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Test Report			Number	V 2183230																																							
Object of analysis:	Tesamorelin + Ipamorelin 14mg																																										
Customer:	Viking Pharma Lab		Sample:	1																																							
Batch:	VP0GR166 Mfg: 10.2025, Exp: 10.2030		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>Tesamorelin:</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><tr><td>Ipamorelin :</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	Tesamorelin:	AVPE	P	Identification	AVPE	P	Content			Ipamorelin :	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>10.2 mg / tab</td><td>10 mg / tab</td><td>-</td></tr><tr><td>Complies</td><td>Complies</td><td>Yes</td></tr><tr><td>4.4 mg / tab</td><td>4 mg / tab</td><td>-</td></tr><tr><td colspan="3">Purity Exceeds 99.7 %</td></tr></tbody></table>		Results	Specifications	complies	Complies	Complies	Yes	10.2 mg / tab	10 mg / tab	-	Complies	Complies	Yes	4.4 mg / tab	4 mg / tab	-	Purity Exceeds 99.7 %		
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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:		Staff:																																								
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.			completed:	18.05.2026																																							