HIPRASUIS AD LIOFILIZADO Y DISOLVENTE PARA SUPENSION INYETABLE PARA PORCINO

Authorised

- Aujeszky's disease virus, strain Bartha K61, Inactivated

Product identification

Medicine name:
HIPRASUIS AD LIOFILIZADO Y DISOLVENTE PARA SUPENSION INYETABLE PARA PORCINO

Active substance:
Aujeszky's disease virus, strain Bartha K61, Inactivated

Target species:
Pig (for fattening)
Pig (female)
Pig (male for reproduction)

Route of administration:
Intramuscular use
Nasal use

Product details

Active substance and strength:

Aujeszky's disease virus, strain Bartha K61, Inactivated
105.50 50% cell culture infectious dose / 2.00 millilitre(s)

Pharmaceutical form:
Lyophilisate and solvent for suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Pig (for fattening)**
  - Meat and offal. 0 day

- **Pig (female)**
  - Meat and offal. 0 day

- **Pig (male for reproduction)**
  - Meat and offal. 0 day

Nasal use:

- **Pig (for fattening)**
  - Meat and offal. 0 day

- **Pig (female)**
  - Meat and offal. 0 day

- **Pig (male for reproduction)**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI09AD01

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

Authorisation status:
Valid

Authorised in:
Spain

Package description:
Available only in Spanish
Available only in Spanish
Available only in Spanish
Available only in Spanish
Available only in Spanish
Available only in Spanish
Available only in Spanish
Available only in Spanish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Informed consent application (Article 13c of Directive No 2001/82/EC)

Marketing authorisation holder:
Laboratorios Hipra S.A.

Marketing authorisation date:
3/12/2014

Manufacturing sites for batch release:
Laboratorios Hipra S.A.

Responsible authority:
Spanish Agency For Medicines And Medical Devices

Authorisation number:
3147 ESP

Date of authorisation status change:
3/12/2014

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

Documents

Summary of Product Characteristics

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Package Leaflet

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Labelling

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