Butador 10 mg/ml Solution for Injection for Horses, Dogs and Cats

Authorised

- Butorphanol tartrate

Product identification

Medicine name:
Butomidor 10 mg/ml Injektionslösung für Pferde, Hunde und Katzen
Butador 10 mg/ml Solution for Injection for Horses, Dogs and Cats

Active substance:
Butorphanol tartrate

Target species:
Dog
Horse
Cat

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
Butorphanol tartrate
14.58 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intravenous use:

- **Horse**
  - Meat and offal. 0 day
  - Milk. 0 hour

Subcutaneous use:

- **Cat**
  - Meat and offal. 0 day
  - Milk. 0 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN02AF01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
United Kingdom (Northern Ireland)

Available in:
United Kingdom (Northern Ireland)

Package description:
Clear glass vials with bromobutyl rubber stoppers and aluminium caps. Package size: 1 x 10 ml
Clear glass vials with bromobutyl rubber stoppers and aluminium caps. Package size: 1 x 50 ml
Clear glass vials with bromobutyl rubber stoppers and aluminium caps. Package size: 10 x 10 ml
Clear glass vials with bromobutyl rubber stoppers and aluminium caps. Package size: 5 x 10 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Vetviva Richter GmbH

Marketing authorisation date:
17/01/2011

Manufacturing sites for batch release:
Vetviva Richter GmbH

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 57446/3006

Date of authorisation status change:
23/11/2024

Reference member state:
Austria

Procedure number:
AT/V/0005/001

Concerned member states:
Belgium Denmark Finland France Germany Greece Iceland Ireland Latvia Lithuania Netherlands Norway Portugal Slovenia Spain Sweden United Kingdom (Northern Ireland)

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