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ADJUSOL TMP SULFA LIQUIDE 83.35 MG/ML + 16.65 MG/ML SOLUTION FOR USE IN DRINKING WATER/MILK

Ima dovoljenje za promet

- Trimethoprim
- Sulfadiazine

Informacije o zdravilu

Ime zdravila:
ADJUSOL TMP SULFA LIQUIDE 83.35 MG/ML + 16.65 MG/ML SOLUTION FOR USE IN DRINKING WATER/MILK

Učinkovina:
Na voljo samo v English
Na voljo samo v English

Ciljne živalske vrste:
brojler
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
Na voljo samo v Bulgarian Spanish Czech Danish Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian

Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
tele
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian

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Pot uporabe:
Dajanje v vodo za pitje/mleko

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
16.65 milligram(s) / 1.00 millilitre(s)

Na voljo samo v English
83.35 milligram(s) / 1.00 millilitre(s)

Farmacevtska oblika:
Raztopina za dajanje v vodo za pitje/mleko

Karenca glede na pot uporabe:
Dajanje v vodo za pitje/mleko:

- **brojler**
  - Meat and offal. 12 day
- **Pig**
  - Meat and offal. 12 day
- **Rabbit**
  - Meat and offal. 12 day
-

Sheep (lamb)
- Meat and offal. 12 day

•

tele
- Meat and offal. 12 day

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Chicken
- Eggs. no withdrawal period

Not for use in birds producing or intended to produce eggs for human consumption.

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QJ01EW10

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

Status dovoljenja za promet z zdravilom:
Valid

Ima dovoljenje za promet v:
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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

Dodatne informacije

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

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

Imetnik dovoljenja za promet z zdravilom:
Virbac

Datum dovoljenja za promet z zdravilom:
15/05/1995

Proizvodna mesta, odgovorna za sproščanje serij:
Virbac
FC France S.A.S.

Pristojni organ:
Directorate General For Food And Veterinary

Številka dovoljenja za promet z zdravilom:
51105

Datum spremembe statusa dovoljenja za promet:
6/10/2026

Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinaski medicini obiščite stran: www.adrreports.eu/vet

Dokumenti

Combined File of all Documents

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