

# MANUFACTURER'S AUTHORISATION<sup>1, 2</sup>

1. Authorisation Number F 20/141
2. Name of authorisation holder PATHOQUEST
3. Address(es) of manufacturing site(s) PATHOQUEST, 11 rue Watt, Biopark bâtiment B, PARIS, 75013, France
4. Legally registered address of authorisation holder 11 rue Watt, Biopark bâtiment B, PARIS, 75013, France
5. Scope of authorisation and dosage forms<sup>2</sup> ANNEX 1 and/ or ANNEX 2
6. Legal Basis of authorisation Art. 40 of Directive 2001/83/EC  
Art. 13 of Directive 2001/20/EC
7. Name of responsible officer of the competent authority of the member state granting the manufacturing authorisation confidential
8. Signature
9. Date 2020-09-03
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)<sup>3</sup>

<sup>1</sup> The authorisation referred to in paragraph 40(1) of Directive 2001/83/EC and 44(1) of Directive 2001/82/EC, as amended, shall also be required for imports coming from third countries into a Member State.

<sup>2</sup> Guidance on the interpretation of this template can be found in the Help menu of EudraGMDP database.

<sup>3</sup> The Competent Authority is responsible for appropriate linking of the authorisation with the manufacturer's application (Art. 42(3) of Directive 2001/83/EC and Art. 46(3) of Directive 2001/82/EC as amended).

**SCOPE OF AUTHORISATION****ANNEX 1**

Name and address of the site : PATHOQUEST, 11 rue Watt, Biopark bâtiment B, PARIS,  
75013, France

Human Medicinal Products
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<b>Authorised Operations</b> MANUFACTURING OPERATIONS (according to part 1)
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<b>Part 1 - MANUFACTURING OPERATIONS</b>	
<b>1.6</b>	<b>Quality control testing</b>
	<i>1.6.3 Chemical/Physical</i>
	<i>1.6.4 Biological</i>

**Any restrictions or clarifying remarks related to the scope of these Manufacturing operations (for Public users)**

Manufacturer (Article R.5124-2 1° of the French Public Health Code) --- Signatory: Mrs Mélanie Cachet, deputy director - Inspection division. The ANSM does not issue hard copy of this authorisation.

**SCOPE OF AUTHORISATION****ANNEX 2**

Name and address of the site : PATHOQUEST, 11 rue Watt, Biopark bâtiment B, PARIS,  
75013, France

Human Investigational Medicinal Products
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**Authorised Operations**

MANUFACTURING OPERATIONS (according to part 1)

**Part 1 - MANUFACTURING OPERATIONS**

<b>1.6</b>	<b>Quality control testing</b>
	1.6.3 Chemical/Physical
	1.6.4 Biological

**Any restrictions or clarifying remarks related to the scope of these Manufacturing operations (for Public users)**

Manufacturer (Article R.5124-2 1° of the French Public Health Code) --- Signatory: Mrs Mélanie Cachet, deputy director - Inspection division. The ANSM does not issue hard copy of this authorisation.