Mercodia

Mall-0514 v3.0

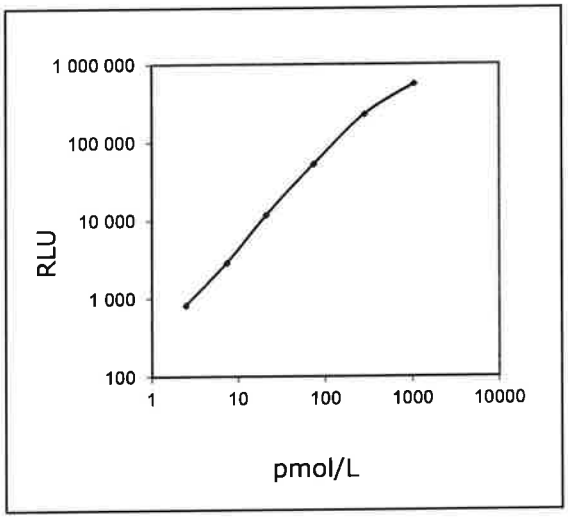
Certificate of Analysis

1. **Manufacturer**

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

2. **Description**

Catalog no:	10-1258-01
Product:	Mercodia Total GIP NL-ELISA
Lot no:	37681
Expiry date:	2028-12-31



Component	*Art no*	*Lot no*	*Exp. date*
Calibrator 0	20-7432	37081	2029-01-21
Calibrator 1, 2,47 pmol/L	20-7439	37074	2029-01-20
Calibrator 2, 7,37 pmol/L	20-7440	37075	2029-01-20
Calibrator 3, 20,9 pmol/L	20-7441	37076	2029-01-20
Calibrator 4, 73,7 pmol/L	20-7442	37077	2029-01-20
Calibrator 5, 281 pmol/L	20-7443	37078	2029-01-20
Calibrator 6, 1050 pmol/L	20-7444	37079	2029-01-20
Coated Plate	20-7454	37072	2029-01-26
Enzyme Conjugate 11X	20-7457	37082	2029-01-28
Enzyme Conjugate Buffer	20-7452	37080	2029-01-16
Wash Buffer 21X	20-6746	37311	2033-06-18
Substrate Reagent A	20-7300	37632	2029-09-15
Substrate Reagent B	20-7301	37633	2029-09-15

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

3. Quality control

Quality control has been performed for lot no 37681 according to standard operating procedures at Mercodia AB, and the product is released based on fulfillment of established acceptance criteria.

4. Calibration

Mercodia Total GIP NL-ELISA is calibrated against a highly purified, commercial GIP(3-42) preparation, validated with amino acid analysis and HPLC-UV LC-MS/MS.

5. Assay method

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

6. Intended use

Mercodia Total GIP NL-ELISA provides a method for the quantitative determination of total GIP in human samples.

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:	Performed by:	Signature:
2026-10-22	Mattias Jart	MJ
Date of approval:	**Approved by:**	**Signature:**
2026-10-22	Elin Wesberg	EW

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