



Report #:
IF-824-QSA-TR006-BTL261022-20

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Certificate of Analysis (CoA) - Finnrick

Date:

Date Tested:

Customer:

Testing material:

Lot Number:

Content/Potency:

Storage:

Visual Description:

Labeled as:

Manufacturer:

Testing Purpose:

8/3/2026

7/30/2026

Finnrick

Retatrutide

EVH7GW4

20 mg

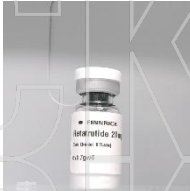
R.T.

Small clear vial: white sample, white label, silver crimp, white plastic cap.

Retatrutide

N/A

FTIR and HPLC analysis for the identification, purity, potency and composition of a peptide product. It does not provide information on particulate matter, microbial contamination or presence of endotoxins.



Test	Method	Specification	Result
General Appearance	USP <630>	white powder	white powder
Mass	USP <41>	As recorded	81.2 mg
FTIR Identification and Composition Analysis	USP <197A>	Sample spectrum should confirm the content of peptide via characteristic bands	FTIR sample spectrum confirms the presence of Retatrutide with addition of excipient(s)/fillers.
HPLC Purity of Peptide Assay	USP <621>	Specifications: ≥ 98 %	99.8 %
HPLC Potency Assay	USP <621>	Specifications: 90 – 110% of 20 mg	21.2 mg (106.1 %)
Peptide-to-Excipients Ratio	USP <1151>	Recommended ratios of (1:2) to (1:10) for (peptide: excipients)	21.2 : 60 mg (1:2.8)

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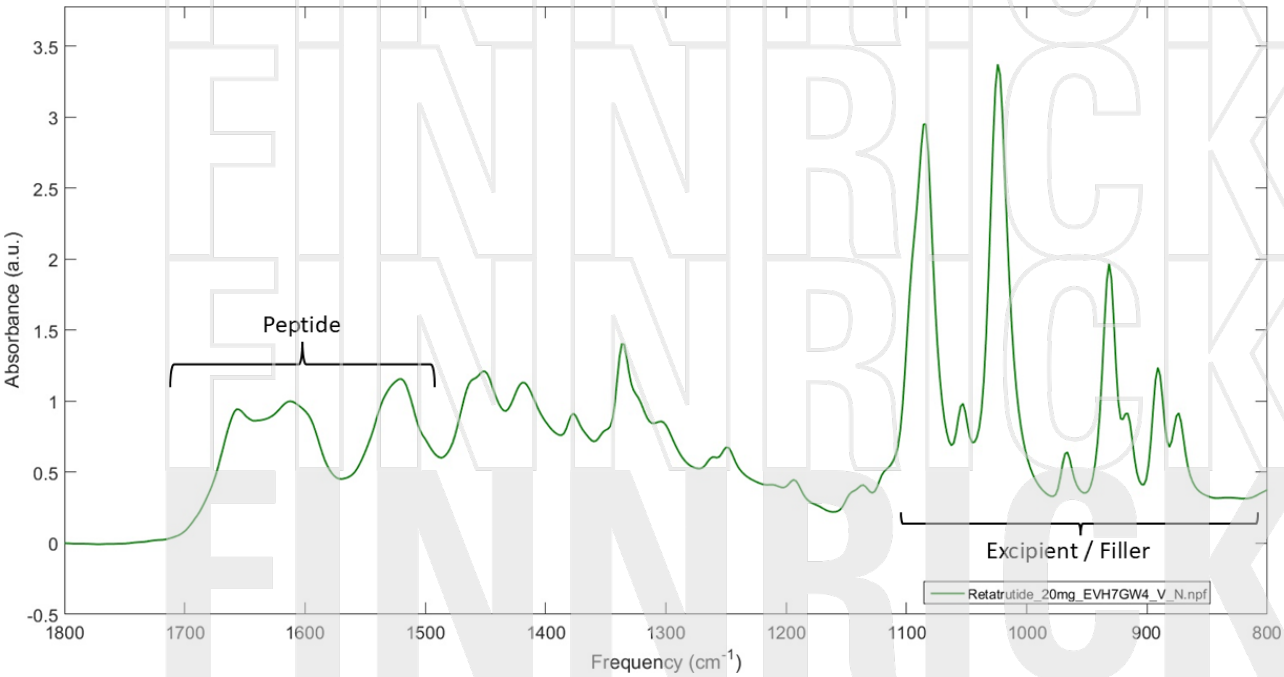


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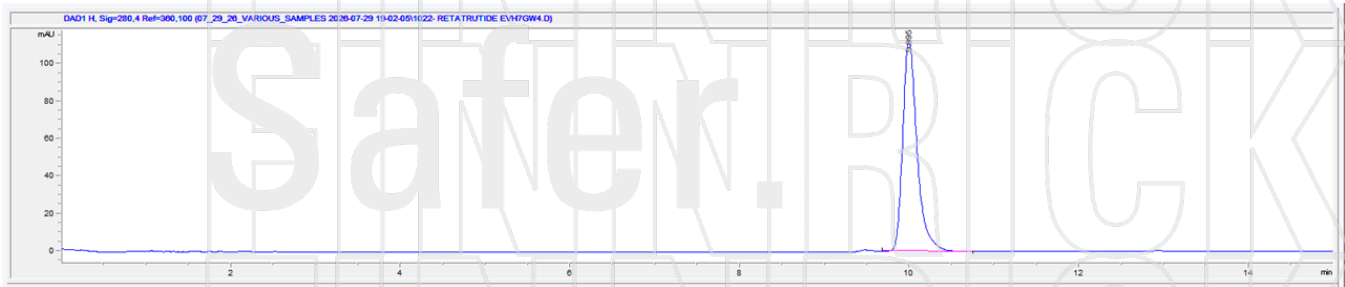
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FTIR ID and Composition Analysis: Retatrutide Lot EVH7GW4



HPLC Purity and Potency Assay @ 280 nm: Retatrutide Lot EVH7GW4



Retatrutide Lot EVH7GW4 @ 280 nm		
Peak #:	Retention Time (min)	Area (mAU*s)
1	9.995	1317.8