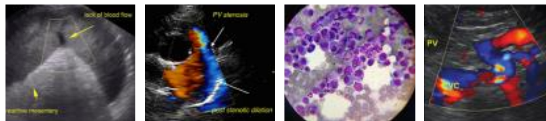

PATIENT
Bruin Barnes

SPECIES
Canine

BREED
Shih Tzu

SEX
MN

AGE
14yr

WEIGHT
18.4lb

INTERPRETED BY
Eric Lindquist, DMV DABVP, Cert. IVUSS

IMAGING PERFORMED BY
Jenna Walsh CVT

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PRESENTING CLINICAL SIGNS

Pet presented for bile acids due to elevated liver values on recent routine blood work at pet's first visit. O notes no vomiting/diarrhea/weight loss. Current Medications trazodone prior to visits. joint supplements. will receive butorphanol +/- alfaxan prior to AUS Primary Question/Differential to Be Answered in This Exam cause(s) of elevated liver values/bile acids

Abnormal PE/Chem/CBC/UA Results: Recent visit (12/7/22): high ALKP>2000 U/L, high ALT H* 657 U/L - r/o endocrine v. hepatobiliary dz v. others. high CHOL H* 412mg/dL - r/o postprandial v. endocrine v. others high GGT H* 28 U/L - r/o artifact v. cholestasis. high GLOB H* 4.6 g/dL TP H* 8.3g/dL - r/o artifact v. inflammation/infection v. others Bile Acids today: Pre-sample taken @ 8:14 am - 159.3 umol/L *H (0.0-14.9) Fed 1/4 can of Hills A/D @ 8:27 am Post-sample taken @ 10:27 am - 125.6 umol/L *H (0.0-29.9)

ULTRASONOGRAPHIC EXAMINATION OF THE ABDOMEN

Urinary System

The urinary bladder, trigone, and pelvic urethra presented normal thicknesses and normal tone to a depth of 2 cm. The ureters were not visible which is normal. No uroliths or sediment were visualized and anechoic urine was present. No evidence of inflammatory or neoplastic changes were noted. Ureteral papillae were normal.

The kidneys revealed largely normal size and structure, corticomedullary definition and ratio (cortex 1/3 of medulla) were essentially maintained with some mild to moderate age-related loss of curvilinear patterns regarding the capsule and C/M junction. A mineralizing infarct was noted in the dorsal cortex of the right kidney. Blood flow to the kidneys appeared to be mildly subnormal. Bilateral cortical mineralization was noted. The cortices presented largely uniform texture with some increased echogenicity expected for his age patient. Medullary structure differed distinctly from that of the cortex and no evidence of pelvic dilation was present. The right kidney measured 4.59 cm in length. The left kidney measured 3.91 cm in length.

Adrenal Glands

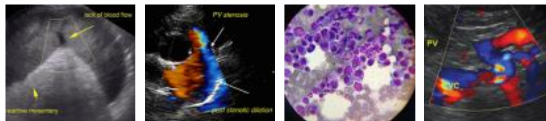
The right adrenal gland revealed an expansive irregular mass deriving from the cranial pole measuring 1.7 cm. Heterogeneous parenchymal changes were noted in the right adrenal gland. The right adrenal mass does not appear to invade into the vena cava however it does impinge against it. The left adrenal gland measured 2.21 cm in length by 0.65 cm caudal pole width by 0.43 cm cranial pole width. The right adrenal gland measured 3.27 cm in length by 0.73 cm caudal pole width.

Spleen

The spleen presented a smooth homogeneous parenchyma hyperechoic to liver and renal cortical parenchyma. The capsule was smooth without noticeable expansion or deviation from within the spleen or adjacent pathology. The splenic vasculature demonstrated normal volume without signs of congestion or thrombosis. No sonographic evidence of acute or chronic inflammatory, neoplastic, or infarctual changes were noted.

Liver

The liver was uniformly swollen with minor, excessive gallbladder debris and over distension with dependent and suspended bile without evidence of overt mucocele formation. However, excessive



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sludge was present. The liver presented coarse architecture with mildly increased portal markings and subtle, mixed echogenic changes. This is consistent with vacuolar hepatopathy and some level of remodeling and history of inflammatory component. There was no overt suspicion of neoplasia.
No pathological hepatic lymphadenopathy was evident. No overt structural evidence of inflammatory, infiltrative or regenerative pathology was evident.

Gastrointestinal

Examination of the gastrointestinal tract revealed a stomach and intestine free of stasis, of normal wall thickness, acceptable curvilinear mural detail, and peristaltic activity. Small and large intestine demonstrated normal luminal chyme and stool consistency respectively. No obstructive or overt infiltrative disease was noted. No associated abnormal lymphatic activity was noted.

Pancreas

The base and limbs of the pancreas were observed to be largely isoechoic to surrounding omental fat. Some parenchymal remodeling, however, with mild deviation from curvilinear normalcy was observed. Pancreatic duct and capsular irregularities were present consistent with age related changes. If pain upon imaging (+ Murphy sign) was present or if the patient is focally painful in subxiphoid palpation then low-grade smoldering chronic pancreatitis should be suspected.

ULTRASONOGRAPHIC FINDINGS

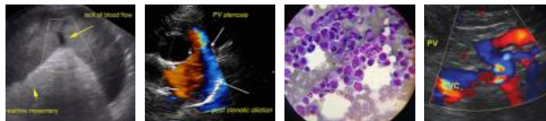
- Right adrenal mass
- Right kidney cortical infarct with bilateral mild to moderate age related renal changes
- Age related pancreatic changes
- Hepatic remodeling
- Gallbladder sludge

INTERPRETATION OF THE FINDINGS & FURTHER RECOMMENDATIONS

Strong concern for right adrenal carcinoma or pheochromocytoma. The right adrenal mass appears resectable. If USG is <1.020 then a workup for adrenal dependent Cushing's is indicated.

A FNA of the liver is indicated for further definition. No evidence of portosystemic shunting was present. Complicating factors such as dysbiosis or similar may be playing a role regarding the bile acid elevation as the liver appears to have mild to moderate remodeling. A liver oriented diet would be appropriate as well as nutraceuticals. If a right adrenalectomy is performed, a hepatic biopsy can be performed at that time.

Notes regarding Cushing's Clinical Presentations:
Nearly all Cushing's dogs have SAP elevations and true PU/PD (USG < 1.025) and most are polyphagic. Cushing's dogs are > 6 years and usually > 9 years old, usually have poor skin coats, body scores > 3/5, and are usually sedentary animals.
Its important to remember that Cushing's dogs usually look and play the part and other diseases cause false + stress related cortisol spikes. On rare occasion a Cushing's dog will not follow the rules but this is truly an exception.
Potential Cushing's patient workups can be costly and frustrating if not definitive and, in my experience, the non-definitive patient usually has something else going on that may be contributing to some of the clinical signs a Cushing's dog will have, especially SAP elevations or PU/PD. Based on this prelude of



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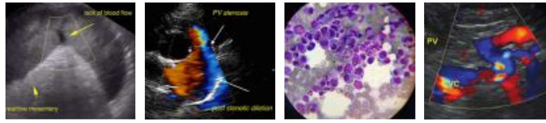
information I came up with the following algorithm in the spirit of diagnostic efficiency.
The following suggested protocol is based on current available literature on Cushing's disease and extensive clinical-sonographic experience evaluation + Cushing's and False + LDDST & ACTH stim. cases in order to maximize the efficiency of a Cushing's workup in practice.

Screen first, workup second
1) UA: Repeatable (2-3 urine samples) Urine specific gravity & urine cortisol/creatinine ratio (UCCR): If repeatable USG< 10.20 and + UCCR move to next step 2.
Note: UA is inexpensive and easy to obtain and if UA criteria is not met for Cushing's then resources can be spent into other more pertinent diagnostics or left on hold until the UA criteria is met in emerging Cushing's cases.
2) Sonogram: Does the patient have concurrent disease clinically or sonographically as non-Cushing's illness will influence the potential false + LDDST or even ACTH stim. The sonogram gives a global perspective of the internal health of the patient to be considered in the Cushing's workup as an assessment of concurrent disease. Is there a concurrent neoplastic process, UTI pancreatitis, mucocele....? Are the adrenals enlarged (Cushing's-PDH, stress, age related or breed variant), or atrophied (Iatrogenic Cushing's or adrenal burnout), have asymmetric enlargement (Adrenal tumor, hyperplasia, adenoma, age related variant), or is there vascular invasion (Invasive pheo with false + UA criteria or adenocarcinoma or phrenic thrombosis)? The sonogram answers these questions proactively. Address & treat concurrent disease first before performing Cushing's testing or testing will be artificially altered increasing false negatives and positives.
3) LDDST (0.01 D-Sodium phosphate mg/kg IV with precise dosing****) (Better screening test but plagued with false + but considered more specific than ACTH stim) Use if there is potential early Cushing's or if adrenal asymmetry present on sonogram suspecting tumor. Use LDDST in cats at a higher dose (0.1 mg/kg IV). Interpetation LDDST: Look at 8-hour post first: If > 1.4 = Cushing's. Then look at 4-hour: if > 1.4 or > 50% baseline = Cushing's. 4-hour do then 8-hour spike most consistent with PDH. Flat line high constant curve without dip more consistent with tumor but can be PDH. See attached graph.

Courtesy: Rebecca Berg DACVIM, DECVIM

4) ACTH stim. (Better confirming test but can have false +) Use if the patient "looks" Cushingoid or if bilateral adrenal enlargement is present, or high normal width on sonogram, or if iatrogenic Cushing's suspected (Cortisone Tx in past). ACTH stim is better for diagnosis of Addisons, Iatragenic Cushing's, and Cushing's therapy monitoring but problematic with initial Cushing's diagnosis. First dx LDDST is suggested.
5) If diabetic then run both LDDST & ACTH stim but stabilize as much as possible first.
5) Run a serial blood pressure in a BP friendly non "white coat effect" atmosphere. Run at least 3 at different times over a few hours or when eating as the patient tends to be calm when eating or give Torbutrol when entering the facility. Cushing's hypertension is usually 150-180 systolic range while pheochromocytoma range is more often > 180 systolic.
6) Perform CT of the pituitary to identify macro adenoma expansion if any lethargy or dullness or other central clinical CNS signs are minimally present. CT for adrenal may be more through for adrenalectomy surgical planning if ultrasound views of the CVC were problematic.
7) Adrenectomy for adrenal mass is prescribed then it is essential to stabilize the patient first regarding secondary disease such as organ dysfunction, hypertension, diabetes mellitus, hypernatremia, thromboembolic risk urinary and other infection in order to minimize potential for operative and postoperative complications as they are common in adrenalectomy. Trilostane stabilization therapy for Cushing's would be the first approach then address surgery and hypertension should be managed ideally < 160 systolic with ace inhibitors, phenoxybenzamine, or amlodipine.

Suggested reading:



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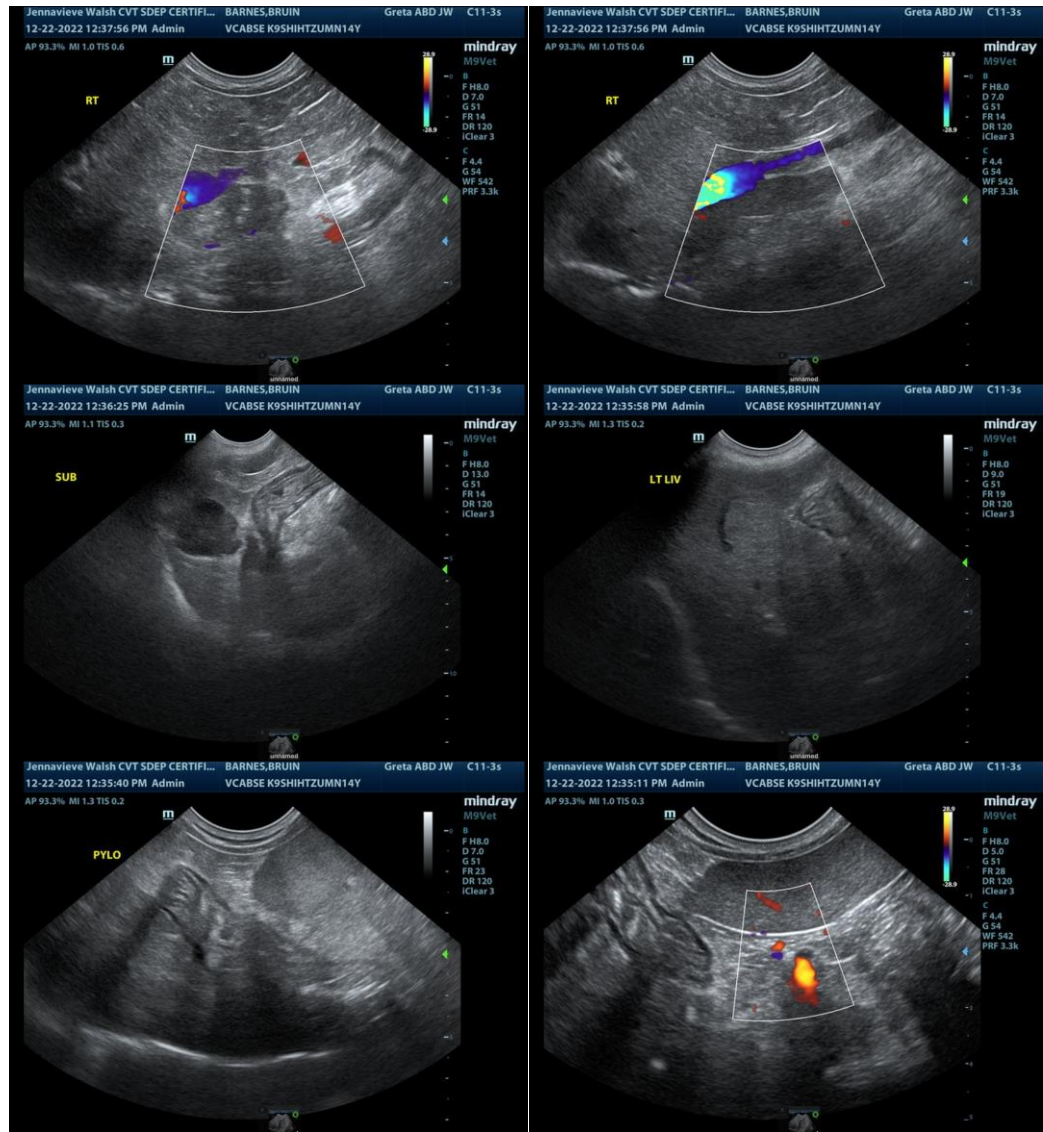
Behrend EN, Kooistra HS, Nelson R, et al. Diagnosis of Spontaneous Canine Hyperadrenocorticism: 2012 ACVIM Consensus Statement (Small Animal). J Vet Intern Med 2013;27:1292–1304 .



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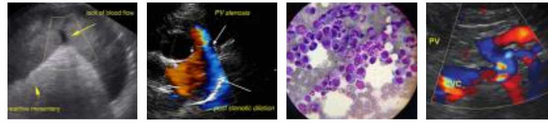

ANIMAL SOUNDS NW
MOBILE VETERINARY ULTRASOUND
IMAGING PERFORMED BY
AnimalSoundsNW.com 541-357-9423

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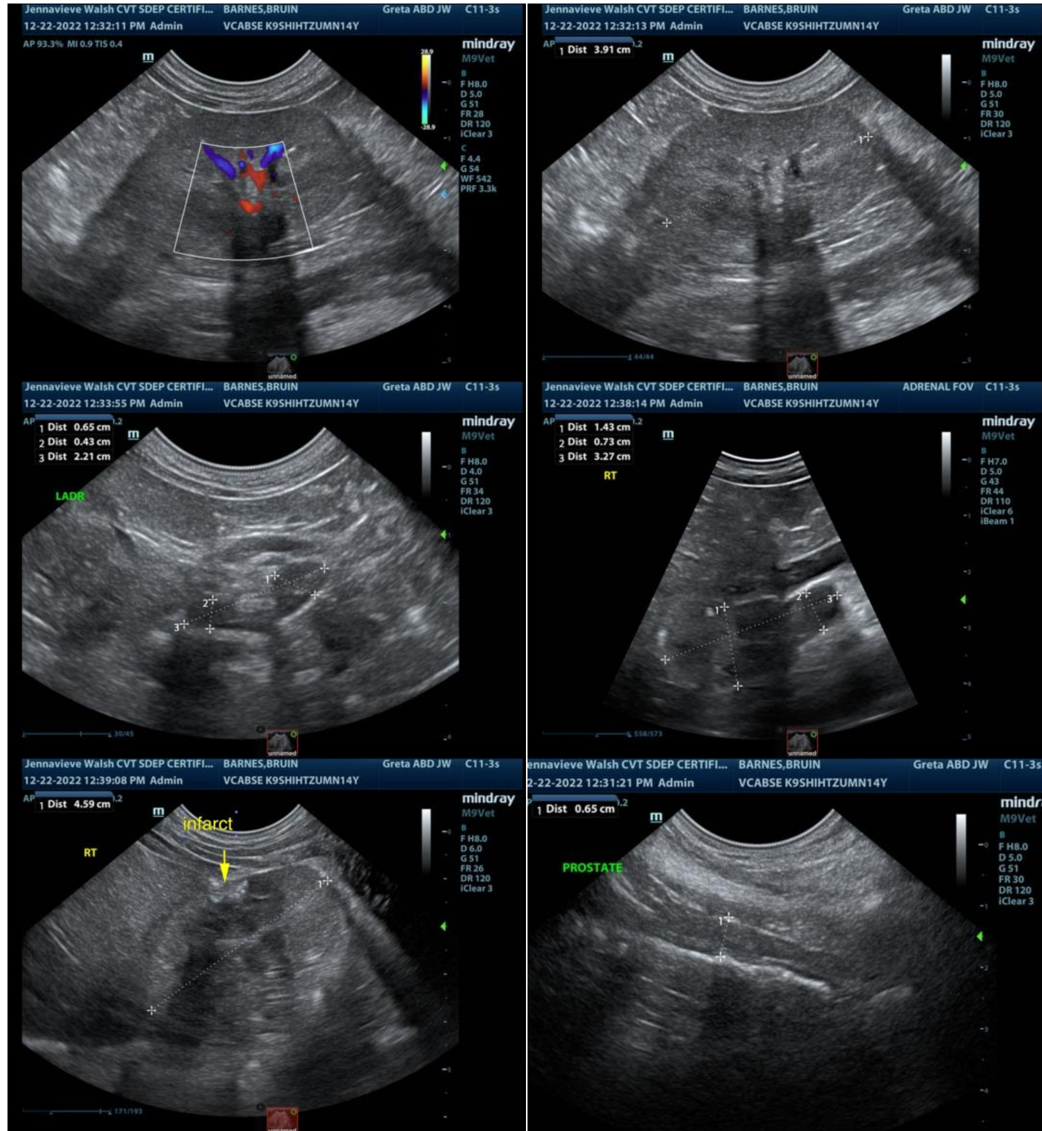


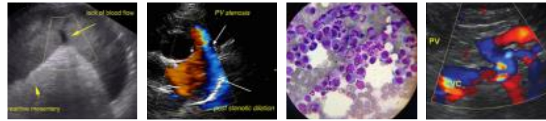
The information and recommendations provided are based on the images presented by the referring veterinarian. No evaluation can be communicated regarding pathology that was not visible in the image/video clips provided.

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.

Eric Lindquist, DMV, DABVP, Cert. IVUSS, CEO of SonoPath.com
Eric.Lindquist@SonoPath.com

Adrenal Tumors

http://www.sonopath.com/AdrenalTumor



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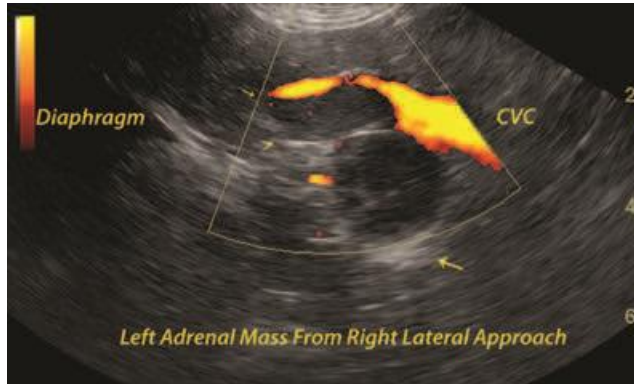


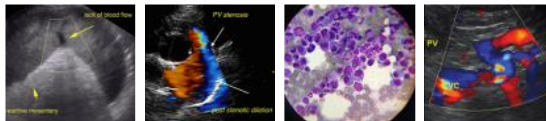
Long axis of the left adrenal gland using a right lateral approach in a dog with an adrenal adenocarcinoma (larger arrow) and caudal vena cava thrombosis. There is a complex mass lesion within the caudal pole of the adrenal gland and hyperechoic thrombus material (either tumor or mounted thrombus) within the caudal vena cava (between arrows). Note the displacement of the power Doppler signal within the caudal vena cava by the thrombus material.

Description: An adrenal mass is suspected when the maximum width of the adrenal gland exceeds 1.5 cm, there is loss of normal architecture or shape, or the shape or size between the affected adrenal gland and the contralateral gland is asymmetrical. The latter comprise the initial criteria for diagnosis; however, a bulbous enlargement of the cranial or caudal pole of the adrenal gland is common in dogs with no adrenal pathology and can be misinterpreted as an adrenal mass. If the suspected mass is not precipitating obvious signs (i.e., aggressive behavior), then an abdominal ultrasound should be repeated to confirm that the mass is a consistent finding before pursuing further diagnostics or surgery. Large breeds (Poodles, German Shepherds, Retrievers, and Terriers) and females appear to be overrepresented in the clinical reviews of adrenal tumors. Adrenal tumors in cats are rare with minimal information to characterize the disease. However, adrenal carcinoma and aldosterone producing tumors are the more common adrenal masses in our archived feline population. More specific information regarding this pathology may be found in the Feline Hyperaldosterone chapter.

Incidental adrenal lesions should be investigated clinically if discovered on ultrasound. Non-neoplastic adrenal lesions, such as cysts or granulomas, are very rare in dogs and cats, and the high incidence of metastatic lesions justifies a thorough hormonal screening as well as evaluation for non-adrenal neoplasms. Although incidental adrenal masses may appear to be nonfunctional at the time of diagnosis, it seems more likely that they are in fact subclinically functional. The diagnosis of functional adrenal tumors is discussed below; however, the identification of a nonfunctional, incidental adrenal mass creates a management dilemma.

Clinical Signs: Clinical signs attributable to adrenal tumors are dependent on hormone secretion type. Please see below.

Diagnostics: Cortical adrenal tumors, such as adenomas and adenocarcinomas, are responsible for 15-20% of hyperadrenocortical cases—what are commonly referred to as adrenal-dependent hyperadrenocortism (ADH)—in dogs. The remaining tumors are the result of pituitary-dependent secretions, which give rise to pituitary-dependent hyperadrenocorticism (PDH). PDH cases tend to demonstrate bilateral hypertrophy with excessive adrenal length and, probably more importantly, width. These enlarged adrenal glands do not invade surrounding vascular structures and are defined by overstimulation resulting from excessive ACTH secretion from the pituitary gland. Yet, ADH cases are usually unilateral (bilateral in 10-20% of cases), may invade the aorta on the left or the vena cava on the right, and metastasize to the liver and lungs most frequently. Practitioners must differentiate ADH masses from hyperplastic, non-functional, benign adrenal tumors, as well as pheochromocytomas. Thus, dynamic function tests (ex. LDDS, HDDS, ACTH stimulation, ACTH baseline, urine cortisol-creatinine ratio) are essential, as is conducting routine biochemistry (ALP is elevated in more than 90% of cases)



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and urinalysis (true polyuria/polydipsia [PU/PD] with USG < 1.020) to determine adequately the need for surgical intervention or aggressive medical therapy. It is important to assess the following: blood pressure for hypertension; oscillating hyper- and hypotensive episodes in cases of pheochromocytomas; urine protein-creatinine ratios; and serum antithrombin III to determine the risk for thromboembolism. Moreover, it is essential to evaluate the entire clinical picture and objective probabilities of possessing a true hyperadrenocorticism case. This further entails ruling out other sources of PU/PD, such as primary polydipsia, renal disease, electrolyte abnormalities, infections, and diabetes insipidus or mellitus.

Malignant or Benign, Functional or Non-Functional: How to Decide?

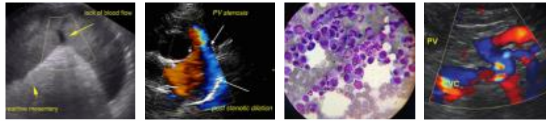
In some cases, it may be difficult to determine whether the mass is malignant or benign, functional or nonfunctional, prior to surgical removal and histopathological examination. A thorough review of the clinical signs, physical examination findings, routine blood work, urine tests, and appropriate hormonal tests should be conducted to determine the functional status of an incidental adrenal mass.

Malignancy is more often associated with larger masses. The larger the mass, the more likely metastasis has already occurred, in spite of a lack of detectable lesions on ultrasound and thoracic radiographs. Invasion of the mass into surrounding organs or blood vessels also supports malignancy, as does the detection of additional mass lesions with abdominal ultrasound and thoracic radiographs. Use of imaging modalities, such as CT and MRI, will likely provide additional data on the characteristics of specific adrenal lesions for use in diagnosis and treatment planning.

Ultrasonography is the primary instrument for assessing tumor size, aggressiveness, non-capsulated versus capsulated appearance, vascular invasion, and hepatic or other metastasis. Ideally, the patient will have fasted prior to the ultrasound; one may choose to administer an enema to enhance visibility around the ascending and descending colon. Ultrasound-guided biopsy or fine needle aspiration (FNA) may be possible on the larger masses, especially on the left side; however, adjacent vascular structures often prevent the feasibility of this procedure.

Diagnosis of the Functional Adrenal Mass:

- Cortisol-Secreting: It is very rare that a patient with hyperadrenocorticism will have a repeatable urine specific gravity greater than 1.020, so it must be determined whether the patient is truly PU/PD. If yes, then dynamic function testing is appropriate. If the patient is not truly PU/PD, then a false positive result must be considered before treatment is initiated, as the resulting hypoadrenocorticism can be life threatening. Other causes of dysuria, such as occult urinary tract infection, must then be considered. The most common functional adrenal tumor identified in dogs and cats results in hyperadrenocorticism. Approximately 15% of hyperadrenocortism cases will be caused by a functional adrenal tumor, of which 50% of these will be malignant.
    - Clinical signs can include: PU/PD; polyphagia; abdominal distention; bilaterally symmetrical truncal alopecia; delayed fur regrowth; hyperpigmentation; comedones; calcinosis cutis; excessive bruising; poor wound healing; ectopic calcification of kidneys and blood vessel walls; pyodermas; muscle weakness; exercise intolerance; hypertension; and panting.
    - Ultrasound usually reveals a small or atrophied contralateral adrenal gland as a result of suppressed pituitary ACTH secretion. Ten to twenty percent of cases have bilateral disease. Adenomas of the adrenal gland are generally less than 2 cm in diameter, and carcinomas can be any size (often they are > 2 cm). Calcification does not appear to be predictive for either adenoma or carcinoma.
    - Specific biochemical tests: Urine cortisol-creatinine ratio, ACTH stimulation test, and LDDS test.
- Catecholamine-Producing: Pheochromocytoma is a tumor derived from the chromaffin cells of the adrenal medulla; it is relatively common in dogs, but quite rare in cats. These cases should be considered malignant until proven otherwise. Invasion/entrapment/compression



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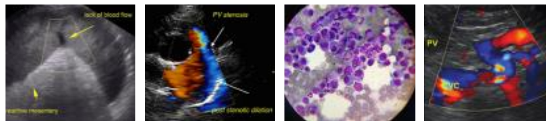
of the caudal vena cava is common. Mural invasion or luminal narrowing of the aorta, renal vessels, adrenal vessels, and hepatic veins may also occur.

- Clinical signs associated with this type of tumor are usually related to the invasion of local structures, metastases, or the secretion of catecholamines. The most common clinical signs of excess catecholamines include generalized weakness, episodic collapse, tachypnea, panting, tachycardia, and cardiac arrhythmias. Catecholamine release and hypertension tends to be episodic; thus, failure to document systemic hypertension does not rule out pheochromocytoma.
- Ultrasound: The contralateral adrenal gland is usually normal in size and shape. Pheochromocytomas do not typically calcify.
- Tests: Many of the clinical signs and blood pressure alterations are similar for pheochromocytoma and ADH. It is therefore important to rule out ADH before focusing on pheochromocytoma. The diagnosis prior to surgery is primarily one of exclusion. Specific hormonal tests, such as those that measure urinary catecholamine concentrations or their metabolites, are not routinely performed.

- Aldosterone-Secreting (rare in dogs and cats):
  - Clinical signs (Conn's Syndrome) are related to excessive secretion of aldosterone, which causes sodium retention and potassium depletion. The resulting symptoms include lethargy, weakness, mild hypernatremia, severe hypokalemia (usually < 3.0 mEq/L), and systemic hypertension.
  - Ultrasound usually reveals a normal contralateral adrenal gland.
  - Tests: Documenting increased plasma aldosterone concentrations before and after ACTH administration is a means of confirming the diagnosis. If weakness and severe hypokalemia are present, plasma aldosterone concentrations can be measured along with plasma cortisol concentrations during the ACTH stimulation test.
- Progesterone-Secreting: Although a functional tumor arising from the zona reticularis of the adrenal cortex could secrete excessive amounts of estrogen, progesterone, or testosterone, to date only progesterone-secreting adrenocortical tumors in cats have been documented.
  - Clinical signs include: diabetes mellitus and feline fragile skin syndrome, which is characterized by progressively worsening dermal and epidermal atrophy, patchy endocrine alopecia, and easily torn skin.
  - Ultrasound usually reveals a normal contralateral adrenal gland.
  - Tests: Diagnosis requires documenting an increased plasma progesterone concentration. The clinical features mimic feline hyperadrenocorticism, which is the primary differential diagnosis. Pituitary-adrenocortical axis test results are normal to suppressed in cats with progesterone-secreting adrenal tumors.
- Deoxycorticosterone-Secreting (rare):
  - Clinical signs are related to mineralocorticoid activity and include weakness, marked hypokalemia, and systemic hypertension.
  - Tests: Increased plasma deoxycorticosterone and non-detectable plasma aldosterone concentrations have been documented in dogs.
- 17-OH-progesterone-Secreting (rare):
  - Clinical signs are similar to hyperadrenocorticism.
  - Tests: Pre- and post-ACTH stimulation plasma 17-OH-progesterone concentrations will be increased.

Treatment: If hormonal tests for ADH and serum electrolytes are normal and clinical signs suggestive of pheochromocytoma are present, one can assume the adrenal mass is a pheochromocytoma and begin treatment with an alpha-adrenergic antagonist (ex. phenoxybenzamine at 0.25 mg/kg PO BID initially) for at least 2 weeks to prevent severe clinical manifestations of hypertension and promote a smooth anesthetic induction if adrenalectomy is planned. Adjustments to the dose are based on clinical response; an increase in the dose should be considered if clinical signs do not improve after 2 weeks of



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treatment. If hormonal tests for ADH and serum electrolyte concentrations are normal, clinical signs suggestive of pheochromocytoma are not present, but an adrenalectomy is nevertheless planned, one should still assume the adrenal mass is a pheochromocytoma and begin phenoxybenzamine treatment prior to adrenalectomy.

When a cortisol-producing adrenal tumor has been documented, medical therapy with trilostane (5-20mg/kg PO Q24hr) or mitotane (25-50 mg/kg PO Q24hr for 10 days, then every 4-7 days) should be considered.

The biggest dilemma is whether to perform an adrenalectomy if hormonal tests for hyperadrenocorticism and serum electrolyte concentrations are normal, and clinical signs and systemic hypertension suggestive of pheochromocytoma are not present.

An aggressive approach—adrenalectomy—is based on the assumption that the mass is malignant until proven otherwise and should be removed before metastasis has occurred. In theory, this approach would offer the best chance for long-term survival; however, the age of the patient, the size of the mass, the presence of concurrent diseases, the level of invasion into other organs, and the probability that metastases already exist should factor into the decision. Poor surgical candidates generally include: dogs compromised from the effects of hypercortisolis; older animals; animals with concurrent disease; those for whom invasion has been aggressive and surgical or post-surgical complications are likely; animals with very large masses that have likely already metastasized; and those with documented potential metastatic disease. In addition, adrenalectomy may not be indicated when the mass is small (< 3 cm diameter) and nonfunctional, and the patient is healthy. Reports suggest that there is an approximate 45% success rate of surgical resection of adrenal masses, with a positive prognosis inversely proportionate to tumor size.

In cases of concurrent hepatic nodular changes, liver biopsy samples can be obtained at surgery in cases of suspicious lesions visualized by ultrasound. Hyperadrenocorticism often causes benign nodular hyperplasia of the liver and should not be automatically interpreted as a sign of hepatic metastasis during ultrasonographic examination. Rather, suspect lesions should be confirmed and biopsied either at surgery or via ultrasound-guided FNA or core biopsy. Post-operative complications include delayed wound healing due to excessive corticoid circulation and wasting, hemorrhage, sepsis, and thromboembolism.

When surgery is a risk and a functional adrenal tumor has been documented, medical therapy, as outlined above, should be considered. Medical therapy will not impede metastatic events. An alternative approach in these cases is to determine the rate of growth of the mass by repeating abdominal ultrasounds initially at 2, 4, and 6 months. If the adrenal mass does not change in size, the time between ultrasound evaluations can be increased to every 4-6 months; however, if the adrenal mass is increasing in size, adrenalectomy should be considered.

Conclusion: Any incidentally discovered adrenal tumor warrants investigation into functionality and metastasis. The course of treatment for each case depends largely on which hormones are secreted by the adrenal tumor. Each case should be carefully evaluated on an individual basis before adrenalectomy is considered for aggressive tumors.

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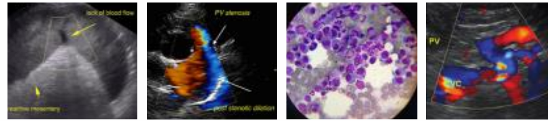


ANIMAL SOUNDS NW
MOBILE VETERINARY ULTRASOUND
IMAGING PERFORMED BY
AnimalSoundsNW.com 541-357-9423

PATIENT

Bruin Barnes

SPECIES

Canine

BREED

Shih Tzu

SEX

MN

AGE

14yr

WEIGHT

18.4lb

INTERPRETED BY

Eric Lindquist, DMV
DABVP, Cert. IVUSS

IMAGING PERFORMED BY

Jenna Walsh CVT

HOSPITAL NAME

Banfield of South Eugene

REFERRING VET

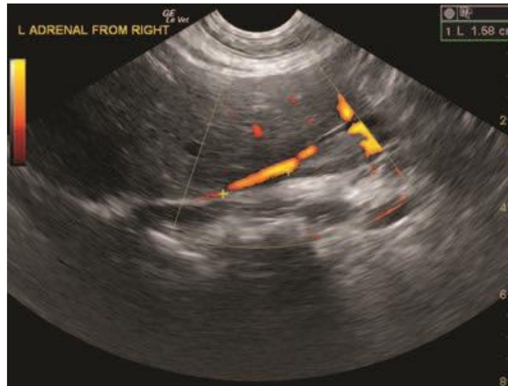
Dr. Wright

INVOICE

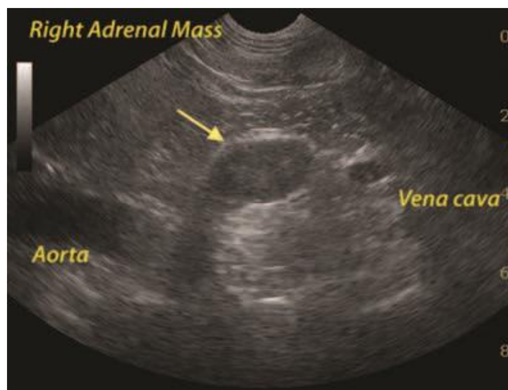
12520ag

DATE

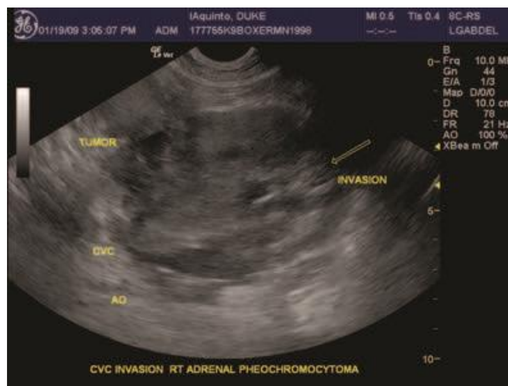
12/22/2022



Long axis of the caudal vena cava using a right lateral approach of the same dog as in the title image. Note the cranial extension of the hyperechoic thrombus material within the caudal vena cava. The liver is seen in the near field. The measurement line represents the distance from the tumor thrombus invasion to the diaphragmatic inlet



Long axis of the right adrenal gland using a right lateral approach in a dog with a surgically resectable pheochromocytoma incidentally identified during a sonogram investigating proteinuria. The adrenal gland is enlarged, rounded and hypoechoic. Note the lack of parenchymal detail with no visible corticomedullary junction as well as the echogenic capsular expansion (arrow). The patient was found to have systemic hypertension after detecting the enlarged adrenal gland sonographically.



Long axis view of the caudal vena cava (CVC) in the prior invasive adrenal mass image. Landmarks are assessed cranial to the right adrenal gland (SDEP scanning position 13 & 14), such as the aorta, in order to identify the invasive adrenal mass occupying the CVC since a normal CVC cannot be found. Color Doppler assessment of the area shows the minor caval blood flow around the invading mass. Invasive adrenal tumors, by definition, are either pheochromocytoma or adenocarcinoma and can become very large and invasive over time, but can sometimes be medically managed to maintain quality of life. A lesion such as this had likely been growing for some time but clinical signs were relatively recent prior to the sonogram.

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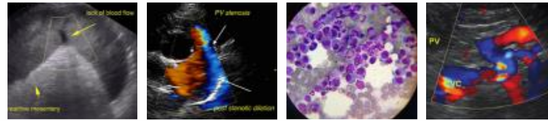

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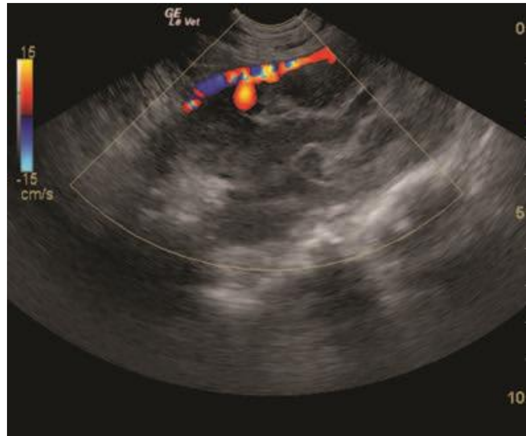

Geriatric Boxer dog presented with vague clinical signs and systemic hypertension. A large cranial abdominal mass is present when imaging the region of the vena cava and right adrenal area. A normal right adrenal could not be found. After scrupulous interrogation of the region and assessment of the regional structures, such as the aorta, the clinical sonographer can discover that the mass is likely of adrenal origin given the ominous vena caval invasion. US-guided 25 g FNA: Pheochromocytoma

References:

Behrend EN, Kooistra HS, Nelson R, et al. Diagnosis of Spontaneous Canine Hyperadrenocorticism: 2012 ACVIM Consensus Statement (Small Animal). J Vet Intern Med 2013;27:1292–1304 .

Heorauf A, Reusch C. Ultrasonographic characteristics of both adrenal glands in 15 dogs with functional adrenocortical tumors. *J Am Anim Hosp Assoc* 1999;35(3):193-99.

Herrara MA, Mehl ML, Kass PH, et al. Predictive factors and the effect of phenoxybenzamine on outcome in dogs undergoing adrenalectomy for pheochromocytoma. *J Vet Intern Med* 2008;22(6):1333-39.

Nelson, RW. Diagnostic approach to the incidental adrenal mass. World Small Animal Veterinary Association World Congress, Granada, Spain, 3-5 October, 2002.

Syme HM, Scott-Moncrioeff JC, Treadwell NG, et al. Hyperadrenocorticism associated with excessive sex hormone production by an adrenocortical tumor in two dogs. *J Am Vet Med Assoc* 2001;219(12):1725-28.

Withrow, S. Management of endocrine neoplasia. World Small Animal Veterinary Association World Congress, Vancouver, BC, 8-11 August, 2001.

von Dehn BJ, Nelson RW, Feldman EC, Griffey SM. Pheochromocytoma and hyperadrenocorticism in dogs: six cases (1982-1992). *J Am Vet Med Assoc* 1995;207(3):322-24.