



Agenzia Italiana del Farmaco

Certificate No: IT/1-1/H/2011

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 CHELAB S.R.L.
Site address VIA FRATTA,25 - 31023 RESANA (TV)

Has been inspected under the national inspection programme in connection with manufacturing authorisation no. aM - 153/2010 dated 10/19/2010 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 01/29/2010 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
SIS : 3116

Mariacristina Orlandi
GMP



Agenzia Italiana del Farmaco

Part 2

Name and address of the site: CHELAB S.R.L. - VIA FRATTA,25 , 31023 RESANA(TV)

Human Medicinal Products

Authorised Operations

Manufacturing Operations (Part 1)

PART 1 - MANUFACTURING OPERATIONS

1.6	Quality control testing
	1.6.1 <i>Microbiological: sterility</i>
	1.6.2 <i>Microbiological: non-sterility</i>
	1.6.3 <i>Chemical/Physical</i>
	1.6.4 <i>Biological</i>

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Name and address of the site: CHELAB S.R.L. - VIA FRATTA,25 , 31023 RESANA(TV)

Human Medicinal Products

Authorised Operations

Manufacturing Operations (Part 1)

PART 1 - MANUFACTURING OPERATIONS OF INVESTIGATIONAL MEDICINAL PRODUCTS

1.6	Quality control testing
	1.6.1 <i>Microbiological: sterility</i>
	1.6.2 <i>Microbiological: non-sterility</i>
	1.6.3 <i>Chemical/Physical</i>
	1.6.4 <i>Biological</i>

Rome, 01/03/2011

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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