

MANUFACTURER'S AUTHORISATION^{1, 2}

1. Authorisation Number 001525/06.08.00.04/2016
2. Name of authorisation holder Oy Medfiles Ltd
3. Address(es) of manufacturing site(s) Oy Medfiles Ltd, Volttikatu 5, Volttikatu 8, Kuopio, FI-70700, Finland
Oy Medfiles Ltd, Neulaniementie 2, Kuopio, 70210, Finland
4. Legally registered address of authorisation holder PL 1450, Kuopio, FI-70701, Finland
5. Scope of authorisation and dosage forms² ANNEX 1 and/ or ANNEX 2
6. Legal Basis of authorisation Art. 40 of Directive 2001/83/EC
Art. 44 of Directive 2001/82/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 2016-07-18
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 and 44(1) of Directive 2001/82/EC, as amended, shall also be required for imports coming from third countries into a Member State.

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

³ 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 : Oy Medfiles Ltd, Volttikatu 5, Volttikatu 8, Kuopio, FI-70700, Finland

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|-------------------------------|
| 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**1.2 Non-sterile products***1.2.2 Batch certification***1.6 Quality control testing***1.6.3 Chemical/Physical***Part 2 - IMPORTATION OF MEDICINAL PRODUCTS****2.1 Quality control testing of imported medicinal products***2.1.3 Chemical/Physical*

SCOPE OF AUTHORISATION**ANNEX 2**

Name and address of the site : Oy Medfiles Ltd, Volttikatu 5, Volttikatu 8, Kuopio, FI-70700, Finland

Human Investigational Medicinal Products

Authorised Operations

MANUFACTURING OPERATIONS (according to part 1)

Part 1 - MANUFACTURING OPERATIONS

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|------------|--|
| 1.2 | Non-sterile products |
| | <i>1.2.1 Non-sterile products (processing operations for the following dosage forms)</i> 1.2.1.5 Liquids for external use 1.2.1.6 Liquids for internal use 1.2.1.8 Other solid dosage forms 1.2.1.15 Other |
| | <i>1.2.2 Batch certification</i> |
| 1.5 | Packaging |
| | <i>1.5.2 Secondary packing</i> |
| 1.6 | Quality control testing |
| | 1.6.3 Chemical/Physical |

SCOPE OF AUTHORISATION**ANNEX 1**

Name and address of the site : Oy Medfiles Ltd, Neulaniementie 2, Kuopio, 70210, Finland

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| Human Medicinal Products Veterinary Medicinal Products |
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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**1.6.4 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**2.1.4 Biological*

SCOPE OF AUTHORISATION**ANNEX 2**

Name and address of the site : Oy Medfiles Ltd, Neulaniementie 2, Kuopio, 70210, Finland

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| Human Investigational Medicinal Products |
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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
- 1.6.4 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
- 2.1.4 Biological