



Certificate No: IT/6/H/2024

## 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 LABANALYSIS LIFE SCIENCE S.R.L.

Site address VIA EUROPA, 5 - 27041 CASANOVA LONATI (PV)

Has been inspected under the national inspection programme in connection with manufacturing authorisation no. aAMM - 10/2024 dated 01/04/2024 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 11/15/2019, 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  
GMP Inspections and Manufacturing Authorizations of Medicinal Products Office  
Via del Tritone, n° 181 - 00187 ROMA (ITALY)  
Tel.+390659784357 Fax +390659784312  
website: [www.agenziafarmaco.it](http://www.agenziafarmaco.it)  
SIS : 3231

RS  
GMP

**Part 2**

Name and address of the site: LABANALYSIS LIFE SCIENCE S.R.L.  
VIA EUROPA, 5  
27041 CASANOVA LONATI (PV)

Human Medicinal Products

**Authorised Operations**

Manufacturing Operations (Part 1)

Importation of medicinal products (Part 2)

**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>

**Any restrictions or clarifying remarks related to the scope of these Manufacturing operations:**

1.6.4 Biological: LAL test.

**PART 2 - IMPORTATION OF MEDICAL PRODUCTS**

2.1	Quality control testing of imported medical products
	2.1.1 <i>Microbiological: sterility</i>
	2.1.2 <i>Microbiological: non-sterility</i>
	2.1.3 <i>Chemical/Physical</i>
	2.1.4 <i>Biological</i>

**Any restrictions or clarifying remarks related to the scope of these Importing operations:**

2.1.4 Biological: LAL test.

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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>

**Any restrictions or clarifying remarks related to the scope of these Manufacturing operations:**

1.6.4 Biological: LAL test.



Rome, 01/04/2024

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

Angela Del Vecchio  
GMP Inspections and Manufacturing  
Authorizations of Medicinal Products Office

*STAMP DUTY PAID ACCORDING TO THE CURRENT ITALIAN LAW*

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