Feature-Tracking Strain Derived from Compressed Sensing Cine Cardiovascular Magnetic Resonance Imaging for Myocardial Infarct Detection: A Feasibility Study

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Abstract

Purpose: This study aimed to use gadolinium-enhanced cardiovascular magnetic resonance (LGE-CMR) scanning to examine the clinical feasibility of feature-tracking strain (FT-strain) analysis on compressed sensing (CS) cine cardiovascular magnetic resonance (CMR) imaging for detecting myocardial infarction (MI).

Methods: We enrolled 37 patients who underwent conventional cine CMR, CS cine CMR, and LGE-CMR scanning to assess cardiovascular disease. FT-strain analysis was used to assess peak circumferential strain (p-CS) based on an 18-segment model in both cine CMR imaging modalities. Based on LGE-CMR imaging findings, myocardial segments were classified as remote, adjacent, subendocardial infarcted, and transmural infarcted. The diagnostic performance of p-CS for detecting MI was compared between CS cine CMR imaging and conventional cine CMR imaging using the receiver operating characteristic (ROC) curve analysis.

Results: A total of 440 remote, 85 adjacent, 76 subendocardial infarcted, and 65 transmural infarcted segments were diagnosed on LGE-CMR imaging. There were significant between-group differences in p-CS on both conventional and CS cine CMR (p < 0.05 in each) imaging. The sensitivity and specificity of p-CS for identifying MI were 85% and 79% for conventional cine CMR imaging, and 82% and 77% for CS cine CMR imaging, respectively. There was no significant difference between conventional and CS cine CMR imaging in the area under the curve of p-CS (0.89 vs. 0.87, p = 0.15).

Conclusion: FT-strain analysis of CS cine CMR imaging may help identify MI; it may be used alongside or instead of conventional CMR imaging.

Introduction

Assessment of cardiac function is important in patients with cardiovascular disease, for which echocardiography is widely used in clinical practice. Left ventricular ejection fraction (LVEF) is a quantitative index of global cardiac function that enables mortality risk prediction and discrimination [1]. Myocardial strain (longitudinal, circumferential, and radial strain) is useful for the assessment of both global and regional cardiac function [2, 3]. It is easily evaluated using two-dimensional speckle-tracking echocardiography (2D-STE), which has a greater prognostic value than LVEF [4, 5]. However, the robustness of myocardial analysis with 2D-STE depends on the skill of the operator. A blind area on echocardiography may prevent the assessment of the entire myocardium. Cardiovascular magnetic resonance (CMR) imaging has been found to have high objectivity, and it allows for the assessment of the whole myocardium.

Feature-tracking strain (FT-strain) analysis is a method of myocardial strain analysis on CMR scans obtained through cine CMR imaging [6, 7]. Lately, compressed sensing (CS) techniques using sparse sampling and iterative reconstruction have reduced the acquisition time of cine CMR images [8]. FT-strain analysis could potentially be used for cine CMR images with CS, but the clinical feasibility of FT-strain
analysis of CS cine CMR images remains unknown. This study aimed to compare FT-strain analysis of CS cine CMR images with that of conventional cine CMR images and evaluate its feasibility for detecting myocardial infarction (MI) using late gadolinium-enhanced (LGE) CMR imaging.

Materials And Methods

This study was a sub-analysis of a dataset obtained in a previous prospective study [8]. It was approved by an ethics review board of Saiseikai Matsuyama Hospital and has therefore been performed in accordance with the ethical standards laid down in the 1964 Declaration of Helsinki and its later amendments. Informed consent was obtained from all patients enrolled in this study. Seventy-four patients who underwent CMR scanning between August 2014 and May 2015 were included in this study. All patients were scheduled for an evaluation of cardiovascular disease; CMR scanning was performed at the discretion of the attending physician. The CMR protocol included conventional cine CMR, CS cine CMR, and LGE-CMR imaging. The exclusion criteria were as follows: (1) non-ischemic cardiovascular disease, (2) data unsuitable for FT-strain analysis (due to arrhythmia or poor breath-holding), and (3) poor LGE-CMR image quality.

Cardiovascular magnetic resonance imaging protocol

A 3T MR scanner (MAGNETOM Skyra; Siemens Healthcare, Erlangen, Germany) with 48 receivers and two parallel transmission channels was used to implement an established protocol [8]. Scout images were acquired to plan cardiac axial views. Using a segmented balanced steady-state free-precession sequence, conventional cine CMR images of the left ventricle (LV) in the short axis direction were acquired with a stack of eight contiguous slices with adequate slice gaps to cover the entire LV. Additionally, LV short-axis CS cine CMR images were acquired using a prototype sequence with sparse incoherent sampling of k-space and non-linear iterative SENSE-type image reconstruction immediately after the conventional cine CMR scans. Temporal resolution, spatial resolution, and slice orientations were identical in both cine protocols. The details of the procedure used for the acquisition and reconstruction of CS cine CMR images have been reported elsewhere [8]. Over 1.5 cardiac cycles of data were acquired during CS cine CMR imaging. LV short-axis LGE-CMR images were acquired 10 min after the injection of gadobutrol (0.1 mmol/kg) using a 2D phase-sensitive inversion recovery sequence. The imaging parameters are presented in Table 1.

Feature-tracking strain analysis

All FT-strain analyses were performed using designated software (CMR42; Circle Cardiovascular Imaging Inc, Calgary, Canada). Three representative LV short-axial slices at the basal-, mid-, and apical levels were selected from both short-axial cine CMR and LGE-CMR images by an independent observer. Subsequently, a software algorithm semi-automatically segmented the LV myocardium into six equidistant segments according to the 18-segment model [9], and peak circumferential myocardium (p-CS) values were independently analyzed in both types of cine CMR images. Inter-observer agreement was assessed using images from
10 randomly selected patients (180 myocardial segments) who were examined by two experienced radiologists with 7 and 15 years of experience in CMR imaging analysis, respectively.

**LGE-CMR analysis**

Two experienced radiologists with 8 and 10 years of experience in CMR imaging, respectively, who were blinded to all other data were tasked with independently classifying myocardial characteristics into four groups at the segment level [10, 11]; the groups were defined as follows: i) a segment without an LGE area that was not adjacent to the border or the infarcted segment was termed the remote segment; ii) a segment without an LGE area that was adjacent to the infarcted segment was termed the adjacent segment; iii) a segment with an LGE transmural extent of ≤ 50% was termed the subendocardial infarcted segment; and iv) a segment with LGE transmural extent of > 50% was termed the transmural infarcted segment. The subendocardial and transmural infarcted segments were defined as MI segments. Any discrepancies between two observers were resolved by discussion and consensus.

**Statistical analysis**

Continuous data were expressed as mean ± standard deviation or median (first quartile, third quartile), depending on the type of distribution. Inter-observer agreement for LGE-CMR analysis was evaluated using Cohen’s κ. Inter-observer agreement for p-CS analysis was evaluated using intra-class correlation coefficients (ICC). The paired t-test was used to compare scanning time between conventional and CS cine CMR scans. The Wilcoxon signed-rank test was used to compare the p-CS values between conventional and CS cine CMR scans. Correlation and agreement between the p-CS values obtained from conventional CMR and CS cine CMR images were evaluated using linear regression and the Bland–Altman analyses. Differences in p-CS values among remote, adjacent, and infarcted regions were assessed using the Steel–Dwass test. To analyze diagnostic performance by segment, we used generalized estimating equations, assumed values for exchangeable correlation structures, and adjusted for the clustered nature of the data using robust estimators of variance [12]. The cutoff values and area under the curve (AUC) of p-CS on conventional and CS cine CMR images for detecting myocardial infarction (subendocardial and transmural) were determined using receiver operating characteristic curve (ROC) analysis. The optimal cut-off values were determined using Youden’s index. Sensitivity, specificity, and positive and negative predictive values (PPV and NPV, respectively) with 95% confidence intervals (CI) were calculated using the cut-off values. In all tests, statistical significance was set at \( p < 0.05 \). Statistical analyses were performed with JMP14 (SAS Institute, Cary, NC, USA).

**Results**

Of the 74 patients initially included in this study, 37 were excluded because of (1) non-ischemic cardiovascular disease (n = 21), (2) data unsuitable for FT-strain analysis (n = 9), and (3) poor LGE image quality (n = 7). Finally, 37 eligible patients were enrolled (Table 2). All patients had a regular sinus rhythm and a mean heart rate of 60.2 ± 10.3 bpm during both cine CMR scans. The total examination time was 114.0 ± 7.0 s for conventional cine CMR scanning and 24.6 ± 4.2 s for CS cine CMR scanning (\( p < 0.0001 \).
Characteristics of myocardial segments on LGE-CMR imaging

A total of 666 segments from 37 patients were classified as remote (n = 440), adjacent (n = 85), subendocardial infarcted (n = 76), or transmural infarcted (n = 65). Among them, 141 (21%) were defined as MI segments. The inter-observer agreement for LGE-CMR imaging was 0.84, indicating satisfactory reproducibility (κ > 0.70).

p-CS on conventional and CS cine CMR images

The p-CS values for conventional and CS cine CMR images were −19.2% (-23.5%, -14.0%), and −19.7% (-23.9%, -14.1%), respectively (p = 0.21). The p-CS values obtained from conventional and CS cine CMR images were significantly correlated (r = 0.75, p < 0.05; Fig. 1A). The mean difference between the conventional and CS cine CMR images based on the Bland–Altman method was 0.5% (95% limits of agreement: -8.4–9.4%; Fig. 1B). The ICC values were 0.89 (95%CI: 0.85–0.92) and 0.85 (95%CI: 0.80–0.89) for conventional and CS cine CMR images, respectively.

Diagnostic performance of p-CS

Figure 2 shows the results of p-CS analysis on conventional and CS cine CMR images in a representative case of MI. Figure 3A shows the p-CS values in the conventional cine CMR image; they were −21.6% (-24.9%, -18.2%), -15.3% (-19.0%, -11.5%), -12.3% (-15.6%, -8.8%), and −7.0% (-12.2%, -3.8%) for the remote, adjacent, subendocardial infarcted, and transmural infarcted segments, respectively. Figure 3B shows the corresponding values in the CS cine CMR image; they were −22.3% (-25.2%, -18.4%), -15.8% (-19.3%, -11.6%), -13.0% (-17.4%, -9.4%), and −8.8% (-12.9%, -5.9%) for the remote, adjacent, subendocardial infarcted, and transmural infarcted segments, respectively. The p-CS values were decreased in all the groups in both conventional and CS cine CMR images (all p < 0.05).

The diagnostic performance estimates of p-CS for detecting MI (subendocardial and transmural infarcted segments) on conventional and CS cine CMR images are shown in Table 3. ROC curve analysis revealed p-CS AUC values of 0.89 (95% CI: 0.85–0.91) and 0.87 (95% CI: 0.83–0.90) for conventional and CS cine CMR images, respectively (p = 0.15; Fig. 4). The optimal cut-off values of p-CS were −16.4% and −16.5% for conventional and CS cine CMR images, respectively.

Discussion

The main findings of this study include (1) the significant correlation between the p-CS values of conventional and CS cine CMR images, (2) high diagnostic performance of p-CS for identifying MI segments on conventional and CS cine CMR images, and (3) evidence that p-CS on CS cine CMR imaging may help classify myocardial characterization on LGE-CMR into remote, adjacent, subendocardial infarcted, and transmural infarcted segments.
Myocardial strain analysis on CMR images is a quantitative method of cardiac function assessment. It is conventionally performed using tagging CMR scans or displacement- or strain-encoded imaging [11, 13, 14]. However, additional data must be acquired for strain analysis using these methods. FT-strain analysis can be performed after processing cine CMR images, providing additional diagnostic value to the visual assessment of cardiac function [15]. Recently, the CS technique has been used with cine CMR images, enabling quick data acquisition while maintaining the robustness of cardiac function assessment [8]. In the present study, the p-CS values of CS cine CMR images were correlated with those of conventional cine CMR images. Furthermore, there was no significant difference in p-CS values between conventional and CS cine CMR images, which was previously reported [16]. The present findings indicate that data acquisition of the full cardiac cycle is important for accurate FT-strain analysis using CS cine CMR images, as a prospective real-time cine CMR scan may lack complete end-diastolic phase data, resulting in the underestimation of peak strain values relative to those obtained from retrospective conventional cine CMR scans [16]. Consequently, in the present study, we acquired the data of over 1.5 cardiac cycles on CS real-time cine CMR scanning to increase estimate accuracy.

There was no significant difference in the AUC values for detecting MI obtained using conventional and CS cine CMR images. In previous studies, p-CS values obtained using conventional CMR strain analysis for detecting MI had a sensitivity estimate of 71–81% and specificity estimate of 74–75%. These estimates are consistent with the estimated from this study [10, 13]. In the present study, the diagnostic performance of p-CS of CS cine CMR imaging for detecting MI was not inferior to that of conventional cine CMR scanning. Moreover, p-CS values on CS cine CMR images were correlated with the severity and location of MI, as reflected in the categories of remote, adjacent, subendocardial infarcted, and transmural infarcted segments. A previous study has shown that p-CS values obtained from conventional CMR images varied depending on the severity and location of MI on an LGE-CMR image [11]. The use of p-CS on CS cine CMR imaging may help differentiate subendocardial from transmural MI as assessed on LGE-CMR scanning, which is clinically relevant to prognosis and viability after revascularization [17, 18]. Buss et al. have shown that, as with the values obtained from LGE-CMR images, p-CS values from FT-strain analysis may be an independent predictor of cardiac function improvement after revascularization [19].

FT-strain analysis of CS cine CMR images is a quantitative technique of cardiac function assessment that is quicker than conventional CMR strain analysis. CS cine CMR imaging is becoming increasingly available, and FT-strain analysis of CS cine CMR scans may be applied to various cardiovascular diseases, as is presently used on conventional cine CMR scans [20–22]. This technique, which does not require a contrast medium, may benefit patients who cannot tolerate lengthy examination times and multiple breath-holdings and to those with advanced renal dysfunction and high risk of nephrogenic systemic fibrosis or renal failure.

There are some limitations of the present study. First, this study was conducted at a single center and with a small sample. Second, the present study only used p-CS values, because they are better for detecting MI than other strain parameters (e.g., longitudinal strain) [13, 23]. Further studies using other
myocardial strain parameters with CS cine CMR images are required. Third, the reconstruction time for the CS cine CMR image was approximately 3 min. Ongoing technological developments will reduce reconstruction times, thus helping to overcome this limitation.

Fourth, the effects of different types of CMR-FT analysis software on obtained estimates were not assessed. As different software types use different algorithms, they may result in different values; these inherent differences should be considered when comparing findings obtained using different software types [24]. Fifth, the present study did not account for differences in temporal and spatial resolution and regularization during FT strain analysis, the impact of which has been reported [25]. Finally, low temporal resolution or high regularization may result in underestimation of strain values [16].

In conclusion, FT-strain analysis of CS cine CMR images may help identify and classify MI segments; it may be used alongside or instead of that of conventional cine CMR images.

Declarations

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Conflict of interest

M.S. is an employee of Siemens Healthcare. The other authors declare no conflict of interest.

Availability of data and material

The date that supports the findings of this study are available from the corresponding author, T.K., upon reasonable request.

Authors’ contributions

TM conceived the study, quantitatively measured the strain, performed the statistical analysis, and drafted the manuscript. YT, TK participated in design of the study, assisted in the interpretation of the results, and helped to revise the manuscript. RO, MN quantitatively measured the strain and image contrast. KW participated in the design of the study and its coordination. MS contributed to the sequence development and the implementation on the scanner, as well as the revision of the manuscript. SA, OY and TK assisted with the interpretation of the results and helped revise the manuscript. All authors read and approved the final manuscript.

Ethics approval

This study was approved by an ethics review board of Saiseikai Matsuyama Hospital.
Consent to participate

Informed consent to participate was obtained from all patients enrolled in this study.

Consent for publication

All the participants in this study provided written consent for publication.

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Tables
**Table 1. Imaging parameters**

|                           | Conventional cine CMR | CS cine CMR | LGE-CMR |
|---------------------------|------------------------|-------------|---------|
| **Sequence type**         | 2D cine true FISP      | 2D cine true FISP | 2D PSIR |
| **ECG mode**              | Retrospective gating   | Prospective triggering | —       |
| **TE/TR (ms)**            | 1.4/3.2                | 1.4/3.2     | 1.3/597 |
| **FOV (mm)**              | 350×350                | 350×350     | 350×350 |
| **Image matrix**          | 208×166                | 208×166     | 133×224 |
| **Spatial resolution (mm)**| 1.7×1.7               | 1.7×1.7     | 1.6×1.6 |
| **Temporal resolution (ms)**| 41                    | 41          | 199     |
| **Slice thickness (mm)**  | 6                      | 6           | 6       |
| **Numbers of slices**     | 8                      | 8           | 12      |
| **Slice gap (mm)**        | 3.6–4.8                | 3.6–4.8     | 0.9     |
| **Flip angle (°)**        | 50                     | 50          | 55      |
| **Bandwidth (Hz/pixel)**  | 1145                   | 1145        | 744     |
| **Cardiac phases**        | 25                     | 19–31       | —       |
| **Numbers of breath-holds**| 4                     | 1           | 2-3     |
| **Acceleration factor**   | 3                      | 12.8        | 2       |
| **Number of iterative reconstructions** | — | 80 | — |

CMR, cardiac magnetic resonance; CS, compressed sensing; ECG, electrocardiogram; FOV, field of view; LGE-CMR, late gadolinium-enhanced magnetic resonance; FISP, fast imaging with steady-state free precession; PSIR, phase-sensitive inversion recovery; TE, echo time; TR, repetition time

**Table 2. Patient characteristics**

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| Age (years)                  | 70.4±9.3 |
|----------------------------|----------|
| Men (% of total)           | 20 (54%) |
| Body mass index (kg/m²)    | 24.1±3.5 |
| HR (beats/min)             | 60.2±10.3|
| Cardiovascular risk factors|          |
| Hypertension               | 21 (56%) |
| Dyslipidemia               | 18 (48%) |
| Diabetes mellitus          | 12 (32%) |
| Smoking habit              | 9 (24%)  |
| Family history of CAD      | 3 (8%)   |

Data are expressed as mean ±SD, or N (%).

CAD, coronary artery disease; HR, heart rate; SD, standard deviation

Table 3. Diagnostic performance of p-CS on conventional and CS cine CMR images for detecting myocardial infarction

|                      | Cut off value (%) | Sensitivity (%) | Specificity (%) | PPV (%) | NPV (%) |
|----------------------|-------------------|-----------------|-----------------|---------|---------|
| Conventional cine CMR| -16.4%            | 85 (79–91)      | 79 (76–83)      | 53 (46–59) | 95 (93–97) |
| CS cine CMR          | -16.5%            | 82 (75–88)      | 77 (73–81)      | 49 (43–55) | 94 (92–96) |

All values are presented as % (95% confidence interval)

CMR, cardiac magnetic resonance; CS, compressed sensing; p-CS, peak circumferential strain; NPV, negative predictive value; PPV, positive predictive value

Figures
Figure 1

Scatter plots (A) and Bland–Altman plots (B) of p-CS on conventional cine CMR and CS cine CMR images. In the Bland-Altman plots, the mean difference between estimates obtained by two techniques is indicated by a solid line; the corresponding double standard deviation intervals are indicated by a dashed line. CMR, cardiac magnetic resonance; CS, compressed sensing; p-CS, peak circumferential strain.

Figure 2

p-CS analysis of conventional cine CMR and CS cine CMR images of a 72-year-old man with myocardial infarction. LGE-CMR shows the late gadolinium enhancement in the LV inferior wall, which is suspected to have undergone infarction (A). The p-CS as assessed using LGE-CMR was decreased in the LV inferior wall on both conventional cine CMR (B) and CS cine CMR (C) images. CMR, cardiac magnetic resonance;
Figure 3

Comparisons of p-CS values among remote, adjacent, subendocardial infarcted, and transmural infarcted segments on conventional cine CMR (A) and CS cine CMR (B) images CMR, cardiac magnetic resonance; CS, compressed sensing; p-CS, peak circumferential strain. *p < 0.05
Figure 4

Receiver operating characteristic curve of p-CS for detecting myocardial infarction on conventional cine CMR and CS cine CMR images CMR, cardiac magnetic resonance; CS, compressed sensing; p-CS, peak circumferential strain