

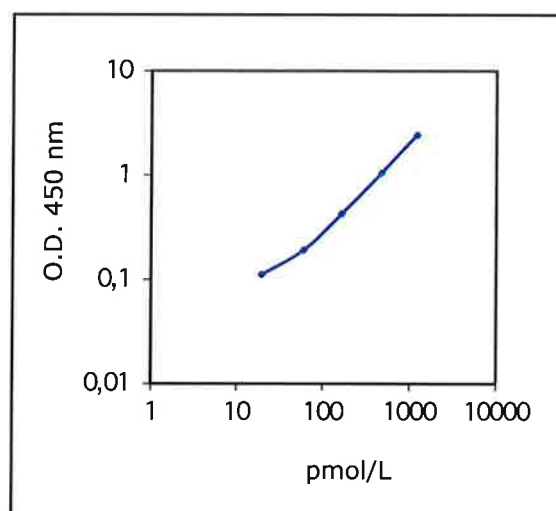
Certificate of Analysis

1. Manufacturer

Mercodia AB, Sylveniusgatan 8A, 754 50 Uppsala, SWEDEN

2. Description

Catalog no: 10-1256-01
Product: Mercodia Porcine C-peptide
ELISA
Lot no: 37092
Expiry date: 2028-03-31



Component	Art no	Lot no	O.D. 450 nm	Exp. date
Calibrator 0	20-6977	36690	0,063	2028-04-06
Calibrator 19,4 pmol/L	20-6983	36694	0,111	2028-04-04
Calibrator 60,1 pmol/L	20-6984	36695	0,191	2028-04-04
Calibrator 164 pmol/L	20-6985	36696	0,426	2028-04-04
Calibrator 470 pmol/L	20-6986	36697	1,047	2028-04-04
Calibrator 1210 pmol/L	20-6987	36698	2,428	2028-04-04
Coated Plate	20-7001	35225		2028-04-11
Assay Buffer	20-6999	36693		2028-04-06
Enzyme Conjugate 11X	20-6995	36691		2028-04-12
Enzyme Conjugate Buffer	20-6997	36692		2028-04-06
Wash Buffer 21X	20-6746	36958		2032-12-03
Substrate TMB	20-2629	36313		2029-02-28
Stop Solution	20-2693	36746		2032-07-01

3. Quality control

Quality control has been performed for lot no 37092 according to standard operating procedures at Mercodia AB, and the product is released based on fulfillment of established acceptance criteria.

4. Calibration

The Mercodia Porcine C-peptide ELISA is calibrated against an in-house reference preparation of porcine c-peptide.

5. Assay method

Test procedure used is according to current Direction for Use for the product and lot.

6. Intended use

Mercodia Porcine C-peptide ELISA provides a method for the quantitative determination of porcine c-peptide in serum, plasma and cell culture medium.

7. Storage and handling

Recommended storage of kit is 2-8°C.

Storage of unused or diluted kit components is stated in the Direction for Use.

8. Hazardous information

Please refer to the Material Safety Data Sheet for hazard identification.

9. Quality standard documentation

The Mercodia Quality System fulfills the QSR and the requirements for the European CE mark, the Directive 98/79/CE on *in vitro* diagnostic medical devices. Mercodia is ISO 13485 certified, for compliance with the International Quality Management System for medical devices.

10. Names and signatures of certifying officers

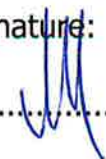
Date of analysis:

2025-06-17

Performed by:

Mattias SL

Signature:



Date of approval:

2026-01-22

Approved by:

Elin Westberg

Signature:

