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ULTRA B

Not authorised

- Thiamine hydrochloride
- Pyridoxine hydrochloride

Product identification

Medicine name:
ULTRA B

Active substance:
Thiamine hydrochloride
Pyridoxine hydrochloride

Target species:
Cattle
Pig
Equid
Sheep
Goat
Dog

Route of administration:
Intramuscular use
Oral use
Intravenous use

Product details

Active substance and strength:
Thiamine hydrochloride
100.00 milligram(s) / 1.00 millilitre(s)

Pyridoxine hydrochloride
8.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Pig**
  - Meat and offal. 0 day
- **Equid**
  - Meat and offal. 0 day
- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Goat**
  - Meat and offal. 0 day
  - Milk. 0 day

Oral use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Pig**
  - Meat and offal. 0 day
- **Equid**
  - Meat and offal. 0 day
- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Goat**
  - Meat and offal. 0 day
  - Milk. 0 day

Intravenous use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Equid**
  - Meat and offal. 0 day
- **Sheep**
  - Meat and offal. 0 day

- Milk. 0 day

- **Goat**
  - Meat and offal. 0 day
  - Milk. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA11DB

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

Authorisation status:
Surrendered

Authorised in:
France

Package description:
Available only in French
Available only in French
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Intervet

Marketing authorisation date:
11/07/1984

Manufacturing sites for batch release:
Trirx Segre

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/4554948 5/1984

Date of authorisation status change:
4/06/2026

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

Documents

Summary of Product Characteristics

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