Rispoval IBR-Marker InactivatumSuspension for injection for cattle

Pooblaščeno

- Infectious Bovine rhinotracheitis virus, strain Difivac gE gene-deleted, Inactivated

Product identification

Ime zdravila:
Rispoval IBR-Marker InactivatumSuspension for injection for cattle
RISPOVAL IBR MARKER INACTIVATUM

Učinkovina:
Na voljo samo v English

Ciljne živalske vrste:
govedo
pitovno govedo
tele
telica

Pot uporabe:
Subkutana uporaba

Product details

Učinkovina / Jakost:
Na voljo samo v English

0.01 titre / 2.00 millilitre(s)

Farmacevtska oblika:
Suspenzija za injiciranje

Withdrawal period by route of administration:

Subkutana uporaba:

- **govedo**
  - Milk. 0 day
  - Meat and offal. 0 day
- **pitovno govedo**
  - Meat and offal. 0 day
- **tele**
  - Meat and offal. 0 day
- **telica**
  - Meat and offal. 0 day

Anatomsko-terapevtsko-kemična veterinarska oznaka (ATCvet):
QI02AA03

Pravni status za dobavo / izdajo zdravila:
Zdravilo, ki se izdaja na veterinarski recept

Status dovoljenja:
Valid

Authorised in:
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Opis ovojnine:
Na voljo samo v English
Na voljo samo v English

Additional information

Entitlement type:

Na voljo samo v English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

Pravna podlaga za izdajo dovoljenja za promet z zdravilom:
Na voljo samo v English Italian Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Zoetis France

Marketing authorisation date:
30/03/1995

Proizvodna mesta, odgovorna za sproščanje serij:
Zoetis Belgium

Pristojni organ:
French Agency For Food, Environmental And Occupational Health & Safety

Številka dovoljenja :
FR/V/6193958 1/1995

Datum spremembe statusa dovoljenja:
30/03/2010

Referenčna država članica:
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Številka postopka:
DE/V/0021/001

Zadevne države članice:
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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Na voljo samo v Estonian English French Swedish Icelandic Norwegian

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

Documents

Povzetek glavnih značilnosti zdravila

Ta dokument ne obstaja v tem jeziku (slovenščina). Spodaj ga najdete v drugem jeziku.

Package Leaflet and Labelling

Ta dokument ne obstaja v tem jeziku (slovenščina). Spodaj ga najdete v drugem jeziku.
