Manufacturer/Importer Authorisation

1. Authorisation Number 2767-F

2. Name of authorisation holder Eurofins Proxy Laboratories B.V. (ORG-100015745)

LOC-100024482)

3. Address(es) of manufacturing site(s) Eurofins PROXY Laboratories B.V. (ORG-100011886 /

LOC-100021255), Darwinweg 24, Leiden, 2333 CR, Netherlands

Eurofins Proxy Laboratories B.V. (ORG-100015745 / LOC-100024482), Archimedesweg 25, Leiden, 2333 CM,

Netherlands

4. Legally registered address of authorisation

holder

Archimedesweg 25, Leiden, 2333 CM, Netherlands

4.a Additional details on units inspected of

legally registered address

5. Scope of authorisation and dosage forms²

ANNEX 1 and/ or ANNEX 2

6. Legal Basis of authorisation

Art. 88 of Regulation (EU) 2019/6

7. Name of responsible officer of the competent authority of the member state granting the

manufacturing authorisation

confidential

8. Signature

9. Date

2025-07-28

10. Annexes attached

Annex 1 and/or Annex 2

Optional Annexes as required:

Annex 3(Addresses of Contract Manufacturing Site(s))

Annex 4(Addresses of Contract laboratories)

Annex 5(Name of Qualified Person)

Annex 6(Name of responsible persons)

Annex 7(Date of inspection on which authorisation granted, scope of last

inspection)

Annex 8(Manufactured/imported products authorised)³

 $^{^{1}}$ The authorisation referred to in paragraph 40(1) of Directive 2001/83/EC as amended and Article 88(1) of Regulation (EU) 2019/6, shall also be required for imports coming from third countries into a Member State.

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Manufacturer/Importer Authorisation: Page 1 of 3

SCOPE OF AUTHORISATION

ANNEX 1

Name and address of the site: Eurofins PROXY Laboratories B.V., Darwinweg 24, Leiden,

2333 CR, Netherlands

Additional Details:

Veterinary Medicinal Products

Authorised Operations

MANUFACTURING OPERATIONS(according to part 1)

IMPORTATION OF MEDICINAL PRODUCTS (according to part 2)

Part 1 - MANUFACTURING OPERATIONS		
1.6	Quality control testing	
	1.6.1 Microbiological: sterility	
	1.6.2 Microbiological: non-sterility	
	1.6.3 Chemical/Physical	
	1.6.4 Biological	
Part 2 - IMPORTATION OF MEDICINAL PRODUCTS		
2.1	Quality control testing of imported medicinal products	
	2.1.1 Microbiological: sterility	
	2.1.2 Microbiological: non-sterility	
	2.1.3 Chemical/Physical	
	2.1.4 Biological	

²Guidance on the interpretation of this template can be found in the Interpretation of the Union format for Manufacturer/Importer Authorisation.

³The Competent Authority is responsible for the appropriate linking of the authorisation with the manufacturer's application (Article 42(3) of Directive 2001/83/EC as amended and Article 90(3) of Regulation (EU) 2019/6).

SCOPE OF AUTHORISATION

ANNEX 1

Name and address of the site: Eurofins Proxy Laboratories B.V., Archimedesweg 25, Leiden,

2333 CM, Netherlands

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IMPORTATION OF MEDICINAL PRODUCTS(according to part 2)

Part 1 - MANUFACTURING OPERATIONS	
1.6	Quality control testing
	1.6.3 Chemical/Physical
	1.6.4 Biological
Part 2	- IMPORTATION OF MEDICINAL PRODUCTS
Part 2 2.1	- IMPORTATION OF MEDICINAL PRODUCTS Quality control testing of imported medicinal products