BLAP HELP

Authorised

- Magnesium acetate
- Acetylmethionine
- Glucose
- L-ARGININE
- Inositol
- Calcium gluconate
- Cyanocobalamin
- Lysine
- Thiamine hydrochloride
- SORBITOL (E420)
- Nicotinamide

Product identification

Medicine name:
BLAP HELP

Active substance:
Magnesium acetate

Acetylmethionine

Glucose

L-ARGININE

Inositol

Calcium gluconate

Cyanocobalamin

Lysine

Thiamine hydrochloride

SORBITOL (E420)

Nicotinamide

Target species:
Cattle

Route of administration:
Intraperitoneal use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
Magnesium acetate
0.30 gram(s) / 500.00 millilitre(s)

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

Glucose
10.00 gram(s) / 500.00 millilitre(s)

L-ARGININE
0.60 gram(s) / 500.00 millilitre(s)

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

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

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

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

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

SORBITOL (E420)
50.00 gram(s) / 500.00 millilitre(s)

Nicotinamide
2.50 gram(s) / 500.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intraperitoneal use:

- **Cattle**
  - Milk. 0 day

Intravenous use:

- **Cattle**
  - Milk. 0 day

Subcutaneous use:

- **Cattle**
  - Milk. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QV06DE

Legal status of supply:
Veterinary medicinal product not subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Italy

Package description:
Available only in Italian
Available only in Italian

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:
25/05/1993

Manufacturing sites for batch release:
Izo S.r.l.

Responsible authority:
Ministry Of Health

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
25/05/1993

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

Documents

Combined File of all Documents

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