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The full details of the article are as follows:

TITLE: Plasma Cardiac Troponin I Concentration and Cardiac Death in Cats with Hypertrophic Cardiomyopathy

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JOURNAL TITLE: JOURNAL OF VETERINARY INTERNAL MEDICINE

VOLUME/EDITION: 28/6

PUBLISHER: Wiley

PUBLICATION DATE: 15 October 2014 (online)

DOI: 10.1111/jvim.12459

Plasma Cardiac Troponin I Concentration and Cardiac Death in Cats with Hypertrophic Cardiomyopathy

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Background: The use of cardiac biomarkers to assist in the diagnosis of occult and symptomatic hypertrophic cardiomyopathy (HCM) in cats has been established. There is limited data describing their prognostic utility in cats with HCM.

Hypothesis: Circulating concentrations of N-terminal B-type natriuretic peptide (NTproBNP) and cardiac troponin I (cTnI) predict cardiac death in cats with HCM.

Animals: Forty-one cats diagnosed with HCM at a veterinary teaching hospital, between February 2010 and May 2011.

Methods: Prospective investigational study. Plasma samples were collected from cats diagnosed with HCM and concentrations of NTproBNP and cTnI were analyzed at a commercial laboratory. Echocardiographic measurements from the day of blood sampling were recorded. Long-term outcome data were obtained. Associations with time to cardiac death were analyzed using Cox proportional hazards models.

Results: When controlling for the presence/absence of heart failure and echocardiographic measures of left atrial size and function, cTnI > 0.7 ng/mL was independently associated with time to cardiac death. In univariable analysis, NTproBNP > 250 pmol/L was associated with cardiac death ($P = .023$), but this did not remain significant ($P = .951$) when controlling for the effect of clinical signs or left atrial size/function.

Conclusions and Clinical Importance: Plasma concentration of cTnI (cutoff >0.7 ng/mL) is a predictor of cardiac death in cats with HCM that is independent of the presence of heart failure or left atrial dilatation.

Key words: Biomarkers; Cardiology; Cardiovascular; Monitoring; Outcome.

Hypertrophic cardiomyopathy (HCM) is the most common cardiac disease in cats.¹⁻⁶ It also affects 1 in 500 people and is the most frequent cause of cardiac death in young adults.⁷ Prognostic indicators have been identified in both humans and cats.^{4,8-10} In humans, independent predictors of death because of cardiac disease include family history of cardiac death, nonsustained ventricular tachycardia, severe hypertrophy (≥ 30 mm) of the left ventricular (LV) wall, LV outflow tract obstruction, late gadolinium enhancement on magnetic resonance imaging and younger age at diagnosis.¹¹⁻¹³ Poor prognostic indicators for cats with HCM include arterial thromboembolism (ATE), congestive heart failure (CHF), increased left atrial (LA) size, decreased LA and LV systolic function, and LV hypertrophy ≥ 9 mm.^{4,10,14,15}

The use of circulating cardiac troponin I (cTnI) and N-terminal pro-B-type natriuretic peptide (NTproBNP) concentrations to assist diagnosis, guide treatment, and screen individuals for HCM has recently received much attention in human and veterinary cardiology research.¹⁶⁻²⁵ However, little information is available on the prognostic value of circulating biomarker concentrations in HCM. A single measure-

Abbreviations:

ATE	arterial thromboembolism
CHF	congestive heart failure
cTnI	cardiac troponin I
HCM	hypertrophic cardiomyopathy
HR	heart rate
LA : Ao	ratio of left atrium to aortic root diameter (short-axis)
LAD	maximum left atrial diameter (long-axis)
LAEF	left atrial ejection fraction
LAFS	left atrial fractional shortening
LA	left atrial/atrium
LVIDd	left ventricular internal diameter in diastole
LV	left ventricular
MST	median survival time
NTproBNP	N-terminal pro-B type natriuretic peptide
RR	respiratory rate
SAM	systolic anterior motion of the mitral valve
SBP	systolic blood pressure (Doppler)
T4	total thyroxine

ment of NTproBNP, cardiac troponin, or a combination of both, act as independent predictors of adverse cardiovascular events, including death because of cardiac disease, in human patients with HCM.²⁶⁻²⁹ One study in cats with heart failure suggested that increased NTproBNP concentrations predicted survival to 1 month after diagnosis of CHF,^a while another recent study suggested that troponins I and T are associated with cardiac death in pedigree cats with HCM.³⁰

The primary aim of this study was to determine if a single measurement of circulating NTproBNP, cTnI, or both, in cats of any breed diagnosed with HCM could provide useful prognostic information. We hypothesized that increases in both NTproBNP and

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Submitted January 12, 2014; Revised July 28, 2014; Accepted August 21, 2014.

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DOI: 10.1111/jvim.12459

cTnI would be significantly associated with reduced time to cardiac death.

Materials and Methods

Animals

Cats presented to a veterinary teaching hospital^b for investigation of heart disease from February 2010 to May 2011 were eligible for prospective enrollment. The inclusion criteria were a diagnosis of HCM on 2-dimensional (2D) echocardiography and a blood sample taken within 24 hours of presentation, from which residual EDTA plasma was available. Echocardiography was performed by a Diplomate in Cardiology or Resident under supervision. Exclusion criteria were any diagnosis other than HCM; chronic kidney disease (IRIS Stage >II) or other significant systemic disease; cats ≥ 8 years old without a recorded blood pressure or where total thyroxine (T4) concentration was unavailable. In addition, cats with other possible causes of LV hypertrophy such as hyperthyroidism, hypertension, or clinical evidence of hypovolemia (“pseudohypertrophy”) were excluded.

Clinical Data

At presentation age, sex, breed, heart rate (HR), respiratory rate (RR), weight, and systolic blood pressure (SBP) were recorded. Echocardiographic measurements were performed at the time of presentation to the clinic and later reviewed by a single cardiologist (DJC). A maximum diastolic LV free wall or interventricular septum measurement of ≥ 6 mm was considered diagnostic for HCM.^{1,6} The ratio of LA diameter to aortic root (LA : Ao) was measured from a 2D short axis view at the heart base, optimized for the LA and aortic valve, in the frame before aortic valve opening (end-ventricular diastole) using an inner edge to inner edge technique.³¹ Maximum LA diameter (LAD) was measured from the 2D right parasternal long-axis view (RPLA), in the frame before mitral valve opening (end-ventricular systole).³² LV internal diameter in diastole (LVIDd) was measured in 2D from the RPLA, in the first frame in which the mitral valve was closed. M-mode measurements LV fractional shortening (LVFS) and LA fractional shortening (LAFS) were made using a leading edge to leading edge method.^{10,33,34} LA ejection fraction (LAEF) was measured in the 2D RPLA, optimized for the LA, using a modified Simpson’s method.^{10,33,34} Transmitral flow velocities (E and A waves) were recorded from pulsed wave Doppler interrogation of diastolic mitral inflow in the left-apical 4-chamber view.³⁵ Waves were considered summated at an E-at-A intersection velocity >0.25 m/s. The presence of mitral systolic cranial motion, regional LV wall hypokinesia, and restrictive mitral inflow pattern (E : A ratio > 2 , nonsummed³⁶) were recorded as categorical variables (yes/no). Regional hypokinesia was defined as localized subjective lack of systolic excursion with thinning of a segment of LV wall.¹⁰

Congestive heart failure was defined as present if the cat had radiographic or ultrasonographic evidence of cardiogenic pulmonary edema or pleural effusion in the presence of LA dilatation (LA : Ao ≥ 1.5 ,³¹ LAD ≥ 16 mm,³² or both), or tachypnea responsive to furosemide in the presence of LA dilatation (LA : Ao ≥ 1.5 ,³¹ LAD ≥ 16 mm,³² or both). If another potential cause of pleural effusion (eg, intrathoracic neoplasia) was present, the cat was excluded from the study.

Cardiac Biomarkers

Residual EDTA plasma samples were obtained after jugular venipuncture for clinical purposes. Samples were processed within

20 minutes of collection. Plasma was separated by centrifugation at $800 \times g$ for 3 minutes, immediately before storage at -80°C . After transport on dry ice to a commercial veterinary laboratory,^c batch analysis of NTproBNP and cTnI (using a high sensitivity assay^d) was performed. The lower limit of detection was 24 pmol/L for the NTproBNP assay³⁷ and 0.01 ng/mL for the cTnI assay. For the purposes of statistical analysis, results below these limits were recorded as 24 pmol/L and 0.01 ng/mL respectively.

Outcome

The end of the study period was September 1, 2013. Owners of cats were contacted annually, according to a protocol approved by the institutional ethics committee. Owners were sent a letter and questionnaire, followed by a telephone call if no response was received within 30 days. Primary veterinarians were contacted regarding the circumstances of death if euthanasia was performed. The end-point of the study was death, categorized as death because of cardiac disease or noncardiac causes. Cardiac causes included death or euthanasia because of clinical signs of CHF or ATE, or sudden death unrelated to known systemic disease or trauma. Noncardiac causes were death or euthanasia after clinical signs not attributable to heart disease (eg, vomiting, diarrhea, trauma, or diagnosed neoplasia).

Statistical Analysis

Statistical analysis was performed using commercially available software.^e For continuous data, normality was assessed graphically and using a Shapiro–Wilk test. Normally distributed continuous data (represented mean \pm SD) were compared using an independent samples *t*-test and nonnormally distributed data (represented median and range) were compared using a Mann–Whitney *U*-test. To look for linear correlation between two continuous variables, Spearman’s rank correlation (ρ_s) was used.

The end-point for survival analysis was death because of cardiac disease. Animals dying of noncardiac disease or alive at the end of the study period were right-censored for survival analysis. Univariable time-to-event models were constructed using Cox proportional hazards analysis. Continuous variables were initially analyzed as quintiles (cTnI) or quartiles (all others) using Kaplan–Meier curves and pairwise Log-rank tests. Variables where hazard increased ordinally were eligible for inclusion in time-to-event analysis as continuous variables. Where one or more groups had disproportionately different hazards to the others, these variables were presumed to exhibit a threshold effect, and were included in Cox models as categorical variables using a cut-off established by the exceptional quartile/quintile.

Multivariable analysis was performed in a forwards stepwise manner, to investigate the association of cardiac biomarkers with time to death because of cardiac disease. Variables significant at $P < .2$ at the univariable level were carried forward to multivariable Cox analysis. The assumptions of Cox proportional hazards were tested using log cumulative hazard plots and Schoenfeld residuals, and overall model fit was evaluated using Cox Snells. Where variables showing statistically significant correlation were included in a model, a variance inflation factor (VIF) were calculated for each pair of correlated variables. Colinearity was considered insignificant for the purposes of Cox multivariable analysis if the VIF was <10 . Akaike’s information criteria were calculated for each of the final multivariable Cox models to assess ability of the overall model to explain difference in time to cardiac death. Statistical significance was set at $P < .05$.

Results

Population

Fifty-nine cats were eligible for inclusion. Eighteen were excluded because of hyperthyroidism (2), non-HCM cardiac disease (8), or no diagnosis of heart disease (8), leaving 41 cats that met the inclusion criteria. Population characteristics, clinical data and echocardiographic measurements are summarized in Table 1.

At the end of the study period, 24/41 cats (59%) had died, 17 (41%) were alive and none were lost to follow-up. Death was considered to be because of cardiac disease in 21/24 cats (ie, 51% of the entire study population reached the primary end-point). Cardiac causes of death were euthanasia because of CHF (14), euthanasia because of clinical signs of ATE (2), sudden death associated with dyspnea (3) and sudden death after clinical signs of limb ATE (2). Noncardiac causes of death were euthanasia because of advanced chronic kidney disease (2) and mediastinal lymphoma (1).

Associations between Biomarkers and Clinical/Echocardiographic Findings

There was significant but weak correlation between cTnI and LA : Ao (ρ_s 0.50), LAD (ρ_s 0.42), LAFS (ρ_s -0.49), LAEF (ρ_s -0.48), LVFS (ρ_s -0.47), weight (ρ_s 0.44), and NTproBNP (ρ_s 0.46). VIF for each of these

significant correlations was <10 (1.10–7.58), suggesting that colinearity was not great enough to impact upon multivariable Cox proportional hazards analysis. There was no correlation between age, HR, RR, maximum LV thickness, or LVIDd. cTnI (but not NTproBNP) concentrations were significantly higher in cats with regional LV wall hypokinesis than those without (0.99 ng/mL, 0.3–4.56 ng/mL; versus 0.2 ng/mL, 0.01–3.8 ng/mL; $P = .027$). All 6 cats with regional hypokinesis had the inferior LV free wall affected.

NTproBNP was significantly correlated with LA : Ao (ρ_s 0.63), LAD (ρ_s 0.57), LAFS (ρ_s -0.77), LAEF (ρ_s -0.69), LVFS (ρ_s -0.42), and SBP (ρ_s -0.47). VIF for each of these significant correlations was <10 (1.64–5.73), suggesting that colinearity was not great enough to impact upon multivariable Cox proportional hazards analysis. There was no correlation between age, weight, HR, RR, maximum LV thickness, or LVIDd. Cats with CHF had significantly higher median NTproBNP concentration than those without (569 pmol/L, 126–1,451 pmol/L; versus 249 pmol/L, 24–786 pmol/L; $P = .016$).

Among echocardiographic variables, significant correlation was present between LA : Ao and LAD (ρ_s 0.87), LAEF (ρ_s 0.81), LAFS (ρ_s 0.85), and LVFS (ρ_s 0.60). In addition, LAD was correlated with LAEF (ρ_s -0.74), LAFS (ρ_s -0.75), and LVFS (ρ_s -0.58); LAFS with LAEF (ρ_s 0.84) and LVFS (ρ_s 0.54); and

Table 1. Population characteristics, clinical data, and echocardiographic findings of the 41 cats enrolled in the study.

Parameter	Alive or Noncardiac Death	Cardiac Death	P-Value
Number of cats	20	21	–
Follow-up period (days)	1,050 (162–1,274)	241 (2–946)	<.001*
Age (years)	6.2 (0.96–16.4)	6.9 (0.72–16.1)	.873
Male, n (%)	12/20 (60)	16/21 (76)	.244
Purebred, n (%)	4/20 (20)	8/21 (38.1)	.177
Body weight (kg)	4.2 (2.6–5.6)	4.43 (2.77–6.34)	.589
Heart rate (per minute)	184 ± 36	180 ± 35	.666
Respiratory rate (per minute)	39 ± 14	46 ± 16	.259
CHF confirmed, n (%)	5/20 (25)	19/21 (90.5)	<.001*
Systolic blood pressure (mmHg)	140 ± 22	121 ± 20	.013*
NTproBNP (pmol/L)	266 (24–786)	573 (24–1,451)	.008*
cTnI (ng/mL)	0.11 (0.01–0.73)	0.27 (0.05–4.56)	.527
LA : Ao ratio	1.52 ± 0.34	2.51 ± 0.76	<.001*
LA maximum diameter (mm)	16.2 ± 3.98	23.7 ± 3.61	<.001*
LA fractional shortening (%)	29 (7.69–46.2)	8.54 (2.94–31.3)	<.001*
LA ejection fraction (%)	53.5 (1.18–67.6)	16.4 (5.13–60.1)	<.001*
LV fractional shortening (%)	54.5 ± 12.9	36.3 ± 13.1	<.001*
IVSd (mm)	6.70 ± 1.37	5.84 ± 2.00	.198
LVFWd (mm)	7.13 ± 1.60	6.91 ± 2.50	.794
LV maximum thickness (mm)	7.74 ± 1.14	7.69 ± 1.35	.915
SAM present, n (%)	13/20 (65)	8/21 (38.1)	.121
Restrictive mitral inflow present, n (%)	3/17 (17.6)	7/10 (70)	.013*
Regional LV hypokinesis present, n (%)	0/20 (0)	6/21 (28.6)	.021*

CHF, congestive heart failure; NTproBNP, N-terminal of pro-B-type natriuretic peptide; cTnI, cardiac troponin I; LA : Ao, left atrium to aortic root ratio; LA, left atrial/atrium; LV, left ventricular; IVSd, interventricular septum measured at end-diastole; LVFWd, left ventricular free wall measured at end-diastole; SAM, systolic anterior motion of the mitral valve.

*Indicates variables significantly different between cats suffering a cardiac death and those that died a noncardiac death or alive at the end of the study period. Normally distributed data are represented as mean ± SD, and nonnormally distributed data are represented as median (range).

LAEF with LVFS (ρ_s 0.62, $P < .05$ for each pairwise correlation listed). SBP was correlated with LAFS (ρ_s 0.57). These significant interactions were considered when performing multivariable Cox proportional hazards analysis.

Survival Analysis

Median survival time (MST) to cardiac death for the whole population was >946 days. Median follow-up period was 834 days (2–1,274 days). Cats with cTnI >0.7 ng/mL had a significantly shorter MST (40 days, 95% CI 0–168 days; versus >1,274 days, $P = .001$; Fig 1). Cats with NTproBNP > 250 pmol/L also had a significantly shorter MST than those with NTproBNP \leq 250 pmol/L (764 days, 95% CI 159–1,369 days; versus >1,257 days, $P = .005$; Fig 1). A combined measurement of NTproBNP > 250 pmol/L and cTnI > 0.7 ng/mL did not yield additional information regarding survival time when compared to measurement of cTnI > 0.7 ng/mL alone.

The results of univariable Cox proportional hazards analysis are shown in Table 2. Two models were devised to evaluate the use of NTproBNP and cTnI as predictors of time to cardiac death in multivariable analysis. First, a model employing cardiac biomarkers where the presence/absence of CHF is confirmed, but echocardiography is unavailable. Second, a multivariable model was constructed where all data were eligible for inclusion (biomarkers, CHF present/absent, and echocardiographic measurements).

In the first model, when controlling for the confirmed presence/absence of CHF, NTproBNP did not provide additional prognostic value. However, cTnI

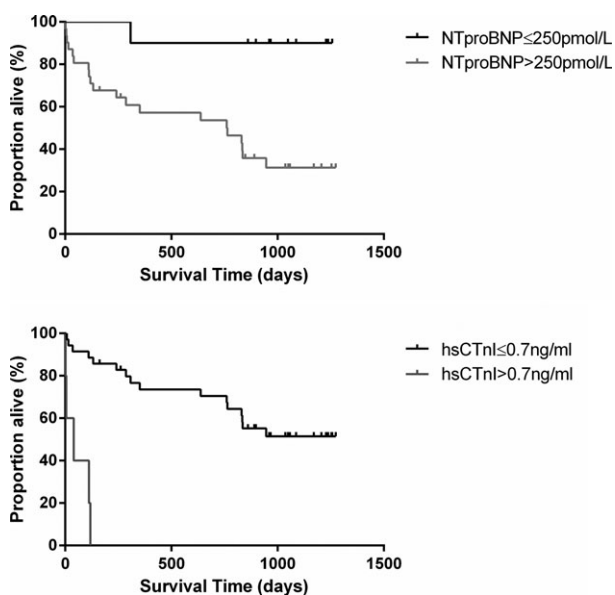


Fig 1. Kaplan–Meier curves to show the effect on survival of cardiac biomarker concentrations at the univariable level. NTproBNP, N-terminal pro-B type natriuretic peptide; cTnI, cardiac troponin I.

Table 2. Variables associated ($P < .2$) with a shorter time to cardiac death in univariable Cox proportional hazards analysis.

Parameter	HR	95% CI	P-Value
Age			
≥3.53 years	Ref		
<3.53 years	4.82	1.12–20.8	.035
LA : Ao			
≤1.76	Ref.		
>1.76	12.3	3.56–42.4	<.001
LA maximum diameter			
≤16 mm	Ref		
>16 mm	16.3	2.18–122.2	.007
LA fractional shortening			
≥14.2%	Ref		
<14.2%	7.8	2.58–23.5	<.001
LA ejection fraction			
≥33.5%	Ref		
<33.5%	12.3	3.56–42.4	<.001
LV fractional shortening			
≥47%	Ref		
<47%	5.49	1.98–15.2	.001
NTproBNP			
≤250 pmol/L	Ref		
>250 pmol/L	10.4	1.39–77.6	.023
cTnI			
≤0.7 ng/mL	Ref		
>0.7 ng/mL	4.77	1.68–13.5	.003
Systolic BP			
≥130 mmHg	Ref		
<130 mmHg	3.08	1.13–8.44	.028
Regional LV hypokinesis			
No	Ref		
Yes	35.9	6.69–192.5	<.001
Nonrestrictive mitral inflow	Ref		
Restrictive mitral inflow present	6.92	1.76–27.2	.006
Congestive heart failure			
No	Ref		
Yes	15	3.4–66.4	<.001

HR, hazard ratio; 95% CI, 95% confidence intervals; LA : Ao, left atrium to aortic root ratio; LA, left atrial; LV, left ventricular; NTproBNP, N-terminal of pro-B-type natriuretic peptide; cTnI, cardiac troponin I.

provided additional, independent predictive value of a significantly shorter time to death because of cardiac disease (Table 3). Other measured variables were not significant.

In the second model, NTproBNP did not remain predictive of shorter time to death because of cardiac disease in any multivariable model featuring confirmed CHF status and echocardiographic measurement of LA size or function. Conversely, cTnI did provide additional prognostic information even when accounting for the effect of LA size, LA function, or LV systolic function. However, if echocardiographic evidence of regional LV hypokinesis was detected, cTnI lost significance, suggesting some interrelationship. Detailed evaluation of the echocardiographic data resulted in the construction of 2 alternate final models (Table 3). In one, CHF status, LA : Ao > 1.76 and cTnI > 0.7 ng/mL were independent predictors of time

Table 3. Results of Cox multivariable analyzes, performed to reflect each of two clinical scenarios. In each, cTnI provided independent predictive value for cardiac death in cats with HCM. The exception was where regional LV hypokinesia was detected by 2D echocardiography.

Multivariable Model	Variable	HR	95% CI	P-Value	AIC value
Model 1: CHF status established, echocardiography unavailable	CHF (yes)	17.4	3.84–78.6	<.001	114.2
	cTnI (>0.7 ng/mL)	6.78	1.98–23.2	.002	
Model 2.1: CHF status established, echocardiography available	CHF (yes)	6.35	1.10–36.7	.039	111.0
	cTnI (>0.7 ng/mL)	5.61	1.61–19.5	.007	
	LA : Ao (>1.76)	4.53	1.04–19.7	.044	
Model 2.2: CHF status established, echocardiography available	CHF (yes)	11.9	2.60–54.4	.001	104.0
	Regional LV hypokinesia (yes)	16.3	2.71–97.6	.002	

HCM, hypertrophic cardiomyopathy; HR, hazard ratio; 95% CI, 95% confidence intervals; AIC, Akaike's information criteria; NTproBNP, N-terminal of pro-B-type natriuretic peptide; cTnI, cardiac troponin I; CHF, congestive heart failure; LA : Ao, left atrium to aortic root ratio; LV, left ventricular.

to cardiac death. In the other, CHF status and presence of regional LV hypokinesia were independent predictors of time to death because of cardiac disease.

Discussion

This prospective study demonstrates the long-term, independent prognostic value of a single measurement of cTnI in cats of any breed with HCM. One recent study by demonstrated that pedigree cats suffering an HCM-related death had higher cTnI at initial diagnosis than survivors.³⁰ In our data, not only was cTnI > 0.7 ng/mL associated with a shorter survival time to cardiac death but also it had predictive value in addition to the presence of CHF and LA dilatation. In our study, higher NTproBNP was not an independent predictor of cardiac death, but both increased NTproBNP and cTnI concentrations were significantly associated with a worse prognosis in univariable analysis (cutoff values >250 pmol/L and >0.7 ng/mL respectively). A combination of both biomarkers did not provide additional prognostic information compared to measurement of cTnI > 0.7 ng/mL alone.

The presence of CHF and LA dilatation are widely acknowledged indicators of risk of cardiac death in cats with HCM.^{4,14,15,38} Echocardiographic measures of poor LA function, extreme LV hypertrophy, and reduced LV systolic function were independent predictors of shorter time to cardiac death in a large population of cats with HCM.¹⁰ In our univariable analysis, CHF status (present/absent), measures of LA size and function, LV systolic function, the presence of restrictive mitral inflow, and regional LV hypokinesia were significantly associated with time to death because of cardiac disease (as previously reported¹⁰).

In multivariable analysis, to investigate whether cardiac biomarker measurement could provide useful additional prognostic information on survival when the effects of CHF status and echocardiographic measurements were accounted for, we derived two alternate models to reflect different clinical situations, where the clinician either has or does not have access

to echocardiographic information (for example, in many general practice clinics). In the first model, multivariable analysis evaluated the prognostic value of biomarkers where CHF was already confirmed as present or absent. In this model, NTproBNP did not remain significant, suggesting that CHF status was a more powerful prognostic predictor. High sensitivity cTnI remained significant; however, likely reflecting its ability to detect myocardial cellular damage.³⁹ Although the ability of cardiac biomarkers to discriminate between cardiac and respiratory causes of tachypnea in cats is well-established,^{17–19,23,40,41} our results show the additional prognostic value of measuring cTnI in cats with HCM and CHF.

In the second model, CHF status and echocardiographic data were included in the multivariable analysis. The model that best accounted for differences in MST included the presence of CHF, LA : Ao and cTnI. If LV regional hypokinesia identified by 2D echocardiography was included in the model, cTnI lost significance. This suggests a close association between LV regional hypokinesia and increased cTnI, similar to elevations in cardiac troponins reported in humans with ischemic myocardial damage.^{42,43} Regional LV hypokinesia is predictive of a shorter survival time in cats with HCM¹⁰ and our results suggest that indicators of myocardial damage (whether increased cTnI concentration or regional LV hypokinesia) provide additional prognostic information independent of the presence of CHF or LA dilatation.

The prognostic value of cTnI identified in our study is similar to reports in humans, and recently cats, with HCM.^{26,27,30} Although our study enrolled a relatively small number of cats, its prospective nature and high event rate add to the statistical power.⁴⁴ Similar studies in humans with HCM, although enrolling a greater number of subjects, have reported a notably lower event rate.^{26,27}

Possible misclassification of cause of death (cardiac/noncardiac) or CHF status is a limitation of this study. Classification of death because of cardiac disease was likely to be relatively accurate for cats euthanized with CHF or ATE by their primary veterinarian,

but may have been less accurate when based on owner reports for cats that died at home. Further limitations include a potential bias of owners or veterinarians in the decision making process surrounding euthanasia, lack of standardization in treatment, and the exclusion of cats with comorbidities that may elevate biomarker concentrations. Although this latter point may limit the applicability of these results to a more heterogeneous feline population in clinical practice, it should make these findings more accurate when considering cats with HCM.

In conclusion, the key finding from this study is that, although NTproBNP did not remain an independent predictor of death because of cardiac disease, a circulating concentration of cTnI >0.7 ng/mL was a significant predictor of cardiac death in cats with HCM that is independent of the presence of heart failure or left atrial dilatation.

Footnotes

^a Fox PR, Ettinger SJ, Lamb KE, et al. Evaluation of NTpro-brain natriuretic peptide (NTproBNP) as a prognostic indicators of short-term outcome in cats with heart failure. *J Vet Int Med* 2013;27:637 (abstract)

^b Queen Mother Hospital for Animals, Royal Veterinary College, Hertfordshire, UK

^c IDEXX Veterinary Laboratories, Wetherby, Yorkshire, UK

^d AccuTnI, a paramagnetic particle chemiluminescent immunoassay from Access Immunoassay Systems (Ref. A78803), Beckman Coulter, High Wycombe, UK

^e IBM SPSS 20 for Windows 7, IBM (UK) Ltd, Portsmouth, UK; GraphPad Prism 6 for Windows 7, GraphPad Software Inc., San Diego, CA

Acknowledgments

The authors thank IDEXX Ltd. for their support of this project.

Conflict of Interest Declaration: This study was not supported by any grant or other source of direct funding. IDEXX Ltd. performed cardiac biomarker assay free of charge, but had no involvement in the preparation of this manuscript. Mr Borgeat and Drs Luis Fuentes and David Connolly have performed consultancy work for Boehringer Ingelheim. Drs Jessie Payne and Luis Fuentes have received research funding from IDEXX Ltd. for previous studies.

Off-label Antimicrobial Declaration: The authors declare no off-label use of antimicrobials.

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