

## Simec AG

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Test report					Number C	4280	8
Object of analysis:	Geneza GP Ha	alotest					
Customer:	200				Samples:	1	
Batch:	GP732T, Exp.	02/2019			Receipt:	07.04.2	2016
Subject:	Fluoxymestero	ne: Assay, I	dentific	ation			
Method(s):	HPLC					07.04.2	2016
Sample / Analysis:		Method	Status	Result	Specification		complies
Fluoxymester Identification Content	one:	HPLC HPLC	N N	complies 9.8 mg/tablet	complies 10 mg/tabl	et	yes -

Method-status:

G = GMP A = accreditated 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.

Signatures:

controlled & released:

Staff:

completed: 13.04.2016

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.

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