





# Certificate of Analysis QCS Pterins in urine

**Product name** Quality comparison material Pterins in urine

Product code	Product code	Colour cap
	PTE-01.1	Green
	PTE-01.2	Red

**Date of issue** 18 March 2026

Batch numbers and Expiry date	Batch number	Exp. date stored at +2°C to +8°C
	 2024.1211	 <b>2024-04</b>
	 2024.1212	 <b>2024-04</b>

**Reconstitution volume** 1.0 mL

Estimated concentrations *	Analyte	Estimated concentrations (µmol/L)	
		Level 1	Level 2
	Neopterin	16.0	127
	Biopterin	11.5	61.7
	Primapterin (7-Biopterin)	3.99	54.9
	Isoxanthopterin	41.4	502

\* See ERNDIM Quality Comparison System at the reverse

## Intended purpose

Not for diagnostic use.

This material is intended solely for the verification of the correct calibration of measurement methods. It is not permitted to use this material (without CE mark) in establishing or assessing patient/client results.

## Disclaimer

It is the responsibility of the customer to limit the use of this material to the intended use described above. Streekeziekenhuis Koningin Beatrix (department MCA laboratory) is not responsible or liable for application outside the intended use.

## Contents

Lyophilized human urine to which pterins have been added to achieve an analytically and physiologically relevant level of the pterins.

## Storage and stability

The product in lyophilized form is stable for 2 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

- Remove cap and stopper.
- Add 1.0 mL aqua destillata
- Replace stopper
- Let stand for 15 minutes at room temperature
- Mix carefully during 20 minutes at room temperature
- Let stand for 15 minutes at room temperature (do not expose to sunlight)
- Process product as patient sample.
- 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 Quality Comparison System: the Concept**

The ERNDIM Quality Comparison System (QCS) 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 these 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 comparison materials.

In essence users can submit results every time they do an analytical run with the comparison 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.erndimqa.nl/General information/Use Website](http://www.erndimqa.nl/General%20information/Use%20Website).

#### *Remark*

On delivery of the materials, the certificate in the package insert shows the values (estimated concentrations) 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* use only.
2. 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](http://www.ERNDIMQA.nl)