Mercodia

Mall-0514 v3.0

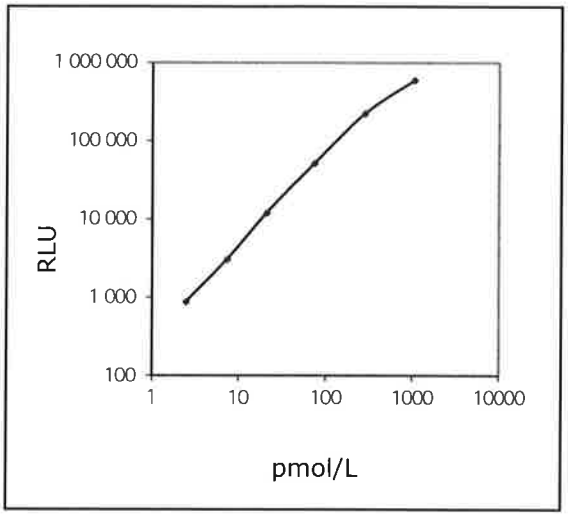
Certificate of Analysis

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: 37187
Expiry date: 2028-01-31



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	36958	2032-12-03
Substrate Reagent A	20-7300	36595	2028-02-14
Substrate Reagent B	20-7301	36596	2028-02-14

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3. **Quality control**

Quality control has been performed for lot no 37187 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:
2026-03-03	Eira Lindqvist	
Date of approval:	**Approved by:**	**Signature:**
2026-03-23	Mathias M	

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