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MYPRA VAC SUIS

Authorised

- Mycoplasma hyopneumoniae, Inactivated

Product identification

Medicine name:

MYPRA VAC SUIS

Active substance:

Mycoplasma hyopneumoniae, Inactivated

Target species:

Pig

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Mycoplasma hyopneumoniae, Inactivated
50.00 Allergy Unit(s) Modified / 1.00 Dose

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Pig

- Meat and offal. 2 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI09AB13

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Bulgaria

Package description:

Available only in [Bulgarian](#)

Available only in [Bulgarian](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Laboratorios Hipra S.A.

Marketing authorisation date:

9/03/2010

Manufacturing sites for batch release:

Laboratorios Hipra S.A.

Responsible authority:

Bulgarian Food Safety Authority

Authorisation number:

0022-2436/09.12.2014

Date of authorisation status change:

8/12/2014

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 and Labelling

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