PROGRESSIS EMULSION FOR INJECTION FOR PIGS (SOWS AND GILTS)

Authorised

- Porcine reproductive and respiratory syndrome virus, type 1, strain P120, Inactivated

Product identification

Medicine name:
PROGRESSIS EMULSION FOR INJECTION FOR PIGS (SOWS AND GILTS)
Progressis Emulsie voor injectie
Progressis Emulsion injectable
Progressis Emulsion zur Injektion

Active substance:
Porcine reproductive and respiratory syndrome virus, type 1, strain P120, Inactivated

Target species:
Pig (for reproduction)

Route of administration:
Intramuscular use

Product details

Active substance and strength:

Porcine reproductive and respiratory syndrome virus, type 1, strain P120, Inactivated 2.50 log10 immunofluorescence unit(s) / 1.00 Dose

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Pig (for reproduction)**
  - All relevant tissues. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI09AA05

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

Authorisation status:
Valid

Authorised in:
Belgium

Package description:
Box of 1 bottle of 5 doses
Box of 10 plastic bottles of 50 doses
Box of 1 plastic bottle of 50 doses
Box of 10 bottles of 25 doses
Box of 1 bottle of 25 doses
Box of 10 bottles of 10 doses
Box of 1 bottle of 10 doses
Box of 10 bottles of 5 doses

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Ceva Sante Animale

Marketing authorisation date:
14/05/2020

Manufacturing sites for batch release:
Ceva-Phylaxia Zrt.

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
14/05/2020

Reference member state:
France

Procedure number:
FR/V/0115/001

Concerned member states:
Austria Belgium Denmark Finland Germany Greece Ireland Italy Luxembourg Netherlands Portugal Spain United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.