

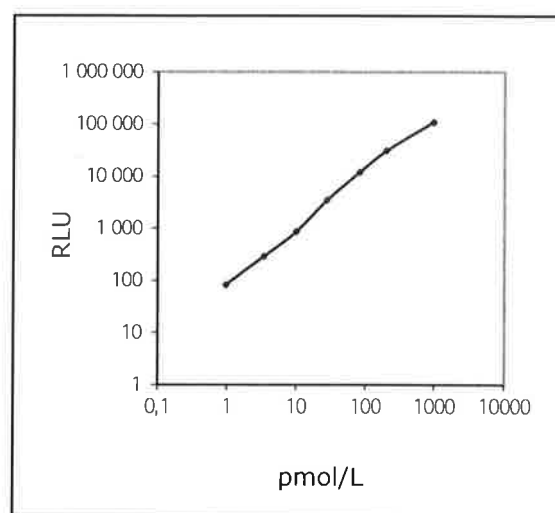
Certificate of Analysis

1. Manufacturer

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

2. Description

Catalog no: 10-1278-01
 Product: Mercodia Total GLP-1 NL-ELISA
 Lot no: 35652
 Expiry date: 2026-04-30



<i>Component</i>	<i>Art no</i>	<i>Lot no</i>	<i>RLU</i>	<i>Exp. date</i>
Calibrator 0	20-7259	35532	20	2026-08-16
Calibrator 0,953 pmol/L	20-7266	35525	82	2026-08-22
Calibrator 3,35 pmol/L	20-7267	35526	289	2026-08-22
Calibrator 9,97 pmol/L	20-7268	35527	858	2026-08-22
Calibrator 27,6 pmol/L	20-7269	35528	3472	2026-08-22
Calibrator 82,0 pmol/L	20-7270	35529	11922	2026-08-22
Calibrator 203 pmol/L	20-7271	35530	30954	2026-08-22
Calibrator 975 pmol/L	20-7299	35531	108204	2026-08-22
Coated Plate	20-7277	35534		2026-08-28
Enzyme Conjugate 11X	20-7275	35536		2026-08-23
Enzyme Conjugate Buffer	20-7273	35533		2026-08-18
Wash Buffer 21X	20-6746	35568		2030-09-05
Substrate Reagent A	20-7300	35463		2026-05-26
Substrate Reagent B	20-7301	35464		2026-05-26

3. Quality control

Quality control has been performed for lot no 35652 according to standard operating procedures at Mercodia AB, and the product is released based on fulfillment of established acceptance criteria.

4. Calibration

Mercodia Total GLP-1 NL-ELISA is calibrated against an in-house reference preparation of GLP-1 (9-36)amide.

5. Assay method

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

6. Intended use

Mercodia Total GLP-1 ELISA provides a method for the quantitative determination of Total GLP-1 in EDTA-plasma.

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 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:

2023-10-16

Performed by:

Fonäs Blück

Signature:



Date of approval:

2023-10-19

Approved by:

Mattias Lil

Signature:

