PARTICULARS TO APPEAR ON THE OUTER PACKAGE CARTON

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Johnson's Wormer Plus 150/144/50mg tablets for dogs & puppies

2. STATEMENT OF ACTIVE SUBSTANCES

Each tablet contains:
150 mg febantel
144 mg pyrantel embonate
50 mg praziquantel

3. PACKAGE SIZE

Cartons containing 1, 2, 4, 6 and 8, tablets

4. TARGET SPECIES

Dogs

5. INDICATIONS

For the treatment of the following common worms affecting dogs and puppies:
Roundworms (ascarids, hookworms, whipworms) Tapeworms

6. ROUTES OF ADMINISTRATION

Read the package leaflet before use

For oral administration only.

Give 1 tablet per 10kg bodyweight.

Weigh your dog carefully before calculating the dose. Do not give any other dose than recommended without consulting your veterinary surgeon or suitably qualified professional. The tablets should be given directly in the mouth of your dog but if necessary, can be disguised in food.

The tablets are scored into halves to enable a range of weights to be treated.

Puppies and Small Dogs:
3-5 kg bodyweight = ½ tablet
Over 5-10 kg bodyweight = 1 tablet

Medium Dogs:
Over 10-15 kg bodyweight = 1 ½ tablets
Over 15-20 kg bodyweight = 2 tablets
Over 20-25 kg bodyweight = 2 ½ tablets
Over 25-30 kg bodyweight = 3 tablets

Large Dogs:
Over 30-35 kg bodyweight = 3 ½ tablets
Over 35-40 kg bodyweight = 4 tablets

For use in puppies from 2 weeks of age and minimum weight of 3 kg

7. WITHDRAWAL PERIODS

8. EXPIRY DATE

Exp. {mm/yyyy}

9. SPECIAL STORAGE PRECAUTIONS

Keep blister in outer carton. Do not store above 25°C. Any part-used tablets should be discarded.

10. THE WORDS "READ THE PACKAGE LEAFLET BEFORE USE"

In case of accidental ingestion, seek medical advice and show the package leaflet or the label to the physician.
In the interests of good hygiene, people administering the tablet directly to the dog or by adding it to the dog's food, should wash their hands afterwards.

Read the package leaflet before use.

11. THE WORDS "FOR ANIMAL TREATMENT ONLY"

For animal treatment only.

12. THE WORDS "KEEP OUT OF THE SIGHT AND REACH OF CHILDREN"

Keep out of the sight and reach of children.

13. NAME OF THE MARKETING AUTHORISATION HOLDER

Vetoquinol UK Limited

14.MARKETING AUTHORISATION NUMBERS

Vm 08007/5037

15. BATCH NUMBER

Lot {number}

16. OTHER INFORMATION

Not for use in dogs weighing less than 3 kg and under 2 weeks of age. The safety of the product has not been assessed in puppies younger than 2 weeks old.

Do not use simultaneously with other deworming products without veterinary advice

Do not use in any animal with hypersensitivity to the active substance or to any of the excipients

The safety of the product has not been assessed in pregnant dogs. Consult a veterinarian before treating pregnant animals for roundworms. The product may be used during lactation

Revised: October 2025
MA split from NI Expiry following AN: 02635/2025

MINIMUM PARTICULARS TO APPEAR SMALL IMMEDIATE PACKAGING - BLISTER STRIPS

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Johnson's Wormer Plus 150/144/50mg tablets for dogs & puppies

2. QUANTITATIVE PARTICULARS OF THE ACTIVE SUBSTANCES

150.0 mg	Febantel
144.0 mg	Pyrantel embonate
50.0 mg	Praziquantel

3. BATCH NUMBER

Lot {number}

4. EXPIRY DATE

Exp. {mm/yyyy}

PARTICULARS TO APPEAR ON THE PACKAGE LEAFLET:

PACKAGE LEAFLET

1. Name of the veterinary medicinal product

Johnson's Wormer Plus 150/144/50mg tablets for dogs & puppies

2. Composition

Each tablet contains:

Active substances:

150.0 mg	Febantel
144.0 mg	Pyrantel embonate
50.0 mg	Praziquantel

Light brown to brown round flat tablet, cross scored on one side.

3. Target species

Dogs.

4. Indications for use

For the treatment of the following gastrointestinal tapeworms and roundworms (ascarids, hookworms, whipworms) in dogs and puppies:

Ascarids:	*Toxocara canis, Toxascaris leonina* (adult and late immature forms).
Hookworms:	*Uncinaria stenocephala, Ancylostoma caninum* (adults)
Whipworms:	*Trichuris vulpis* (adults)
Tapeworms:	*Echinococcus* spp. *Taenia* spp. and *Dipylidium* species (adult and immature forms).

5. Contraindications

Do not use in cases of hypersensitivity to the active substances or to any of the excipients.

Do not use simultaneously with piperazine compounds as piperazine may block the action of pyrantel embonate contained in this veterinary medicinal product. Other worming products may contain piperazine.

Do not use simultaneously with other deworming products without veterinary advice.

Not for use in dogs weighing less than 3 kg.

6. Special warnings

Special warnings:

Dogs may become infected with worms by eating insects (including fleas and lice), birds, small rodents, rabbits or raw offal from affected sheep, goats and cattle. Dogs will continue to be re-infected unless the route of infection is controlled e.g. treating a flea infestation or preventing a dog from scavenging or hunting.

Fleas serve as intermediate hosts for one common type of tapeworm - *Dipylidium caninum*. Tapeworm infestation is certain to re-occur unless control of intermediate hosts such as fleas, mice etc. is undertaken.

Echinococcus multilocularis is a tapeworm of foxes, dogs and cats which is common in parts of Europe. It can also infect humans potentially causing severe liver disease which can be fatal. To prevent *Echinococcus multilocularis* establishing in the UK and Ireland it is a requirement that all dogs must be treated with a product containing praziquantel before entering the country.

Special precautions for safe use in the target species:

The safety of the product has not been assessed in puppies younger than 2 weeks old

Parasite resistance to any particular class of anthelmintic (de-wormer) may develop following frequent, repeated use of an anthelmintic of that class.

Underdosing could result in ineffective use and may favour resistance development.

To ensure a correct dosage, body weight should be determined as accurately as possible

Do not exceed the stated dose.

The safety of the product has not been assessed in pregnant dogs. Consult a veterinarian before treating pregnant animals for roundworms.

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

In case of accidental ingestion, seek medical advice and show the package leaflet or the label to the physician.

In the interests of good hygiene, people administering the tablet directly to the dog or by adding it to the dog’s food, should wash their hands afterwards.

Other precautions:

Echinococcosis represents a hazard for humans. As Echinococcosis is a notifiable disease to the World Organisation for Animal Health (WOAH), specific guidelines on the treatment and follow-up, and on the safeguard of persons, need to be obtained from the relevant competent authority.

Pregnancy and lactation:

Consult a veterinary surgeon before treating pregnant animals for roundworms. The product may be used during lactation.

Interaction with other medicinal products and other forms of interaction:

Overdose:

The product is well tolerated in dogs. In safety studies doses of 5 x or greater gave rise to occasional vomiting.

Special restrictions for use and special conditions for use:

None.

Major incompatibilities:

None known.

7. Adverse events

Very rare (<1 animal / 10,000 animals treated, including isolated reports):	Digestive tract disorders (e.g., vomiting and diarrhoea)[1], Anorexia, Lethargy, Hyperactivity.

[1]Mild and transient.

Reporting adverse events is important. It allows continuous safety monitoring of a product. If you notice any side effects, even those not already listed in this package leaflet, or you think that the medicine has not worked, please contact, in the first instance, your veterinarian. You can also report any adverse events to the marketing authorisation holder or the local representative of the marketing authorisation holder using the contact details at the end of this leaflet, or via your national reporting system at:

Website: https://www.gov.uk/report-veterinary-medicine-problem/animal-reacts-medicine

e-mail: adverse.events@vmd.gov.uk

8. Dosage for each species, routes and method of administration

Dosage

Give as a single oral administration
1 tablet per 10 kg bodyweight. The tablet can be given directly to the dog or disguised in food. No starvation is needed before, or after, treatment.

Puppies and Small Dogs:

3-5 kg bodyweight	= ½ tablet
Over 5-10 kg bodyweight	= 1 tablet

Medium Dogs:

Over 10-15 kg bodyweight	= 1½ tablets
Over 15-20 kg bodyweight	= 2 tablets
Over 20-25 kg bodyweight	= 2½ tablets
Over 25-30 kg bodyweight	= 3 tablets

Large Dogs:

Over 30-35 kg bodyweight	= 3½ tablets
Over 35-40 kg bodyweight	= 4 tablets

9. Advice on correct administration

For the control of *Toxocara,* nursing bitches should be dosed 2 weeks after giving birth and every two weeks until weaning. Puppies should be treated at 2 weeks of age and every 2 weeks until 12 weeks of age. Thereafter, the need for and frequency of re-treatment(s) should be based on professional advice (from a veterinarian or suitably qualified professional) and should take into account the local epidemiological situation and the animal's lifestyle. It is advisable to treat the bitch at the same time as the puppies.

In case of suspected heavy roundworm infestation, please contact your veterinary surgeon for diagnosis and treatment recommendation.

10. Withdrawal periods

Not applicable.

11. Special storage precautions

Do not store above 25ºC. Keep blister in outer carton. Part used tablets should be discarded. Do not use this veterinary medicinal product after the expiry date stated on the blister after Exp. The expiry date refers to the last day of that month.

12. Special precautions for disposal

Medicines should not be disposed of via wastewater.

Use take-back schemes for the disposal of any unused veterinary medicinal product or waste materials derived thereof in accordance with local requirements and with any applicable national collection systems. These measures should help to protect the environment.

Ask your veterinary surgeon or pharmacist how to dispose of medicines no longer required.

13. Classification of veterinary medicinal products

Veterinary medicinal product not subject to prescription.

14. MARKETING AUTHORISATION NUMBERS AND PACK SIZES

Vm 08007/5037
Aluminium foil blister or polyethylene-coated aluminium blister
Cartons containing 1, 2, 4, 6 and 8 tablets
Not all pack sizes may be marketed.

15. PID LINK (Do not print heading)

[The following statement must be included where reference to the European Union Product Database is included on the product information. This statement is relevant to both UK(GB) and UK(NI) products:]

Find more product information by searching for the 'Product Information Database' on www.gov.uk.

16. Contact details

Marketing authorisation holder:

Vetoquinol UK Limited
Steadings Barn
Pury Hill Business Park
Nr. Alderton
Towcester
Northamptonshire
NN12 7LS

Manufacturer responsible for batch release:

KVP Pharma + Veterinär Produkte GmbH, Projensdorfer Str. 324, 24106 Kiel, Germany

and

EUROPHARTECH, 34 rue Henri Matisse, 63370 Lempdes, France

Local representatives and contact details to report suspected adverse reactions:

Johnson's Veterinary Products Ltd.
Sutton Coldfield
West Midlands
B75 7DF, UK

Tel: +44 (0)121 378 1684
Email: info@johnsons-vet.com

For any information about this veterinary medicinal product, please contact the local representative of the marketing authorisation holder.

17. Other information

AVM-GSL

Gavin Hall
Approved: 30 October 2025