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Infant formula and adult nutritionals — Simultaneous determination of total vitamins B1, B2, B3 and B6 — Enzymatic digestion and LC-MS/MS

Formules infantiles et produits nutritionnels pour adultes — Détermination simultanée de la teneur en vitamines B1, B2, B3 et B6 - Digestion enzymatique et CL-SM/SM

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Foreword

ISO (the International Organization for Standardization) is a worldwide federation of national standards bodies (ISO member bodies). The work of preparing International Standards is normally carried out through ISO technical committees. Each member body interested in a subject for which a technical committee has been established has the right to be represented on that committee. International organizations, governmental and non-governmental, in liaison with ISO, also take part in the work. ISO collaborates closely with the International Electrotechnical Commission (IEC) on all matters of electrotechnical standardization.

The procedures used to develop this document and those intended for its further maintenance are described in the ISO/IEC Directives, Part 1. In particular, the different approval criteria needed for the different types of ISO documents should be noted. This document was drafted in accordance with the editorial rules of the ISO/IEC Directives, Part 2 (see www.iso.org/directives).

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For an explanation of the voluntary nature of standards, the meaning of ISO specific terms and expressions related to conformity assessment, as well as information about ISO's adherence to the World Trade Organization (WTO) principles in the Technical Barriers to Trade (TBT) see www.iso.org/iso/foreword.html.

This document was prepared by Technical Committee ISO/TC 34, *Food products* in collaboration with AOAC INTERNATIONAL. It is being published by ISO and separately by AOAC INTERNATIONAL. The method described in this document is equivalent to the AOAC Official Method 2015.14: *Simultaneous Determination of Total Vitamins* B_1, B_2, B_3, *and* B_6 *in Infant Formula and Related Nutritionals by Enzymatic Digestion and LC-MS/MS.*

Any feedback or questions on this document should be directed to the user's national standards body. A complete listing of these bodies can be found at www.iso/org/members.html.

Infant formula and adult nutritionals — Simultaneous determination of total vitamins B1, B2, B3 and B6 — Enzymatic digestion and LC-MS/MS

1 Scope

This document specifies a method for the simultaneous quantitative determination of four water-soluble vitamins in infant formula and related nutritional products, including relevant forms of vitamins B_1, B_2, B_3 and B_6 by enzymatic digestion and UHPLC-MS/MS.

2 Normative references

There are no normative references in this document.

3 Terms and definitions

For the purposes of this document, the following terms and definitions apply.

ISO and IEC maintain terminological databases for use in standardization at the following addresses:

— ISO Online browsing platform: available at https://www.iso.org/obp

— IEC Electropedia: available at http://www.electropedia.org/

3.1
adult nutritional
nutritionally complete, specially formulated food, consumed in liquid form, which may constitute the sole source of nourishment, made from any combination of milk, soy, rice, whey, hydrolysed protein, starch and amino acids, with and without intact protein

3.2
infant formula
breast-milk substitute specially manufactured to satisfy, by itself, the nutritional requirements of infants during the first months of life up to the introduction of appropriate complementary feeding

[SOURCE: Codex Standard 72-1981]

4 Principle

Samples are prepared by enzymatic digestion with papain and α-amylase to hydrolyze protein and complex carbohydrate and acid phosphatase to free phosphorylated vitamin forms. Stable-isotope labelled internal standards are incorporated into the sample preparation to correct for variability in both the sample preparation and instrument response. A series of six mixed working standard solutions spanning two orders of magnitude in vitamin concentration are used to generate calibration curves based on the peak response ratio of the analyte to its stable-isotope labelled internal standard.

Prepared samples and working standard solutions are injected onto ultra-high pressure liquid chromatograph (UPLC) interfaced to a triple-quadrupole mass spectrometer (MS/MS) for analysis. The MS/MS is configured to monitor parent-daughter (precursor-fragment) ion pairs for each analyte and internal standard. This reaction forms the basis for method selectivity. Analytes are quantified by least squares regression using the response ratio of the analyte to its internal standard.

5 Reagents and materials

During the analysis, unless otherwise stated, use only reagents of recognized analytical grade and distilled or demineralized water or water of equivalent purity.

5.1 Nicotinamide, primary reference standard, for example USP Reference Standard, Catalog # 1462006[1)].

Store as indicated on label.

5.2 Niacin (nicotinic acid), primary reference standard, for example USP Reference Standard, Catalog # 1461003[1].

Stored as indicated on label.

5.3 Pyridoxine hydrochloride, primary reference standard, for example USP Reference Standard, Catalog # 1587001[1].

Store in desiccator protected from white light. Dry according to manufacturer's instructions prior to use.

5.4 Riboflavin, primary reference standard, for example USP Reference Standard, Catalog # 1603006[1].

Store in desiccator protected from white light. Dry according to manufacturer's instructions prior to use.

5.5 Thiamine hydrochloride, primary reference standard, for example USP Reference Standard, Catalog # 1656002[1].

Store in desiccator protected from white light. Measure the moisture content of the powder prior to use.

5.6 Pyridoxamine dihydrochloride

Fluka Analytical Standard, catalog# P9380.[1]

5.7 Pyridoxal hydrochloride

Sigma, catalog# P9130.[1]

5.8 $^{2}H_{4}$-Nicotinamide

CDN Isotopes, Catalog # D-3457[1].

5.9 $^{2}H_{4}$-Nicotinic acid

CDN Isotopes; Catalog # D-4368[1].

5.10 $^{13}C_{4}$-Pyridoxine: pyridoxine:HCl (4,5-bis(hydroxymethyl)-$^{13}C_{4}$)

Cambridge Isotope Laboratory;Catalog # CLM-7563[1].

5.11 $^{2}H_{3}$-Pyridoxal.

IsoSciences; Catalog # 7098[1].

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1) This is an example of a suitable product available commercially. This information is given for the convenience of users of this document and does not constitute an endorsement by ISO of the product named. Equivalent products may be used if they can be shown to lead to the same results.

5.12 $^{2}H_{3}$-Pyridoxamine.

IsoSciences; Catalog # 7099[1].

5.13 $^{13}C_{4}$-Thiamine chloride.

IsoSciences; Catalog # 9209[1].

5.14 $^{13}C_{4}$,$^{15}N_{2}$-Riboflavin.

IsoSciences, Catalog# 7072[1].

5.15 Acid phosphatase, type II from potato, 0,5 U/mg to 3,0 U/mg.

Sigma, catalog# P3752[1].

5.16 Papain from Carica papaya, ≥ 3 U/mg.

Sigma, catalog# 76220[1].

5.17 α–amylase from aspergillus oryzae, 150 U/mg.

Sigma, catalog# A9857[1].

5.18 Hydrochloric acid concentrated (substance concentration c = 12 mol/l).

ACS grade, or equivalent.

5.19 Ammonium formate, for mass spectrometry (purity ≥ 99,0%).

Fluka 70221 or equivalent[1].

5.20 Glacial acetic acid.

Sigma ACS Reagent Grade, or equivalent[1].

5.21 Formic acid.

Sigma ACS Reagent Grade, or equivalent[1].

5.22 Laboratory water.

18,0 MΩ, < 10 ppb TOC, or equivalent.

5.23 Methanol.

Fisher LC-MS/MS Optima grade or EMD Omni-Solve LC-MS grade[1].

5.24 Ethylenediaminetetracetic acid, disodium salt dihydrate (EDTA).

ACS grade (99 % to 101 %), or equivalent.

5.25 Potassium phosphate dibasic.

ACS grade (purity > 98 %), or equivalent.

5.26 meta-Phosphoric acid.

ACS grade (33,5 % to 36,5 %), or equivalent.

5.27 Buffer solutions for pH meter calibration, pH = 4,0, 7,0, and 10,0.

5.28 Phosphoric acid, 85 %.

ACS grade, or equivalent.

5.29 Potassium hydroxide, 40 %.

ACS grade, or equivalent.

6 Standard and solution preparation

6.1 Mobile phase A, substance concentration c = 0,020 mol/l ammonium formate in water.

Using a graduated cylinder, transfer 500 ml laboratory water to a mobile phase reservoir. Add 0,631 g of ammonium formate (5.19) and mix well. Expiration is 3 days.

6.2 HCl solution, c = 0,12 mol/l.

Add approximately 300 ml of water to a 500 ml graduated cylinder. Add 5,0 ml ± 0,1 ml of concentrated HCl solution (5.18) and swirl to mix. Bring to 500 ml with laboratory water and mix well.

6.3 Acetic acid solution, 1,0 %.

Add approximately 30 ml of water to a 500 ml graduated cylinder. Add 5,0 ml ± 0,1 ml of glacial acetic acid (5.20) and swirl to mix. Bring to 500 ml with laboratory water and mix well.

6.4 Mobile phase B, methanol.

6.5 Weak needle wash, 10 % methanol in water, expiration 3 months.

6.6 Strong needle wash, methanol.

6.7 Ammonium formate solution, c = 0,050 mol/l.

Using a graduated cylinder, transfer 1 400 ml of laboratory water to an appropriate reservoir. Add 4,41 g of ammonium formate (5.19) and mix well. 1 400 ml is adequate for 6 working standards and 32 samples. Scale as needed. Expiration is 3 days.

6.8 Mixed enzyme solution

Using a graduated cylinder, transfer 200 ml of ammonium formate buffer (6.7) to an appropriate reservoir. Add 200 mg ± 10 mg of acid phosphatase (5.15), 80 mg ± 5 mg of α-amylase (5.17) and 400 mg ± 10 mg of papain (5.16). Mix for 10 min with a magnetic stir plate and stir bar. Check pH and adjust to 4,25 ± 0,25 with formic acid (5.21, ~ 100 µl). 200 ml is adequate for 6 working standards and 32 samples. Scale as needed. Prepare fresh daily.

6.9 Stable isotope labelled compounds, individual, internal standard stock solutions

6.9.1 General

Internal standard stock solutions have an expiration of 6 months. However, the following guidelines can be used to troubleshoot internal standard and, when documented as part of routine system suitability checks, extend the expiration dates indefinitely.

Based on U.S. FDA bioanalytical method validation guidelines that the lowest-level calibration shall be 5 times the analyte response of the blank, see,[9] the channel of the non-labelled analyte of interest shall be monitored to ensure the stable isotope-labelled internal standard does not contribute more than 20 % of the area count of the lowest-level calibration standard. No response should be generated in any other channels being monitored in the method, as this is a sign of contamination, in which case fresh solution should be prepared or fresh lot of material should be ordered.

The area count of the internal standard should be at least 3 times the area count of the analyte in the lowest-level calibration standard and the lowest level matrix-based QC sample.

6.9.2 $^{2}H_{4}$-Nicotinamide stock solution, mass concentration $\rho \approx 560$ µg/ml.

Weigh 14,0 mg ± 0,1 mg into a tarred weighing vessel. Quantitatively transfer to a 25 ml volumetric flask with laboratory water and fill to the mark with laboratory water. Mix well and transfer to a 50 ml amber bottle and store refrigerated (2 °C to 8 °C). Expiration, see 6.9.1.

6.9.3 $^{2}H_{4}$-Nicotinic acid stock solution, $\rho \approx 500$ µg/ml.

Weigh 12,5 mg ± 0,1 mg into a tarred weighing vessel. Quantitatively transfer to a 25 ml volumetric flask with laboratory water and fill to the mark with laboratory water. Mix well and transfer to a 50 ml amber bottle and store refrigerated (2 °C to 8 °C). Expiration, see 6.9.1.

6.9.4 $^{13}C_{4}$-Pyridoxine stock solution, $\rho \approx 70$ µg/ml.

Weigh 7,0 mg ± 0,1 mg into a tarred weighing vessel. Quantitatively transfer to a 100 ml volumetric flask with laboratory water and fill to the mark with laboratory water. Mix well and transfer to a 100 ml amber bottle and store refrigerated (2 °C to 8 °C). Expiration, ee 6.9.1.

6.9.5 $^{2}H_{3}$-Pyridoxal stock solution, $\rho \approx 40$ µg/ml.

Weigh 4,0 mg ± 0,1 mg into a tarred weighing vessel. Quantitatively transfer to a 100 ml volumetric flask with laboratory water and fill to the mark with laboratory water. Mix well and transfer to a 100 ml amber bottle and store refrigerated (2 °C to 8 °C). Expiration, ee 6.9.1.

6.9.6 $^{2}H_{3}$-Pyridoxamine stock solution, $\rho \approx 40$ µg/ml.

Weigh 4,0 mg ± 0,1 mg into a tarred weighing vessel. Quantitatively transfer to a 100 ml volumetric flask with laboratory water and fill to the mark with laboratory water. Mix well and transfer to a 100 ml amber bottle and store refrigerated (2 °C to 8 °C). Expiration, ee 6.9.1.

6.9.7 $^{13}C_{4}$-Thiamine chloride stock solution, $\rho \approx 100$ µg/ml.

Weigh 5,0 mg ± 0,1 mg of $^{13}C_{4}$-thiamine into a tarred weighing vessel. Quantitatively transfer to a 50 ml volumetric flask with HCl solution (6.2) and fill to the mark with HCl solution (6.2). Mix well and transfer to a 100 ml amber bottle and store refrigerated (2 °C to 8 °C). Expiration, ee 6.9.1.

6.9.8 $^{13}C_4$,$^{15}N_2$-Riboflavin stock solution, $\rho \approx 73$ µg/ml.

Weigh 7,3 mg ± 0,1 mg of $^{13}C_4$,$^{15}N_2$-riboflavin into a tarred weighing vessel. Quantitatively transfer to a 100 ml volumetric flask with acetic acid solution (6.3) and fill to the mark with acetic acid solution (6.3). Mix well and transfer to a 100 ml amber bottle and store refrigerated (2 °C to 8 °C). Expiration, ee 6.9.1.

6.10 Internal standard stock mixture (ISSM).

Combine 2 500 µl of ammonium formate solution (6.7) with 250 µl of $^{2}H_4$-nicotinamide stock solution (6.9.2), 250 µl of $^{2}H_4$-nicotinic acid stock solution (6.9.3), 250 µl of $^{13}C_4$-pyridoxine stock solution (6.9.4), 200 µl of $^{2}H_3$-pyridoxal stock solution (6.9.5), 50 µl of $^{2}H_3$-pyridoxamine stock solution (6.9.6), 250 µl of $^{13}C_4$-thiamine stock solution (6.9.7) and 250 µl of $^{13}C_4$,$^{15}N_2$-riboflavin stock solution (6.9.8). Volume provides sufficient ISSM for 6 working standards and 32 samples. Scale as needed. Prepare fresh daily.

6.11 Phosphate buffer solution, pH = 5,0 (0,010 mol/l potassium phosphate dibasic, 1 % EDTA, 2 % metaphosphoric acid.

Weigh 20,0 g ± 0,2 g of EDTA into a tarred weighing vessel and quantitatively transfer to a 2 000 ml beaker containing approximately 1 800 ml laboratory water and add a magnetic stir bar.

Weigh 34,8 g ± 0,1 g of potassium phosphate dibasic into a tarred weighing vessel and quantitatively transfer to the 2 000 ml beaker already containing approximately 1 800 ml laboratory water and EDTA. Mix by stirring on a magnetic stir plate until both the EDTA and potassium phosphate dibasic is completely dissolved.

Weigh 40,0 g ± 0,2 g of metaphosphoric acid into a tarred weighing vessel and quantitatively transfer to the 2 000 ml beaker containing approximately 1 800 ml laboratory water, EDTA, and potassium phosphate dibasic. Mix by stirring on a magnetic stir plate until the metaphosphoric acid is completely dissolved.

Adjust the pH of the solution to pH = 5,00 ± 0,02 using 40 % potassium hydroxide or 85 % phosphoric acid. Quantitatively transfer the solution to a 2 000 ml volumetric flask and dilute to volume with laboratory water. Expiration: 48 hours.

6.12 Stock standard solutions of native compounds

6.12.1 Vitamin standard stock mixture (VSSM)

Accurately weigh the indicated amounts for the following standards using separate weighing funnels or other appropriate weighing vessel and quantitatively transfer to a 100 ml volumetric flask using phosphate buffer (pH = 5):

a) niacinamide: 70,5 mg ± 0,5 mg;

b) thiamine hydrochloride: 10,5 mg ± 0,2 mg ;

   Determine the moisture of the thiamine hydrochloride reference standard (5.5) as directed on the container immediately prior to weighing. The percent moisture determined for the reference standard is used to calculate the concentration of thiamine in the VSSM.

c) riboflavin: 7,0 mg ± 0,2 mg;

   Dry an appropriate amount of the riboflavin reference standard (5.4) at 105°C ± 1 °C for 2 h (± 10 min) prior to weighing.

d) pyridoxine hydrochloride: 10,8 mg ± 0,2 mg;

   Dry an appropriate amount of the pyridoxine hydrochloride reference standard (5.3) over indicating absorbent in vacuo for 4 h prior to weighing.

