Mercodia

Mall-0264 v2.0

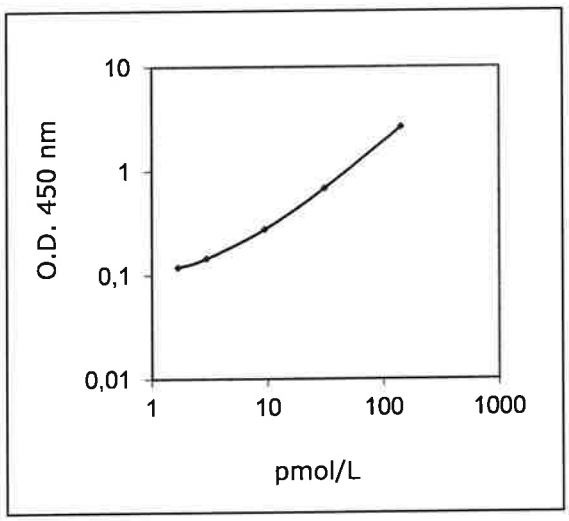
Certificate of Analysis

1. Manufacturer

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

2. Description

Catalog no:	10-1271-01
Product:	Mercodia Glucagon ELISA
Lot no:	37390
Expiry date:	2029-07-31



Component		*Art no*	*Lot no*	*O.D. 450 nm*	*Exp. date*
Calibrator 0		20-7040	37380	0,085	2029-09-03
Calibrator 1,67	pmol/L	20-7042	37381	0,119	2029-08-26
Calibrator 2,94	pmol/L	20-7044	37382	0,145	2029-08-26
Calibrator 9,36	pmol/L	20-7046	37383	0,276	2029-08-26
Calibrator 30,8	pmol/L	20-7048	37384	0,686	2029-08-26
Calibrator 141	pmol/L	20-7050	37385	2,653	2029-08-26
Assay Buffer		20-7551	37614		2029-09-28
Coated Plate		20-7057	37387		2029-09-14
Enzyme Conjugate 11X		20-7053	37388		2029-08-25
Enzyme Conjugate Buffer		20-7055	37389		2029-08-12
Wash Buffer 21X		20-6746	37528		2033-08-28
Substrate TMB		20-2629	37312		2030-08-31
Stop Solution		20-2693	37299		2033-05-29

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

3. Quality control

Quality control has been performed for lot no 37390 according to standard operating procedures at Mercodia AB, and the product is released based on fulfillment of established acceptance criteria.

4. Calibration

Mercodia Glucagon ELISA is calibrated against WHO 1st international reference preparation 69/194.

5. Assay method

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

6. Intended use

Mercodia Glucagon ELISA provides a method for the quantitative determination of glucagon in EDTA-plasma, serum 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:	Performed by:	Signature:
2026-11-04	Mattias Jide	

Date of approval:	Approved by:	Signature:
2026-11-09	Jonas Kvick	

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