# Equip Artervac Emulsion for Injection for Horses and Ponies

Authorised

- Equine arteritis virus, strain Bucyrus, Inactivated
- Equine arteritis virus, strain Bucyrus, Inactivated

## Product identification

**Medicine name:**
Equip Artervac Emulsion for Injection for Horses and Ponies

**Active substance:**
Equine arteritis virus, strain Bucyrus, Inactivated
Equine arteritis virus, strain Bucyrus, Inactivated

**Target species:**
Horse
Horse (pony)

**Route of administration:**
Intramuscular use
Intramuscular use

## Product details

**Active substance and strength:**
Equine arteritis virus, strain Bucyrus, Inactivated
1.80 relative unit(s) / 1.00 millilitre(s)

Equine arteritis virus, strain Bucyrus, Inactivated
1.80 relative unit(s) / 1.00 millilitre(s)

**Pharmaceutical form:**
Emulsion for injection

**Withdrawal period by route of administration:**
**Intramuscular use:**

- **Horse**
  - Meat and offal. 0 day
  - Milk. 0 day

**Intramuscular use:**

- **Horse (pony)**
  - Meat and offal. 0 day
  - Milk. 0 day

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QI05AA07

**Legal status of supply:**
Veterinary medicinal product subject to veterinary prescription

**Authorisation status:**
Valid

**Authorised in:**
Germany

**Package description:**
(ID3) 2 millilitre(s): Box (Cardboard) with 2 Syringe (Glass) each with 1 millilitre(s)
(ID2) 10 millilitre(s): Box (Cardboard) with 10 Syringe (Glass) each with 1 millilitre(s)
(ID1) 1 millilitre(s): Box (Cardboard) with 1 Syringe (Glass) with 1 millilitre(s)

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

**Marketing authorisation holder:**
Zoetis Deutschland GmbH

**Marketing authorisation date:**
23/05/2002

**Manufacturing sites for batch release:**
Zoetis Belgium

**Responsible authority:**
Paul-Ehrlich-Institut

**Authorisation number:**
PEI.V.01240.01.1

**Date of authorisation status change:**
29/03/2010

**Reference member state:**
Germany

**Procedure number:**
DE/V/0235/001

**Concerned member states:**
France Ireland United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

# Documents

Combined File of all Documents

English (PDF)
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