

Form SP-PR-LIB-F-011 Status : APPLICABLE

Certificate of Analysis DIVERSEY for ClearKlens Bi-Spore RTU VH26S 4x5L

Date of application: 13/07/2023
Index: 01

Diversey	earKlens Bi-Spore RTU VH26S 4x5L SKU 100852440
Batch Number	SPH25072023
Production date	08/2023
Expiration Date	06/2025

This is to certify that the above batch of product has been tested and conforms to the manufacturing Quality Control specification for the product.

	ClearKlens Bi-Sp	ore Base Solution	RTU	
Test	Test Method	Specification	Unit	Results
Appearance	PS-CON-I-029	Clear colorless liquid	N/A	clear colorless liquid
Relative density (20°C)	PS-CON-I-027	1.000 - 1.010	-	1.001
pH (neat Solution)	PS-CON-I-040	2.5 - 5.0	-	2.7
	ClearKlens Bi-S	ore Activator Solu	tion	
Test	Test Method	Specification	Unit	Results
Appearance	PS-CON-I-029	Clear colorless liquid	N/A	clear colorless liquid
Relative density (20°C)	PS-CON-I-027	1.004 - 1.020	-	1.008
pH (neat Solution 20°C)	PS-CON-I-040	9.0 - 12.5	-	11.4
Potential chlorine dioxide content	PS-CON-I-049	100 - 180	ppm	151

ClearKlens Bi-Spore Activator Solution + ClearKlens Base Solution RTU				
Test	Test Method	Specification	Unit	Results
Appearance	PS-CON-I-029	Clear yellow liquid	N/A	Clear yellow liquid
Relative density (20°C)	PS-CON-I-027	1.000 - 1.010	-	1.001
pH (neat Solution 20°C)	PS-CON-I-040	2.5 - 5.0	-	3
Potential chlorine dioxide content	PS-CON-I-049	100 - 180	ppm	122

	Water	routine test		
Microbiological Quality of WFI Water	Water PPI VRAC European pharmacopoeia	<10 cfu/100ml	CFU / 100mL	<1



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Control of the Sterility of the product:

Test	Test Method	Specification	Conformity	Reference number
Sterilization	Gamma Irradiation	25.0 - 45.0 (kGy)	С	FR02S12748381 -1-1 + 23T02260D + 2013-1188A
Sterility certificate	Membrane filtration sterility testing EP 11.0 2.6.1	No Growth	С	23/CBH/STE/212

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

We certify that, except for the exceptions or deviations listed above, the quoted supply has been manufactured and tested in accordance with the requirements of the specifications in force

SIMA-PHARMA Control (Responsible for the production process of the final product) :	Stamp and signature
Name: E. JACQUIN Position: General Hauager	
Produced on behalf of Diversey at: Sima Pharma: Z.I. de Rousset / Peynier, 54 Av. de la Plaine, 13790 Rousset, France	
Date: 11109123	

End of certificate of analysis

For further information, please contact Diversey at pharma@diversey.com