Mercodia

Mall-0674 v 1.0

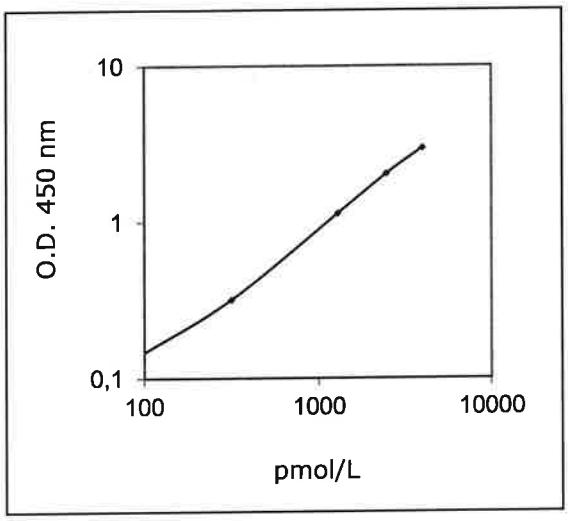
Certificate of Analysis

1. **Manufacturer**

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

2. **Description**

Catalog no:	10-1136-03
Product:	Mercodia C-peptide ELISA
Lot no:	37697
Expiry date:	2028-07-31



Component		*Art no*	*Lot no*	*O.D. 450 nm*	*Exp. date*
Calibrator 0		20-3415	36814	0,062	2028-09-04
Calibrator 97,8	pmol/L	20-3416	36815	0,145	2028-08-26
Calibrator 315	pmol/L	20-3417	36816	0,319	2028-08-26
Calibrator 1280	pmol/L	20-3418	36817	1,137	2028-08-26
Calibrator 2450	pmol/L	20-3419	36818	2,044	2028-08-26
Calibrator 3980	pmol/L	20-3420	36819	2,983	2028-08-26
Coated Plate		20-3422	36810		2028-09-08
Assay Buffer		20-2933	36813		2028-08-28
Enzyme Conjugate 11X		20-3425	36820		2028-09-18
Enzyme Conjugate Buffer		20-3427	36812		2028-08-27
Wash Buffer 21X		20-6746	37528		2033-08-28
Substrate TMB		20-2629	36313		2029-02-28
Stop Solution		20-2693	37299		2033-05-29

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E-mail info@mercodia.com | **Website** www.mercodia.com

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3. Quality control

Quality control has been performed for lot no 37697 according to standard operating procedures at Mercodia AB, and the product is released based on fulfillment of established acceptance criteria.

4. Calibration

The Mercodia C-peptide ELISA is calibrated against the International Reference Reagent for C-peptide, IRR C-peptide 84/510.

5. Assay method

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

6. Intended use

Mercodia C-peptide ELISA provides a method for the quantitative determination of human c-peptide in serum, plasma or urine.

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

Mercodia is ISO 13485 certified, demonstrating compliance with the international quality management system standard for medical devices. This product is supplied for research use only and is not intended for use as an in vitro diagnostic (IVD) medical device.

10. Names and signatures of certifying officers

Date of analysis:	Performed by:	Signature:
2025-11-03	Mattias Jad	

Date of approval:	Approved by:	Signature:
2026-10-28	Jonas Krick	

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