Speckle tracking echocardiography in cats with preclinical hypertrophic cardiomyopathy

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**Abstract**

**Background:** Cats with hypertrophic cardiomyopathy (HCM) have decreased left ventricular (LV) longitudinal deformation detected by mitral annular plane systolic excursion (MAPSE) and speckle tracking echocardiography. People with preclinical HCM have decreased systolic LV longitudinal and radial strain (S) and strain rate (SR), with preserved circumferential S and SR.

**Hypothesis/Objectives:** Cats with preclinical HCM have decreased systolic LV deformation compared to normal cats.

**Animals:** Seventy-three client-owned cats with (n = 37) and without (n = 36) preclinical HCM.

**Methods:** Retrospective echocardiographic study. Left and right ventricular longitudinal and radial strain and strain rate, LV radial and circumferential strain and strain rate were calculated by STE. Left ventricular mass was also calculated. Correlation between STE variables and LV hypertrophy was determined and receiver-operating characteristic (ROC) curves were plotted for prediction of HCM.

**Results:** Cats with HCM had smaller absolute longitudinal S (−14.8 ± 3.3% vs −19.7 ± 2.7%, P < .001), longitudinal SR (−2.36 ± 0.62 vs −2.95 ± 0.68 second⁻¹, P < .001), radial S (46.2 ± 21.3% vs 66.7 ± 17.6%, P < .001), and radial SR (5.60 ± 2.08 vs 6.67 ± 1.8 second⁻¹, P < .001) compared to healthy controls. No difference was observed for circumferential S and SR. Cats with HCM had greater LV mass (13.2 ± 3.7 g vs 8.6 ± 2.7 g, P < .001). The ROC with the greatest area under the curve (AUC) for the identification of HCM (0.974) was plotted from a logistic regression equation combining LV mass, MAPSE at the free wall, and LV internal diameter in diastole (LVIDd).

**Conclusions and clinical importance:** Cats with preclinical HCM have decreased long axis and radial deformation. Decreased longitudinal deformation and decreased LVIDd are factors that would support a diagnosis of HCM.

**KEYWORDS**

echocardiography, feline, hypertrophic cardiomyopathy

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**Abbreviations:** IVSd, mean end-diastolic interventricular septal thickness; LA:Ao, left atrium to aorta ratio; LAD, left atrial diameter in long axis; LAFS, left atrial fractional shortening; LV, left ventricular; LVEF, left ventricular ejection fraction; LVFS, left ventricular fractional shortening; LVFWd, mean end-diastolic left ventricular wall thickness; LVIDd, end diastolic left ventricular internal diameter; MAPSE FW, mitral annular plane systolic excursion measured at the free wall; MAPSE IVS, mitral annular plane systolic excursion measured at the interventricular septum; maxIVSd, maximal end-diastolic interventricular septal thickness; maxLVFWd, maximal end-diastolic left ventricular wall thickness; RVd, end diastolic right ventricular wall thickness; S, strain; SR, strain rate; STE, speckle tracking echocardiography; TAPSE, tricuspid annular plane systolic excursion.
Hypertrophic cardiomyopathy (HCM) is the most common heart disease in cats, and is characterized by left ventricular (LV) hypertrophy in the absence of abnormal loading conditions. Cats showing clinical signs have high cardiac mortality rates, but the prognosis in cats with preclinical HCM is highly variable.

HCM is generally considered to be a disease of diastolic impairment, with decreased systolic function (left ventricular fractional shortening [LVFS]) being identified as a prognostic factor in cats with HCM as a marker for late systolic impairment; however, early systolic impairment occurs with reduced LV longitudinal displacement (mitral annular plane systolic excursion) and longitudinal myocardial deformation (longitudinal strain) both in people and cats with HCM.

Speckle tracking echocardiography (STE) allows the quantification of myocardial deformation in the longitudinal, radial, and circumferential plane and can provide information on regional and global myocardial function. STE appears to be a sensitive technique for the early diagnosis of HCM in people, and it has been used to differentiate left ventricular hypertrophy because of HCM from athletes’ heart, cardiac amyloidosis, or storage disease. STE abnormalities have been documented in people with pathogenic HCM mutations who are phenotype-negative based on conventional echocardiographic imaging.

Reported applications for left ventricular STE in veterinary medicine have included the investigation of dilated cardiomyopathy, tachycardia-induced cardiomyopathy, degenerative mitral valve disease, and patent ductus arteriosus. Three STE studies have reported investigations of cardiac mechanics in cats with HCM. One study showed no difference in peak systolic longitudinal strain (S) and strain rate (SR) in cats with asymptomatic HCM compared to healthy controls, whereas another study showed reduced longitudinal and radial S in cats with asymptomatic HCM compared with controls. Left ventricular wall thickness was inversely correlated with longitudinal S, radial S and SR, and circumferential S and SR. The most recent study investigated multilayer STE in asymptomatic cats with dynamic outflow obstruction and found lower longitudinal S and SR on all layers and an increased epicardial-to-endocardial ratio in circumferential S and SR with preserved endocardial circumferential S when compared to healthy cats.

Left ventricular (LV) wall thickness is the measurement most commonly used to diagnose HCM in both people and cats; however, it is a relatively crude measure of hypertrophy and does not take into account the overall distribution. Left ventricular mass can be calculated from echocardiography and 1 study in people reported good correlation between longitudinal S and LV mass. Left ventricular mass is not routinely obtained in cats, although 1 study reported moderate correlation between LV mass measured post-mortem (considered the gold standard) with LV mass calculated by the truncated ellipse method from 2-dimensional (2D) echocardiography in cats with HCM.

Because previous STE studies in cats have enrolled relatively small, heterogeneous populations and found discordant results, we sought to investigate whether cardiac mechanics in cats with preclinical HCM were different compared to healthy control cats, and to evaluate whether the magnitude of left ventricular hypertrophy correlated with STE-derived variables. An additional aim of the study was to identify the best logistic regression model that could aid in differentiating cats with HCM from healthy controls using echocardiographic indexes other than LV wall thickness.

Our hypotheses were that longitudinal and radial S and SR would be lower in cats with HCM compared to healthy control cats, whereas circumferential S and SR would not differ. We also hypothesized that STE-derived variables would correlate with LV mass and that measurements of longitudinal deformation and LV mass would be useful in supporting a diagnosis of HCM.

2 | MATERIALS AND METHODS

The study received ethical approval from the Royal Veterinary College (SR 2018-1559). The electronic database and echocardiographic records were reviewed for cats diagnosed with preclinical HCM between April 2015 and September 2017. The control group comprised mainly healthy cats undergoing cardiac assessment as part of a blood donor program or cats in which a heart murmur had been auscultated but no structural abnormalities had been identified on echocardiography.

Enrolment criteria were a complete case record (owner data, cat signalment and history, complete physical examination) and a complete echocardiographic examination. Cats were included in the preclinical HCM group if the mean left ventricular wall thickness measured at end diastole was ≥6 mm for at least 1 myocardial segment (at the interventricular septum or left ventricular free wall). Cats were included in the healthy control group if no abnormalities were detected on the echocardiographic examination and no systemic illnesses that might affect hemodynamics or the cardiovascular system were observed. Additional inclusion criteria were the presence of good quality images with a minimum frame rate of 100 fps, and cine loops to allow STE evaluation from a right parasternal short axis view at the level of the papillary muscles, and a left apical 4 chamber view. Additional requirements included a symmetric, circular LV cross-section with minimal translational motion (“out-of-plane motion”) for the short axis views, and long axis views excluding the LV outflow tract, optimized for LV length and endocardial borders. When available, a cine loop from the left apical view optimized for longitudinal right ventricular strain and strain rate was preferable but not necessary for inclusion in the study.

Cats were excluded if there was a history compatible with congestive heart failure; aortic thromboembolism; arrhythmias; receiving medications; diagnosis of other conditions that could affect LV wall thickness such as hyperthyroidism, systemic hypertension, acromegaly, cardiomyopathies other than HCM, congenital heart disease, or neoplastic disease. Cats were excluded for incomplete case records or an inadequate echocardiographic examination. Left atrial enlargement was not an exclusion criterion.

All echocardiographic examinations were performed using a commercial ultrasound machine with 8-12 MHz phased array ultrasound probes (Vivid E95, GE systems, Hatfield, United Kingdom).
All echocardiographic measurements represented an average of 3 measurements taken from different cardiac cine loops/still images unless stated otherwise. Mean interventricular (IVSd) and free wall (FWd) septal thickness were measured from a 2-dimensional right parasternal long axis (2D RPLax) 4-chamber view at end diastole, based on averaged thickness taken from 3 consecutive cardiac cycles. Maximal interventricular (maxIVSd) and free wall (maxFWd) thickness at end-diastole were measured as the mean value of 3 different cardiac cycles taken from the thickest segment of 3 different views: 2D RPLax 4-or 5-chambered view or a short-axis view at the papillary muscle level. End diastolic right ventricular wall thickness (RVd) was measured from the RPLax 4-chamber view at end-diastole. Left ventricular internal diameter in diastole (LVDDd) was also measured at end diastole as an average from 3 2D RPLax 4-chamber view cine loops. A leading edge to leading edge technique was used for IVSd, FWd, maxIVSd, maxFWd, RVd, and LVDDd. Left atrial size was assessed as left atrium to aorta ratio (LA/Ao), which was measured from the right parasternal short axis view at the level of the heart base optimized for the left atrium with a trailing edge to leading edge technique in the first frame after aortic valve closure.7,10 Additionally, left atrial diameter in long axis was measured from a right parasternal 4-chamber long axis view by drawing a line parallel with the mitral annulus bisecting the left atrium in the last frame before mitral valve opening.7 Left atrial fractional shortening (LAFS) and left ventricular fractional shortening (LVFS) were measured with a leading edge to leading edge method from M-Mode views of right parasternal short axis view at the level of the heart base and papillary muscle, respectively, as previously described.7,38

The area-length method from the left apical 4-chamber view was used to calculate left ventricular ejection fraction.38,39 LV mass was calculated by the truncated ellipse method recommended in people and previously applied in cats,37 with the apical 4-chamber view as the reference long axis view.39 Mitral annular plane systolic excursion at the level of the free wall (MAPSE FW) and interventricular septum (MAPSE IVS), as well as tricuspid annular plane systolic excursion (TAPSE) were measured as described.11 Speckle tracking postprocessing was performed off-line by 1 single, trained observer (IS). Commercial software was used for off-line analysis (Echo Pac off-line measurement software, GE systems, Hatfield, United Kingdom). The cine loop was evaluated frame by frame to select a single frame as a starting point. Generally, the software would automatically select end-systole, but the observer assessed the whole cycle before selecting the best still frame. The endocardium was tracked manually, after which a region of interest was generated by the software, and was adjusted to fit the LV wall thickness by the observer. The software automatically divided the myocardium into 6 segments and evaluated tracking quality during the cardiac cycle. The whole cine loop was further evaluated by the operator and adjusted, if needed; only 1 manual attempt to adjust endocardial points was considered acceptable. After this procedure, S and SR curves were obtained by the software. Data were then exported to an Excel data sheet and prepared for statistical analysis.

All echocardiographic values were measured 3 times on a single day by the same operator based on a single-cycle clip/image and the results shown are averaged. For advanced echocardiographic values, a global peak systolic S and SR are provided, which are the averaged value of the 6 regional segments identified by the software. Global IVS and global LVFW S and SR were also calculated as the mean value of the 3 segments belonging to the IVS or FW. Right ventricular longitudinal S and SR were calculated only at the right ventricle wall (3 segments) by using a LV 4 chamber template and the interventricular values were discarded. Circumferential and radial S and SR were measured from the right parasternal short axis view, whereas LV longitudinal S and SR were measured at the apical 4-chamber view, RV longitudinal S and SR, when available, from a left apical view optimized for the RV. All S and SR values were obtained from a mid-myocardium level. A within-day and between-day coefficient of variation (CV) of both global and regional strain and strain rate was assessed in 10% of the cats.

2.1 Statistical analysis

Data were analyzed by a commercially available statistical software (IBM SPSS Statistics version 22, IBM [UK] Ltd, Portsmouth, UK) and statistical significance was set at P < 0.05. Continuous data were normally distributed according to the Shapiro-Wilk test. Data were therefore presented as mean ± SD, except for body condition score, which is an ordinal variable and was presented as median (range). In order to compare healthy control cats with affected, a t-test was used for continuous variables, a chi-squared test was performed for categorical variables, and Mann Whitney U test was applied for ordinal variables.

For the repeatability study, within-day and between-day coefficients of variation (CV) were calculated, including global and regional CV. CV was calculated as the ratio between SD and mean × 100.

Correlations between STE-derived variables and left ventricular wall thickness were tested by the Pearson's correlation coefficient. The correlation was considered perfect, strong, moderate or weak when the value of the correlation coefficient (r) was 1, 0.7-0.9, 0.4-0.6, or 0.1-0.3, respectively.40 A receiver-operating characteristic (ROC) curve with HCM (Y/N) as the predicting factor was performed to identify the echocardiographic variables that would best classify cats with HCM by methods other than left ventricular wall thickness (gold standard). The area under the curve (AUC) was considered excellent if the AUC was between 0.90 and 1, good for 0.80-0.90, fair for 0.70-0.80 and poor for 0.60-0.70.41 Cutoffs were also calculated and the Youden Index: ([sensitivity + specificity] – 1) was calculated to aid in the identification of the optimal cutoff.

A binary logistic regression model with the presence of HCM (Y/N) as the dependent variable was constructed. Potential explanatory variables were chosen using the values obtained from the ROC analysis of all echocardiographic variables for which there were no missing values. Variables with 95% confidence intervals for the ROC that did not include the value 0.5 were considered for entry into the model. The model was constructed in a forward stepwise manner. Variables with the highest AUC were entered into the model first and subsequent variables entered in descending order of AUC.
Improvement of the model at each step was indicated by an increase in the proportion of cases classified correctly. No more than 4 explanatory variables were entered in the model at any one time. In the final model, only variables with P < .05 were retained. The sensitivity of the final model to inclusion of previously excluded variables was evaluated by individually adding back all excluded variables into the final model. All variables entered into the model were checked for the risk of collinearity. No pair of independent variables had an r-value of greater than 0.8 in the correlation matrix. Goodness of fit of the model was assessed using the Hosmer and Lemeshow test.

Predicted probabilities derived from the equation generated by the final logistic regression model were used to plot an additional ROC curve.42

3 | RESULTS

After case selection based on the inclusion criteria (Figure 1), the final study population consisted of 73 cats: 36 healthy controls and 37 cats with preclinical HCM. Mean age was 5.1 ± 3.5 years and mean body

**FIGURE 1** Flow diagram showing the selection process for cats with HCM and healthy controls based on the inclusion criteria. HCM, hypertrophic cardiomyopathy

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**TABLE 1** Baseline demographics of asymptomatic cats with HCM and healthy controls

|                      | Healthy controls (n = 36) | HCM (n = 37) | P value |
|----------------------|---------------------------|--------------|---------|
| Age (years)          | 5 ± 3.4                   | 5.3 ± 3.7    | 0.75    |
| Sex (% male)         | 75% (27/36)               | 70% (26/37)  | 0.47    |
| Breed (% non-pedigree)| 69.4% (25/36)             | 56.7% (21/37)| 0.27    |
| Weight (kg)          | 4.7 ± 1.1                 | 4.5 ± 1.1    | 0.48    |
| Body condition score (1-9) | 5 (3-8) | 5 (2.5-7) | 0.96    |

Data presented as mean ± SD or as a percentage (for categorical data), or median (range) for ordinal variable (body condition score). To identify any difference between groups, T-test was performed for continuous variables, chi-square for categorical variables, and Mann-Whitney-U test for ordinal variables.

**TABLE 2** Coefficient of variation for global and regional speckle tracking echocardiography variables

|                    | Circ S | Circ SR | Rad S | Rad SR | Long S | Long SR | RV Long S | RV Long SR |
|--------------------|--------|---------|-------|--------|--------|---------|-----------|------------|
| Global within day  | 4%     | 6%      | 11%   | 13%    | 4%     | 6%      | 5%        | 10%        |
| Global between day | 5%     | 8%      | 9%    | 7%     | 4%     | 6%      | 4%        | 12%        |
| Regional within day| 9%     | 12%     | 12%   | 11%    | 8%     | 14%     | 6%        | 15%        |
| Regional between day| 9%    | 13%     | 13%   | 15%    | 11%    | 16%     | 11%       | 17%        |

Abbreviations: Circ S, circumferential strain; Circ SR, circumferential strain rate; Long S, longitudinal strain; Long SR, longitudinal strain rate; Rad S, radial strain; Rad SR, radial strain rate; RV Long S, right ventricular longitudinal strain; RV Long SR, right ventricular longitudinal strain rate.

Coefficient of variation for advanced echocardiographic parameters.
TABLE 3  Standard echocardiographic variables in asymptomatic cats with HCM and healthy controls

| Clinical and echocardiographic parameters | Healthy controls (n = 36) | HCM (n = 37) | P value for comparison between groups |
|------------------------------------------|--------------------------|--------------|--------------------------------------|
| Heart Rate (bpm)                          | n = 73                   |              |                                      |
| IVSd (mm)                                 | 4.3 ± 0.4                | 6.8 ± 0.8    | N/A                                  |
| LVFWd (mm)                                | 4.3 ± 0.5                | 6.3 ± 1.3    | N/A                                  |
| Max IVSd (mm)                             | 4.5 ± 0.5                | 7.3 ± 0.8    | N/A                                  |
| Max LVFWd (mm)                            | 4.3 ± 0.6                | 6.5 ± 1.3    | N/A                                  |
| LVIDD (mm)                                | 14.9 ± 1.9               | 12.4 ± 1.9   | <.001                                |
| RVd (mm)                                  | 2.1 ± 0.4                | 2.5 ± 0.4    | <.001                                |
| LAD (mm)                                  | 14.1 ± 1.59              | 15.4 ± 2.7   | .022                                 |
| LA/Ao                                     | 1.23 ± 0.20              | 1.32 ± 0.27  | .093                                 |
| LAFS (%)                                  | 33.2 ± 5.6               | 28.8 ± 7.5   | .006                                 |
| LVFS (%)                                  | 48.5 ± 7.4               | 54.2 ± 10.1  | .008                                 |
| LVEF (%)                                  | 63.4 ± 8.5               | 65.3 ± 12.1  | .45                                  |
| MAPSE FW (mm)                             | 5.5 ± 1.0                | 4.1 ± 0.8    | <.001                                |
| MAPSE IVS (mm)                            | 5.1 ± 0.9                | 4.2 ± 0.9    | <.001                                |
| TAPSE (mm)                                | 8.7 ± 1.7                | 7.9 ± 1.4    | .028                                 |
| LV mass (gr)                              | 8.6 ± 2.7                | 13.2 ± 3.7   | <.001                                |

Abbreviations: IVSd, mean end-diastolic interventricular septal thickness; LA/Ao, left atrium to aorta ratio; LAD, left atrial diameter in long axis; LAFS, left atrial fractional shortening; LVFS, left ventricular fractional shortening; LVFWd, mean end-diastolic left ventricular wall thickness; LVIDD, end diastolic left ventricular internal diameter; MAPSE FW, mitral annular plane systolic excursion measured at the free wall; MAPSE IVS, mitral annular plane systolic excursion measured at the interventricular septum; maxLVSD, maximal end-diastolic interventricular septal thickness; maxLVFWd, maximal end-diastolic left ventricular wall thickness; N/A, statistical analysis not applicable (criteria used for the classification into healthy/HCM); RVd, end diastolic right ventricular wall thickness; TAPSE, tricuspid annular plane systolic excursion; LV mass, end diastolic left ventricular mass.

Data are presented as mean ± SD. Independent t-test analysis was performed to identify difference between groups.

TABLE 4  Speckle-tracking echocardiography (STE) variables

| STE                         | Healthy controls (n = 36 for all observations except for RV S/SR) | HCM (n = 37 for all observations except for RV S/SR) | P value for between group comparisons |
|-----------------------------|-----------------------------------------------------------------|--------------------------------------------------------|--------------------------------------|
| Circumferential S (%)       | −23.2 ± 3.9                                                      | −21.2 ± 6.2                                            | .12                                  |
| Circumferential S (%)       | −22.4 ± 5.2                                                      | −19.2 ± 6.5                                            |                                      |
| Circumferential SR (s⁻¹)    | −3.84 ± 0.94                                                     | −3.73 ± 1.01                                           | .25                                  |
| Radial S (%)                | 66.7 ± 17.6                                                     | 44.0 ± 21.3                                            | <.001                                |
| Radial S (%)                | 67.6 ± 18.5                                                     | 44.3 ± 19.7                                            |                                      |
| Radial SR (s⁻¹)             | 6.67 ± 1.8                                                      | 5.6 ± 2.08                                             | .022                                 |
| Longitudinal S (%)          | −19.7 ± 2.74                                                    | −14.8 ± 3.3                                            | <.001                                |
| Longitudinal S (%)          | −23.1 ± 3.3                                                      | −17.9 ± 3.9                                            |                                      |
| Longitudinal SR (s⁻¹)       | −2.9 ± 0.68                                                      | −2.36 ± 0.62                                           | <.001                                |
| Longitudinal S (s⁻¹)        | −3.32 ± 0.81                                                     | −2.79 ± 0.75                                           | .30                                  |

Abbreviations: Circumferential S, circumferential strain; Circumferential SR, circumferential strain rate; Longitudinal S, left ventricular longitudinal strain; Longitudinal SR, left ventricular longitudinal strain rate; Radial S, radial strain; Radial SR, radial strain rate; RV longitudinal S, right ventricular strain; RV longitudinal SR, right ventricular strain rate.

Data are presented as mean ± SD. Independent t-test analysis was performed to identify difference between groups.
Twenty-eight cats with preclinical HCM had dynamic left ventricular outflow tract obstruction. When evaluating echocardiographic data, cats with HCM had increased left atrial diameter in long axis (LAD, \( P = .002 \)), LVFS (\( P = .008 \)), and LV mass (\( P < .001 \)) compared to healthy controls. Furthermore, cats with HCM had lower LAFS (\( P = .006 \)), MAPSE IVS (\( P < .001 \)), MAPSE FW (\( P < .001 \)), and TAPSE (\( P = .028 \)) compared to healthy control cats (Table 3). When compared to normal cats, cats with HCM had lower radial S (\( P < .001 \)) and SR (\( P = .022 \)) and less negative longitudinal S (\( P < .001 \)) and SR (\( P < .001 \)) (Table 4), whereas no difference was observed for circumferential S (\( P = .12 \)) and SR (\( P = .25 \)) or RV longitudinal S (\( P = .76 \)) and SR (\( P = .30 \)).

When analyzing correlations between STE and left ventricular hypertrophy, Pearson correlation’s coefficient showed moderate correlation between maximal IVS and FW thickness and the respective global IVS and FW longitudinal S (\( r = 0.659, P < .001 \) and \( r = 0.524, P < .001 \)), moderate correlation between maximal IVS and FW and the respective global IVS and FW radial strain (\( r = −0.494, P < .001 \) and \( r = −0.413, P < .001 \)). Weak correlation was observed for maximal IVS and FW thickness and the respective global IVS and FW longitudinal SR (\( r = 0.327, P = .005 \) and \( r = 0.371, P = .001 \)) as well as maximal IVS and global IVS circumferential S (\( r = 0.307, P = .008 \)).

Left ventricular mass was strongly correlated with IVSd and LVFWd (\( r = 0.701, P < .001 \) and \( r = 0.715, P < .001 \) respectively). Additionally, LV mass moderately correlated with longitudinal S and SR (\( r = 0.511, P < .001 \) and \( r = 0.311, P = .007 \), respectively), but not with other STE-derived variables. No significant correlation was found between body weight and LV mass or longitudinal S or SR. No significant correlation was found between heart rate and STE-derived variables.

When assessing the predictive ability of STE-derived variables, LV mass and standard echocardiographic variables, the ROC curves showed that LV mass, and longitudinal S had some of the highest AUCs of 0.867 (CI: 0.783-0.951) and 0.862, respectively (CI:0.781-0.943) (Figure 2, Table 5). Based on the Youden Index, the best cutoff for LV mass was 10 g (Youden Index 0.62) and −15.9% for Long S (Youden index 0.56).

The final logistic regression model showed that LV mass, MAPSE FW, and LVIDd in combination best predicted the presence of HCM (Table 6).

The combination of these tests increased the probability of a cat with HCM to be correctly classified as affected (predicted ROC curve, AUC 0.974, CI 0.945-1.000) (Figure 2).

![FIGURE 2](attachment:image.png) ROC curve constructed using probabilities derived from the binary logistic regression and selected echocardiographic variables to differentiate asymptomatic cats from those with HCM. The 4 individual echocardiographic variables with the greatest areas under the curve are plotted in addition to the predicted probabilities from the regression equation. The gold standard was 2-dimensional left ventricular wall thickness. The area under the curve with 95% confidence interval values is displayed in Table 5. Predicted probability, derived from binary logistic regression model; LV mass truncated ellipsoid, end-diastolic left ventricular mass measured by the truncated ellipse method; longitudinal S, left ventricular longitudinal strain; MAPSE FW, mitral annular plane systolic excursion measured at the free wall; LVIDd, end diastolic left ventricular internal diameter

| Variable                  | Area under the curve (AUC) | 95% CI          |
|---------------------------|----------------------------|-----------------|
| Predicted model           | 0.974                      | 0.945-1.000     |
| LV mass                   | 0.867                      | 0.783-0.951     |
| MAPSE FW                  | 0.864                      | 0.781-0.947     |
| Longitudinal S            | 0.862                      | 0.781-0.943     |
| LVIDd                     | 0.827                      | 0.734-0.921     |
| RVd                       | 0.803                      | 0.699-0.907     |
| Radial S                  | 0.788                      | 0.682-0.895     |
| MAPSE IVS                 | 0.784                      | 0.638-0.858     |
| Longitudinal SR           | 0.751                      | 0.639-0.862     |
| LAFS                      | 0.681                      | 0.556-0.805     |
| LVFS                      | 0.673                      | 0.548-0.798     |
| Radial SR                 | 0.655                      | 0.528-0.782     |
| LAD                       | 0.651                      | 0.523-0.779     |
| TAPSE                     | 0.627                      | 0.495-0.758     |
| LA/Ao                     | 0.610                      | 0.480-0.741     |
| EF                        | 0.584                      | 0.450-0.718     |
| Circumferential S         | 0.577                      | 0.444-0.710     |
| Circumferential SR        | 0.521                      | 0.387-0.655     |
| RV Longitudinal S         | 0.483                      | 0.319-0.648     |
| RV Longitudinal SR        | 0.446                      | 0.282-0.611     |

Output from ROC curves for echocardiographic variables and predicted probabilities obtained from multivariable logistic regression analysis. Area under the curve (AUC) and 95% confident interval (CI) are provided.
planes and is an echocardiography-based technique. As with counter-rotation to allow twisting of the left ventricle to aid ejection, additional feature of the contracting heart, as the heart base and apex tend to be preserved or increased, with some studies reporting deterioration of the circumferential deformation as a possible explanation for the development of clinical signs.

The results of the present study showed that in 2 groups of cats of similar age, sex, weight, body condition score, and heart rate, cats with preclinical HCM had lower absolute values of both longitudinal and radial S and SR, and greater LV mass. Furthermore, a correlation was found between regional IVS or FW longitudinal S and regional LV wall thickness, as well as between longitudinal S or SR and LV mass. In the study population, these variables performed well individually at differentiating cats with preclinical HCM from controls when compared to standard echocardiographic variables. The logistic regression model that best predicted the presence of HCM included a combination of LV mass, MAPSE FW, and LVIDd. Within-day and between-day CV was acceptable and similar to previous studies, with global S and SR values achieving better reproducibility than regional measurements, as identified in a previous study on S/SR.

Studies on myocardial fiber distribution showed that the myocardium has longitudinally and circumferentially oriented fibers, which support cardiac contraction in different deformation planes. Short axis, long axis, and twist represent deformation planes that contribute to cardiac ejection. LVFS can be considered a surrogate of short axis deformation, as well as circumferential and radial S and SR. Long axis function can be identified by atroventricular systolic plane excursion (MAPSE), as during contraction the cardiac base descends toward the cardiac apex; longitudinal S and SR are also additional long axis deformation variables. The right ventricle has mainly longitudinally oriented fibers, with minimal short axis deformation, therefore TAPSE and longitudinal S and SR are the deformation variables that can be used to assess right ventricular deformation. Torsion and twist are an additional feature of the contracting heart, as the heart base and apex counter-rotate to allow twisting of the left ventricle to aid ejection.

STE allows the evaluation of myocardial deformation in different planes and is an echocardiography-based technique. As with every technique, there are limitations, which for STE are mainly related to image quality, out of plane motion, and load dependency. Additional limitations include a steep learning curve, inter-vendor software analysis variation and an overall acceptable CV, mainly with global results, yet regional segment analysis can have greater variation (up to 16% of variation). The present study included cases seen at the authors’ outpatient clinic, and the required cine loops were not different from the standard views required for a standard echocardiogram. It is important to take particular care with image quality, symmetry and in limiting out-of-plane motion to allow STE analysis and this technique might therefore be difficult in uncooperative or critically ill cats.

The results of our study showed that longitudinal deformation is impaired early in the disease, similar to previous reports in people with preclinical HCM where early impairment in longitudinal and radial S and SR has been observed, whereas circumferential S and SR tend to be preserved or increased, with some studies reporting deterioration of the circumferential deformation as a possible explanation for the development of clinical signs.

Previous studies assessing STE in cats with HCM showed discordant results. One study did not identify any change in longitudinal systolic S or SR, with the only abnormality detected being a diastolic change in longitudinal E and A SR, but not its ratio. The cat population was relatively homogeneous and no cat received treatment. However, sample size was smaller than our study, and the degree of hypertrophy was mild, which might account for the differences in the results. Also, no segmental S or SR data were shown, and only global values were presented. In the present study, diastolic SR analysis was not performed so it is not possible to fully compare our study to those previously published. Another study reported decreased radial and longitudinal systolic S and SR in cats with HCM not showing clinical signs and with congestive heart failure, but not in those showing clinical signs other than CHF (exercise intolerance, dyspnea or tachypnea). The multiple subgroups with low number of cats in each group could have limited statistical power. No characterization on hypertrophy distribution and regional mechanics was mentioned in that study, and cardiac therapy was allowed. Our results confirm the previous observation that cats with asymptomatic HCM have lower absolute values of longitudinal and radial deformation compared to healthy cats. A more recent study identified preserved endocardial circumferential strain but lower epicardial values resulting in greater epicardial to endocardial ratio in cats with dynamic outflow tract obstruction and HCM, so it could be that more detailed analysis of circumferential deformation would be required in the assessment of cats with HCM. Compared with healthy control cats, the cats with preclinical HCM in our study had decreased longitudinal and radial deformation with similar circumferential S and SR, that was assessed at the mid-myocardium level, in contrast to the previously cited study, where the whole layer circumferential results were lower in asymptomatic cats with HCM, but were not statistically different on a previous study from the same group. There are similar discrepancies.

### TABLE 6 Binary logistic regression analysis results

| Parameter     | B value | OR   | CI      | Wald | P value |
|---------------|---------|------|---------|------|---------|
| LV mass       | 0.792   | 2.207| 1.435-3.395 | 12.99| <.001   |
| LVIDd         | −0.681  | 0.506| 0.287-0.891 | 5.569| =.018   |
| MAPSE FW      | −1.524  | 0.218| 0.060-0.792 | 5.352| =.021   |
| Constant      | 8.385   | 4381.982| 5.057 |      | =.025   |

Abbreviations: B value, beta value coefficient; CI, confident interval; LV mass, end diastolic left ventricular mass; LVIDd, end diastolic left ventricular internal diameter; OR, odds ratio; MAPSE FW, mitral annular plane systolic excursion measured at the free wall, which were all statistically significant in the model; Wald, wald test result.

### 4 DISCUSSION

The results of the present study showed that in 2 groups of cats of similar age, sex, weight, body condition score, and heart rate, cats with preclinical HCM 具有较低的绝对值的纵向和径向S和SR，并有较大的LV质量。此外，与区域IVS或FW的纵向S和区域LV壁厚度，以及纵向S或SR和LV质量的成正比相关。在研究人群中，这些变量在个体水平上表现良好，与先前的研究相似，其中全局S和SR值达到更好的可重复性，而区域测量值，在先前的S/SR研究中被识别。
in circumferential S and SR in humans\textsuperscript{12,14,16} and there is currently interest in the assessment of a transmural strain difference both on MRI and echocardiography,\textsuperscript{66,47} in human cardiology.

The best correlation with regional hypertrophy and left ventricular mass was observed with longitudinal S and SR. No weight-dependence was identified in the present study in STE-derived values, although the weight range in our group of cats was relatively narrow, with few cats weighing more than 5 kg or less than 3 kg. A similar lack of correlation between LV mass and bodyweight was also observed. The present findings suggest that in this population of cats, STE and LV mass can provide a weightless index to aid in the characterization of HCM; however, further studies are required to elucidate if a wider body weight range would impact LV mass and STE variables based on the findings of increasing LV wall thickness with increasing body weight.\textsuperscript{48}

LV mass has been quantified by MRI, echocardiography and post-mortem in cats\textsuperscript{37}; however, no extensive use of this technique has been applied in veterinary medicine. To the author's knowledge, no repeatability studies in the veterinary literature are available. Different LV mass calculations in people are available, including M-mode and 2D based calculations. The truncated ellipse method is the 2-dimensional method recommended in human guidelines\textsuperscript{39} and previously applied in cats.\textsuperscript{37} It requires both short- and long-axis views, but an overall acceptable correlation was found compared to post-mortem LV mass assessment in cats, although MRI-based mass calculation correlated better to actual LV mass.\textsuperscript{37} Healthy control cats in our study had similar values to the ones reported previously.\textsuperscript{37} Longitudinal S and SR seemed to correlate well with LV mass, with less negative longitudinal S values associated with increasing LV mass. There is similar correlation in people with HCM.\textsuperscript{12,14}

Early diagnosis and risk stratification are important targets in human medicine in order to deliver the best treatment options. A decrease in longitudinal S has been associated with a poorer prognosis in people, with the highest incidence of adverse events (including congestive heart failure, ventricular tachycardia, sudden cardiac death or the implantation of a cardiodefibrillator device) with lower absolute values (ie, less negative results).\textsuperscript{69} A longitudinal S greater than −16% (ie, less negative) was an independent predictor of adverse effects in people.\textsuperscript{50} Longitudinal S could be of help in classifying equivocal or challenging cases or allow a closer monitoring over time; however, longitudinal studies are required to investigate this aspect. Because of the previously reported limitations in the technique and the results of our regression analysis, MAPSE seems to be a more widely applicable alternative in the assessment of longitudinal deformation, with the main limitation of this technique associated with the specific regional sampling required (atrioventricular annulus and not a global index of longitudinal deformation).

The logistic regression model that showed the best association with a diagnosis of HCM in our population combined LV mass, MAPSE FW, and LVIDd. When left ventricular hypertrophy is manifest, left ventricular wall thickness is likely sufficient to discriminate between affected and non-affected. However, combining different variables could help in settings where the hypertrophy is not as obvious, such as in equivocal cases or in cats with a family history of HCM. Interestingly, the 3 variables that were shown to increase the probability of a cat to have HCM include a marker of global wall thickness (LV mass), LV chamber size, and longitudinal deformation. This might indicate that alongside wall thickness, LV geometry and function are additional tools that could aid identifying HCM or improve monitoring. This potential application requires further investigation. Longitudinal observational studies in cats with preclinical HCM and with equivocal HCM might provide insight into the applicability of longitudinal S or MAPSE and LV mass as an aid in risk stratification for HCM development or progression. Genotype-positive and phenotype-negative people represent a challenge in human cardiology, and few studies have investigated abnormalities using advanced echocardiographic techniques.\textsuperscript{27,28} Maine Coon cats heterozygous for the mutation A31P do not always show LV hypertrophy\textsuperscript{51–55} but the prevalence of hypertrophy increases with age\textsuperscript{53,55} so use of a combined predictive model would be worth evaluating in this context.

As expected, cats with HCM had thicker end-diastolic RV wall thicknesses values as previously identified.\textsuperscript{56,57} Interestingly, in the asymptomatic phase, no difference in RV Long S and SR was observed. TAPSE values are lower in cats with CHF, so it could be possible that RV global long axis deformation is impaired later on, when clinical signs develop or when there is an increase in pulmonary arterial pressure or pulmonary hypertension.\textsuperscript{58}

Limitations of the present study include its retrospective nature; however, a standardized protocol for clinical cats and healthy controls was applied. It was not possible to blind the operator to the presence of LV hypertrophy when making echocardiographic measurements, and our study did not address diastolic function deformation via standard echocardiography or STE, or torsion or twist. Variability of right ventricular shape makes RV imaging more challenging; whereas available, the authors used RV focused views; otherwise, RV S and SR were not assessed. Our study was not designed to assess genotype. The classification and ROC curve output was based on a 2D end-diastolic LV wall thickness cutoff of 6 mm, which is considered a conservative cutoff for the diagnosis of HCM in cats, it is possible that other cutoffs could be better at classifying HCM in cats. Furthermore, the authors cannot exclude that the healthy cats group might have included cats with preclinical HCM not meeting the current criteria for HCM. Additionally, the regression model was performed in a dichotomous population of cats, where LV hypertrophy was manifest or absent based on the selected cutoff, so the results of this model might not be readily applied in a population of cats with equivocal/genotype positive-phenotype negative cats, for which a longitudinal study would be required.

In conclusion, cats with HCM had decreased systolic long axis deformation and radial deformation, but overall preserved circumferential deformation. Left ventricular mass was increased in cats with HCM. LV mass and longitudinal S correlated with the degree of left ventricular hypertrophy and performed best when compared to standard echocardiographic variables. STE CV was acceptable, with the lowest CV with global S and SR. Longitudinal S and LV mass might be useful additional variables in the evaluation of cats with HCM.
CONFLICT OF INTEREST DECLARATION

Authors declare that they have no conflict of interest with the contents of this article.

OFF-LABEL ANTIMICROBIAL DECLARATION

Authors declare no off-label use of antimicrobials.

INSTITUTIONAL ANIMAL CARE AND USE COMMITTEE (IACUC) OR OTHER APPROVAL DECLARATION

Royal Veterinary College IACUC approval, SR 2018-1559.

HUMAN ETHICS APPROVAL DECLARATION

Authors declare that human ethics approval was not needed for this study.

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