












Virtual pacing of a patient's digital twin to predict left ventricular reverse remodelling after cardiac resynchronization therapy

Tijmen Koopsen ¹, Willem Gerrits², Nick van Osta ¹, Tim van Loon ¹, Philippe Wouters ², Frits W. Prinzen ³, Kevin Vernooy ^{4,5,6}, Tammo Delhaas ¹, Arco J. Teske ², Mathias Meine ², Maarten J. Cramer ², and Joost Lumens ^{1*}

¹Department of Biomedical Engineering, CARIM Cardiovascular Research Institute Maastricht, Maastricht University, Universiteitssingel 40, 6200 MD, The Netherlands; ²Department of Cardiology, Division of Heart and Lungs, University Medical Center Utrecht (UMCU), Utrecht, The Netherlands; ³Department of Physiology, CARIM Cardiovascular Research Institute Maastricht, Maastricht University, Maastricht, The Netherlands; ⁴Department of Cardiology, CARIM Cardiovascular Research Institute Maastricht, Maastricht University, Maastricht, The Netherlands; ⁵Department of Cardiology, Maastricht University Medical Center (MUMC), Maastricht, The Netherlands; and ⁶Department of Cardiology, Radboud University Medical Center, Nijmegen, The Netherlands

Received 26 October 2023; accepted after revision 9 January 2024; online publish-ahead-of-print 30 January 2024

Aims

Identifying heart failure (HF) patients who will benefit from cardiac resynchronization therapy (CRT) remains challenging. We evaluated whether virtual pacing in a digital twin (DT) of the patient's heart could be used to predict the degree of left ventricular (LV) reverse remodelling post-CRT.

Methods and results

Forty-five HF patients with wide QRS complex (≥ 130 ms) and reduced LV ejection fraction ($\leq 35\%$) receiving CRT were retrospectively enrolled. Echocardiography was performed before (baseline) and 6 months after CRT implantation to obtain LV volumes and 18-segment longitudinal strain. A previously developed algorithm was used to generate 45 DTs by personalizing the CircAdapt model to each patient's baseline measurements. From each DT, baseline septal-to-lateral myocardial work difference ($MW_{LW-S,DT}$) and maximum rate of LV systolic pressure rise ($dP/dt_{max,DT}$) were derived. Biventricular pacing was then simulated using patient-specific atrioventricular delay and lead location. Virtual pacing-induced changes $\Delta MW_{LW-S,DT}$ and $\Delta dP/dt_{max,DT}$ were correlated with real-world LV end-systolic volume change at 6-month follow-up ($\Delta LVESV$). The DT's baseline $MW_{LW-S,DT}$ and virtual pacing-induced $\Delta MW_{LW-S,DT}$ were both significantly associated with the real patient's reverse remodelling $\Delta LVESV$ ($r = -0.60$, $P < 0.001$ and $r = 0.62$, $P < 0.001$, respectively), while correlation between $\Delta dP/dt_{max,DT}$ and $\Delta LVESV$ was considerably weaker ($r = -0.34$, $P = 0.02$).

Conclusion

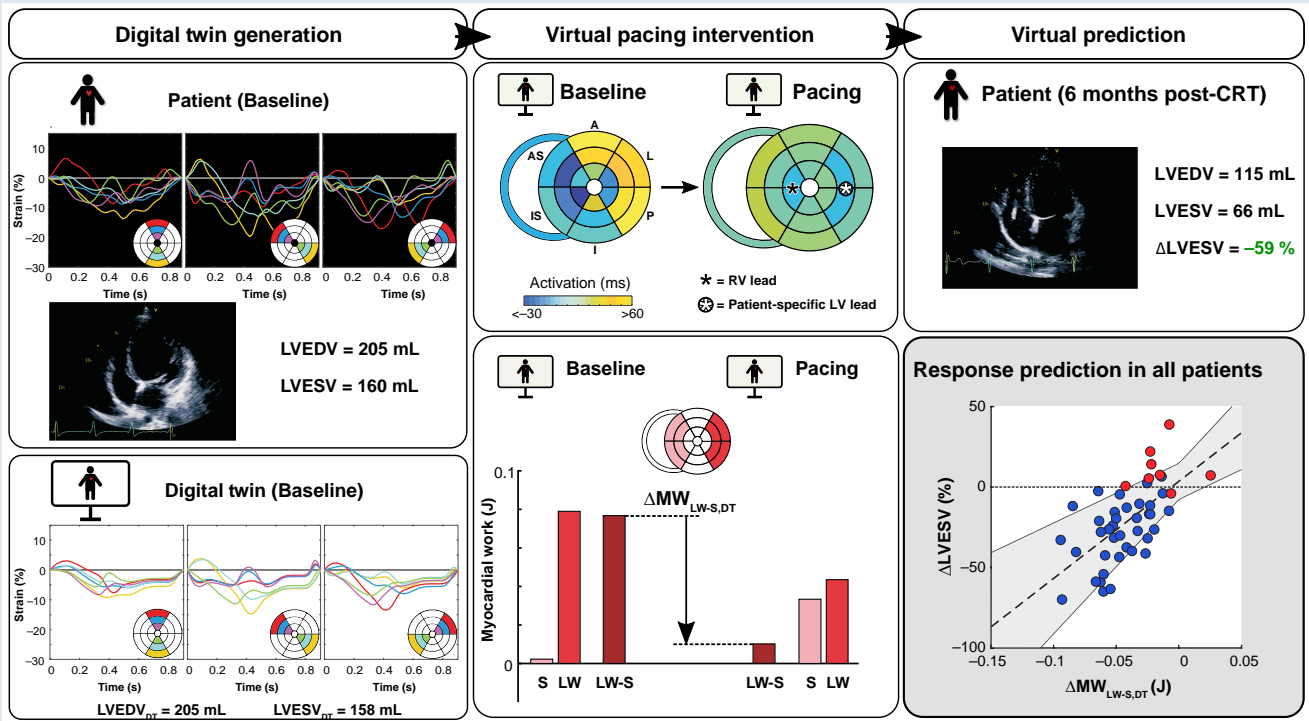
Our results suggest that the reduction of septal-to-lateral work imbalance by virtual pacing in the DT can predict real-world post-CRT LV reverse remodelling. This DT approach could prove to be an additional tool in selecting HF patients for CRT and has the potential to provide valuable insights in optimization of CRT delivery.

* Corresponding author. Tel: +31 43 3881666. E-mail address: joost.lumens@maastrichtuniversity.nl

© The Author(s) 2024. Published by Oxford University Press on behalf of the European Society of Cardiology.

This is an Open Access article distributed under the terms of the Creative Commons Attribution License (<https://creativecommons.org/licenses/by/4.0/>), which permits unrestricted reuse, distribution, and reproduction in any medium, provided the original work is properly cited.

Graphical Abstract



Keywords

In silico clinical trial • Left bundle branch block • CircAdapt • Myocardial work • Patient-specific modelling • Computer modelling and simulation

What's new?

- Imaging-based digital twins were created of 45 heart failure patients treated with cardiac resynchronization therapy.
- Virtual pacing intervention in the digital twins revealed a direct correlation between its mechanical response and the actual degree of left ventricular reverse remodelling observed in patients during follow-up.
- Future investigations should explore the capacity of this digital twin methodology to enhance patient outcomes by predicting the optimal strategy for pacing delivery tailored to each individual.
- The study underscores the potential of the CircAdapt model of the human heart and circulation as a virtual platform for *in silico* (pre-) clinical trials in virtual heart failure patients.

Introduction

The goal of cardiac resynchronization therapy (CRT) is to improve left ventricular ejection fraction (LVEF), reduce symptoms, and improve survival of patients with dyssynchronous heart failure (HF).^{1,2} However, despite the importance of CRT as a HF treatment, a substantial number of implanted patients show no improvements.³ At the same time, CRT is widely underutilized in eligible patients.⁴ Uncertainty of which patients will benefit remains a challenge that affects the full potential of this HF therapy.^{4,5} To improve patient selection, by taking into account more than the standard LVEF and electrocardiographic

characteristics,⁶ a personalized modelling approach using the patient's digital twin (DT) may provide important insights.⁷

The DT is a virtual representation of the patient's heart, obtained from integrating a set of patient-specific measurements into a computer model.⁸ Thereby, the DT describes the unique pathophysiology of the patient and can function as a platform to evaluate the effect of therapeutic interventions. Based on the diagnostic value of myocardial strain measurements in dyssynchronous HF patients,^{9–11} we have previously developed an algorithm that uses left ventricular (LV) strain and volume measurements to generate a DT.¹²

In this study, we evaluate whether virtual pacing of these DTs can be used to predict the degree of post-CRT LV reverse remodelling. Hereto, we retrospectively perform simulations of biventricular pacing as clinically administered in a population of dyssynchronous HF patients, including personalized atrioventricular (AV) delay and lead location. Based on previous findings suggesting that acute recoordination is more strongly associated with long-term LV reverse remodelling than baseline dyssynchrony only,^{13,14} we quantify the pacing-induced reduction of septal-to-lateral work difference in the DT as a potential predictor of LV reverse remodelling, as well as acute haemodynamic improvement.

Methods

Study population

We retrospectively included 45 HF patients with a QRS width ≥ 120 ms and a reduced LVEF ($\leq 35\%$), all previously selected for CRT device implantation. Patients were randomly selected from three different studies^{13,15,16}

based on availability and quality of all three echocardiographic views to allow derivation of 18-segment longitudinal strain. All three studies were approved by the institutional review boards or local medical ethics committees of the participating centres, and informed consent was obtained from all study participants. Characteristics of the patient population are shown in Table 1.

Echocardiography

Routine echocardiographic imaging was performed at baseline and 6 months after CRT device implantation (mean follow-up: 6.4 ± 1.4 months) using a Vivid E9 or E95 Ultrasound system (GE Healthcare, Horten, Norway). All images were digitally stored and analysed offline. Left ventricular end-diastolic and end-systolic volumes were calculated using biplane Simpson's method and were used to calculate the relative change of LV end-systolic volume from baseline to follow-up (Δ LVESV). Good quality baseline two-chamber, three-chamber, and four-chamber echocardiographic acquisitions were used for speckle tracking analysis (EchoPac version 203) to obtain LV 18-segment longitudinal strain. Regions of interest were automatically tracked and manually adjusted to the endo- and epicardial border following the standard recommendations.¹⁸ Timing of mitral valve closure was derived from Doppler flow tracings and selected as the moment of zero-strain reference.

Cardiac resynchronization therapy

Cardiac resynchronization therapy devices were implanted according to local implantation protocols under local anaesthesia.^{13,15,16} Right ventricular (RV) and right atrial leads were placed transvenously at conventional positions. Taking into account adequate pacing thresholds and absence of phrenic nerve stimulation, the LV lead was preferably placed in a lateral or posterolateral vein. Right ventricular leads were positioned preferably in the RV apex. Although no uniform AV and ventriculo-ventricular (VV) delays were mandated, paced AV delay was set to 90–130 ms, and VV delay was set to either 0 or –40 ms (LV first).

Digital twin generation algorithm

For each patient, a DT was automatically generated using a previously developed algorithm that personalized the CircAdapt cardiovascular model.¹² Details on the DT generation algorithm and on the model equations can be found in the [Supplementary material](#). In short, this algorithm used the patient's baseline measurements of LV 18-segment longitudinal strain, end-diastolic volume (EDV), and end-systolic volume (ESV) and was blinded to follow-up data. The algorithm utilized dynamic multi-swarm particle swarm optimization (DMS-PSO) to minimize the mean squared error between measured and simulated LV strain and volumes. To calculate this mean squared error, the difference between measured and simulated strain and strain rate as well as the difference between measured and simulated LVEDV and LVEF were normalized to their expected measurement uncertainties. In total, 75 model parameters describing cardiac output, global LV activation duration, AV delay, and LV regional myocardial constitutive properties were selected based on an objective parameter subset reduction pipeline¹² and optimized to personalize parameter values (*Figure 1*). As a result, the patient's unique DT was obtained.

Digital twin: baseline indices

Maximum rate of LV systolic pressure rise ($dP/dt_{\max,DT}$) and septal-to-lateral myocardial work difference ($MW_{LW-S,DT}$) of the baseline DT were calculated. Segmental MW was calculated as the area of the regional stress-strain loop multiplied with segmental wall volume. Value of $MW_{LW-S,DT}$ was calculated as the absolute difference between average MW of the four basal and mid-ventricular lateral and posterior wall segments and average MW of the four basal and mid-ventricular antero-septal and inferoseptal segments.

Virtual pacing and derived indices

Starting from the baseline DT, a patient-specific biventricular pacing intervention was simulated in which activation propagated radially from the location of the RV and LV leads with an intersegmental delay of 15 ms (*Figure 2*). While the location of the RV lead was similar in all patients and

Table 1 Clinical characteristics of the patient population ($n = 45$)

Age (years)	66 ± 10
Male gender (% , n)	62%, 28
QRS duration (ms)	171 ± 21
LBBB morphology ^a (% , n)	84%, 38
CRT Class I indication (% , n)	82%, 37
Atrial fibrillation (% , n)	11%, 5
Ischaemic heart disease (% , n)	33%, 15
LVEDV (mL)	217 ± 83
LVESV (mL)	172 ± 81
LVEF (%)	23 ± 9
ACE-inhibitor/AT2 (% , n)	93%, 42
Beta-blocker (% , n)	67%, 30
Diuretics (% , n)	96%, 43
Spirinolactone/eplerenone (% , n)	51%, 23

ACE, angiotensin converting enzyme; AT2, angiotensin-2; CRT, cardiac resynchronization therapy; LBBB, left bundle branch block; LVEDV, left ventricular end-diastolic volume; LVEF, left ventricular ejection fraction; LVESV, left ventricular end-systolic volume.

^aBased on the Strauss criteria for LBBB.¹⁷

therefore fixed, LV lead location differed between patients. Therefore, four different patient-specific LV lead locations were simulated, i.e. anterolateral (AL), lateral (L), posterolateral (PL), and posterior (P). Furthermore, AV delay was set to either 130¹⁶ or 100 ms^{13,15} depending on the pacing protocol of the original study. During pacing, activation of the RV wall was assumed to equal the average activation time of the septum. Absolute changes of $MW_{LW-S,DT}$ ($\Delta MW_{LW-S,DT}$) and $dP/dt_{\max,DT}$ ($\Delta dP/dt_{\max,DT}$) were calculated.

Statistical analysis

Analysis was performed using MATLAB R2022b (MathWorks, Natick, MA, USA). The relations between baseline $MW_{LW-S,DT}$, $\Delta MW_{LW-S,DT}$, $\Delta dP/dt_{\max,DT}$, and Δ LVESV were assessed by univariate linear regression analysis with calculation of Pearson's correlation coefficient (r) and associated P -values.

Results

Digital twin generation

The patient's echocardiographic volume and strain data were well reproduced by their DTs. In summary, LVEDV and LVESV differed from the measurements by 8 ± 7 and 5 ± 5 mL, respectively, while average deviation of strain per measurement point during the cardiac cycle was $1.5 \pm 0.7\%$.

Virtual pacing and work redistribution

For a single patient, *Figure 3* shows the LV volumes and regional strain of the baseline DT, as well as its $MW_{LW-S,DT}$ at baseline and after simulating pacing. It can be observed that the septum produced almost zero work at baseline, while the LV lateral wall generated a large amount of positive work. Consequently, $MW_{LW-S,DT}$ had a large positive value at baseline. During pacing, septal work was increased while LV lateral wall work was reduced, leading to a relatively large reduction of $MW_{LW-S,DT}$. During clinical follow-up, the patient demonstrated a Δ LVESV of –59%.

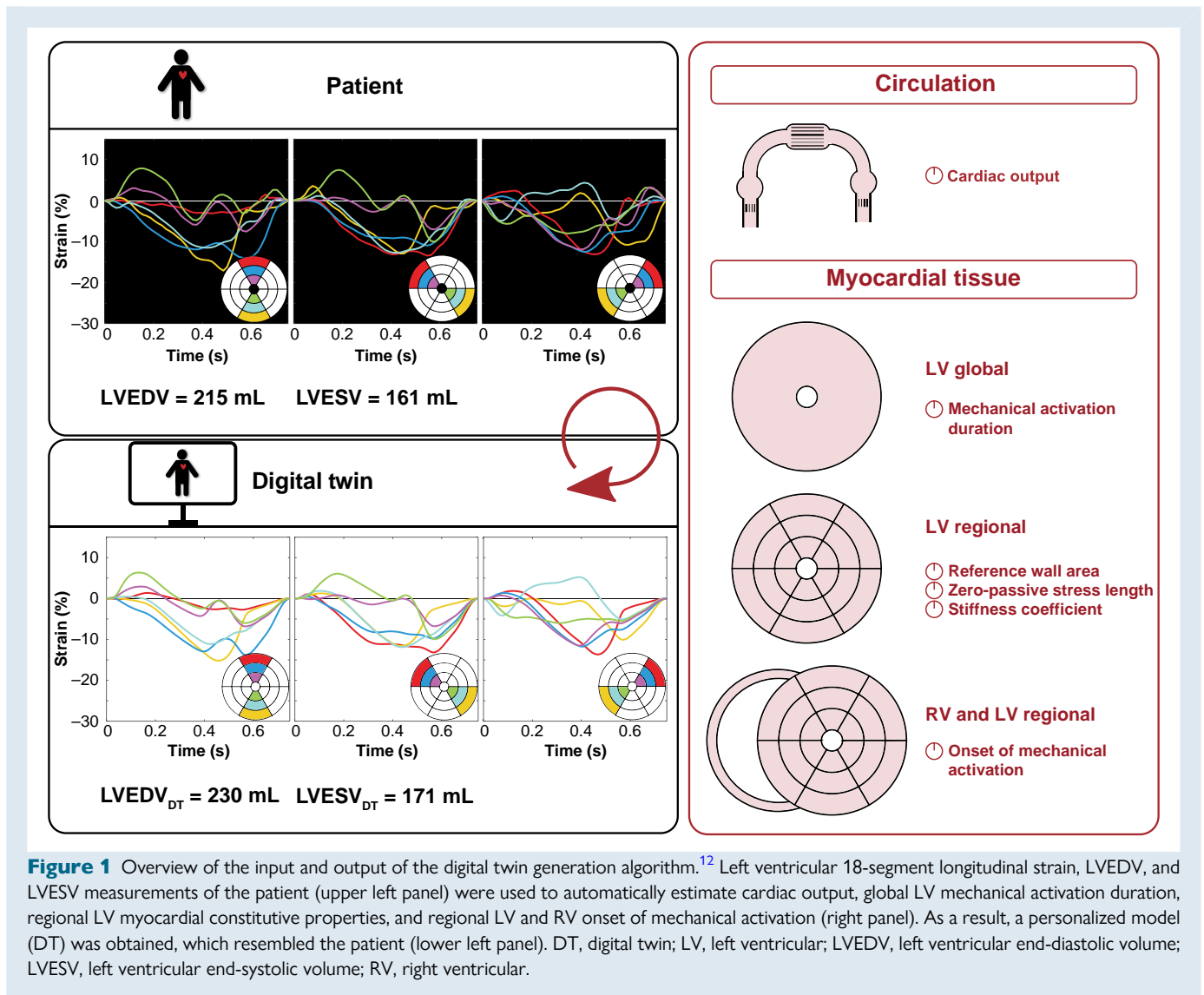


Figure 1 Overview of the input and output of the digital twin generation algorithm.¹² Left ventricular 18-segment longitudinal strain, LVEDV, and LVESV measurements of the patient (upper left panel) were used to automatically estimate cardiac output, global LV mechanical activation duration, regional LV myocardial constitutive properties, and regional LV and RV onset of mechanical activation (right panel). As a result, a personalized model (DT) was obtained, which resembled the patient (lower left panel). DT, digital twin; LV, left ventricular; LVEDV, left ventricular end-diastolic volume; LVESV, left ventricular end-systolic volume; RV, right ventricular.

For the whole patient population, baseline $MW_{LW-S,DT}$ and $\Delta MW_{LW-S,DT}$ correlated moderately with $\Delta LVESV$ ($r = -0.60$, $P < 0.001$ and $r = 0.62$, $P < 0.001$, respectively; Figure 4). At the same time, $\Delta dp/dt_{max,DT}$ correlated weaker with $\Delta LVESV$ ($r = -0.34$, $P = 0.02$).

Discussion

The present study investigated whether virtual pacing in DTs of HF patients, generated using baseline echocardiographic imaging data, could be used to predict the degree of LV reverse remodelling 6 months after CRT. It was found that the virtual pacing-induced reduction of septal-to-lateral work difference in the DT was significantly associated with the patient's LVESV reduction at clinical follow-up. This DT technology could prove to be an additional tool in selecting HF patients for CRT and has the potential to provide valuable insights in optimization of CRT delivery.

Virtual pacing of our DTs revealed that the DT's acute mechanical recoordination was more strongly associated with the patient's LVESV reduction than the DT's acute haemodynamic improvement. This result agrees with the findings of recent clinical studies^{13,14} and suggests that mechanical recoordination can be an effective target for optimizing pacing delivery, also in the DT. Acute haemodynamic improvement represents

the acute success of resynchronization but its predictive value for long-term LV reverse remodelling varies among studies.^{19,20}

The DT's baseline septal-to-lateral work imbalance as well as its change after virtual pacing correlated well with real-world LV reverse remodelling (Figure 4). Firstly, this corroborates previous clinical studies showing that baseline septal-to-lateral work imbalance is a marker of CRT response potential.^{14,21} Secondly, this finding suggests that our virtual pacing interventions were representative for clinical CRT, thereby strengthening the credibility of our DT technology.

Interestingly, the DTs of the few patients with Class II indications according to the 2013 ESC Guidelines on cardiac pacing and CRT²² showed a small reduction of septal-to-lateral work difference by virtual pacing as compared with the DTs of the patients with a Class I indication. Even more interestingly, the virtual pacing effect appeared to have additional prognostic value in patients with a Class I indication for CRT. Within this subgroup, there was significant heterogeneity in the degree of LV reverse remodelling, which was strongly associated with the virtual pacing-induced work redistribution in their DTs.

The comparison of DT-derived indices with measures of long-term LV reverse remodelling is supported by the good correlation of LVESV reduction with mortality.²³ Furthermore, a similar association between acute MW redistribution by CRT and LV reverse remodelling has

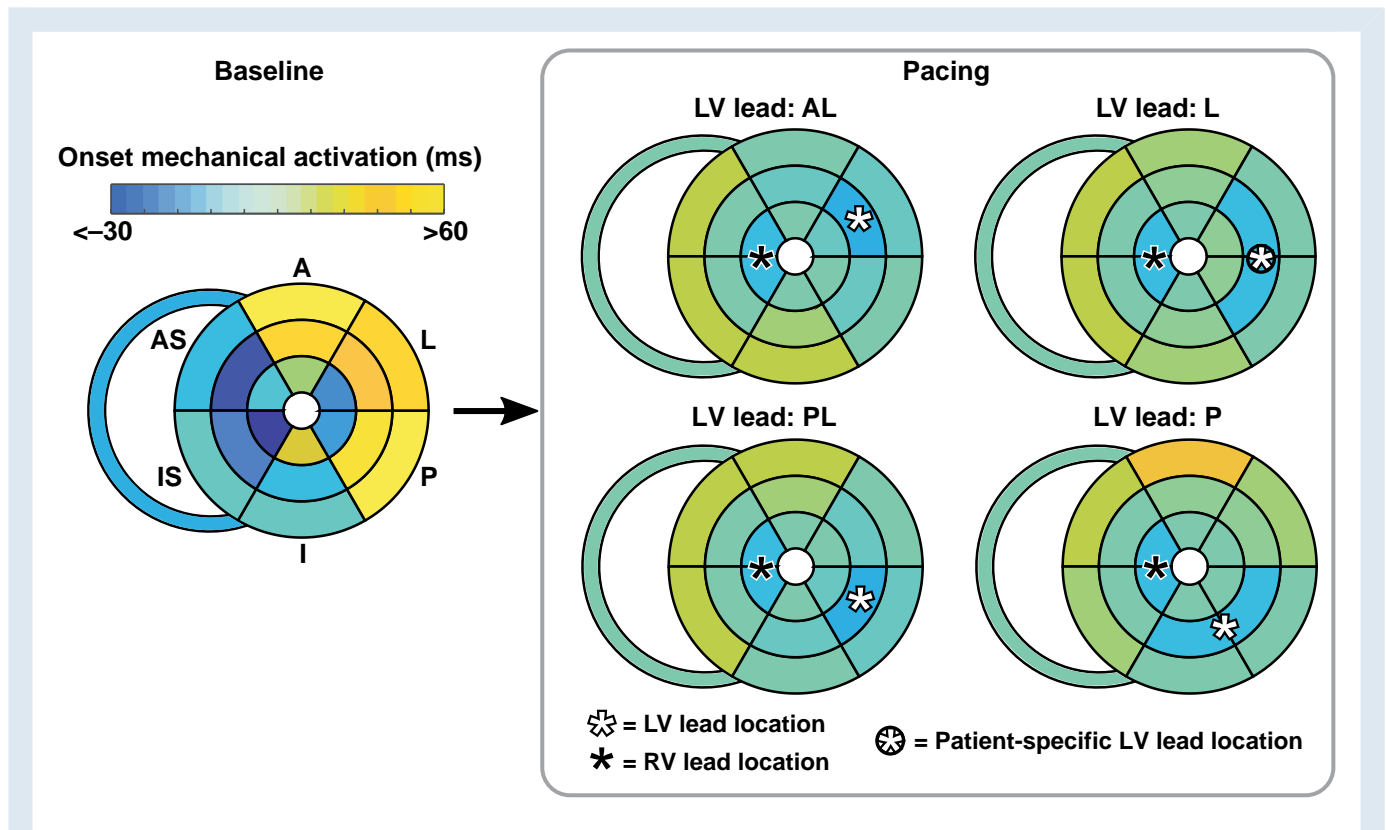


Figure 2 Bullseye plots of RV and LV regional mechanical activation at baseline and during virtual biventricular pacing in the DT. Starting from the baseline DT with automatically estimated onset of activation in the RV wall and LV wall segments (Baseline), a personalized pacing configuration was simulated in which activation was initiated at the location of the RV and patient-specific LV lead, indicated by the black and white asterisks, respectively (Pacing). From the pacing lead locations, activation propagated radially with an intersegmental delay of 15 ms. The patient illustrated here had a L LV lead location, indicated by the circled asterisk. Furthermore, patient-specific AV delay was set to 130 ms based on the pacing protocol of the original study. Note that definition of myocardial segment location differs slightly from that of LV lead location. A, anterior; AL, anterolateral; AS, anteroseptal; I, inferior; IS, inferoseptal; L, lateral; LV, left ventricular; P, posterior; PL, posterolateral; RV, right ventricular.

previously been established in clinical studies.^{13,14} These observations strengthen the current validation approach that was based on LV remodelling indices. However, the correlation of the DT with other response measures, such as hard clinical endpoints, remains to be investigated in future studies.

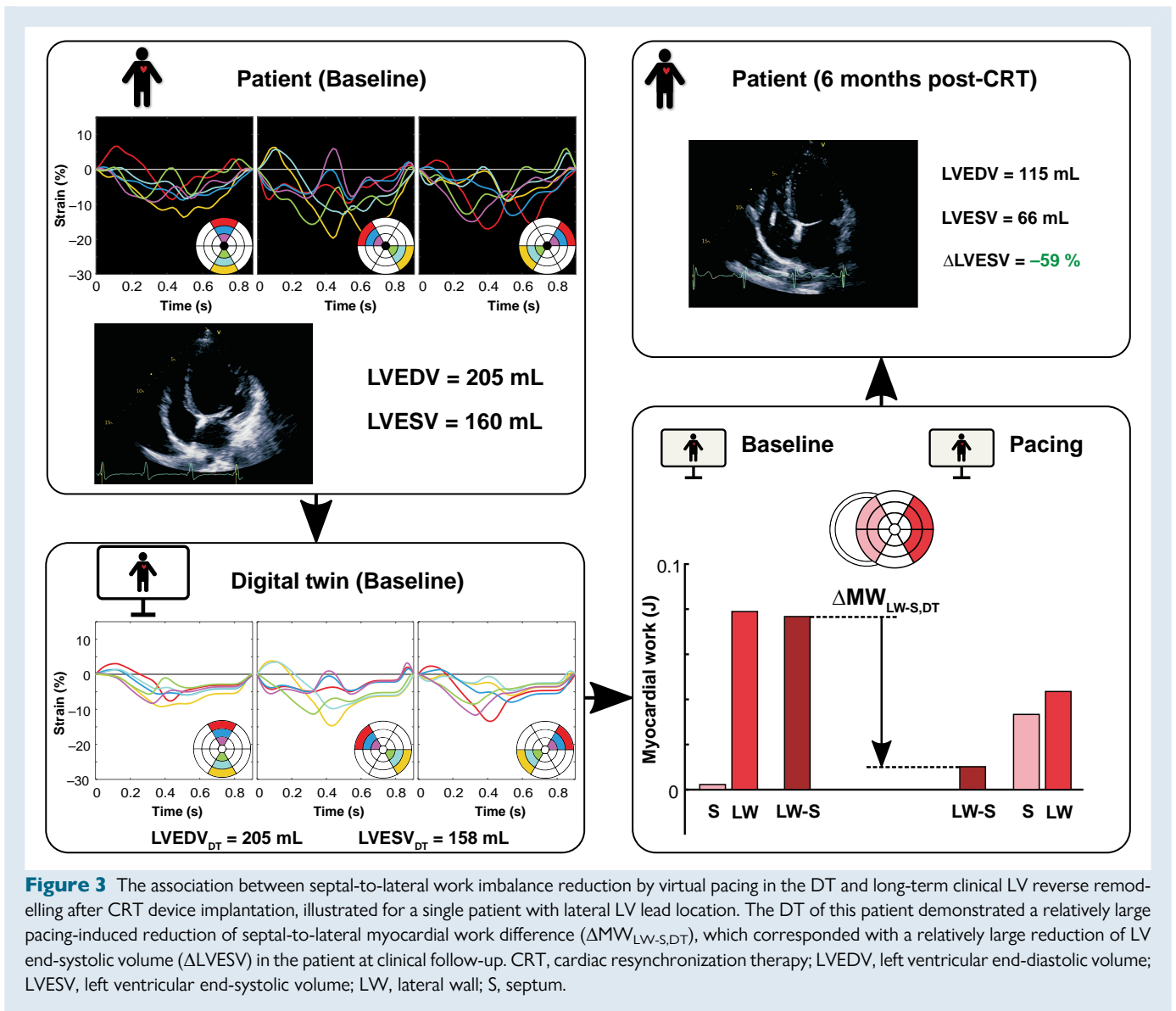
Future studies will also investigate the predictive value of the DT at individual patient level, i.e. by virtual optimization of pacing settings (e.g. sites and delays). Investigating the effects of LV lead location was beyond the scope of the current manuscript and may require a prospective study setup where multiple LV lead locations are tested in the same patient. A previous modelling study suggested that LV lead position sensitively determines the distribution of MW, particularly in hearts with left bundle branch block (LBBB) and myocardial scar, where a balanced position remote from both the scar and the RV lead was associated with the most homogeneous distribution of MW.²⁴

As far as we know, this is the first patient-specific modelling study that calculates the pacing-induced change of septal-to-lateral work imbalance as a predictor of CRT response. Calculation of acute haemodynamic response has been more common and shows conflicting results between modelling studies. In contrast to our findings, acute improvement of LV dP/dt_{max} was predictive for long-term LV reverse remodelling in other personalized heart models.^{25,26} However, another study showed that the simulated shortening of QRS duration, which may be closely related to acute LV dP/dt_{max} improvement,²⁷

correlated poorly with LV reverse remodelling in ischaemic LBBB patients.²⁸

While our current DT approach relied on mechanical input information only, this approach should be considered in combination with available electrical measurements of the patient. Baseline QRS area has been shown to have a strong association with echocardiographic response to CRT^{16,29} and may e.g. be combined in a multivariate analysis with the degree of septal-to-lateral work reduction in the DT. Another index that quantifies electrical dyssynchrony at baseline is the standard deviation of electrical activation times, which has been shown to be predictive for post-CRT LVEF improvement.³⁰ Although regional RV and LV activation times were automatically estimated by the DT algorithm, it should be noted that this activation represents mechanical rather than electrical activation.

Besides the use of electrical measurements to supplement the DT's response prediction, these measurements could also be integrated into the DT itself. While the current model implementation is not capable to do so, integration of electrical information into the DT to better characterize the patient's electrical activation pattern could further improve prediction of the effects of pacing. This electrical characterization may especially be important in ischaemic patients that have abnormal conduction in scarred areas.³¹ When also combining such a DT with models of electrical wave propagation to simulate different pacing conditions,³² the interaction between the patient's electrical and non-electrical tissue characteristics may be more closely investigated.



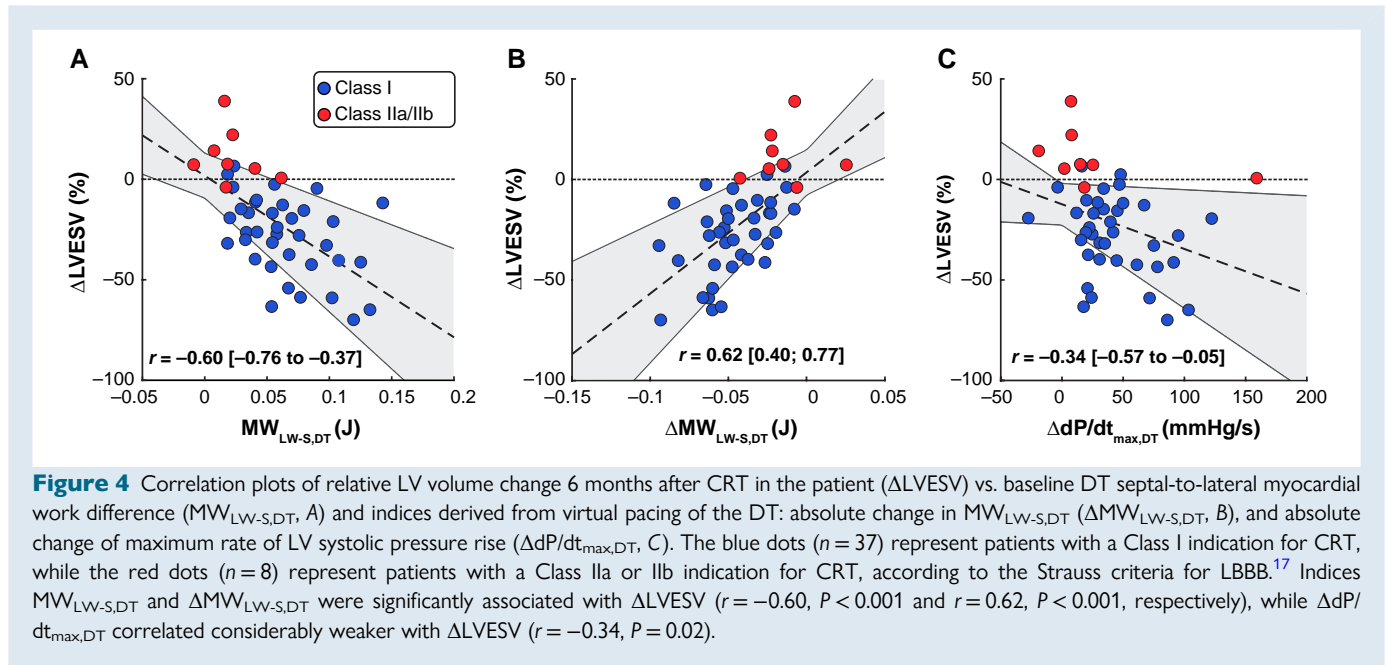
Clinical implications

The DTs generated by our algorithm reveal the unique myocardial disease substrates of the patient and may thereby support CRT patient selection as well as therapy delivery. In the first place, the DT's baseline septal-to-lateral work imbalance provides a marker of CRT response potential. Furthermore, identification of hypocontractile segments in the DT that may indicate ischaemia or scar could guide LV lead location during implantation and potentially present an alternative to the use of late gadolinium enhancement magnetic resonance imaging (MRI) techniques.^{21,33} This study focused on the effects of MW since these have been associated with response to CRT. However, more mechanistic insight might be concealed in the estimated DT tissue characteristics. Future research should try to reveal these insights, which may further improve the predictive power of the DT technology for CRT response.

Based on the interaction of the DT with virtual pacing, moreover, more insight into the mechanisms by which a patient does or does not positively respond to CRT can be obtained. This interaction between the DT and pacing intervention may at the same time provide a platform for therapy optimization in individual HF patients. For example, manipulation of

virtually paced AV and VV delay settings and lead location in the DT could allow personalization of pacing delivery. In contrast to using default pacemaker settings, or guiding lead location by e.g. haemodynamic optimization during the implantation procedure,²⁰ virtual pacing may identify one or several adequate lead locations prior to pacemaker device implantation. Similarly, the technology could facilitate optimization of pacemaker settings during later stages of CRT delivery, potentially based on generation of a new DT at clinical follow-up.

Application of this DT technology requires data obtained using routine echocardiography and could therefore be relatively easily implemented in clinical practice. Prior to being eligible as a clinical tool to support patient selection and therapy delivery, however, further validation of this technology is essential. This validation could explore how potential errors of currently used clinical data propagate in the DT technology and whether other imaging modalities, such as Doppler echo or cine MRI, are of additional or alternative diagnostic value. Generation of the patient's DT currently consumed between 12 and 24 h and was thereby by far more time consuming than the virtual pacing, which was performed within seconds. Evaluation of different pacing conditions using the current approach would therefore be highly feasible.



Limitations

The current study did not compare predictivity of pacing-induced septal-to-lateral work imbalance reduction in the DT with already existing indices to predict the effect of CRT. Our primary focus currently was to present a novel methodology and to demonstrate its potential for clinical application. However, future research should investigate whether this DT technology improves clinical decision-making in dyssynchronous HF patients.

Furthermore, our patient population included eight patients with a Class IIa or IIb indication for CRT. These patients demonstrated a relatively low response to CRT in terms of LV reverse remodelling. Although we found a similarly negative result for these patients by virtual pacing of their DT, i.e. a relatively low workload imbalance reduction, it remains essential to test this technology in more patients with a Class IIa or IIb indication for CRT.

Due to the incomplete documentation of personalized AV and VV delay settings as applied in these patients during clinical CRT, we chose to simulate representative population-based AV and VV delay settings as derived from the original studies. This choice may have influenced the correlations obtained in this study. However, we hypothesize that simulation of these general AV and VV delay settings will rather have worsened than improved these correlations.

Conclusions

We successfully used virtual biventricular pacing interventions in DTs of dyssynchronous HF patients to predict response to CRT. Our results suggest that the reduction of septal-to-lateral work imbalance by virtual pacing in the DT can predict the patient's post-CRT LV reverse remodelling. This DT approach could prove to be an additional tool in selecting HF patients for CRT and has the potential to provide valuable insights in optimization of CRT delivery.

Supplementary material

Supplementary material is available at *Europace* online.

Funding

This study was funded by the Dutch Heart Foundation (grant no. 2015T082 to J.L.), the Dutch Research Council (ZonMw-VIDI grant 016.176.340 to J.L.), and the European Union's Horizon 2020 research and innovation programme under the Marie Skłodowska-Curie grant agreement no. 860745.

Conflict of interest: The authors of this manuscript report the following competing interests: K.V. has been a consultant for Medtronic, Abbott, Boston Scientific, Philips, and Biosense Webster (fees paid to institute) and has received research/educational grants from Medtronic, Abbott, and Biosense Webster (grants paid to institute); J.L. has received research grants from Medtronic and speaker fees from Abbott. Others have nothing to declare.

Data availability

The data underlying this article will be shared on reasonable request to the corresponding author.

References

- Cleland JGF, Daubert J-C, Erdmann E, Freemantle N, Gras D, Kappenberger L *et al.* The effect of cardiac resynchronization on morbidity and mortality in heart failure. *N Engl J Med* 2005;**352**:1539–49.
- Moss AJ, Hall WJ, Cannom DS, Klein H, Brown MW, Daubert JP *et al.* Cardiac-resynchronization therapy for the prevention of heart-failure events. *N Engl J Med* 2009;**361**:1329–38.
- Daubert C, Behar N, Martins RP, Mabo P, Leclercq C. Avoiding non-responders to cardiac resynchronization therapy: a practical guide. *Eur Heart J* 2017;**38**:1463–72.
- Mullens W, Auricchio A, Martens P, Witte K, Cowie MR, Delgado V *et al.* Optimized implementation of cardiac resynchronization therapy: a call for action for referral and optimization of care: a joint position statement from the Heart Failure Association (HFA), European Heart Rhythm Association (EHRA), and European Association. *Eur J Heart Fail* 2020;**22**:2349–69.
- Auricchio A, Prinzen FW. Enhancing response in the cardiac resynchronization therapy patient: the 3B perspective—bench, bits, and bedside. *JACC Clin Electrophysiol* 2017;**3**:1203–19.
- McDonagh TA, Metra M, Adamo M, Gardner RS, Baumbach A, Böhm M *et al.* 2021 ESC Guidelines for the diagnosis and treatment of acute and chronic heart failure. *Eur Heart J* 2021;**42**:3599–726.
- Niederer SA, Lumens J, Trayanova NA. Computational models in cardiology. *Nat Rev Cardiol* 2019;**16**:100–11.
- Corral-Acero J, Margara F, Marciniak M, Rodero C, Loncaric F, Feng Y *et al.* The 'digital twin' to enable the vision of precision cardiology. *Eur Heart J* 2020;**41**:4556–4564.

9. Gorcsan J, Anderson CP, Tayal B, Sugahara M, Walmsley J, Starling RC et al. Systolic stretch characterizes the electromechanical substrate responsive to cardiac resynchronization therapy. *JACC Cardiovasc Imaging* 2019;**12**:1741–52.
10. Leenders GE, Lumens J, Cramer MJ, De Boeck BWL, Doevendans PA, Delhaas T et al. Septal deformation patterns delineate mechanical dyssynchrony and regional differences in contractility: analysis of patient data using a computer model. *Circ Heart Fail* 2012;**5**:87–96.
11. Aalen JM, Remme EW, Larsen CK, Andersen OS, Krogh M, Duchenne J et al. Mechanism of abnormal septal motion in left bundle branch block: role of left ventricular wall interactions and myocardial scar. *JACC Cardiovasc Imaging* 2019;**12**:2402–13.
12. Koopsen T, van Osta N, van Loon T, Meiburg R, Huberts VW, Beela AS et al. Parameter Subset Reduction for Imaging-Based Digital Twin Generation of Patients with Left Ventricular Mechanical Discoordination. <https://www.researchsquare.com/article/rs-3442063/v1>.
13. Wouters PC, Leenders GE, Cramer MJ, Meine M, Prinzen FW, Doevendans PA et al. Acute recoordination rather than functional hemodynamic improvement determines reverse remodelling by cardiac resynchronisation therapy. *International Journal of Cardiovascular Imaging* 2021;**37**:1903–11.
14. Duchenne J, Aalen JM, Cvijic M, Larsen CK, Galli E, Bézy S et al. Acute redistribution of regional left ventricular work by cardiac resynchronization therapy determines long-term remodelling. *Eur Heart J Cardiovasc Imaging* 2020;**21**:619–28.
15. Van Everdingen WM, Zweerink A, Cramer MJ, Doevendans PA, Nguyễn UC, Van Rossum AC et al. Can we use the intrinsic left ventricular delay (QLV) to optimize the pacing configuration for cardiac resynchronization therapy with a quadripolar left ventricular lead? *Circ Arrhythm Electrophysiol* 2018;**11**:1–12.
16. Maass AH, Vernooy K, Wijers SC, Van't Sant J, Cramer MJ, Meine M et al. Refining success of cardiac resynchronization therapy using a simple score predicting the amount of reverse ventricular remodelling: results from the Markers and Response to CRT (MARC) study. *Europace* 2018;**20**:393–393.
17. Strauss DG, Selvester RH, Wagner GS. Defining left bundle branch block in the era of cardiac resynchronization therapy. *American Journal of Cardiology* 2011;**107**:927–34.
18. Voigt JU, Pedrizzetti G, Lysyansky P, Marwick TH, Houle H, Baumann R et al. Definitions for a common standard for 2D speckle tracking echocardiography: consensus document of the EACVI/ASE/Industry Task Force to standardize deformation imaging. *Eur Heart J Cardiovasc Imaging* 2015;**16**:1–11.
19. Bogaard MD, Houthuizen P, Bracke FA, Doevendans PA, Prinzen FW, Meine M et al. Baseline left ventricular dP/dt max rather than the acute improvement in dP/dt max predicts clinical outcome in patients with cardiac resynchronization therapy. *Eur J Heart Fail* 2011;**13**:1126–32.
20. De Roest GJ, Allaart CP, Kleijn SA, Delnoy PPHM, Wu L, Hendriks ML et al. Prediction of long-term outcome of cardiac resynchronization therapy by acute pressure-volume loop measurements. *Eur J Heart Fail* 2013;**15**:299–307.
21. Aalen JM, Donal E, Larsen CK, Duchenne J, Lederlin M, Cvijic M et al. Imaging predictors of response to cardiac resynchronization therapy: left ventricular work asymmetry by echocardiography and septal viability by cardiac magnetic resonance. *Eur Heart J* 2020;**41**:3813–23.
22. Brignole M, Auricchio A, Baron-Esquivias G, Bordachar P, Boriani G, Breithardt O-A et al. 2013 ESC Guidelines on cardiac pacing and cardiac resynchronization therapy: the task force on cardiac pacing and resynchronization therapy of the European Society of Cardiology (ESC). Developed in collaboration with the European Heart Rhythm Association (EHRA). *Europace* 2013;**15**:1070–118.
23. Stankovic I, Belmans A, Prinz C, Ciarka A, Daraban AM, Kotrc M et al. The association of volumetric response and long-term survival after cardiac resynchronization therapy. *Eur Heart J Cardiovasc Imaging* 2017;**18**:1109–17.
24. Huntjens PR, Walmsley J, Ploux S, Bordachar P, Prinzen FW, Delhaas T et al. Influence of left ventricular lead position relative to scar location on response to cardiac resynchronization therapy: a model study. *Europace* 2014;**16**:iv62–8.
25. ichi Okada J, Washio T, Nakagawa M, Watanabe M, Kadooka Y, Kariya T et al. Multi-scale, tailor-made heart simulation can predict the effect of cardiac resynchronization therapy. *J Mol Cell Cardiol* 2017;**108**:17–23.
26. Isotani A, Yoneda K, Iwamura T, Watanabe M, ichi Okada J, Washio T et al. Patient-specific heart simulation can identify non-responders to cardiac resynchronization therapy. *Heart Vessels* 2020;**35**:1135–47.
27. Crozier A, Blazevic B, Lamata P, Plank G, Ginks M, Duckett S et al. Analysis of lead placement optimization metrics in cardiac resynchronization therapy with computational modelling. *Europace* 2016;**18**:IV113–20.
28. Oomen PJA, Khoi T, Seth NP, Kenneth HW, Jeffrey CB. A rapid electromechanical model to predict reverse remodeling following cardiac resynchronization therapy 1 Introduction. *Biomech Model Mechanobiol* 2022;**21**:231–47.
29. van Stipdonk AMW, ter Horst I, Kloosterman M, Engels EB, Rienstra M, Crijns HJGM et al. QRS area is a strong determinant of outcome in cardiac resynchronization. *Circ Arrhythm Electrophysiol* 2018;**11**:e006497.
30. Gage RM, Curtin AE, Burns K V, Ghosh S, Gillberg JM, Bank AJ. Changes in electrical dyssynchrony by body surface mapping predict left ventricular remodeling in patients with cardiac resynchronization therapy. *Heart Rhythm* 2017;**14**:392–9.
31. Lee AWC, Razeghi O, Solis-Lemus JA, Strocchi M, Sidhu B, Gould J et al. Non-invasive simulated electrical and measured mechanical indices predict response to cardiac resynchronization therapy. *Comput Biol Med* 2021;**138**:104872.
32. Meiburg R, Rijks JHJ, Beela AS, Bressi E, Grieco D, Delhaas T et al. Comparison of novel ventricular pacing strategies using an electro-mechanical simulation platform. *Europace* 2023;**25**:1–11.
33. Taylor RJ, Umar F, Panting JR, Stegemann B, Leyva F. Left ventricular lead position, mechanical activation, and myocardial scar in relation to left ventricular reverse remodeling and clinical outcomes after cardiac resynchronization therapy: a feature-tracking and contrast-enhanced cardiovascular magnetic resonance study. *Heart Rhythm* 2016;**13**:481–9.