Manufacturer/Importer Authorisation

1. Authorisation Number 0000012546/23/1

Qualimetrix S.A. (ORG-100021115 / LOC-100029818) 2. Name of authorisation holder

3. Address(es) of manufacturing site(s) Qualimetrix S.A. (ORG-100021115 / LOC-100029818), Mesogeion

Avenue 579, Agia Paraskevi, Athens, 153 43, Greece, GPS: 38.01221,

23.815573

4. Legally registered address of authorisation

holder

Mesogeion Avenue 579, Agia Paraskevi, Athens, 153 43, Greece,

GPS: 38.01221, 23.815573

5. Scope of authorisation and dosage forms²

ANNEX 1 and/ or ANNEX 2

Art. 40 of Directive 2001/83/EC 6. Legal Basis of authorisation

> Art. 13 of Directive 2001/20/EC 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 2023-12-14

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

¹The 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.

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

 $^{^3}$ The 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: Qualimetrix S.A., Mesogeion Avenue 579, Agia Paraskevi,

Athens, 153 43, Greece, GPS: 38.01221, 23.815573

Additional Details:

Human Medicinal Products Veterinary Medicinal Products

Authorised Operations

MANUFACTURING OPERATIONS(according to part 1)
IMPORTATION OF MEDICINAL PRODUCTS(according to part 2)

Part 1 - MANUFACTURING OPERATIONS		
1.1	Sterile products	
	1.1.3 Batch certification	
1.2	Non-sterile products	
	1.2.2 Batch certification	
1.3	Biological medicinal products (list of product types)	
	1.3.2 Batch Certification (list of product types)	
	1.3.2.6 Human or animal extracted products	
1.6	Quality control testing	
	1.6.2 Microbiological: non-sterility	
	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)

Part 2 - IMPORTATION OF MEDICINAL PRODUCTS		
2.1	Quality control testing of imported medicinal products	
	2.1.2 Microbiological: non-sterility	
	2.1.3 Chemical/Physical	
	2.1.4 Biological	
2.2	Batch certification of imported medicinal products	
	2.2.1 Sterile products	
	2.2.1.1 Aseptically prepared	

2.2.1.2 Terminally sterilised
2.2.2 Non-sterile products
2.2.3 Biological medicinal products 2.2.3.6 Human or animal extracted products

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

SCOPE OF AUTHORISATION

ANNEX 2

Name and address of the site: Qualimetrix S.A., Mesogeion Avenue 579, Agia Paraskevi,

Athens, 153 43, Greece, GPS: 38.01221, 23.815573

Human Investigational Medicinal Products

Authorised Operations

MANUFACTURING OPERATIONS (according to part 1)
IMPORTATION OF MEDICINAL PRODUCTS (according to part 2)

Part 1 - MANUFACTURING OPERATIONS		
1.1	Sterile products	
	1.1.3 Batch certification	
1.2	Non-sterile products	
	1.2.2 Batch certification	
1.3	Biological medicinal products (list of product types)	
	1.3.2 Batch Certification (list of product types)	
	1.3.2.6 Human or animal extracted products	
1.6	Quality control testing	
	1.6.2 Microbiological: non-sterility	
	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)

Part 2 - IMPORTATION OF MEDICINAL PRODUCTS		
2.1	Quality control testing of imported medicinal products	
	2.1.2 Microbiological: non-sterility	
	2.1.3 Chemical/Physical	
	2.1.4 Biological	
2.2	Batch certification of imported medicinal products	
	2.2.1 Sterile products	
	2.2.1.1 Aseptically prepared	
	2.2.1.2 Terminally sterilised	

2.2.2	Non-sterile products
2.2.3	Biological medicinal products
	2.2.3.6 Human or animal extracted products

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