the product-specific applications. **Note:** Analyte concentrations are not provided for these extracts.

Overview

Catalog No.	Description	No. of Analytes	Kit Contents
L-ISO1	Crude Lipid Yeast Extract (U- ¹³ C, 99%)	100s	1 vial of U-¹³C crude lipid yeast extract - document package (user manual, CoA, SDS, product flyer)
L-ISO1-UNL	Crude Lipid Yeast Extract (unlabeled)	100s	1 vial of unlabeled crude lipid yeast extract - document package (user manual, CoA, SDS, product flyer)
ISO1	Metabolite Yeast Extract (U- ¹³ C, 99%)	100s	1 vial of U-¹³C metabolite yeast extract - document package (user manual, CoA, SDS, product flyer)
ISO1-UNL	Metabolite Yeast Extract (unlabeled)	100s	1 vial of unlabeled metabolite yeast extract - document package (user manual, CoA, SDS, product flyer)
ISO1-KIT	Metabolite Yeast Extract Kit	100s	1 vial of U-¹³C metabolite yeast extract 1 vial of unlabeled metabolite yeast extract - document package (user manual, CoA, SDS, product flyer)
MSK-CRED-DD-KIT	Credentialed <i>E. coli</i> Cell Extract Kit (dried down)	100s	1 vial (blue cap) of ¹³C-labeled <i>E. coli</i> cell extract (dried down) 1 vial (yellow cap) of unlabeled <i>E. coli</i> cell extract (dried down) - document package (user manual, CoA, SDS, product flyer)
MSK-CRED-KIT	Credentialed <i>E. coli</i> Cell Extract Kit (solution)	100s	1 vial (black) of ¹³C-labeled <i>E. coli</i> cell extract (solution) 1 vial (red) of unlabeled <i>E. coli</i> cell extract (solution) - document package (user manual, CoA, SDS, product flyer)



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Lipid Yeast Extracts

Compositions

Tabulated is a reproducibly identified panel of fatty acids and lipids measured in the lipid yeast extracts (U-¹³C, L-ISO1; unlabeled, L-ISO1-UNL) using one general RPLC-MS method. **Note:** Additional analytes have been measured with alternate methods and analysis techniques. Analyte concentrations are not provided for these extracts.

Category	Class	Examples (Fatty Acid Level)			
Glycerolipids	Diglyceride (DG)	34:1 (16:0/18:1)	34:2 (16:1/18:1)	34:3 (16:1/18:2)	
		34:4 (16:2/18:2)	36:1 (18:0/18:1)	36:2 (18:0/18:2)	
		36:3 (18:1/18:2)	36:4 (18:1/18:3)	36:5 (18:2/18:3)	
	Triglyceride (TG)	50:1 (16:0/16:0/18:1)	50:2 (16:0/18:2/16:0)	50:3 (16:1/16:1/18:1)	
		50:4 (16:1/16:1/18:2)	50:5 (16:1/16:1/18:3)	52:1 (18:0/16:0/18:1)	
		52:2 (16:0/18:1/18:1)	52:3 (16:1/18:1/18:1)	52:4 (18:2/18:2/16:0)	
		52:5 (16:1/18:2/18:2)	52:6 (16:1/18:2/18:3)	52:7 (18:3/18:3/16:1)	
		54:1 (18:0/18:1/18:0)	54:2 (18:0/18:1/18:1)	54:3 (18:1/18:1/18:1)	
		54:4 (18:1/18:1/18:2)	54:5 (18:1/18:2/18:2)	54:6 (18:2/18:2/18:2)	
		54:7 (18:1/18:3/18:3)	54:8 (18:3/18:2/18:3)	54:9 (18:3/18:3/18:3)	
Glycerophospholipids	Phosphatidic acid (PA)	34:1 (16:0/18:1)	34:2 (16:0/18:2)	34:3 (16:0/18:3)	
		36:1 (18:0/18:1)	36:2 (18:0/18:2)	36:3 (18:1/18:2)	
	Phosphatidylcholine (PC)	34:1 (16:0/18:1)	34:2 (16:0/18:2)	34:3 (16:0/18:3)	
		34:4 (16:1/18:3)	36:2 (18:0/18:2)	36:3 (18:1/18:2)	
		36:4 (18:1/18:3)	36:5 (18:2/18:3)	36:6 (18:3/18:3)	
	Phosphatidylethanolamine (PE)	34:1 (16:0/18:1)	34:2 (16:0/18:2)	34:3 (16:0/18:3)	
		36:1 (18:0/18:1)	36:2 (18:0/18:2)	36:3 (18:1/18:2)	
		36:4 (18:1/18:3)	36:5 (18:2/18:3)	36:6 (18:3/18:3)	
	Phosphatidylglycerol (PG)	34:1 (16:0/18:1)	34:2 (16:1/18:1)		
	Phosphatidylinositol (PI)	34:1 (16:0/18:1)	34:2 (16:0/18:2)	36:1 (18:0/18:1)	
		36:2 (18:1/18:1)			
Lysophospholipids	Lysophosphatidylcholine (LPC)	16:1	18:1	18:2	18:3
	Lysophosphatidylethanolamine (LPE)	18:1	18:2	18:3	
Sphingolipids	Ceramide (Cer)	d34:1 (d18:1/16:0)	d36:1 (d18:1/18:0)	t34:1 (t18:1/16:0)	
	Hexosyl ceramide (HexCer)	t36:0 (t18:0/18:0)	t36:2 (t18:1/18:1)		
Other	Acylcarnitines (AC)	16:1			
	Coenzyme (Co)	Q8			
	Fatty acid (FA)	18:2	18:3		

Usage Specifications

Extract Type	L-ISO1	L-ISO1-UNL
No. of uses	~50*	
<i>Before reconstitution:</i>		
Storage	-80°C; protect from light	
Expiration	4 years from date of manufacture	

*Conservative estimate, with numbers varying depending on method and application.

Solution Preparation Procedure Example

1. Reconstitute the lipid yeast extract (L-ISO1 or L-ISO1-UNL) in 1 mL solvent (e.g., isopropanol).
2. Following a high-speed vortex and brief centrifuge, the resulting clear solution can then be matrix-spiked, diluted (1/10 v/v), or prepared further for calibrant, QC, and/or sample addition.

Example Results

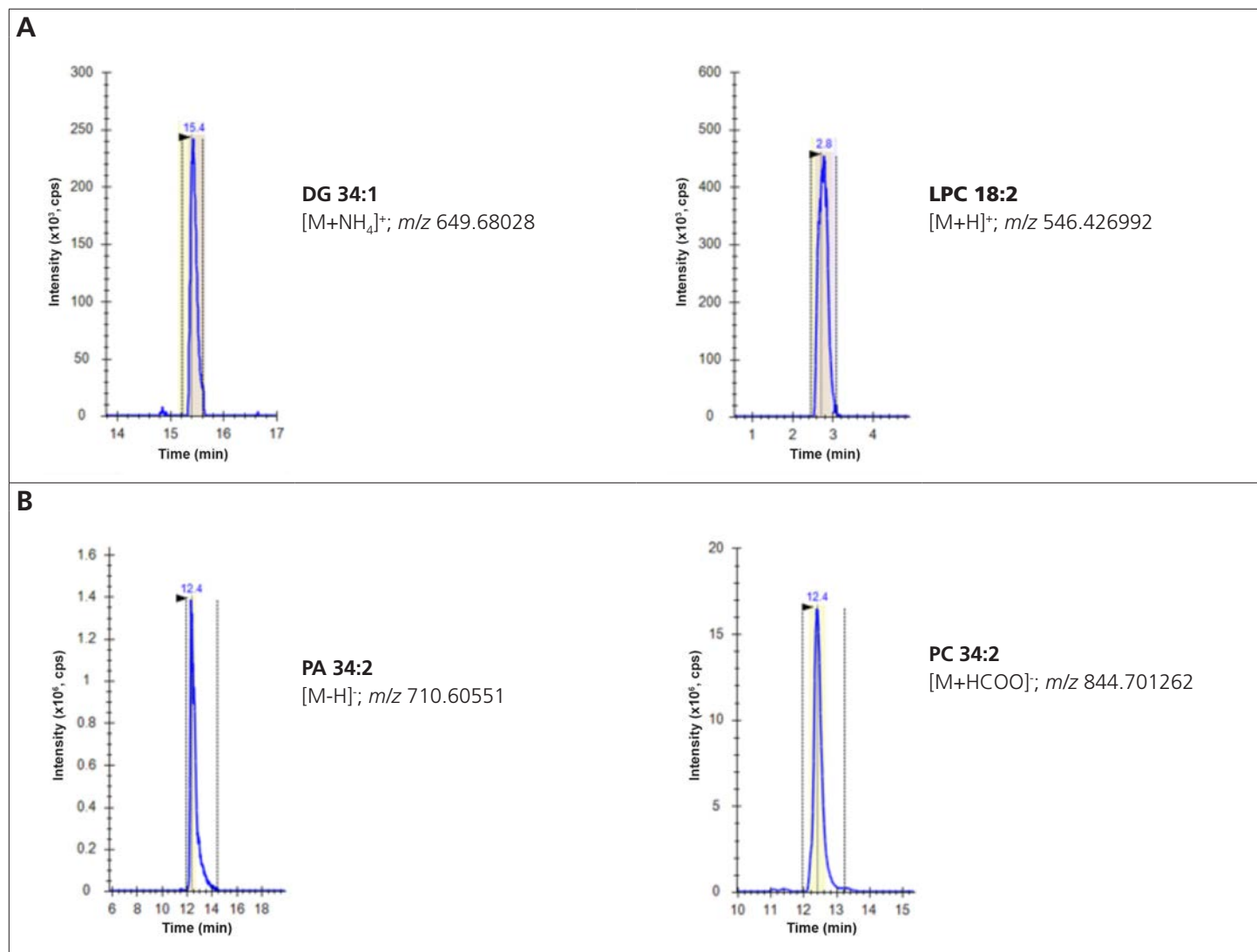


Figure. Representative XICs of example lipids measured in the U-¹³C lipid yeast extract by RPLC-MS (ACQUITY HSS T3, Orbitrap ID-X). The ionization mode was ESI+ in **A** and ESI- in **B**. Data is courtesy of the Patti lab at Washington University.

Table. Example set of identified fatty acids and lipids measured by RPLC-IM-MS (ACQUITY™ Premier CSH™ C18 column, SYNAPT™ XS). Tabulated is analytical data for unlabeled and U-¹³C lipids acquired under ESI+ and ESI- (in blue) polarities. Error tolerances on the ion mobility-derived collision cross-sections (CCS) are ±3%. Data is courtesy of Waters (Manchester, UK).

Lipid (Fatty Acid Level)	RT (min)	Precursor <i>m/z</i>		Adduct	CCS (Å ²)	
		Unlabeled Lipids	U- ¹³ C Lipids		Unlabeled Lipids	U- ¹³ C Lipids
Cer d34:1 (d18:1/16:0)	3.31	560.5023	594.6124	[M+Na] ⁺	257.79	258.44
Cer d34:1 (d18:1/16:0)	3.31	536.5059	570.6159	[M-H] ⁻	252.85	253.30
PC 34:2 (16:0/18:2)	2.85	758.5700	800.7074	[M+H] ⁺	293.16	294.46
PC 36:4 (18:1/18:3)	2.53	782.5697	826.7160	[M+H] ⁺	300.69	301.52
PE 34:2 (16:0/18:2)	2.96	716.5220	755.6510	[M+H] ⁺	279.92	280.33
PE 34:2 (16:0/18:2)	2.96	738.5032	777.6310	[M+Na] ⁺	286.03	286.47
LPC(16:1)	0.82	494.3223	518.4045	[M+H] ⁺	229.29	229.61
LPC(16:1)	0.82	538.3145	562.3950	[M+HCOO] ⁻	238.29	239.23

Metabolite Yeast Extracts

Compositions

Tabulated are the routinely identified analytes in the metabolite yeast extracts (U-¹³C, ISO1; unlabeled, ISO1-UNL). **Note:** Additional metabolites have been measured with alternate methods and analysis techniques. This includes cofactors (e.g., NMN, ADPR, NADPH) and coenzymes (e.g., acetyl and malonyl coenzyme A). **Note:** Analyte concentrations are not provided for these extracts.

Amino Acids and Derivatives (L enantiomer where applicable)		
S-Adenosyl-homocysteine (SAH)	Glutamate (Glu)	Methionine (Met)
Alanine (Ala)	Glutamine (Gln)	(±)-3-Methyl-2-oxovalerate (K-IVal)
α-Aminoadipate (AAD)	Glycine (Gly)	Ornithine (Orn)
Arginine (Arg)	Guanidineacetate (GAA)	Phenylalanine (Phe)
Argininosuccinate (ASA)	Histidine (His)	Proline (Pro)
Asparagine (Asn)	Homoserine (Hse)	Sarcosine (Sar)
Aspartate (Asp)	Isoleucine (Ile)	Serine (Ser)
Betaine (BET)	α-Ketoisovalerate (KIV)	Threonine (Thr)
Citrulline (Cit)	Kynurenine (KYN)	Tryptophan (Trp)
Cystathionine (CYS)	Leucine (Leu)	Tyrosine (Tyr)
Dihydroxyisovalerate (DIHV)	Lysine (Lys)	Valine (Val)
Nucleobases, Nucleosides, and Nucleotides		
Adenine (Ade)	Cytidine triphosphate (CTP)	Inosine (Isin)
Adenosine (Asin)	Deoxyadenosine monophosphate (dAMP)	Inosine monophosphate (IMP)
Adenosine diphosphate (ADP)	5'-Deoxy-5'-methylthioadenosine (MTAP)	5-Methyluridine (m ⁵ U)
Adenosine monophosphate (AMP)	Guanine (Gnin)	Pseudouridine (PsU)
Adenosine triphosphate (ATP)	Guanosine (Gsin)	Uridine (Uri)
Cyclic adenosine monophosphate (cAMP)	Guanosine diphosphate (GDP)	Uridine diphosphate (UDP)
Cyclic guanosine monophosphate (cGMP)	Guanosine monophosphate (GMP)	Uridine monophosphate (UMP)
Cytidine monophosphate (CMP)	Guanosine triphosphate (GTP)	Uridine triphosphate (UTP)
Organic Acids		
cis-Aconitate (cis-Ac)	DL-2-Hydroxyglutarate (2-HG)	Malate (Mal)
Citrate (CA)	Isocitrate (Iso)	Pyruvate (Pyr)
Fumarate (Fum)	α-Ketoglutarate (α-KG)	Succinate (SA)
Gluconate (GA)	Lactate (Lac)	
Sugar and Sugar Phosphates (D enantiomer where applicable)		
Dihydroxyacetone phosphate (DHAP)	Glucose (Glc)	6-Phosphogluconate (6PGA)
Erythritol (Erthrit)	Glucose-6-phosphate (G6P)	Ribose (RIB)
Fructose (Fuc)	Mannitol (Man-OI)	Ribose-5-phosphate (R5P)
Fructose-1,6-bisphosphate (FBP)	Mannose (Man)	Sedoheptulose-7-phosphate (S7P)
Fructose-6-phosphate (F6P)	Mannose-6-phosphate (M6P)	Trehalose (TRE)
Galactose (Gal)	2-Phosphoglycerate (2PG)	
Vitamins and Coenzymes		
Biotin (B ₇)*	Nicotinamide (NAM)	Nicotinamide adenine dinucleotide, reduced (NADH)
Choline (CHOL)	Nicotinamide adenine dinucleotide, oxidized (NAD ⁺)	Nicotinamide adenine dinucleotide phosphate, oxidized (NADP ⁺)
Other Small Molecules		
Glutamylcysteine (Glu-Cys)	Glutathione, reduced (GSH)	
Glutathione, oxidized (GSSG)	Mevalonate (MVA)	

*Identified in ISO1-UNL only.

Usage Specifications

Extract Type	ISO1	ISO1-UNL
No. of uses	~50*	
Before reconstitution:		
Storage	-80°C; protect from light	
Expiration	6 years from date of manufacture	
Upon reconstitution:		
Storage	4°C	
Recommended retest	4 weeks	

*Conservative estimate, with numbers varying depending on method and application.

Solution Preparation Procedure Example

1. Reconstitute the metabolite yeast extracts (ISO1 or ISO1-UNL) with 2 mL solvent (e.g., water, 50% methanol).
 2. Vigorously shake by hand with intermittent high-speed vortexing (2 minute minimum).
 3. Centrifuge at 20°C for 5 min at 4000 rcf.
 4. The clear standard solution can then be diluted (1/10 v/v) for direct use or prepared further for calibration and matrix addition.
- If particulate matter remains after steps 2) and 3), repeat with extra intense vortexing and shaking until complete dissolution.



A video demonstration of this procedure can be obtained here, as well as in the resources section of the Metabolite Yeast Extracts application page (see isotope.com/applications/metabolic-research/metabolomics-mixes-and-kits/metabolite-yeast-extracts).

Example Results

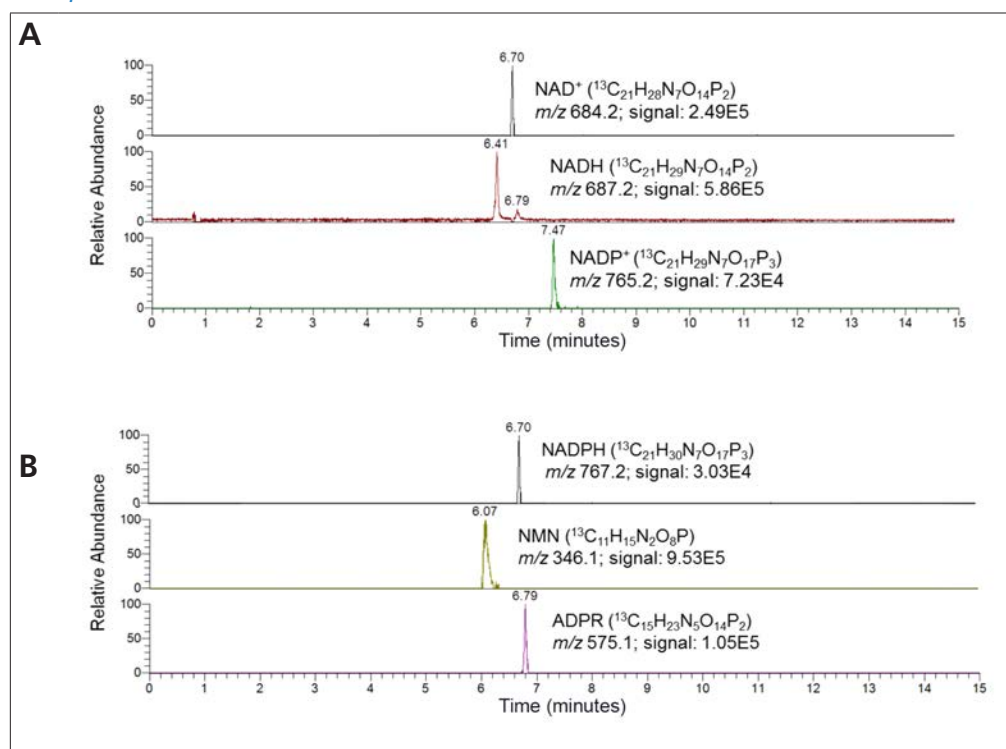


Figure. Representative XICs of ¹³C-labeled cofactors measured by HILIC-MS (positive ESI, Q Exactive HF). Shown in **A** are the routinely identified metabolites and **B** the additionally observed cofactors. Procedurally, the extract was reconstituted in 2 mL water then diluted 1:10 before analysis. The HILIC separation utilized an Acquity UPLC BEH Amide column (100 × 2.1 mm, 1.7 μm particles).

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Application Notes

- Percy, A.J.; Munjoma, N.; Heywood, D.; et al. **2022**. Targeted MRM screening of U-¹³C lipid yeast extracts for robust lipidomics applications. (CIL application note #54)
- Mohsin, S.B.; Batoon, P. **2022**. Absolute quantitation of fragile metabolites by isotope dilution mass spectrometry on the Agilent 6495 Triple Quadrupole LC/MS. (Agilent application no. 5994-4439EN)

E. coli Cell Extracts

Compositions

Condensed table of example metabolites detected in the *E. coli* extract samples (see PMID: 29256075 for application details). The spread of logPs (source: ALOGPS) highlights the breadth of physicochemical diversity. **Note:** Analyte concentrations are not provided for these extracts.

Compound (Abbrev. or Alt. Name)	LogP	Metabolite Class
Adenine (Ade)	-0.38	Nucleobase
Adenosine monophosphate (AMP)	-2.30	Nucleotide
Biotin (vitamin B ₇)	0.30	Vitamin
Coenzyme Q10 (CoQ10; ubiquinone)	9.94	Coenzyme
Diacylglycerol (DG) 17:0/17:0	10.16	Lipid
Diacylglycerol (DG) 18:0/18:2	10.28	Lipid
Diacylglycerol (DG) 18:1/18:1	10.26	Lipid
Elaidic acid (EA; <i>trans</i> 18:1)	7.68	Fatty acid
Glutathione (GSH)	-2.70	Peptide
Oleic acid (OA; 18:1)	7.68	Fatty acid
Palmitic acid (PAL; 16:0)	7.23	Fatty acid
Palmitoyl CoA (PAL-CoA)	2.35	Coenzyme
Phosphatidylcholine (PC) 18:2/18:2	5.68	Lipid
Phosphatidylethanolamine (PE) 16:1/16:1	7.89	Lipid
Phosphatidylethanolamine (PE) 18:1/18:1	8.81	Lipid
Phenethylamine (PEA)	1.41	Neurotransmitter
Proline (Pro)	-2.70	Amino acid
Phosphatidylserine (PS) 18:1/18:1	4.96	Lipid
Retinol (vitamin A)	6.38	Vitamin
Stearic acid (STE; 18:0)	8.02	Fatty acid
Thymidine (Thd)	-1.30	Nucleoside
Tryptophan (Trp)	1.04	Amino acid
Uracil (U)	-1.20	Nucleobase
Uridine (Uri)	-1.80	Nucleoside

Usage Specifications

Criteria	Recommendation
No of sample runs	≤20 (with 5 µL LC-MS injections)
Storage	-80°C; protect from light, air, and moisture
Recommended retest	4 years

Solution Preparation Procedure Example

1. Reconstitute the dried-down extracts in 100 µL of ACN:water (e.g., 1:1 v/v). **Note:** Alternate solvent ratios are possible without impact to the credentialing approach. Changing the ratio to increase ACN or water will simply increase the coverage of organic or water-soluble metabolites, respectively.
2. Sonicate briefly the solutions (equates to 40 µL/mg of *E. coli* extract) then centrifuge (e.g., for 15 min at 13,000 RPM and 4°C) and incubate overnight at 4°C. This results in a clear solution.
3. Mix aliquots of the unlabeled and ¹³C-labeled extracts (at 99% isotopic enrichment) into autosampler vials at distinct ratios (e.g., 1:2 and 1:1 v/v). Vortex briefly.
4. Load the vials onto an autosampler rack for the benchmarking optimizations.

[illegible]

PeptiQuant™ Plus Assay Kits

Researchers in academia and life science industries continue to implement a bottom-up MS-based workflow for protein biomarker screening. Biomarker verification/validation requires absolute quantification of surrogate peptides in a sample matrix, a requirement that is best achieved using well-characterized stable isotope-labeled standards. To ensure robust quantitative measurements, QC checks should be routinely performed. CIL offers a collection of PeptiQuant Assay Kits (from MRM Proteomics Inc.) for QC and biomarker assessment using bottom-up LC-MS/MS methodologies. The QC kits are designed to evaluate the performance of an LC-MS platform, either alone or in combination with a human plasma proteomics workflow (see corresponding flyer for panel details). The biomarker assessment kits (BAKs) are intended to help researchers screen target panels of candidate protein disease biomarkers in human or mouse plasma samples (see corresponding flyer for panel details). The current platform-specific offerings for each kit type are listed in the overview below.



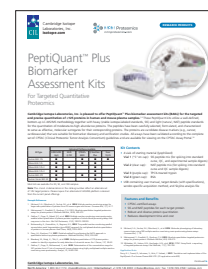
Overview

QC Kits					
Catalog No.	Description	Kit Contents	No. of Peptides	Unit Size	Optimized Instrument
LCMSP-QC	PeptiQuant Plus Human Plasma Daily QC Kit	$^{13}\text{C}/^{15}\text{N}$-labeled peptide mix - USB (e.g., user manual, acquisition and analysis files)	35	10, 20, or 50 injections	6490/6495 QqQ - QTRAP® 6500 - Q Exactive™ Plus
WFPK	PeptiQuant Plus Human Plasma Workflow QC Kit	$^{13}\text{C}/^{15}\text{N}$-labeled peptide mix - unlabeled peptide mix - trypsin - BSA - human plasma - USB (e.g., user manual, acquisition and analysis files)	35	1 or 2 runs	6490/6495 QqQ - QTRAP 6500 - Q Exactive Plus

Biomarker Assessment Kits (BAKs)					
Catalog No.	Description	Kit Contents	No. of Quant. Proteins	Unit Size	Optimized Instrument
Human					
BAK-125	PeptiQuant Plus Human Plasma Proteomics Kit	$^{13}\text{C}/^{15}\text{N}$-labeled peptide mix - unlabeled peptide mix - trypsin - BSA - USB (e.g., user manual, acquisition and analysis files)	125	20, 50, or 100 samples	6490/6495 QqQ - QTRAP 6500 - Q Exactive Plus - Xevo TQ-XS
BAK-270	Expanded PeptiQuant Plus Human Plasma Proteomics Kit	$^{13}\text{C}/^{15}\text{N}$-labeled peptide mix - unlabeled peptide mix - trypsin - BSA - USB (e.g., user manual, acquisition and analysis files)	270	20, 50, or 100 samples	6490/6495 QqQ - QTRAP 6500 - Q Exactive Plus
BAK-CNCR50	DiseaseQuant Human Tissue Cancer Pathway Proteomics Kit	$^{13}\text{C}/^{15}\text{N}$-labeled peptide mix - unlabeled peptide mix - trypsin - BSA - USB (e.g., user manual, acquisition and analysis files)	50	50 or 100 samples	6490/6495 QqQ - Q Exactive Plus
Mouse					
M-BAK-125*	PeptiQuant Plus Mouse Plasma Proteomics Kit	$^{13}\text{C}/^{15}\text{N}$-labeled peptide mix - unlabeled peptide mix - trypsin - BSA - USB (e.g., user manual, acquisition and analysis files)	125	20, 50, or 100 samples	6490/6495 QqQ - QTRAP 6500 - Q Exactive Plus 6545 Q-TOF

*Alternate sets of 125 target proteins are available (see PeptiQuant Plus BAK flyer for details).

PeptiQuant is a trademark of MRM Proteomics Inc.



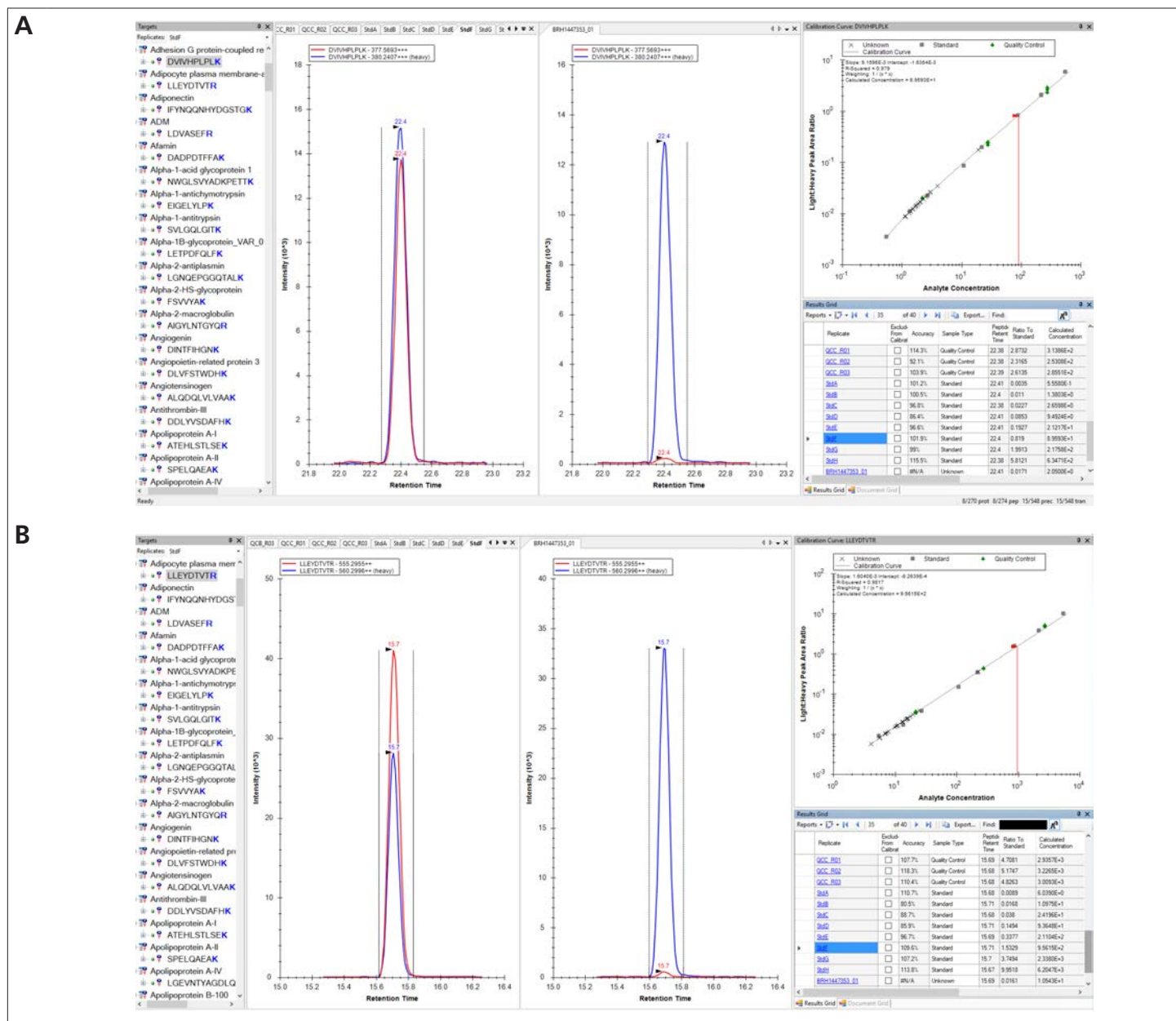
Click on the thumbnails or visit isotope.com for more information.

Usage Specifications

Before reconstitution (for peptide mixes):	
Storage	-80°C
Recommended retest	6 months from date of delivery

Note: The dried-down trypsin and BSA materials supplied with the BAKs are not stable at room temperature and must be stored at 4°C. Trypsin is to be prepared immediately prior to use and stored on ice until then.

Example Results



Example References

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Application Note

Percy, A.J.; Trouvé, R.; Lehmann, S.; Hirtz, C. Vialaret, J. **2021**. Translation and implementation of the PeptiQuant™ Plus Human Plasma BAK-270. (CIL application note #50)

“The PeptiQuant Plus Platform Performance Kit has proven to be a vital component of our everyday quality assurance that enables us to deliver high-quality targeted proteomics data in an accurate and timely manner. This kit has a ‘dilute and shoot’ operation and comes with vendor-specific LC-MRM/MS parameters and a Skyline analysis file for quick input and results output. Altogether, the performance kit is an excellent means to rapidly assess LC-MS performance that should become a routine staple in a proteomic user’s toolbox.”

– Tasso Miliotis, PhD

Associate Principal Scientist at AstraZeneca Gothenburg (Sweden)

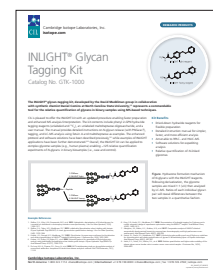
“PeptiQuant Plus Assay Kits contain all the essential materials, including the standards and methods, for performing absolute protein quantification by LC-MRM/MS in a standardized way. The standard protocol helped us reduce the assay development time, while improve the reproducibility and precision of multiplex protein quantification. In addition to the biomarker assessment kits, the quality control kits enable the instrument performance and assay reproducibility to be monitored and assessed, which ultimately provided us confidence in the reliability of the quantification results.”

– Elaine Wong, PhD

Scientific Officer at Queen Mary Hospital, Fu Lam (Hong Kong)

INLIGHT® Glycan Tagging Kit

Glycans participate in a large number of cellular, molecular, and biological processes and are implicated in a number of diseases (e.g., Alzheimer's, cancer). To aid the identification and relative quantification of glycans by LC-MS, CIL offers the innovative INLIGHT® (Individuality Normalization when Labeling with Isotopic Glycan Hydrazide Tag) glycan-tagging kit. This kit employs both natural (NAT – $^{12}\text{C}_6$) and stable isotope-labeled (SIL – $^{13}\text{C}_6$) phenyl 2-GPN reagents in the hydrazide derivatization of free *N*-glycans. While this tagging strategy was developed for *N*-glycans, it has also been adapted to sample analysis of *O*-glycan and heparin oligomer profiles. The INLIGHT kit contains five vials of NAT reagent and five vials of SIL reagent, which in total provides sufficient tagging for approximately 125 relative quantification experiments. The user manual provides step-by-step instructions for executing the modified INLIGHT strategy using maltoheptaose and fetuin A as examples. Data processing and analysis of derivatized glycans can be facilitated in GlycoHunter or Skyline.



Click on the thumbnail or visit isotope.com for more information.

Overview

Catalog No.	Description	Kit Contents
GTK-1000	INLIGHT® Glycan Tagging Kit	• Light phenyl 2-GPN INLIGHT reagent (5 × 0.25 mg) • Heavy phenyl 2-GPN INLIGHT reagent ($^{13}\text{C}_6$, 5 × 0.25 mg) • Unlabeled maltoheptaose (5 × 10 µg)

Usage Specifications

Criteria	Recommendation
No. of Uses	25 per vial
Before reconstitution:	
Storage	ambient temperature; protect from light and moisture
Recommended retest	5 years from date of manufacture

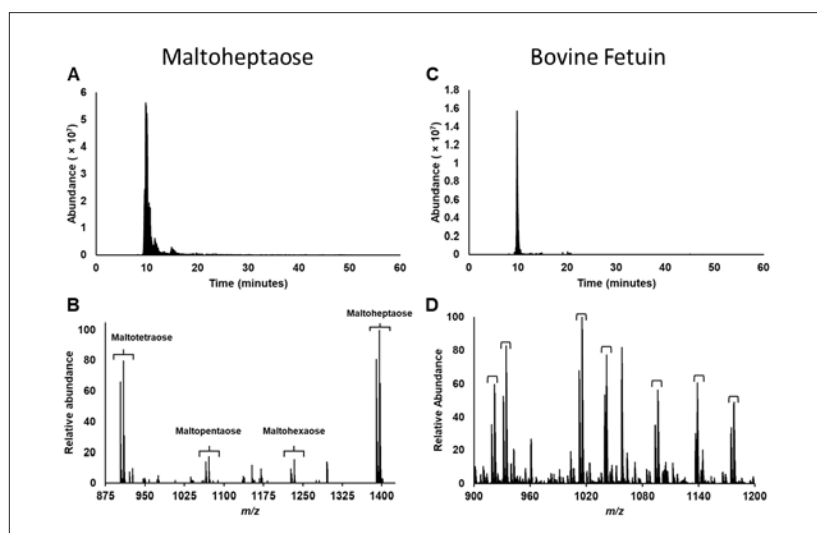


Figure. LC-MS examples of INLIGHT derivatized maltoheptaose and *N*-linked glycans from bovine fetuin.

- A.** Extracted ion chromatogram of $[\text{NAT} + \text{H}]^+$ maltoheptaose.
B. Mass spectrum at 10 minutes illustrating NAT and SIL derivatized maltoheptaose, maltohexaose, maltopentaose, and maltotetraose.
C. Extracted ion chromatogram of a fetuin-based glycan at m/z 1012.3980 corresponding to $[\text{NAT} + 2\text{H} + (\text{Fuc})_1(\text{Gal})_2(\text{GlcNAc})_4(\text{Man})_3]^{2+}$.
D. Average mass spectrum of a fetuin analysis showing the NAT and SIL derivatized *N*-linked glycan pairs in the LC window 8-11 min.

Example References

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Q&As

Listed below are a series of general Q&As for CIL's mixes and kits. Product-specific FAQs can be located on isotope.com's [FAQs](#) page and on their corresponding product application page (e.g., Life Sciences → Metabolomics Mixtures and Kits → [QC Mixes and Kits](#)).

What are the advantages of selecting an off-the-shelf mix vs. a collection of individual isotope standards for self-mixing?

The MSK/NSK products are formulated to exacting standards following detailed batch records developed from over 20 years of formulation experience. Following production, randomly selected vials are analyzed to ensure both accuracy and consistency. The astute attention and process control generates exceptional vial-to-vial and lot-to-lot reproducibility. This high reproducibility return is one of the merits of utilizing CIL prepared mixes. Additional advantages are:

- reduced development time and cost;
- enhanced data quality;
- ease of user implementation; and
- improved confidence in analytical results.

The majority of the mixes are ^{13}C - and/or ^{15}N -labeled.

What are the advantages of $^{13}\text{C}/^{15}\text{N}$ -labeling vs. D-labeled in MS measurements?

The nature of the stable isotope can potentially impact the preanalytical (e.g., storage and handling) and analytical (e.g., sample preparation and processing) phases of an experiment. In comparison to D labels, $^{13}\text{C}/^{15}\text{N}$ labels can have:

- improved isotope stability;
- negligible isotope scrambling issues;
- conserved chromatographic elution (relative to its unlabeled standard); and
- heightened analytical reliability.

In cases where D-labeled compounds were selected, the D-label is located at nonexchangeable positions and was stability tested for preservation as well as product application. For further background, please refer to our [technical note](#) that describes the benefits of ^{13}C vs. D standards in MS-based studies.

How often should QC measurements be performed, and what mixes are most suitable?

Prior to first use, QCs should be conducted several times in succession prior to sample analysis to establish baseline performance and intervention limits. Once established, QCs should be performed routinely (i.e., before, during, and after sample analysis) to monitor the effectiveness of the analytical method and instrument platform over time. While all of CIL's mixes could be used in these types of operations, the ones ideally suited for these type of measurements are the metabolomic QC kits (see [MSK-QC-KIT](#) and [MSK-QReSS-KIT](#)) and the proteomic QC kits (see [PeptiQuant Plus](#) daily and workflow QC kits).

In a targeted quantitative application, how many MRM transitions should be monitored per analyte?

Ideally, a minimum of two MRM transitions per compound (be it metabolite or peptide) should be targeted. This will allow for a quantifier and qualifier(s) assignment, with the ion ratios serving as an additional metric for performance qualification. Nonetheless, this may not be possible for all analytes given the compound's fragmentation chemistry, the employed MS/MS parameters, and the mode of operation utilized. Empirical experiments with isotope-labeled standards should therefore be first performed to optimize the MS/MS parameters prior to conducting the quantitative study with precious experimental samples.

Can custom mixes or add-on vials be formulated?

Yes, we have the ability to customize. We would first review feasibility and then provide a quotation on your specific mix. To start this process, please provide the necessary details on this [Custom Mix Request](#) form or contact your local sales representative.

Please visit isotope.com for a complete list of isotope-labeled compounds.

Research products are distributed and sold worldwide via our extensive network.

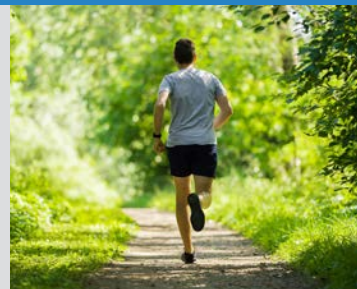
CIL's distributor listing is available at isotope.com.

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