

RESTRICTED VETERINARY MEDICINE
KEEP OUT OF REACH OF CHILDREN
FOR ANIMAL TREATMENT ONLY

Buscopan® compositum



SPASMOLYTIC AND ANALGESIC

Composition

Hyoscine-N-Butylbromide 4 mg/mL
Dipyron 500 mg/mL

Buscopan® compositum is a combination of the spasmolytic agent Buscopan® (Hyoscine-N-Butylbromide) and the analgesic dipyron.

Activity

Buscopan® compositum acts in three ways:

- directly as a spasmolytic
- directly as an analgesic
- indirectly as an anti-inflammatory agent

These three actions are interrelated.

Pharmacology

Buscopan® blocks the transmission of the neural stimuli at the intramural ganglia of the hollow abdominal organs, which are innervated by the parasympathetic nervous system. Absorption and distribution studies established that Buscopan® compositum is concentrated in the walls of the hollow abdominal organs causing a spasmolysis. This accounts for the lack of systemic atropine like side effects. Buscopan® remains mainly in the enterohepatic circulation. Dipyron has central analgesic activity plus a direct effect on the smooth muscle cells, lowering the excitability

of the smooth muscle fibres. This latter effect together with dipyron's anti-inflammatory action enhances the properties of Buscopan®. Dipyron is well tolerated parenterally.

Clinical Action

Buscopan® compositum is recommended in all conditions of severe pain involving spasm in the gastrointestinal, biliary and urogenital tracts. With intravenous administration pain often recedes before the injection is completed and pain relief is clearly apparent. The injection may be repeated after four hours if necessary. The duration of action is approximately six hours.

Buscopan® compositum brings about relief of colic by abolishing the spasm and relieving the pain ensuring a rapid return to normal peristalsis and intestinal function.

Enteritis associated with scouring quickly debilitates the animal through excessive fluid loss. Buscopan® compositum normalises intestinal tone and function, preventing further fluid loss. Concurrent treatment with an appropriate antibacterial or anthelmintic may be instituted. Clinical reports confirm Buscopan® compositum's usefulness of colic in horses

and gastroenteritis in small animals.

Indications

Assists in the treatment of gastroenteritis.

Spasm of the gastro-intestinal, biliary or urogenital tracts associated with severe pain. Buscopan® compositum relieves both pain and smooth muscle spasm.

Horses

Spasmodic colic, scours, oesophageal obstruction, enteritis with diarrhoea.

Cattle/Calves

Pain with enteritis and digestive disturbance, diarrhoea, oesophageal obstruction, postparturient relief of pain, functional tympanitis.

Pigs

All forms of colic, gastroenteritis, diarrhoea.

Dogs

Colic, gastro-enteritis with diarrhoea, vomiting, spasm of the urogenital tract, pain associated with digestive disturbances, tenesmus, urinary calculi.

In gastro-enteritis Buscopan® compositum can be used in combination with antibiotics, sulphonamides or anthelmintics, as required.

DIRECTIONS FOR USE

Horses	20-30 mL intravenously
Cattle	20-30 mL intravenously or intramuscularly
Calves/Pigs	5-10 mL intramuscularly
Piglets	1-2 mL intramuscularly
Dogs	1-2.5 mL subcutaneously or intravenously

Intravenous injections to be given slowly.

The injection may be repeated after 4 hours if necessary.

It is an offence for users of this pro-

duct to cause residues exceeding the relevant MRL in the New Zealand (Maximum Residues Limits of Agricultural Compounds) Food Standards.

WITHHOLDING PERIOD

MEAT: ANIMAL PRODUCING MEAT OR OFFAL FOR HUMAN CONSUMPTION MUST NOT BE SOLD FOR SLAUGHTER EITHER DURING TREATMENT OR WITHIN: HORSES 2 DAYS, CATTLE AND PIGS 14 DAYS OF CESSATION OF THE LAST TREATMENT. MILK: INTENDED FOR HUMAN CONSUMPTION MUST BE DISCARDED DURING TREATMENT AND FOR 12 HOURS FOLLOWING THE LAST TREATMENT.

Disposal

Dispose of empty container by wrapping with paper and putting in garbage.

Presentation

100 mL bottle solution for injection

Restricted Veterinary Medicine. No. A3483

See www.foodsafety.govt.nz for registration conditions.

Storage

Store below 30°C (Room Temperature).

Protect from light.

Avoid freezing.

After opening, the colour of the injection solution may change. This does not influence the efficacy.

Distributed by

Boehringer Ingelheim (NZ) Ltd.

Animal Health Division

Level 1, Unit 9

42 Ormiston Road

East Tamaki Auckland