

CERTIFICATE OF RELEASE

A) BASIC DATA

PRODUCT:

BISOLVON INJ. 100 ML

COUNTRY:

AUSTRIA

PRODUCT CODE LABIANA:

50884

MANUFACTURING SPECIFICATION Nº-

C 0390-01-06

TESTING SPECIFICATION Nº:

101 8502-F87R-01

LABIANA BATCH Nº:

F13407F-13

SPECIFIC BATCH No:

F13407F-13

ANALYSIS No:

49532

MANUFACTURING DATE:

04-11-2013

EXPIRY DATE:

11-2018

UNITS RELEASED:

1741

COMMENTS:

B) RAW MATERIALS

We herewith certify that all raw materials have been tested and are in compliance with the requirements of the testing specifications

C) MANUFACTURING / PACKAGING

The manufacturing and packaging documentation for the above mentioned product have been reviewed and it was determined that this lot number was manufactured and packaged according to the requirements of the manufacturing specifications and master production documents.

D) QUALITY ASSURANCE

All documentation has been found to be in compliance with the requirements of the above mentioned manufacturing specifications and master production documents. Explanations for deviations, if any are attached. This lot has been tested to the requirements of the above mentioned testing specification and the certificate of analysis is attached. All batch documentation is available upon request.

E) QUALIFIED PERSON RELEASE

In-process Control (QA)

The pharmaceutical product has been tested and complies with the requirements of the testing specifications and the relevant marketing authorisation and European cGMP requirements.

The pharmaceutical product is released by a Qualified Person (75/319/EEC or 81/851/EEC).

Reviewed by:

Released by:

Qualified Person (QP

Date: 14 01 14

141 07 174



ANALYSIS CERTIFICATE

Producto Acabado

Producto/Product: BISOLVON			Código/Code : 45580 E	
Lote/Batch: F13407	Caducidad/Expir	y: (*)	Cantidad/Amount: 2000L	
Fecha Fabricación/Manufac.Date:04.11.13		Formato: 100ml		Reporte/Report: 649734
Recibido/Received : 06.11.13 N° Análisis/Analysis : 49532		Fecha Análisis/Q Analysis Date/Q: Fecha Análisis/M Analysis Date/M:		11.11.13 22.11.13
Prescripción análisis : 101 8502 561C-03 (10.03.2004)		2004)	Tipo Lote/Use of batch: Producción/Production	

ENSAYO/TEST	ESPECIFICICACION/SPECIFICATION	RESULTADOS/RESULTS
1. DESCRIPTION . 1.1 LIQUID . COLOUR	Clear and colourless solution, practically free from particles, filled into injection vials Almost imperceptible.	Complies Complies
2. PHYSICO-CHEMICAL DETERMINATIONS 2.2 PH 2.4 RELATIVE DENSITY 2.9 IDENTIFICATION	3.0 - 4.0 0.9945 - 1.0045 (20°C)	3.0 1.0007
2.9.1 IDENTIFICATION BY TLC	N-A 274 CL: Corresponding to standard Methylparahydroxybenzoate:Corresponding to standard Propylparahydroxybenzoate:Corresponding to standard	Complies Complies Complies
. 2.9.2 IDENTIFICATION BY HPLC	N-A 2/4 CL: Corresponding to standard Methylparahydroxybenzoate:Corresponding to standard Propylparahydroxybenzoate:Corresponding to standard	Complies Complies
2.10 CLARITY OF SOLUTION 2.11 COLOUR OF SOLUTION 2.19 EXTRACTABLE VOLUME	≤ I ≤ B9 Format 50 ml : 50.0 - 55.0 ml Format 100 ml : 100.0 - 110.0 ml Format 250 ml : 250.0 - 260.0 ml	< 1 < 89 104.0 - 106.0 ml
2.42 SUB-VISIBLE PARTICLES 3. ASSAY	Compiles with Ph. Eur. /Ed.	Complies
3.5 HPLC 4 MICROBIOLOGICAL DETERMINATIONS 4.1 STERILITY TEST 6. IMPURITIES	N-A 274 CL : 285.0 - 315.0 mg/100.0 ml Complies with Ph. Eur. 7Ed.	300.0 mg/100.0 ml Complies
. 6.16 ACTIVE INGREDIENT DECOMPOSITION	\$ 1.0% N-A 274 CL equivalent to \$ 0.7% N-AB 773 XX \$ 0.3% N-A 274 CL equivalent to \$ 0.3% N-A 1740 CL Each individual unknown impurities:	0.07 % 0.04 %
7. PRESERVATIVES	≤ 0.3% calculated as % decomposed N-A 274 CL Total impurities: ≤ 1.3%	0.02 x 0.13 x
7.5 METHYLPARABEN /PROPYLPARABEN	Methylparahydroxybenzoate: 66.5 - 73.5 mg/100.0 ml Propylparahydroxybenzoate: 28.5-31.5 mg/100.0 ml (*)Specific for climatic zones. *End of report*	70.1 mg/100.0 ml 30.0 mg/100.0 ml

Firma/Signature

Control Calidad Quality Control

Diference Manager

Approbado Rechazado Approvind-Rejected

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