



Product certificate ERNDIM IQCS Amino Acids

Product name Control Amino Acids

Product code

Product code	Colour cap	
AMI-02.1	Green	
AMI-02.2	Red	

Date of issue 31-10-2019

Batch numbers and Expiry date

Batch number	Exp. date stored at +2°C to +8°C	
LOT 2019.1571	2024-05	
LOT 2019.1572	2024-05	

Reconstitution volume

1.0 mL

Estimated concentrations *

Analyte	Estimated concentrations (µmol/L)		
	Level 1	Level 2	
α-aminobutyric acid	26	89	
Alanine	299	878	
Arginine	25	490	
Asparagine	36	186	
Aspartic acid	13	74	
Citrulline	17	404	
Cystathionine	5	28	
Cystine	24	54	
Glutamic acid	59	261	
Glutamine	535	1051	
Glycine	297	940	
Histidine	105	406	
Hydroxyproline	36	84	
Isoleucine	31	376	
Leucine	74	849	
Lysine	78	479	
Methionine	10	230	
Ornithine	59	598	
Phenylalanine	78	907	
Proline	164	563	
Serine	47	409	
Taurine	45	391	
Threonine	99	383	
Tryptophan	137	317	
Tyrosine	51	874	
Valine	152	770	

 $[\]ensuremath{^{*}}$ See ERNDIM Internal Quality Control System at the reverse



PC-AMI-02_EN.v2

Amino Acids ERNDIM IQCS

Intended purpose

These materials are control material (thus no calibrators) for the internal control of analytical systems for the determination of amino acids in serum.

Contents

Lyophilized human serum to which amino acids have been added to achieve an analytically and physiologically relevant level of the amino acids.

Storage and stability

The product in lyophilized form is stable for 5 years when stored at +2°C to + 8°C. Expiration dates are found on the product certificate (reverse). The stability of the reconstituted product is comparable to patient samples.

Instructions for use

- a. Remove cap and stopper.
- b. Add 1 mL aqua destillata
- c. Replace stopper
- d. Let stand for 15 minutes at room temperature
- e. Mix carefully during 20 minutes at room temperature
- f. Process product as patient sample, i.e. it is advisable to immediately deproteinise samples and separate the supernatant to minimise stability problems of certain amino acids.
- g. If not analysed on the same day according to your usual procedure for patient samples in your laboratory, the supernatant should be stored at -24°C to -16°C.

ERNDIM Internal Quality Control System: the Concept

The ERNDIM Internal Quality Control System (IQCS) consists of samples and a website for data management.

Samples

Samples contain analytes specifically selected for laboratories active in the field of inborn errors of metabolism. They come in two levels (1=low and 2=high) with for each analyte a relevant concentration.

Data Management

ERNDIM offers users of control materials a data management system (Note: this is an option to serve users; users do not have the obligation to use it). The strength of this system is that it does not only monitor the data of the laboratory but also compares the labs results with results of labs using the same batch of internal control materials.

In essence users can submit results every time they do an analytical run with the control material and then download two reports.

The Review Day Report shows the results of the last run in comparison to

- a) the acceptance limits set by the lab,
- b) the mean of all previous runs of the lab
- c) the mean of all laboratories.

By clicking on the name of a specific analyte in the report, Shewhart charts of that analyte are shown.

The Cumulative Table report shows the cumulative data of the lab.

Details can be found under www.erndimga.nl/General information/Use Website.

Remark

On delivery of the control materials, the certificate in the package insert shows the values as measured by a peer laboratory. Once in use laboratories submit their results and the reports will show the trimmed mean of all laboratories. This mean is a running mean which changes with every new submission: Thus a dynamic assigned value resulting from "crowd targeting".

Precautions and warnings

- 1. For in vitro diagnostic use only.
- 2. Tested and found negative for Hepatitis B Surface Antigen (HbsAg), antibody to hepatitis C (HCV) and antibody to HIV.
- 3. This product should be handled with care, as appropriate for biological materials. Outdated and left-over material should be discarded as potentially infectious material, according to the procedures in your institute.

References

www.ERNDIMQA.nl

Dr C.W. Weykamp on behalf of the ERNDIM Internal Quality Control System Working Group