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Test Report					Number	V 18	352400	
Object of analysis:	Triple Stack Tab 55mg							
Customer:	Viking Pharma Lab Sample:				1			
Batch:	VD0CDC40 Mfa: 02 2022 Even 09 2029				t: 03.02.2023			
Subject:	Content, Identification							
Method(s):	HPLC			Anzats- datum	12.04.2023			
Samples / Ana	ysis Method S	Status	Results	:	Specifictions	co	omplies	
Oxandrolon Identification Content	e: HPLC HPLC	N N	Complies 26.1 mg / tab		Complies 25 mg / tab		Yes -	
Stanozolol: Identification Content	HPLC HPLC	N N	complies 15.3 mg / tab		complies 15 mg / tab		Yes -	
Methenolon Identification Content	e: HPLC HPLC	N N	complies 15.5 mg / tab		complies 15 mg / tab		Yes -	
ТАМС ТҮМС	SOP M 006 SOP M 006	N N	< 1 KBE / tab < 1 KBE / tab		- -		-	
Method-status G= GMP A= accreditated V= generally validated P= validated on product N= not validated E= external lab								
Remarks:	not controlled externa! processes like manufacturing, labelling, sampling, shlpplng, 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 ralease 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	1: AL	Staff: 1.5	public K	c	omplet	ed:	
The enclosed results retar exclusively to the object of examination described above. The measuring accuracy is available on request. Only handwritten signaturas are					13	13.05.2023		
valid. Without written permission, It is not allowed to publish single parts of this page 1 of 1 report.								
Content Stanozolol: Identification Content Methenolon Identification Content TAMC TYMC Method-status Remarks: Signatures: The enclosed refine measuring valid. Without	HPLC HPLC HPLC HPLC HPLC HPLC SOP M 006 SOP M 006 SOP M 006 SOP M 006 SOP M 006 SOP M 006 The sample(s) mentioned a not controlled external pro- are for information on manufacturer/distributor. G full GMP/GDP regime, inclu used for commercial reaso analysis in the context of do by the tutor/trainer or any c controlled & released esults retar exclusively to the c	N N N N N N N N N /- generally val /- generally val bove have bee cesses like mar ily and do ienerally, pharm iding GMP-com ins incl. produc ping/sports/cco other person.	26.1 mg / tab complies 15.3 mg / tab complies 15.5 mg / tab < 1 KBE / tab < 1 KBE / tab < 1 KBE / tab lidated P= validate analysed as they bufacturing, labelli not compensat naceutical product pliant analyses w ct ralease and/or ompetitions or any Staff:	y have been ng, sampli e for constraints have to lead the validate of the val	25 mg / tab complies 15 mg / tab complies 15 mg / tab - - - duct N= not val en sent to us by ng, shlpplng, s porrect quality be produced ar ed methods. The ntrol. It is not yal actlon, neith	y the clien torage. The contro nd distrib his report allowed her by the complet 3.05.20	- Yes - Yes - - - - - - - - - - - - - - - - - - -	

