MIXOHIPRA - H, Prášek a rozpouštědlo pro injekční suspenzi

Not authorised

- Myxoma virus, strain VMI 30, Live

Product identification

Medicine name:
MIXOHIPRA - H, Prášek a rozpouštědlo pro injekční suspenzi

Active substance:
Myxoma virus, strain VMI 30, Live

Target species:
Rabbit

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Myxoma virus, strain VMI 30, Live
5.30 50% tissue culture infectious dose / 1.00 Dose

Pharmaceutical form:

Powder and solvent for suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Rabbit**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI08AD02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Czechia

Package description:

Available only in Czech
Available only in Czech
Available only in Czech
Available only in Czech

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:

19/06/1997

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

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
97/538/97-C

Date of authorisation status change:
4/12/2024

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.