Canergy 100mg tablets for dogs

Species: Dogs
Therapeutic indication:Pharmaceuticals: Cardiovascular and respiratory preparations
Active ingredient: Propentofylline
Product: Canergy 100mg tablets for dogs
Product index: Canergy 100mg tablets for dogs

Presentation

PHARMACEUTICAL FORM

Tablet.

Light brown with brown spots, round and convex flavoured tablet with a cross-shaped break line on one side.

The tablets can be divided into 2 or 4 equal parts.

Each tablet contains :

Active substance:

Propentofylline 100 mg

Uses

Target species: Dogs

For the improvement of peripheral and cerebral vascular blood circulation.

For improvement in dullness, lethargy and overall demeanour in dogs.

Dosage and administration

The basic dosage is 6-10 mg propentofylline/kg bodyweight daily, divided into two doses as follows:

	100 mg Tablets			
Body weight (kg)	**Morning**	**Evening**	**Daily total tablets**	**Daily total dose (mg/kg)**
5 – 8	1/4	1/4	½	6.25 – 10.0

>8 – 10	1/2	1/4	¾	7.5 – 9.4
>10 – 15	1/2	1/2	1	6.7 – 10.0
>15 – 25	3/4	3/4	1 ½	6.0 – 10.0
>25 – 33	1	1	2	6.1 – 8.0
>33 – 49	1 ½	1 ½	3	6.1 – 9.1
>49 – 66	2	2	4	6.1 – 8.2
>66 – 83	2 ½	2 ½	5	6.0 – 7.6

The tablets can be administered directly in the mouth, onto the back of the dog's tongue or can be mixed in a small ball of food and should be administered at least 30 minutes before feeding.

Tablets can be divided into 2 or 4 equal parts to ensure accurate dosing. Place the tablet on a flat surface, with its scored side facing up and the convex (rounded) side facing the surface.

Halves: press down with your thumbs on both sides of the tablet

Quarters: press down with your thumb in the middle of the tablet

Contra-indications, warnings, etc

Contraindications

Do not use in dogs weighing less than 5 kg.

Do not use in case of hypersensitivity to the active substance or to any of the excipients.

Special precautions for use

Special precautions for use in animals

Specific diseases (e.g. kidney disease) should be treated accordingly.

Consideration should be given to rationalising the medication of dogs already receiving treatment for congestive heart failure or bronchial disease.

In the case of renal failure, the dose should be reduced.

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

Care should be taken to avoid accidental ingestion.

In the event of accidental ingestion of the tablets, seek medical advice immediately and show the package leaflet or the label to the physician. Wash hands after use.

Adverse reactions (frequency and seriousness)

On rare occasions (more than 1 but less than 10 animals in 10,000 animals), allergic skin reactions, vomiting and cardiac disturbances have been reported. In these cases, the

treatment should be stopped.

Use during pregnancy or lactation

The safety of the veterinary medicinal product has not been established during pregnancy and/or lactation. Use in pregnant or lactating bitches or breeding animals is therefore not recommended.

Interaction with other medicinal products and other forms of interaction

None known.

Pharmaceutical precautions

Shelf-life

Shelf-life of the veterinary medicinal product as packaged for sale: 3 years

Shelf life of divided tablets after first opening the immediate packaging: 4 days.

Special precautions for storage

This veterinary medicinal product does not require any special temperature storage conditions.

Any unused tablet portion should be returned to the open blister and inserted back into the carton to be used for the next administration.

Special precautions for the disposal of unused veterinary medicinal product or waste materials derived from the use of such products

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

Legal category

Legal category:POM-V

Packaging quantities

Nature and composition of immediate packaging

Aluminium - PA/ALU/PVC blister

Cardboard box of 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 25 or 50 blisters of 10 tablets

Not all pack sizes may be marketed.

Marketing Authorisation Holder (if different from distributor)

Le Vet Beheer B.V.

Wilgenweg 7

3421 TV Oudewater.

The Netherlands

Further information

Marketing Authorisation Number

Vm 41821/4019

Significant changes

GTIN

GTIN description:--
GTIN:--

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