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Ibaflin (WD) 30 mg - Tablet

Not authorised

- Ibafloxacin

Product identification

Medicine name:
Ibaflin (WD) 30 mg - Tablet

Active substance:
Ibafloxacin

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Ibafloxacin
30.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01MA96

Legal status of supply:

Medicinal product subject to medical prescription

Authorisation status:

Surrendered

Surrendered

Authorised in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland) , Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Additional information

Entitlement type:

Marketing Authorisation

Marketing Authorisation

Legal basis of product authorisation:

Complete application (stand-alone) - Council Directive 81/851/EEC

Complete application (stand-alone) - Council Directive 81/851/EEC

Marketing authorisation holder:

INTERVET INTERNATIONAL B.V.

INTERVET INTERNATIONAL B.V.

Marketing authorisation date:

13/06/2000

13/06/2000

Manufacturing sites for batch release:

Intervet Ges.m.b.H.

Responsible authority:

European Commission
European Commission

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

Date of authorisation status change:
21/06/2013
23/02/2004

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