



## PATIENT

Cleopatra Keefe

## SPECIES

Feline

## BREED

DSH

## SEX

Spayed Female

## AGE

13 Years

## WEIGHT

13.5 lbs

## INTERPRETED BY

Bradley Harris, DVM,  
DACVECC, DACVIM  
(cardiology)

## IMAGING PERFORMED BY

Dr. Andrea Nason

## HOSPITAL NAME

Caravan Vet

## REFERRING VET

Dr. Andrea Nason

## INVOICE

16103

## DATE

05/11/26

## PRESENTING CLINICAL SIGNS

Cleo had an echocardiogram last November due to an elevated proBNP (135). Her echo was consistent with HCM stage B1. As per recommendations, we're re-screening 6 months later to assess for progression and if Felycin CA1 is an option for this patient.

Blood pressure = 158 systolic Crea 0.9, BUN 21, USG 1.041 T4 1.6

## ULTRASONOGRAPHIC EXAMINATION OF THE HEART

FELINE CARDIAC PARAMETERS	BODY WEIGHT (kg)	HR (BPM)	IVSd (cm)	LVIDd (cm)	LVWd (cm)	FS (%)	EF (%)
NORMAL PARAMETER	-----	150-240	0.3-0.6	1.0-2.1	0.25-0.6	35-67	80-100
PATIENT	6.14	200	0.65	1.31	0.62	51	NM
FELINE CARDIAC PARAMETERS	LA/AO (M-mode)	LA/AO HEART BASE (Sisson)	LAD LA MAX 4 Chamber		LVOT VEL (m/s)	RVOT VEL (m/s)	IVRT (m/)
NORMAL PARAMETER	<1.5	1.6	0.7-1.7		<1.6	<1.3	40-60
PATIENT	NM	1.38	1.36		NM	1.0	NM
Adapted from June Boon, Veterinary Echocardiography, 1998							
Sisson D et al. JVIM 1991; 5: 232, Jacobs et al. Am J Vet Res 1985; 46:1705							

## Cardiac Presentation

The left atrium is normal in dimension. There are no distinct left atrial thrombi/clots or spontaneous echo contrast appreciated. The left ventricle is normal in dimension, with \_\_\_ hypertrophy, and no evidence of restriction. Left ventricular systolic function is normal, with adequate contractility. The right atrium and ventricle are subjectively normal in dimension and systolic function. The anterior and posterior mitral and tricuspid valve leaflets presented normal linear structure, extension in systole, and union in diastole without regurgitation. There is no evidence of systolic anterior mitral valve motion documented. The left ventricular outflow tract demonstrated normal laminar flow and subjective structural valvular integrity. The visible aorta is unremarkable. Pulmonary outflow tract assessment revealed normal valve structure, laminar flow, and appropriate diameter and distensibility. There is no evidence of semilunar valve insufficiency or pulmonary hypertension documented. There is no visible pericardial, pleural, or free peritoneal fluid noted.

## ECG

The underlying rhythm is sinus in origin with an average rate of 200bpm. There is a leftward mean electrical axis deviation. The R-R intervals are regular, with a uniform P-R interval that is within normal limits. There are rare premature complexes with a wide QRS (>40ms), consistent with a ventricular origin. There are no ventricular couplets or runs of tachycardia documented. There is no evidence of atrioventricular block or atrial ectopy documented.



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## ULTRASONOGRAPHIC FINDINGS

- These findings identify left ventricular hypertrophy in the absence of an overt outflow tract obstruction, consistent with hypertrophic cardiomyopathy (HCM). A ventricular arrhythmia is also noted. In cats, ventricular arrhythmias are usually secondary to underlying structural heart disease. Causes include cardiomyopathy (e.g., hypertrophic, restrictive, arrhythmogenic, dilated) or secondary myocardial disease (e.g., hyperthyroidism, hypertension). Rarely, ventricular arrhythmias develop secondary to extracardiac conditions (e.g., neurologic disease, metabolic disease, fever, anemia, trauma, GI disease, DIC and sepsis). The left axis deviation is reflective of the mild left ventricular hypertrophy, which is the presumed cause of the ventricular ectopy in this case.

## INTERPRETATION OF THE FINDINGS & FURTHER RECOMMENDATIONS

A systemic blood pressure and thyroid level are recommended to rule out systemic hypertension and hyperthyroidism as a cause for the left ventricular hypertrophy, respectively. If normal, then the left ventricular hypertrophy is secondary to primary hypertrophic cardiomyopathy. Given that the hypertrophy is mild and there is no left atrial enlargement, no specific treatment is recommended at this time. The clinical course for cats with HCM is incredibly variable. Results of future rechecks, especially the echocardiogram, will help us better determine the long-term prognosis. Cats with mild cardiac changes can live for years with static disease. Complications are more likely to occur in cats with advanced heart disease, and include congestive heart failure, sudden death due to arrhythmias, and thromboembolism.

Recently, the FDA authorized conditional approval of the medication Felcycin (delayed-released rapamycin) in feline patients with sub-clinical HCM which in early studies has shown to reduce left ventricular wall thickening/hypertrophy. However, extensive information regarding the use/indications, potential side effects, and monitoring is still emerging, and the drug is still not largely available (<https://www.fda.gov/animal-veterinary/cvm-updates/fda-conditionally-approves-drug-management-ventricular-hypertrophy-cats>). In the absence of liver disease and/or diabetes mellitus, introduction of this medication is reasonable (0.3 mg/kg PO once weekly). If this is performed monitoring of patient bloodwork is advised at least on a 3-6 month basis unless clinically warranted sooner.

A recheck echocardiogram, thoracic radiographs, and blood pressure are recommended in 12 months to monitor for progression, or sooner, if new clinical signs are noted. Owners should begin monitoring the resting respiratory rate. A normal respiratory rate is less than 30 breaths per minute; however, the trend in breathing rate is most important. If a progressive increase in respiratory rate is seen, then evaluation by a veterinarian is necessary.

Anesthesia considerations:

If anesthesia is necessary, then alpha-2 agonists, ketamine, high dose acepromazine, and Telazol should be avoided. Fluid therapy during anesthesia should be considered at a conservative rate (e.g., 5 ml/kg/hour) if possible (i.e., if not hypotensive). A shorter anesthetic duration will reduce the risk of complications. Pre-oxygenation is advised. Premedication with an opioid (i.e., butorphanol, hydromorphone, oxymorphone) with or without a benzodiazepine is generally the safest protocol. An induction agent such as Propofol, Alfaxalone, or diazepam/etomidate can be used to effect. Maintenance of anesthesia with isoflurane or sevoflurane is reasonable.



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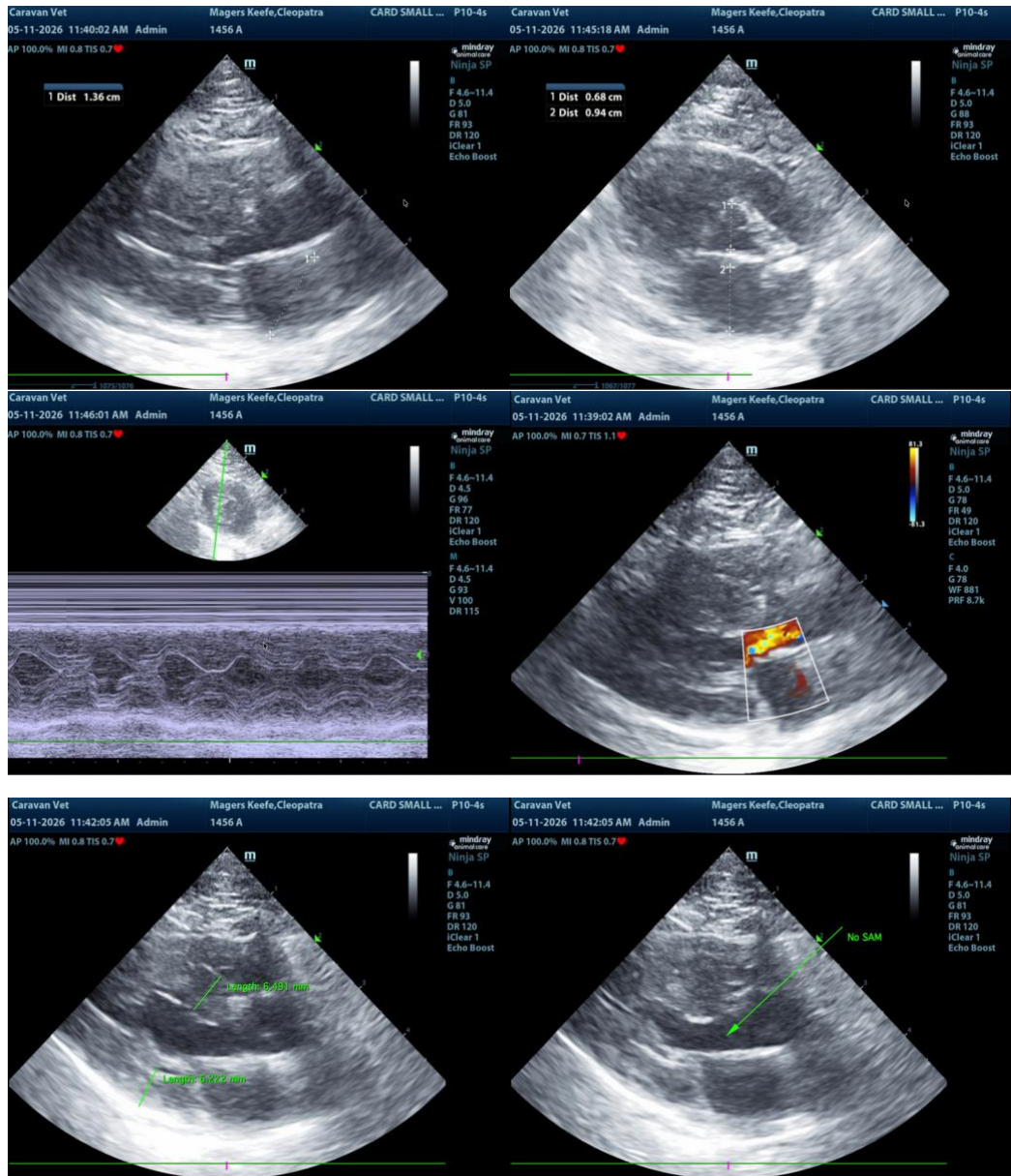
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Diet:

No special considerations are necessary. Any high-quality food from Hills, Royal Canin, Science Diet, Eukanuba, Iams, or Purina is reasonable

Activity:

No special considerations are necessary.





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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.

**Bradley Harris, DVM, DACVECC, DACVIM (cardiology)**

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