Radiographic Projections for Detection of **Pneumoperitoneum**

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In the literature

Ng J, Linn KA, Shmon CL, Parker S, Zwicker LA. The left lateral projection is comparable to horizontal beam radiography for identifying experimental small volume pneumoperitoneum in the canine abdomen. Vet Radiol Ultrasound. 2020;61(2):130-136.

Pneumoperitoneum can be an important indicator of GI perforation.

FROM THE PAGE ...

In the absence of trauma or penetrating wounds, pneumoperitoneum can be an important indicator of GI perforation. However, detection of small amounts of free gas can be difficult to distinguish from intraluminal gas on standard radiographic projections. Obtaining images with horizontal beam projection could enhance detection of free gas. Air may be more likely to accumulate against the body wall and may be more easily distinguished from intestinal gas with either ventrodorsal or lateral horizontal beam projections. However, a small cadaveric study failed to find a difference in the accuracy of horizontal beam projections over standard views in detection of pneumoperitoneum.² To investigate this further, the present study evaluated whether use of horizontal beam projections improves the ability to detect abdominal free gas and whether the level of training of the radiographic interpreter impacts accuracy.

To gauge the sensitivity of radiographic detection of free air, intraperitoneal catheters were placed in 14 research dogs and 0 mL, 2.5 mL, 5 mL, or 10 mL of air was injected in the abdominal cavity. Radiographs were then taken using standard left lateral and ventrodorsal projections and lateral and ventrodorsal horizontal beam projections in a randomized order.

All films were interpreted by 3 board-certified radiologists and 3 nonspecialist clinicians who were blinded as to air volume, dog identity, and time of image acquisition

Osurnia

(florfenicol, terbinafine, betamethasone acetate)

Otic gel

For Otic Use in Dogs Only

Do not use in cats

CAUTION: Federal (USA) law restricts this drug to use by or on the order

or a licensed veternarian. BRIEF SUMMARY (for full prescribing information, see package

DESCRIPTION: OSURNIA contains 10 mg florfenicol, 10 mg terbinafine DESCRIPTION: OSURNIA contains 10 mg florfenicol, 10 mg terbinafine and 1 mg betamethasone acetate per ml. and the inactive ingredients propylene carbonate, glycerol formal, hypromellose, phospholipid, oleic acid and BHT in an off-white to slightly yellow translucent gel.

INDICATION: OSURNIA is indicated for the treatment of otitis externa in dogs associated with susceptible strains of bacteria (Staphylococcus pseudintermedius) and yeast (Malassezia pachydermatis).

DOSAGE AND ADMINISTRATION: OSURNIA should be administered in the clinic. Clean and dry the external ear canal before administering the initial dose of the product. Administer one dose (1 tube) per affected ear(s) and repeat administration in 7 daws. Do not feen the ear canal

ear(s) and repeat administration in 7 days. Do not clean the ear canal for 45 days after the initial administration to allow contact of the gel for 45 days after the initial administration to allow contact of the gel with the ear canal. Cleaning the ear may affect product effectiveness (see Effectiveness in the product insert), if alternative obt the trapiles are required it is recommended to clean the ear(s) before application. Open tube by twisting the soft tip, insert the flexible tip into the affected external ear canal(s). After application, gently massage the base of the ear to allow the gel to penetrate to the lower part of the ear canal. CONTRAINDIGATIONS: Do not use in dogs with known tympanic perforation (see Precautions in the product insert). Do not use in dogs with a burearessibility to florfeniol teripication or confidenticities.

with a hypersensitivity to florfenicol, terbinafine, or corticosteroids.

WARMINGS. Human Safety Warning:

numan safety warning.

OSURNIA may cause eye injury and irritation

Not for use in humans. Keep this and all medications out of reach of children. Consult a physician in case of accidental ingestion by humans. In case of accidental skin contact, wash area thoroughly with water Avoid contact to the eyes. In case of accidental eye contact, flush thoroughly with water for at least 15 minutes. If symptoms develop, seek medical advice.

seek medical advice.

PRECAUTIONS: Wear eye protection when administering OSURNIA and restrain the dog to minimize post-application head shaking. Reducing the potential for splatter of product will help prevent accidental eye exposure in people and dogs and help to prevent occular injury. Do not administer orally. The use of OSURNIA in dogs with perforated wimpanic membranes has not been evaluated. The integrity of the tympanic membranes have the control of the production of the product of the pr membranes has not been evaluated. The integrity of the tympanic membrane should be confirmed before administering this product. Reevaluate the dog if hearing loss or signs of vestibular dysfunction are observed during treatment. Use of topical offic corticosteroids has been associated with adrenocortical suppression and latrogenic hyperadrenocorticalm logo, (see Animal Safety in the product inset Use with caution in dogs with impaired hepatic function (see Animal Safety and Adverse Reactions in the product insert). The safe use of OSURNIA in dogs used for breeding purposes, during pregnancy, or in lactating bitches, has not been evaluated.

ADVERSE REACTIONS: The following adverse reactions were reported.

ADVERSE REACTIONS: The following adverse reactions were reported during the course of a US field study for treatment of otitis externa in dogs treated with OSUBNIA in decreasing order: elevated liver enzymes dogs treated with USUHNIA in decreasing order: elevated liver enzymes, vomiting, weight loss; 6-10% body weight and hearing loss. To report suspected adverse events, for technical assistance or to obtain a copy of the SDS, contact Dechra Veterinary Products at (866) 933-2472. For additional information about adverse drug experience reporting for animal drugs, contact FDA at 1-888-FDA-VETS or online at http://www.fda.gov/AnimalVeterinary/SafetyHealth.

POST-APPROVAL EXPERIENCE (2020): The following adverse events are broad as post-formation and the product of the control of the control

based on post-approval adverse drug experience reporting for OSURNIA. Not all adverse events are reported to FDA/CVM. It is not always possible

to reliably estimate the adverse event frequency or establish a causal relationship to product exposure using this data.

In humans, accidental exposure leading to corneal ulcers and other ocular injuries such as eye irritation, burning, stinging, and itchiness heen reported to occur when the dog shook its head after application of OSURNIA.

In dogs, the adverse events reported for OSURNIA are presented below in decreasing order of reporting frequency: Deafness, ear discharge, ear irritation and pain, vomiting, head shaking, head tilt, ataxia, vocalization, corneal ulcer, keratoconjunctivitis sicca, nystagmus, tympanic rupture,

INFORMATION FOR DOG OWNERS: Owners should be aware that adverse INFORMATION FOR DUG OWNERS SHOULD BE WARE that adverse reactions may occur following administration of OSURNIA and should observe dog for signs such as deafness, ear pain and irritation, vomiting, head shaking, head tilt, incoordination, eye pain and cutal discharge (see Animal Safety and Post-Approval Experience in the product insert). Owners should be advised to contact their veterinarian if any of the above signs are observed.

Owners should also be informed that splatter may occur if the dog

Owners should also be informed that splatter may occur if the log shakes its head following administration of OSURNIA which may lead to coular exposure. As a result, eye injuries in humans and dogs have been reported including comeal ulcers.

FEFECTIVENESS: Effectiveness was evaluated in 235 dogs with othis externa. The study was a double-masked field study with a placebo control (vehicle without the active ingredients). One hundred and fifty-nine dross were treated with OSURIAM and sevenbs-ity dross were treated with OSURIAM and sevenbs-ity dross were treated with dogs were treated with OSURNIA and seventy-six dogs were treated with the placebo control. All dogs were evaluated for safety. Treatment (1 mL) was administered to the affected ear(s) and repeated 7 days later. Prior to was administered to the affected earlys and repeated / ays later. Prior to the first administration, the earlys were cleaned with saline but not prior to the Day 7 administration. Six clinical signs associated with othis externa were evaluated: pain, erythema, exudate, swelling, odor and ulceration. Total clinical scores were assigned for a dog based on the severity of each clinical sign on Days 0, 71, 43, 00 and 45, Success was determined by clinical improvement at Day 45. The success rates of the two groups were simificantly different (no. 10,001; 62,764, of does administrated were significantly different (p=0.0094); 64.78% of dogs administered OSURNIA were successfully treated, compared to 43.42% of the dogs in

Coordinate were successfully dealed, compared to 43-24-26 or life dogs in the placebo control group. **STORAGE CONDITIONS:** CSURNIA should be stored under refrigerated conditions between 36° - 46° F (2° - 8° C). To facilitate comfort during administration, OSURNIA may be brought to room temperature and stored for up to three months.

MANUFACTURED FOR Dechra Veterinary Products 7015 College Boulevard, Suite 525 Overland Park, KS 66211 USA

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but not to the radiographic projection used. Each image was assessed as being definitive for no free air (0), uncertain (1), or having free air (2). Scores were then compared to determine the odds ratio of predicting whether air was present. Overall accuracy was higher with standard left lateral (80.4%) and lateral horizontal (78.9%) projections as compared with standard ventrodorsal (60.7%) and ventrodorsal horizontal (65.5%) beam projections. There was no significant difference in the accuracy of standard left-lateral and lateral horizontal beam projections. This suggests that horizontal beam projection did not increase the ability to detect air. The board-certified radiologists had higher overall accuracy (74.9%) than the nonspecialist clinicians (67.9%), particularly with the 5-mL and 10-mL quantities. Results suggest that standard left lateral and/or lateral horizontal beam projections provide the best opportunity for identifying small-volume pneumoperitoneum, and review by a board-certified radiologist can help increase the accuracy of diagnosis.

... TO YOUR PATIENTS

Key pearls to put into practice:

In the absence of trauma or penetrating wounds, pneumoperitoneum typically indicates GI perforation, which can lead to life-threatening

Standard left lateral and lateral horizontal beam projections are associated with the highest accuracy and odds of detecting free gas. In this study, there was no significant difference between these 2 projections, suggesting a horizontal beam does not significantly enhance accuracy compared with standard left lateral projections.

Review by a board-certified radiologist can increase the accuracy of detection of pneumoperitoneum.

References

- Frank PM. The peritoneal space. In: Thrall DE, ed. Textbook of Veterinary Diagnostic Radiology. 6th ed. Elsevier Saunders: 2013:662-663.
- Marolf A. Blaik M. Ackerman N. Watson E. Gibson N. Thompson M. Comparison of computed radiography and conventional radiography in detection of small volume pneumoperitoneum. Vet Radiol Ultrasound. 2008;49(3):227-232.