

PATIENT

Ushi Morrison

SPECIES

Canine

BREED

Mixed

SEX

NM

AGE

15 years

WEIGHT

11.2 lbs

INTERPRETED BY

Beth Johnson, DVM
DACVIM

IMAGING PERFORMED BY

Dr. Sheldon

HOSPITAL NAME

Advanced PetCare of
Oakland

REFERRING VET

Dr. Sheldon

INVOICE

11474

DATE

3/12/2026

PRESENTING CLINICAL SIGNS

- Owner has noticed that he has a decreased appetite and is lethargic. Bloodwork: Very elevated ALT 2898, ALP 5839, increased Tbil 0.5, low T4 0.9(likely non thyroidal), neutrophilia. 1 lb weight loss.

ULTRASONOGRAPHIC EXAMINATION OF THE ABDOMEN

Urinary System

The urinary bladder is adequately distended with anechoic contents. No masses, inflammatory changes, echogenic sediment or cystoliths are observed. The urinary bladder, trigone and visible pelvic urethra are normal in thickness with a smooth mucosal surface.

The prostate is unable to be well visualized in these images.

Kidneys are overall normal in size and shape with smooth peripheral margination. A normal 1:3 cortex to medulla ratio is maintained. The medulla and cortices are uniform in texture with some mild increased cortical echogenicity and mild loss of corticomedullary distinction, expected in this age patient. Trace pyelectasia noted bilaterally. There is no evidence of mineral or infarcts observed. Left kidney measures 4.68 cm, and the right kidney measures 4.04 cm.

Adrenal Glands

The right adrenal gland is normal in size (0.58 cm at cranial pole and 0.48 cm at caudal pole), shape and overall architecture, echogenicity and echotexture. Visible surrounding vasculature appears normal.

The left adrenal gland is normal in size (0.46 cm at cranial pole and 0.58 cm at caudal pole), shape and overall architecture, echogenicity and echotexture. Visible surrounding vasculature appears normal.

Spleen

The spleen is subjectively normal in size with a normal smooth capsular contour. Parenchyma is appropriately finely textured and homogenous with normal echogenicity relative to surrounding tissue (hyperechoic to liver). No focal nodules or masses are observed. Splenic vasculature appears normal.

Liver

Liver is subjectively enlarged with mildly irregular margins. Parenchyma is mildly heterogenous characterized by multiple poorly defined hypoechoic nodules within otherwise hyperechoic liver parenchyma. Visible vasculature and biliary tree appear normal without distension or congestion. In the right caudal liver, an approximately 1.8 cm x 2.4 cm anechoic density/suspect cyst is noted.

Gallbladder is mildly overdistended with a moderate amount of non-dependent, mildly aggregated/inspissated sludge. Hypo to anechoic cystic areas are noted between the gallbladder sludge and luminal wall. The wall is otherwise smooth without visible thickening. There is no evidence of cystic or CBD dilation. There is no evidence of effusion.

Gastrointestinal

The visible stomach wall is normal in thickness and layering. The lumen of the stomach is empty with no evidence of obstruction, foreign material or infiltrative disease. Pyloric outflow tract appears patent.



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The visible small intestines are normal in wall thickness and layering. Small intestinal motility appears adequate (1-3 contractions per min). The lumen of the small intestine is empty with no evidence of obstruction, foreign material or infiltrative disease.

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The visible colon is normal in wall thickness (< 0.2 cm) and layering. Contents are consistent with normal formed feces and gas.

Pancreas

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The pancreas that is observed appears appropriately isoechoic to surrounding omental fat. Visible capsule is smooth and normal in contour. Visible pancreatic parenchyma is homogenous and unremarkable. There is no visible pancreatic duct dilation. There is no evidence of active peripancreatic inflammation.

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Free Abdomen

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There is no visible free peritoneal effusion noted in these images.

There is no apparent pathologic lymphadenopathy noted in these images.

PRIMARY FINDINGS

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- Emerging gallbladder mucocele – Cholecystic debris is of unknown clinical significance. It can be seen with biliary stasis from fasting or illness. Cholecystic debris is not necessarily related to hepatobiliary disease. The non-dependent nature of this sludge combined with the cystic areas are suggestive, however, of possible emerging cystic mucosal hyperplasia or early gallbladder mucocele.
- Suspect incidental hepatic cyst.
- Otherwise, an obvious cause for the subtle liver changes is not identified in these images. Microscopic disease such as Leptospirosis, bacterial cholangiohepatitis, chronic active hepatitis, copper-associated hepatotoxicity, other hepatotoxicity, other reactive hepatopathy, infiltrative neoplasia, etc. cannot be definitively ruled out.

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SECONDARY FINDINGS

- Age related kidney changes with trace bilateral pyelectasia.

INTERPRETATION OF THE FINDINGS & FURTHER RECOMMENDATIONS

Patient's reported clinical signs and liver enzymes could be secondary to the suspected emerging gallbladder mucocele, although workup of a concurrent underlying hepatopathy is also recommended, including bile acids if patient's total bilirubin is not increased.

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Testing for leptospirosis could be considered.

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Fine needle aspirates of the liver could be considered if patient's coagulation status is appropriate.

In the meantime, in addition to supportive/symptomatic medical management of clinical signs, empirical hepatic nutraceuticals including ursodiol, empirical broad-spectrum antibiotics, etc. could be considered, but ultimately, if clinical signs don't improve and/or progress, an exploratory laparotomy for further evaluation of the gallbladder and likely cholecystectomy and liver biopsy, may be indicated after assessing patient's coagulation status.



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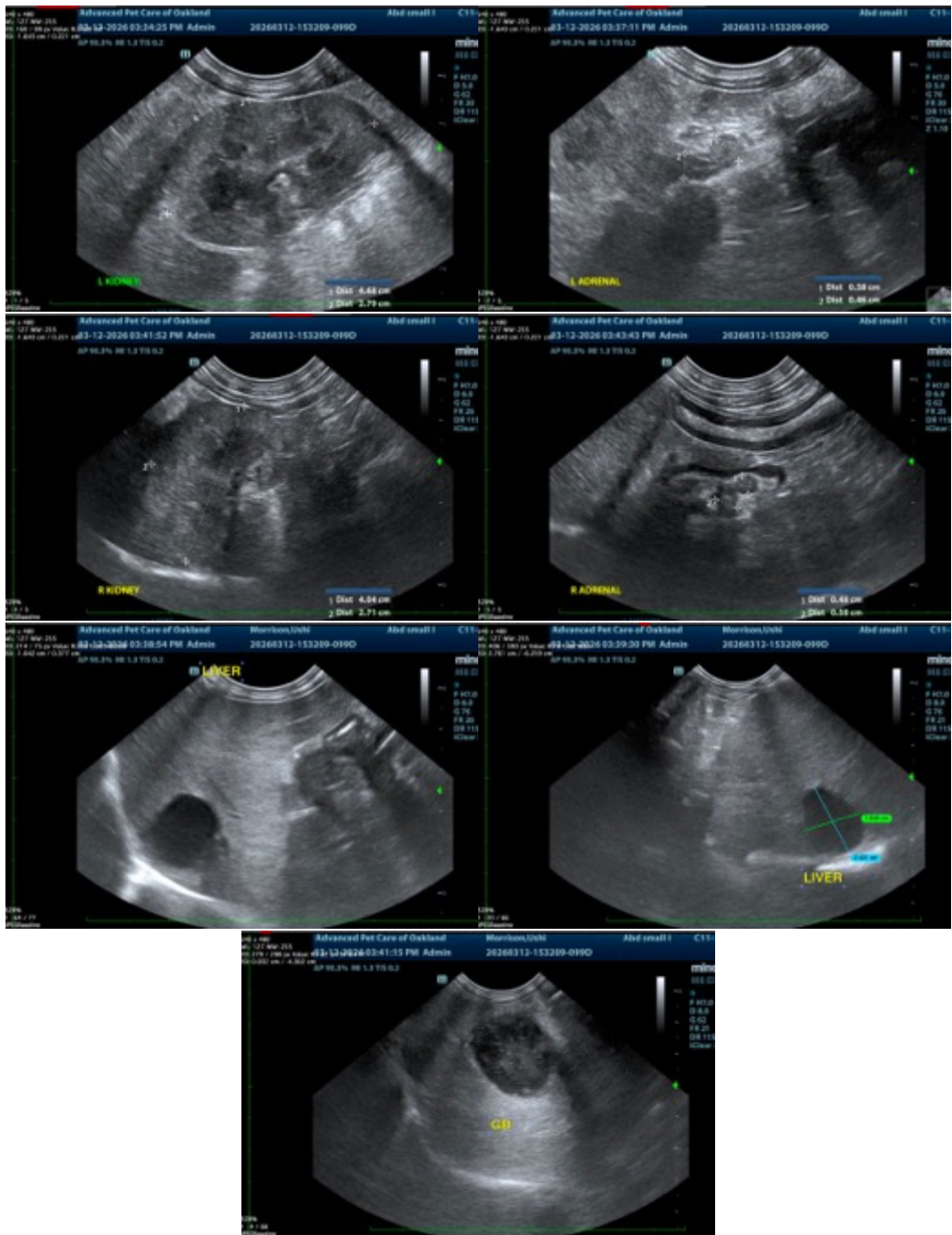
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The information and recommendations provided are based on the images presented by the referring veterinarian/sonographer. No evaluation can be communicated regarding pathology that was not visible in the image/video clips provided.

Thank you for this referral. If the clinical or image interpretation does not parallel your findings or if I can be of any further assistance please contact me.



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info@sonopath.com

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