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BLAP HELP, injekcinis tirpalas karvėms

Authorised

- Nicotinamide adenine dinucleotide phosphate
- Cyanocobalamin
- Thiamine hydrochloride
- Magnesium acetate
- Calcium gluconate
- Lysine hydrochloride
- Arginine hydrochloride
- Acetylmethionine
- Inositol nicotinate
- GLUCOSE BP
- SORBITOL BP

Product identification

Medicine name:
BLAP HELP, injekcinis tirpalas karvėms

Active substance:
Nicotinamide adenine dinucleotide phosphate

Cyanocobalamin

Thiamine hydrochloride

Magnesium acetate

Calcium gluconate

Lysine hydrochloride

Arginine hydrochloride

Acetylmethionine

Inositol nicotinate

GLUCOSE BP

SORBITOL BP

Target species:
Cattle (cow)

Route of administration:
Intravenous use
Subcutaneous use
Intraperitoneal use

Product details

Active substance and strength:
Nicotinamide adenine dinucleotide phosphate
2.50 gram(s) / 500.00 millilitre(s)

Cyanocobalamin
0.50 milligram(s) / 500.00 millilitre(s)

Thiamine hydrochloride
0.50 gram(s) / 500.00 millilitre(s)

Magnesium acetate
0.30 gram(s) / 500.00 millilitre(s)

Calcium gluconate
10.00 gram(s) / 500.00 millilitre(s)

Lysine hydrochloride
0.60 gram(s) / 500.00 millilitre(s)

Arginine hydrochloride
0.60 gram(s) / 500.00 millilitre(s)

Acetylmethionine
12.50 gram(s) / 500.00 millilitre(s)

Inositol nicotinate
5.00 gram(s) / 500.00 millilitre(s)

GLUCOSE BP
10.00 gram(s) / 500.00 millilitre(s)

SORBITOL BP
50.00 gram(s) / 500.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QV06DE

Legal status of supply:
This information is not available for this product.

Authorisation status:
Valid

Authorised in:
Lithuania

Package description:
Available only in Lithuanian
Available only in Lithuanian

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:
Izo S.r.l.

Marketing authorisation date:
19/05/2008

Manufacturing sites for batch release:

Izo S.r.l.

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/08/1792/001

Date of authorisation status change:
2/06/2013

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

Documents

RV1792.pdf