

DCM, LVNC and transplant

Preparatory course to certification in CMR

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Disclosures of interest



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3.2. Dilated cardiomyopathy (DCM)

- 3.2.1. LGE in DCM
 - 3.2.1.1. Frequency
 - 3.2.1.2. Typical spatial distribution
 - 3.2.1.3. Prognostic importance
- 3.2.2. Relative diagnostic yields for DCM by echo and CMR
- 3.2.3. Potential of CMR to reveal underlying cause of DCM
- 3.2.4. Differential diagnosis in heart failure of unclear aetiology



Dilated cardiomyopathy: DCM



- LV dilatation and globally impaired systolic function
- Diagnosis by exclusion of secondary causes, e.g.
 - Valvular disease
 - Ischaemic heart disease
 - Hypertension
- 20-50% familial



DCM: CMR protocol



- Anatomy
- LV function, RV function (Cine CMR)
- Oedema-sensivite CMR imaging
- Late gadolinium enhancement
- ± mapping techniques



DCM: CMR reporting



- LV and RV volumes, function (global and regional), and mass/wall thickness
- Evidence of mitral regurgitation
- Presence and pattern of late gadolinium enhancement
- Evidence of LV thrombus



Frequency and typical spatial distribution of LGE in DCM



TABLE 3. Findings of Contrast-Enhanced CMR

•			
	DCM	Ischemic LV	P
Enhancement, n (%)	26 (41)	27 (100)	< 0.001
Average gadolinium score	15 ± 33	66 ± 46	< 0.001
Gadolinium score, % of patients			
0	59	0	
1–50	31	37	
50–100	8	39	
>100	2	24	
Enhancement location, n (%)			
Absent	37/63 (59)	0	
Endocardial	8/26 (13)	27/27 (100)	
Midwall	18/26 (28)	0	

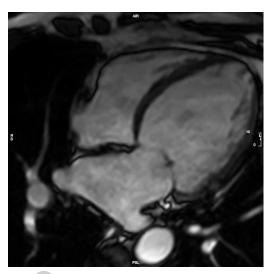
McCrohon JA et al. Circulation. 2003

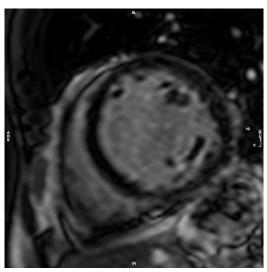


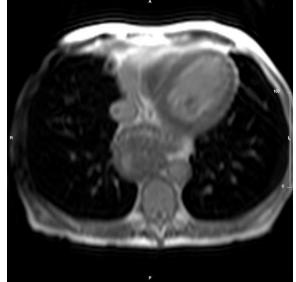
DCM: CMR and prognosis



- LVEF <35%
- Presence and extent of mid-wall fibrosis







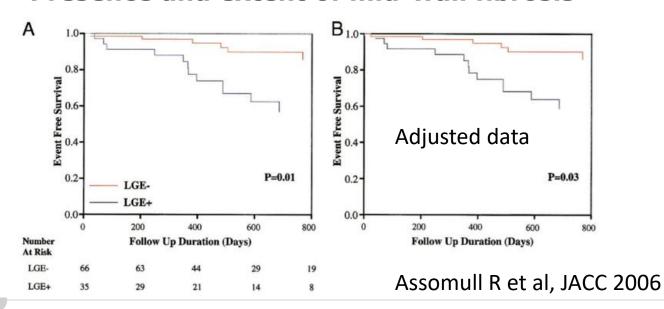


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DCM: CMR and prognosis



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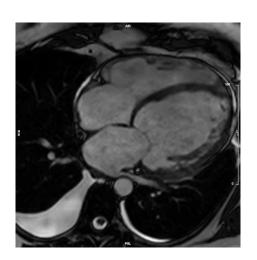


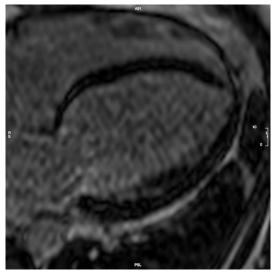


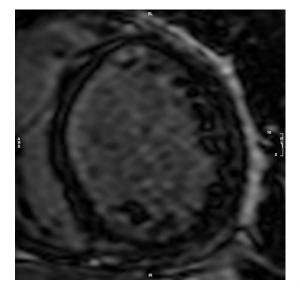
DCM: CMR and prognosis



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Potential of CMR to reveal underlying cause of DCM



Assomull et al LGE-CMR as a Gatekeeper to Angiography in HF 1353

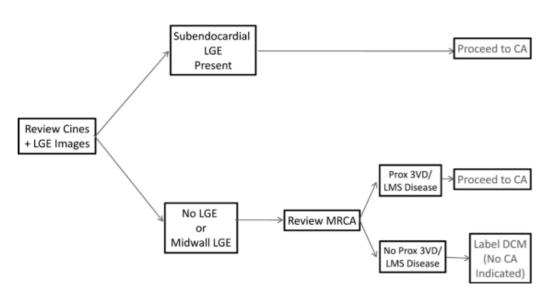


Figure 1. Predefined decision algorithm for the cardiac magnetic resonance consensus panel to decide whether to proceed to invasive x-ray coronary angiography (CA). The algorithm states that the presence of subendocardial late gadolinium enhancement (LGE) should trigger the decision to proceed to CA. In cases when subendocardial LGE is not present, magnetic resonance coronary angiography (MRCA) images should be reviewed before deciding whether CA is required. The review of MRCA should exclude proximal severe 3-vessel disease (Prox 3VD) or left main stem (LMS) disease before a scan be labeled dilated cardiomyopathy (DCM).

Assomull R et al. Circulation. 2011



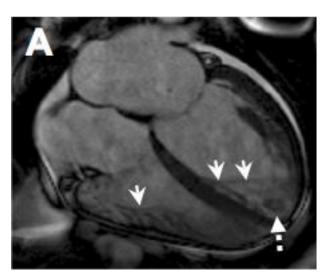


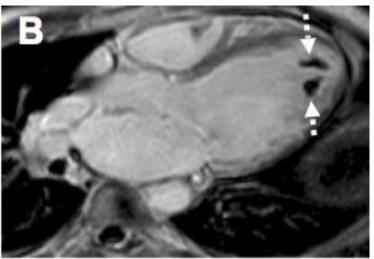
- 3.5. Non-compaction cardiomyopathy (NCP)
 - 3.5.1. Criteria for NCP (including differences in echo and CMR criteria)
 - 3.5.2. Relative diagnostic yields for NCP by echo and CMR
 - 3.5.3. LGE in NCP



LVNC: Left ventricular non-compaction







Petersen et al Heart 2005

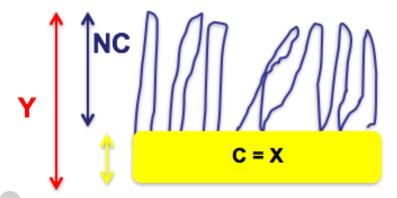
- Characterised by prominent myocardial trabeculations and deep inter-trabecular recesses that communicate with the ventricular cavity
- Can lead to chronic cardiac failure, malignant arrhythmias and thromboembolic events



Diagnostic echocardiography criteria: LVNC



- Jenni criteria: NC/C>2 in systole
- Chin criteria: X/Y=C/(NC+C) ≤0.5
- Stollberger criteria: > 3 trabeculations protruding the left ventricular wall, apically to the papillary muscles, visible in a single image plane



Jenni R et al. Heart 2001 Chin TK et al. Circulation 1990 Stollberger C et al. AM J Cardiol 2002

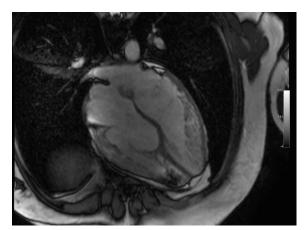


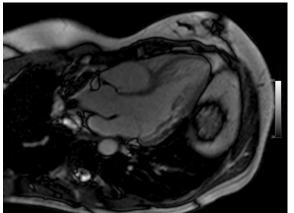
Diagnostic CMR criteria: LVNC

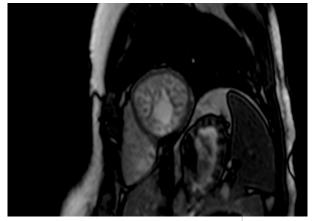


	Jenni	Petersen	Jacquier	Stacey	Captur
Modality	echo	CMR	CMR	CMR	CMR
Criteria	N/C in parasternal short- axis views Colour Doppler evidence of deep perfused intertrabecular recesses Decreased thickening and hypokinesia present within, but not limited to, the non-compacted segments	two layered myocardium long axis SSFP cine measured at the most pronounced trabeculations measurement perpendicular to compacted myocardium	short axis SSFP fines to obtain total LV mass same used to obtain compacted myocardial mass difference between first and second provides the trabecular mass papilary muscles included in the myocardial mass	-apical short-axis views 16 to 24 mm from the true apical slice -region with the largest NC/C ratio	- short axis views - loss of base-to-apex FD gradient
Cardiac phase	End-systole	End-diastole	End-diastole	End-systole	End-diastole
Definition	NC/C ratio > 2	NC/C ratio > 2.3	Trabecular mass > 20%	NC/C ratio ≥ 2	FD ≥1.30





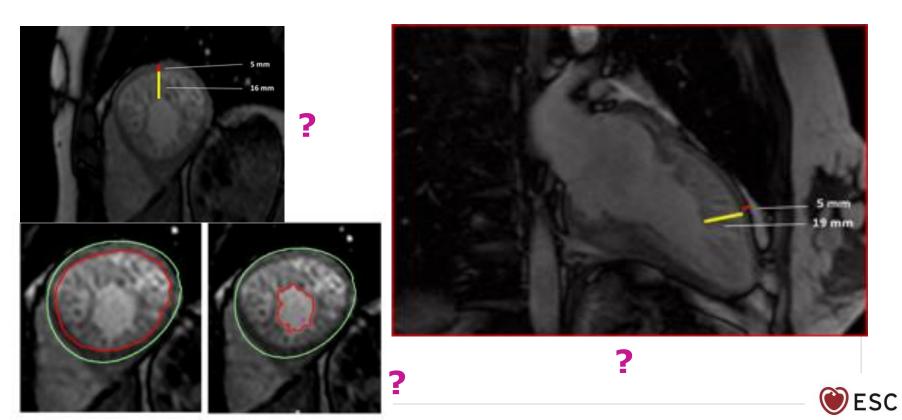






LVNC: CMR analysis approaches

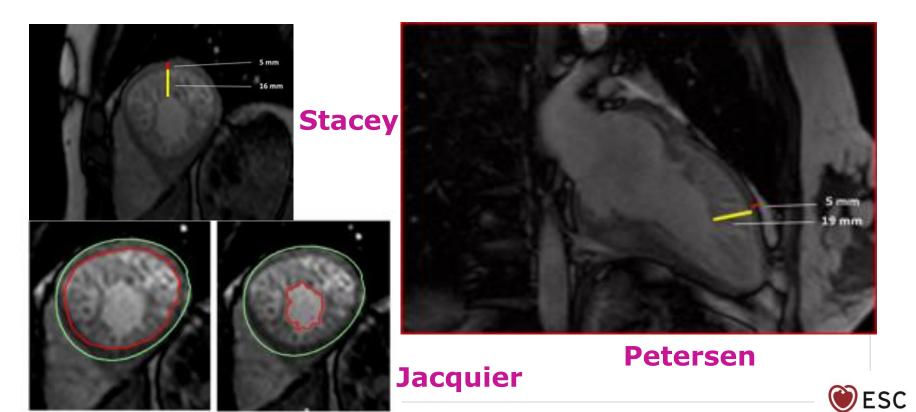




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LVNC: CMR analysis approaches





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Measure of endocardial complexity: Fractal dimension



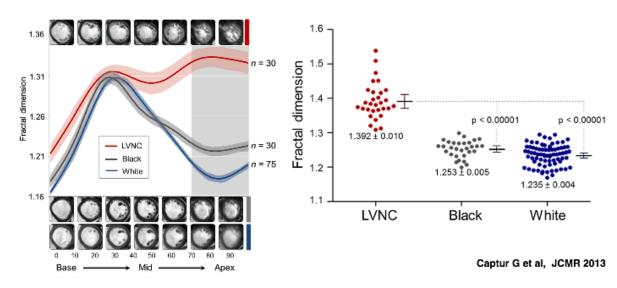


Table 2 Accuracy statistics for the fractal method and comparator CMR criteria.

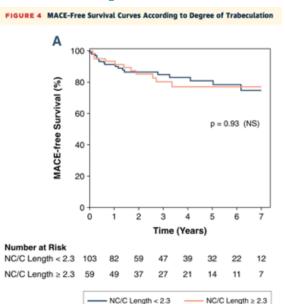
Accuracy statistic*	Max. Apical FD	Global LV FD	Perimetry	Petersen	Jacquier
Sensitivity (95% CI, UL-LL)	100 (88-100)	83 (61-92)	70 (48-85)	70 (46–87)	95 (73–100)
Specificity (95% CI, UL-LL)	100 (97-100)	86 (78-92)	91 (86-97)	98 (85-100)	53 (36-68)

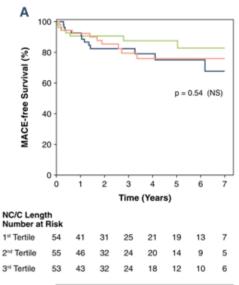
Captur G et al, JCMR 2013



Prognosis: trabeculations and DCM/LVNC







— 3rd Tertile

Amzulescu M-S et al. Prognostic Impact of Hypertrabeculation and Noncompaction Phenotype in Dilated Cardiomyopathy: A CMR Study. JACC Cardiovasc Imaging. 2015 Jul 8;8(8):934–46.

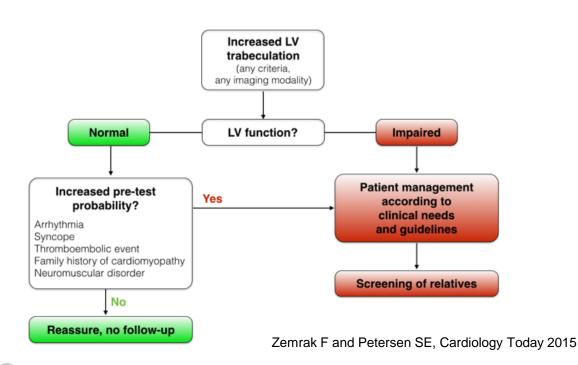


NC/C Length

— 1st Tertile

Management algorithm for LVNC







LGE and LVNC



- Evidence free zone
- Non-ischaemic pattern of LGE
- Ischaemic pattern of LGE
 - if coexisting coronary artery disease
 - If thromboembolic myocardial infarction





- 3.7. Transplant Cardiomyopathy
 - 3.7.1 Sequences
 - 3.7.2 Clinical importance of CMR imaging



Heart transplant: Role of CMR in follow-up



Anatomy:

- 'Double left atrium' appearance with older surgical technique
- Pericardial effusion common in early phase

Function:

- EF is usually preserved but abnormalities in strain and long axis function.
- Significant TR (valve injury due to repeated RV biopsy as part of surveillance for rejection)
- LVH and diastolic impairment (long term)

Tissue characterisation:

- Evidence of myocardial oedema and globally elevated native T1 and T2 times (during first 5 months)
- LGE (usually non-ischaemic pattern) -> likely due to cumulative injury from rejection episodes



Heart transplant: Role of CMR in follow-up



Fig. 1 a, b Left atrial anastomosis. c Bicaval anastomosis. Stars represent superior vena cava anastomosis; triangles represent inferior vena cava anastomosis



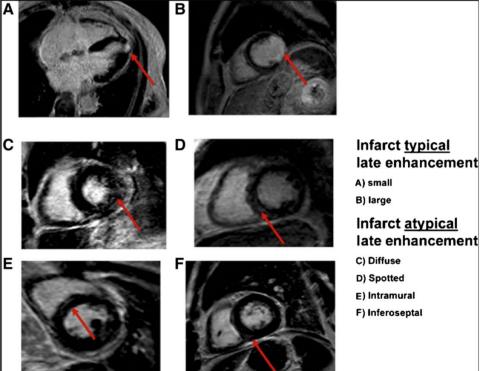
Reid, A. B., Waldron, N., Schmitt, M., & Miller, C. A. (2015). The Value of Cardiovascular Magnetic Resonance in Heart Transplant Patients. Current Cardiology Reports, 17(7), 612. http://doi.org/10.1007/s11886-015-0612-x



Heart Transplant:

Role of CMR in follow-up

Fig. 2 Patterns of late gadolinium enhancement seen in cardiac transplants. (Reproduced with permission from Steen H, Merten C, Refle S, et al: Prevalence of different gadolinium enhancement patterns in patients after heart transplantation. J Am Coll Cardiol 2008, 52:1160-1167) [8•]



Reid, A. B., Waldron, N., Schmitt, M., & Miller, C. A. (2015). The Value of Cardiovascular Magnetic Resonance in Heart Transplant Patients, Current Cardiology Reports, 17(7), 612. http://doi.org/10.1007/s11886-015-0612-x



EACVI

European Association of Cardiovascular Imaging

Graft disease: Role of CMR



Primary graft failure

Severe LVSD (within hours/days)

Acute rejection

- Elevated T2 time (oedema); Increased EGE relative signal intensity
- Caution with interpretation T2 time usually elevated in first 6/12

Allograft vasculopathy

- CMR useful in detecting silent infarcts (ischaemic LGE pattern)
- Worse strain rate (from tagging sequences)

Other consideration

- Erratic response to adenosine in stress perfusion (due to denervation)
- Increased incidence of solid organ tumours and haematological malignancy (Careful scrutiny of extra-cardiac images)





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