

**PARTICULARS TO APPEAR ON THE OUTER PACKAGE {Carton, when a pull-out label is not used} AND THE IMMEDIATE PACKAGE {Tub label. When concertina label is used, the front face is also duplicated and attached to the container}**

**1. NAME OF THE VETERINARY MEDICINAL PRODUCT**

Furosemide Tablets BP (Vet) 40 mg

Furosemide

**2. STATEMENT OF ACTIVE SUBSTANCES**

Furosemide 40 mg

**3. PHARMACEUTICAL FORM**

Tablet

**4. PACKAGE SIZE**

1000 Tablets

**5. TARGET SPECIES**

Dogs and cats

**6. INDICATION(S)**

**7. METHOD AND ROUTE(S) OF ADMINISTRATION**

Oral use. Read the package leaflet before use.

**8. WITHDRAWAL PERIOD**

**9. SPECIAL WARNING(S), IF NECESSARY**

Wear gloves or wash hands immediately after handling tablets. In case of accidental ingestion seek medical attention and show product label and/or package leaflet to the doctor.

Read the package leaflet before use.

**10. EXPIRY DATE**

EXP {MM/YYYY}

## 11. SPECIAL STORAGE CONDITIONS

[When a carton is used instead of a pull-out label:] Keep the container in the outer carton.

Store in the original container in order to protect from light.

## 12. SPECIFIC PRECAUTIONS FOR THE DISPOSAL OF UNUSED PRODUCTS OR WASTE MATERIALS, IF ANY

Dispose of waste material in accordance with local requirements.

## 13. THE WORDS “FOR ANIMAL TREATMENT ONLY” AND CONDITIONS OR RESTRICTIONS REGARDING SUPPLY AND USE IF APPLICABLE *[Distribution category]*

For animal treatment only. To be supplied only on veterinary prescription.

POM-V

## 14. THE WORDS “KEEP OUT OF THE SIGHT AND REACH OF CHILDREN”

KEEP OUT OF THE SIGHT AND REACH OF CHILDREN

## 15. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER

Millpledge Ltd  
Whinleys Estate  
Clarborough  
Retford  
Nottinghamshire  
DN22 9NA

## 16. MARKETING AUTHORISATION NUMBER

Vm 04409/4000

## 17. MANUFACTURER’S BATCH NUMBER

**PACKAGE LEAFLET FOR:**

**Furosemide Tablets BP (Vet) 40 mg**

**1. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER AND OF THE MANUFACTURING AUTHORISATION HOLDER RESPONSIBLE FOR BATCH RELEASE, IF DIFFERENT**

Marketing Authorisation holder:

Millpledge Ltd  
Whinleys Estate  
Clarborough  
Retford  
Nottinghamshire  
DN22 9NA

Manufacturer responsible for batch release:

Millpledge Ltd  
Unit 6/8 Heapham Road Industrial Estate  
Gainsborough  
Lincolnshire DN21 1RZ

**2. NAME OF THE VETERINARY MEDICINAL PRODUCT**

Furosemide Tablets BP (Vet) 40 mg

Furosemide

**3. STATEMENT OF THE ACTIVE SUBSTANCE (S) AND OTHER INGREDIENTS**

Furosemide 40mg

Oral Tablets containing the stated amount of Furosemide. Flat faced, white circular with bevelled edges and a scored half break line, embossed F40 and plain on the reverse

**4. INDICATION(S)**

For the treatment of oedema, the product may be used in ascites, hydrothorax, pulmonary oedema of the mammary glands or legs, as well as oedema resulting from cardiac insufficiency, hepatic or renal dysfunction, parasitism or of a traumatic origin.

**5. CONTRAINDICATIONS**

Do not use in acute glomerular nephritis, in electrolyte diseases, in patients with anuria, or patients that have received excessive doses of cardiac glycosides. Because of the danger of potentiating their toxic effects do no use with aminoglycoside or cephalosporin antibiotics. Allergic reactions have been associated with use with sulphonamides.

## **6. ADVERSE REACTIONS**

The patient may increase its water intake to compensate for the diuresis. Consideration should be given to restricting water intake if the patient's condition makes such a course appropriate.

## **7. TARGET SPECIES**

Cat and dog

## **8. DOSAGE FOR EACH SPECIES, ROUTE(S) AND METHOD OF ADMINISTRATION**

Oral use. 5mg/KgBW, one or two times per day. For patients weighing less than 8Kg dosage with the 20mg tablet (which may be halved) is recommended. Avoid overdosage in weak and old patients.

## **9. ADVICE ON CORRECT ADMINISTRATION**

## **10. WITHDRAWAL PERIOD(S)**

Not applicable.

## **11. SPECIAL STORAGE PRECAUTIONS**

Keep out of the sight and reach of children.

Store in the original container in order to protect from light.

[When a carton is used instead of a pull-out label:] Keep the container in the outer carton.

Do not use this veterinary medicinal product after the expiry date which is stated on the tub. The expiry date refers to the last day of that month.

## **12. SPECIAL WARNING(S)**

### Special precautions for use in animals

Prolonged dosage may on occasions justify potassium supplementation and thus monitoring for hypokalaemia should be considered, especially if the product is used in conjunction with cardiac glycosides.

Pregnancy and lactation: The safety of use in pregnancy is not well established and a careful assessment of the likely benefits and potential risks should be made. A deleterious effect on lactation is to be expected, particularly if drinking water is restricted. Furosemide passes into milk, but not to a great extent.

Interactions: Potential interactions with other drugs include ototoxicity with aminoglycosides and nephrotoxicity with cephalosporins. Use in combination with sulphonamide treatment may lead to sulphonamide allergy. There is a possibility of interaction with cardiac glycosides.

Overdose: Dehydration and electrolyte depletion may occur. Monitor and correct as necessary. Dosage higher than that which is recommended, may cause transitory

deafness. Cardiovascular side effects may be observed in weak and old patients following overdose.

Special precautions to be taken by the person administering the medicinal product to animals:

Wear gloves or wash hands immediately after handling tablets. In case of accidental ingestion seek medical attention and show product label and/or pack insert to the doctor.

For Animal Treatment Only

**13. SPECIAL PRECAUTIONS FOR THE DISPOSAL OF UNUSED PRODUCT OR WASTE MATERIALS, IF ANY**

Any unused veterinary medicinal product or waste materials derived from such veterinary medicinal product should be disposed of in accordance with local requirements.

**14. DATE ON WHICH THE PACKAGE LEAFLET WAS LAST APPROVED**

April 2021

**15. OTHER INFORMATION**

POM – V To be supplied only on veterinary prescription.  
Vm 04409/4000

Approved: 22/04/21

