

Certificate of Analysis

Date 18/12/2017

Article
SKU 100934589 ClearKlens oxifast 1L sterile

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batch : 0001340140
expiry date : 2018/11
date of manufacture : 06/11/2017

Characteristic Method	Unit	Value	lower limit	upper limit	conformity
peracetic acid MAN LCQ 011	%	0,09	0,07	0,10	Yes
hydrogen peroxide MAN LCQ 011	%	3,12	2,90	3,40	Yes
loss TAO 16h at 96°C MAN LCQ 009	%	0,38	0,00	5,00	Yes
Sterility certificate European Pharmacopeia 2.6.1					Yes

released on : 18/12/2017

by : TRIBONDEAU-JAOUI

Comments:
Sterility certificate n°171201-0063-001




The analysis results above could change over time and in function of the storage temperature. It is imperative to store the products according to the recommended conditions indicated in the MSDS.

End of certificate of analysis

Labor L+S AG Mangelstfeld 4, 5, 6 | 97708 Bad Bocklet-Großenbrach | Germany

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validé le 18/12/2017


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Akkreditiert nach ISO / IEC 17025

Durch die DAKKS Deutsche Akkreditierungsstelle GmbH nach DIN EN ISO/IEC 17025 akkreditiertes Prüflaboratorium. Die Akkreditierung gilt für die in der Urkunde aufgeführten Verfahren.



Bad Bocklet 18 Dec 2017 / AP / BIOXCh

Certificate of Analysis

L+S No:	171201-0063-001	L+S Code:	1005537 / S
Product name:	CLEARLENs OXIFAST VH49		
Lot No:	SEP/0001340140/17/307		
Entry temperature:	room temperature		
Order dated:	27 Nov 2017	Sample receipt:	01 Dec 2017
Start of test:	01 Dec 2017	End of test:	18 Dec 2017

Test for Sterility

Parameter	Specification / Demands	Result
Test procedure according to:	Ph. Eur. 2.6.1 (Produktspez. Eignungsprüfung/Methodentransfer liegt für dieses Volumen nicht vor)	
Reference document:	Growth promotion test (140225-0139-001)	
Test method:	Membrane filtration	
Test environment:	Clean room	
Number of samples tested:	8	8
Division of samples:	Prüfung auf 2 Testsysteme aufteilen	
Tested volume per sample:	200 ml	200 ml
Type of filter material:	PVDF (Durapore®)	PVDF (Durapore®)
Test system:	TZHVAB210	TZHVAB210
Nutrient media:	2 x TSB- & 2 x FTM with neutralization agents (2 x C+/2 x T+)	2 x C+/2 x T+
Start of incubation:		04 Dec 2017
End of incubation:		18 Dec 2017
Period of incubation:	At least 14 days	14 days
Evaluation:	No macroscopic growth visible	complies

The test was conducted in compliance with GMP guidelines. There were no test-related deviations.

This document was created by a GMP-supervised LIMS and approved by electronic signature.

Approved on 18 Dec 2017 at 08:29 by Katharina Schlereth, Division Manager.

CERTIFICATE OF STERILITY

BIOXAL SA

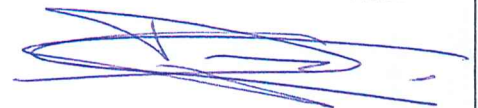
Route des Varennes
BP 30072
F 71103 CHALON / SAONE
FRANCE,

Certifies that the product:

CLEARKLENS OXIFAST STERILE
Item code: 70001369
Batch number: 0001340140
Quantity: 1136 bottles,

is sterile, according to sterility test Ph.Eur. 2.6.1.

Samuel LALEVÉE
Quality Manager



PO : Diane-Claire TRIBONDEAU-JAOUI
Regulatory Affairs Manager