

Parvosuin MR/AD emulsão injectável para suínos

Authorised

- Porcine parvovirus, strain NADL-2, Inactivated
- Aujeszky's disease virus, strain Bartha K61, Live
- Erysipelothrix rhusiopathiae, strain R32E11, Inactivated

Product identification

Medicine name:

Parvosuin MR/AD emulsão injectável para suínos

Active substance:

Porcine parvovirus, strain NADL-2, Inactivated

Aujeszky's disease virus, strain Bartha K61, Live

Erysipelothrix rhusiopathiae, strain R32E11, Inactivated

Target species:

Pig

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Porcine parvovirus, strain NADL-2, Inactivated

2048.00 haemagglutination inhibiting unit(s) / 2.00 millilitre(s)

Aujeszky's disease virus, strain Bartha K61, Live

7.50 50% cell culture infectious dose / 2.00 millilitre(s)

Erysipelothrix rhusiopathiae, strain R32E11, Inactivated

20000000000.00 Colony forming unit / 2.00 millilitre(s)

Pharmaceutical form:

Emulsion for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI09AL

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Portugal

Package description:

Available only in Portuguese

Available only in Portuguese

Available only in Portuguese

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

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

Marketing authorisation holder:

Laboratorios Hipra S.A.

Marketing authorisation date:

29/05/2000

Manufacturing sites for batch release:

Laboratorios Hipra S.A.

Responsible authority:

Directorate General For Food And Veterinary

Authorisation number:

634/99 DGV

Date of authorisation status change:

1/10/2014

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

Documents

Combined File of all Documents

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