



SimecAG
Gartnereiweg 4 B CH-4665 Oftringen
Telefon +41 62 762 83 08 Fax +41 62 732 83 09

Test Report			Number	V 2183225																								
Object of analysis:	Cagrilintide 10mg																											
Customer:	Viking Pharma Lab		Sample:	1																								
Batch:	VPOGR171 Mfg: 11.2025, Exp: 11.2030		Receipt:	03.02.2026																								
Subject:	Content, Identification																											
Method(s):	AVPE		Anzats-datum	17.04.2026																								
<table border="1"><thead><tr><th>Samples / Analysis</th><th>Method</th><th>Status</th></tr></thead><tbody><tr><td>Cagrilintide:</td><td>AVPE</td><td>P</td></tr><tr><td>Identification</td><td>AVPE</td><td>P</td></tr><tr><td>Content</td><td></td><td></td></tr></tbody></table>			Samples / Analysis	Method	Status	Cagrilintide:	AVPE	P	Identification	AVPE	P	Content			<table border="1"><thead><tr><th>Results</th><th>Specifications</th><th>complies</th></tr></thead><tbody><tr><td>Complies</td><td>Complies</td><td>Yes</td></tr><tr><td>10.3 mg / tab</td><td>10 mg / tab</td><td>-</td></tr><tr><td colspan="3">Purity Exceeds 99.1 %</td></tr></tbody></table>		Results	Specifications	complies	Complies	Complies	Yes	10.3 mg / tab	10 mg / tab	-	Purity Exceeds 99.1 %		
Samples / Analysis	Method	Status																										
Cagrilintide:	AVPE	P																										
Identification	AVPE	P																										
Content																												
Results	Specifications	complies																										
Complies	Complies	Yes																										
10.3 mg / tab	10 mg / tab	-																										
Purity Exceeds 99.1 %																												
Method-status G= GMP A= accredited V= generally validated P= validated on product N= not validated E= external lab Remarks: The sample(s) mentioned above have been analysed as they have been sent to us by the client. We have not controlled external processes like manufacturing, labelling, sampling, shipping, storage. These results are for information only and do not compensate for correct quality control by the manufacturer/distributor. Generally, pharmaceutical products have to be produced and distributed under full GMP/GDP regime, including GMP-compliant analyses with validated methods. This report cannot be used for commercial reasons incl. product release and/or quality control. It is not allowed to use this analysis in the context of doping/sports/competitions or any other illegal action, neither by the athlete nor by the tutor/trainer or any other person. Signatures: controlled & released: <i>A. M.</i> Staff: <i>J. S. M.</i> completed: 18.05.2026 The enclosed results refer exclusively to the object of examination described above. The measuring accuracy is available on request. Only handwritten signatures are valid. Without written permission, it is not allowed to publish single parts of this report. page 1 of 1																												