



*Agenzia Italiana del Farmaco*

**AIFA**



Certificate No: IT/190-1/H/2014

**CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER**

**Part 1**

**Issued following an inspection in accordance with Art. 111(5) of Directive 2001/83/EC**

The competent authority of Italy confirms the following:

The manufacturer S.A.L.F. SPA LABORATORIO FARMACOLOGICO  
Site address VIA G. MAZZINI, 9 - 24069 CENATE SOTTO (BG)

Has been inspected under the national inspection programme in connection with manufacturing authorisation no. aM - 108/2014 dated 07/11/2014 in accordance with Art. 40 of Directive 2001/83/EC/ transposed in the following national legislation: D.Lvo 219/2006 art. 50.

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on 03/21/2014 it is considered that it complies with the Good Manufacturing Practice requirements referred to in The principles and guidelines of Good Manufacturing Practice laid down in Directive 2003/94/EC.

This certificate reflects the status of the manufacturing site at the time of the inspection noted above and should not be relied upon to reflect the compliance status if more than three years have elapsed since the date of that inspection, after which time the issuing authority should be consulted.

The authenticity of this certificate may be verified with the issuing authority.

AIFA Italian Medicines Agency  
Manufacturing Authorization Office  
Via del Tritone, n° 181 - 00187 ROMA (ITALY)  
Tel.+390659784489 Fax +390659784312  
website: [www.agenziafarmaco.it](http://www.agenziafarmaco.it)  
SIS : 141

PC  
GMP



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**Part 2**

Name and address of the site: S.A.L.F. SPA LABORATORIO FARMACOLOGICO - VIA G. MAZZINI, 9 , 24069 CENATE SOTTO(BG)

Human Medicinal Products

**Authorised Operations**

Manufacturing Operations (Part 1)

**PART 1 - MANUFACTURING OPERATIONS**

<b>1.1</b>	<b>Sterile Products</b>	
	1.1.2	<i>Terminally sterilised</i>
	1.1.2.1	Large volume liquids
	1.1.2.3	Small volume liquids

Rome, 07/24/2014

Name and signature of the authorised person of the Competent Authority of Republic of Italy

Dott. Renato Massimi  
AIFA – Manufacturing Authorization Unit



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