

# Aftovaxpur DOE (12) O1 Manisa + A Turkey 14/98

Not  
authorised

- Foot-and-mouth disease virus, serotype O, strain O1 Manisa, Inactivated
- Foot-and-mouth disease virus, serotype A, strain Turkey 14/98, Inactivated

## Product identification

**Medicine name:**

Aftovaxpur DOE (12) O1 Manisa + A Turkey 14/98

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**Active substance:**

Foot-and-mouth disease virus, serotype O, strain O1 Manisa, Inactivated

Foot-and-mouth disease virus, serotype A, strain Turkey 14/98, Inactivated

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**Target species:**

This information is not available for this product.

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**Route of administration:**

Intramuscular use

Subcutaneous use

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## Product details

**Active substance and strength:**

Foot-and-mouth disease virus, serotype O, strain O1 Manisa, Inactivated

Presentation\_strength:? 6 PD50 Reference:Hse Index:0

Foot-and-mouth disease virus, serotype A, strain Turkey 14/98, Inactivated  
Presentation\_strength:? 6 PD50 Index:11

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**Pharmaceutical form:**

Emulsion for injection

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI02AA04

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Surrendered

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**Authorised in:**

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Boehringer Ingelheim Vetmedica GmbH

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**Marketing authorisation date:**

15/07/2013

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**Manufacturing sites for batch release:**

Boehringer Ingelheim Animal Health France  
Boehringer Ingelheim Animal Health France

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**Responsible authority:**

European Commission

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

16/05/2023

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

English (PDF)

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