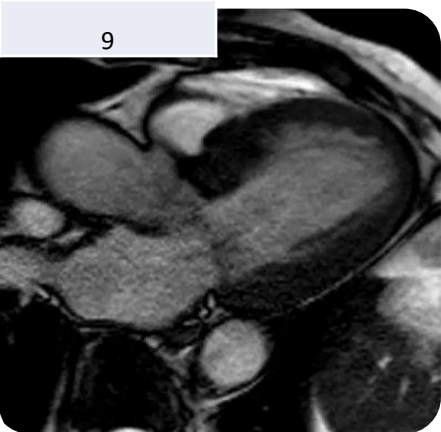


# Yo pediría una cardioRM para...

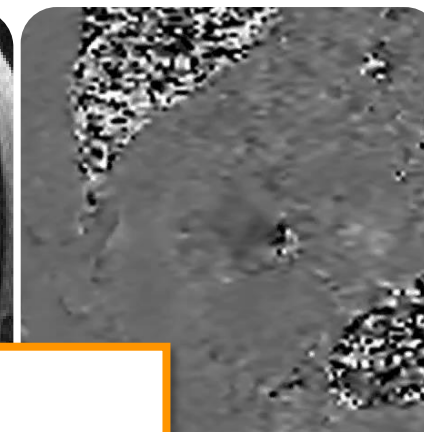
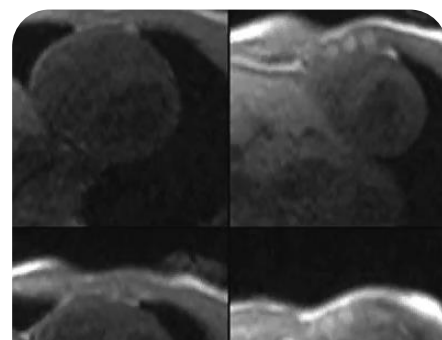
Dr. Alicia M. Maceira, MD, PhD, FESC, FEACVI  
Unidad Cardiovascular, ASCIRES Grupo Biomédico  
Dpto de Medicina, Facultad de CCSS, UCH-CEU  
Valencia



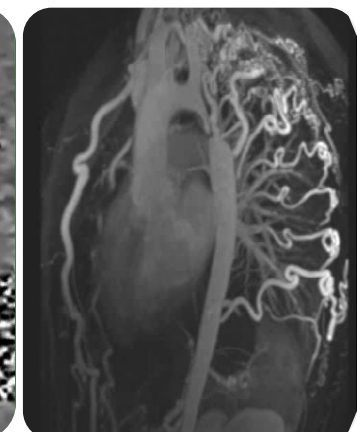
SSFP Cine



TSE



Contraste de fase

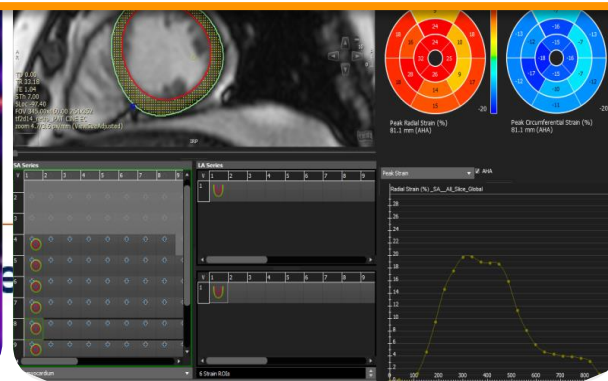
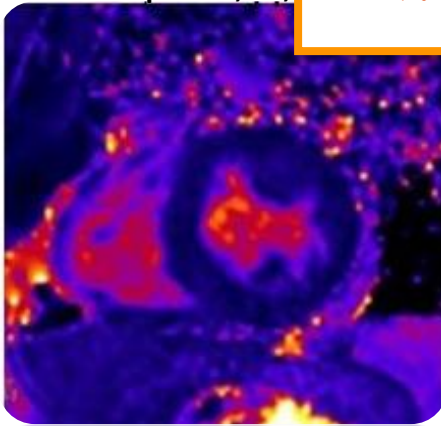


Angiografía RM 3D

## YO PEDIRÍA CRM PARA....

1. Medición de dimensiones y función sistólica VI y VD
2. Estudio de cardiopatía isquémica
3. Miocardiopatías (MCD, MCH, MCR, MCA), miocarditis, d.d. corazón de atleta
4. Pericardiopatías, tumores, trombos
5. C. Congénitas
6. Valvulopatías complejas
7. Patología de grandes vasos

Mapeo T1, T2,



Leiner et al. *J Cardiovasc Magn Reson* (2020) 22:76  
<https://doi.org/10.1186/s12968-020-00682-4>

Journal of Cardiovascular  
Magnetic Resonance

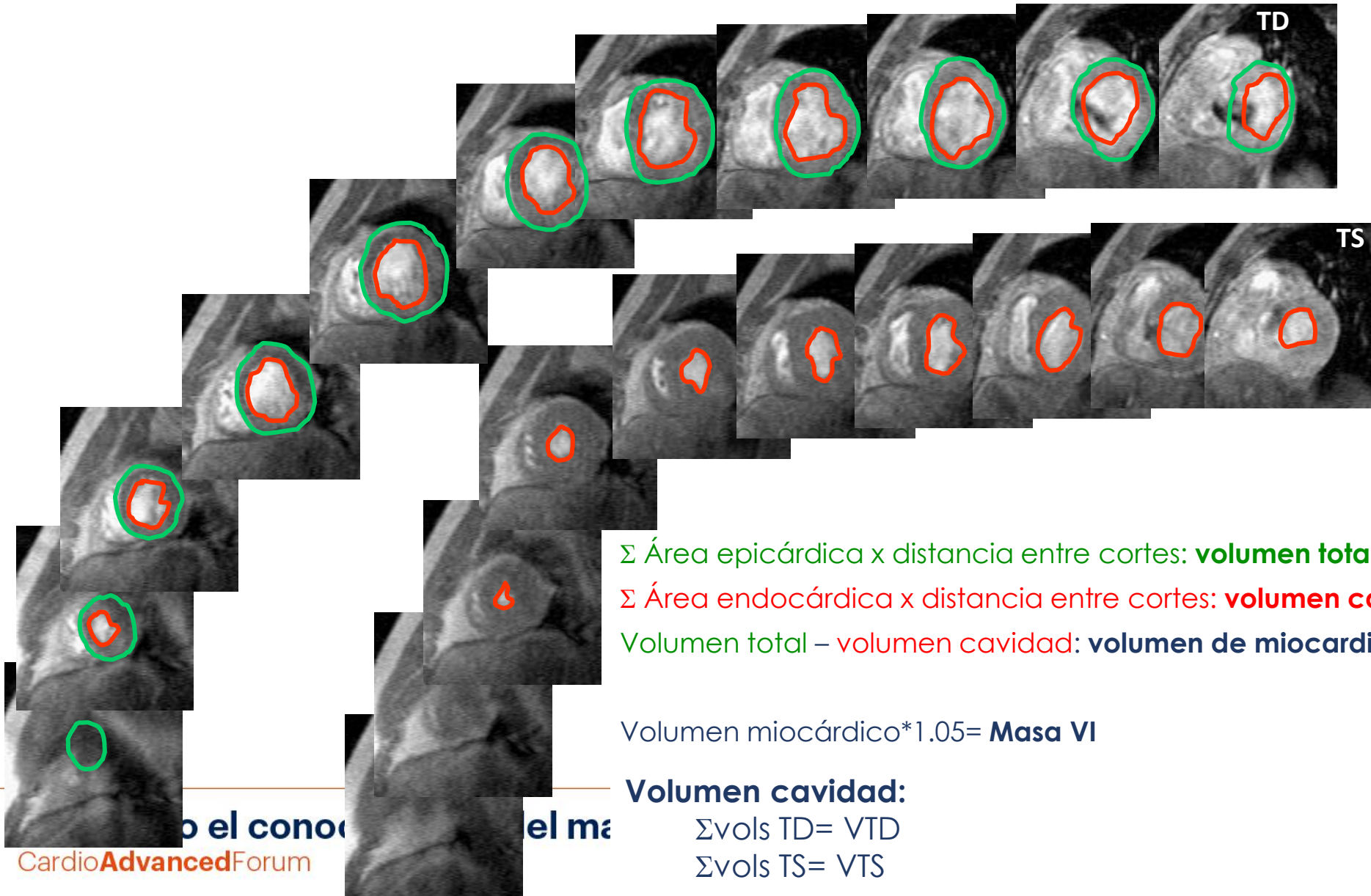
REVIEW

Open Access

SCMR Position Paper (2020) on clinical indications for cardiovascular magnetic resonance



# 1. Estudio de dimensiones y función ventricular



$\Sigma$  Área epicárdica x distancia entre cortes: **volumen total**

$\Sigma$  Área endocárdica x distancia entre cortes: **volumen cavidad**

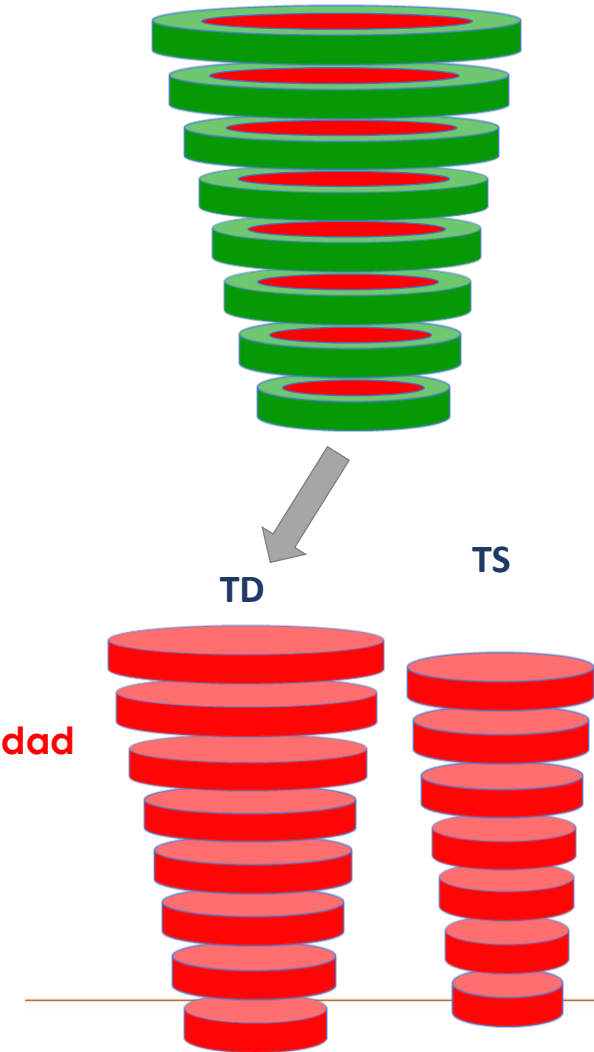
Volumen total – volumen cavidad: **volumen de miocardio**

Volumen miocárdico \* 1.05 = **Masa VI**

**Volumen cavidad:**

$\Sigma$ vols TD = VTD

$\Sigma$ vols TS = VTS



Maceira, JCMR 2006

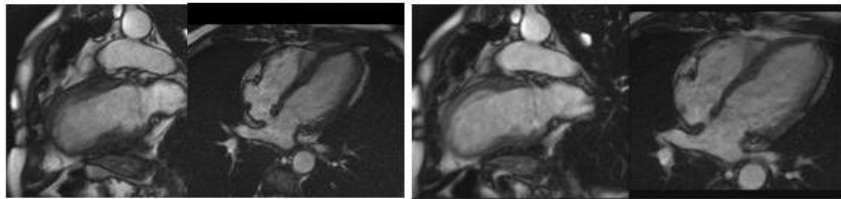


# 2. Cardiopatía isquémica

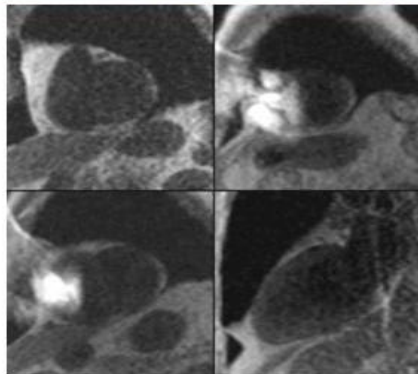
68% de nuestras indicaciones

## Detección de isquemia

Dobutamina



Vasodiladores (dipiridamol/adenosina/regadenoson)

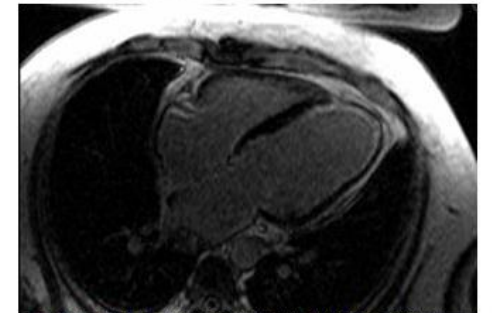
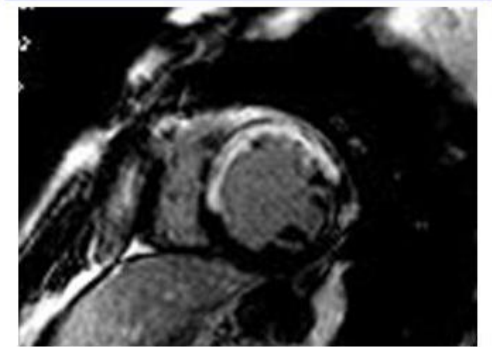


**Table 3 Indications for CMR in coronary artery disease**

Indication	Class
1. Acute coronary syndromes	I
2. Chronic coronary artery disease	I
3. Myocardial infarction with non-obstructive coronary arteries (MINOCA)	I
4. Coronary artery anomalies	II

## Detección de necrosis

## Captación tardía contraste

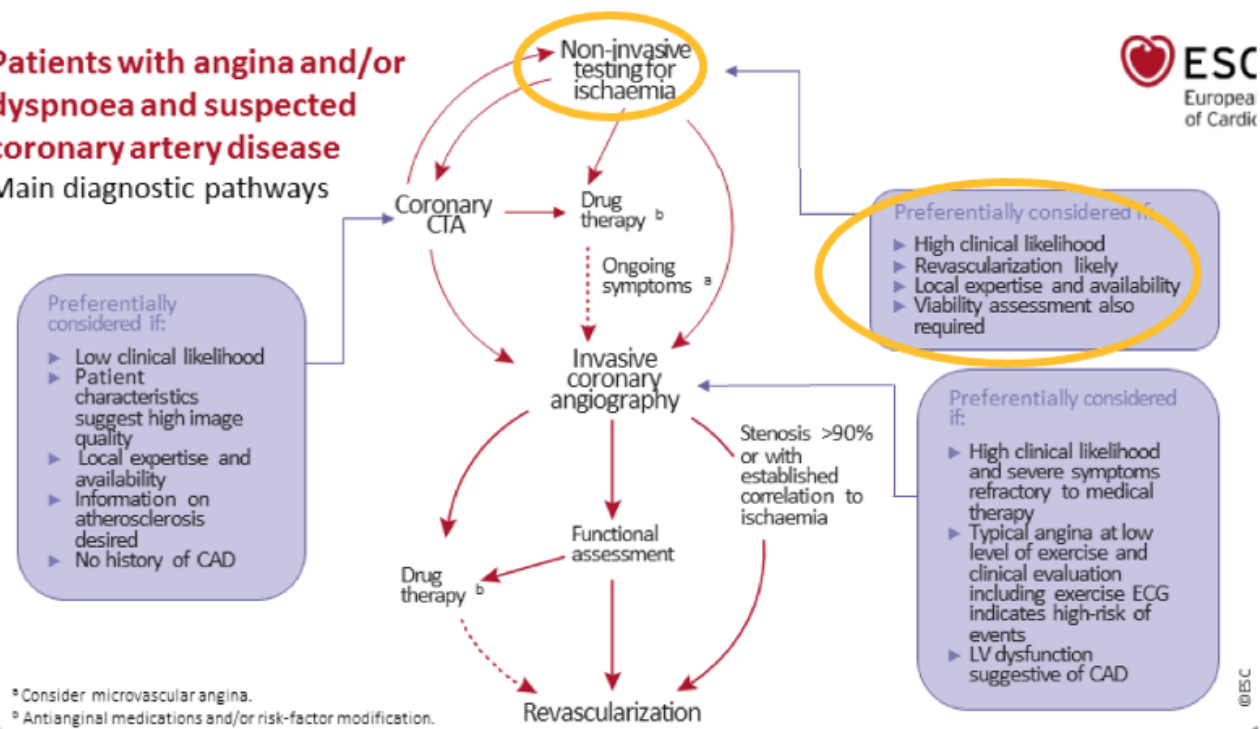


## Selección tipo PRUEBA INICIAL NO INVASIVA según:

1. Probabilidad clínica EAC
2. Precisión Dx del test
3. Características del paciente (FC, obesidad, colaboración)
4. Caract. del medio (experiencia local, disponibilidad)



**Patients with angina and/or dyspnoea and suspected coronary artery disease**  
Main diagnostic pathways



<sup>a</sup> Consider microvascular angina.

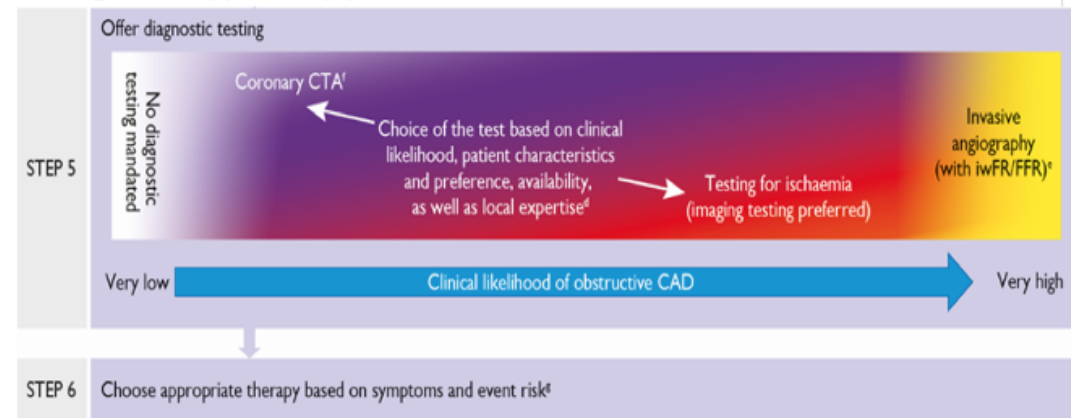
<sup>b</sup> Antianginal medications and/or risk-factor modification.

[www.escardio.org/guidelines](http://www.escardio.org/guidelines)

ESC Guidelines on the diagnosis and management of chronic coronary syndromes  
(European Heart Journal 2019; 10.1093/eurheartj/ehz425)

**Patients with angina and/or dyspnoea and suspected coronary artery disease**

Diagnostic approach (2)



ESC guidelines CCS. EHJ 2019

Liderando el conocimiento del mañana

CardioAdvancedForum

# DETECCIÓN DE ISQUEMIA\_CRM DE ESTRÉS



## 2018 ESC/EACTS Guidelines on myocardial revascularization

### Indications for revascularization in patients with stable angina or silent ischaemia

Extent of CAD (anatomical and/or functional)		Class <sup>a</sup>	Level <sup>b</sup>
<b>For prognosis</b>	Left main disease with stenosis >50%. <sup>c 68-71</sup>	I	A
	Proximal LAD stenosis >50%. <sup>c 62,68,70,72</sup>	I	A
	Two- or three-vessel disease with stenosis >50% with impaired LV function (LVEF ≤35%). <sup>c 61,62,68,70,73-83</sup>	I	A
	Large area of ischaemia detected by functional testing (>10% LV) or abnormal invasive FFR. <sup>d 24,59,84-90</sup>	I	B
	Single remaining patent coronary artery with stenosis >50%. <sup>c</sup>	I	C
<b>For symptoms</b>	Haemodynamically significant coronary stenosis <sup>c</sup> in the presence of limiting angina or angina equivalent, with insufficient response to optimized medical therapy. <sup>e 24,63,91-97</sup>	I	A

**Table 6** Definitions of high event risk for different test modalities in patients with established chronic coronary syndromes<sup>a 102-104</sup>

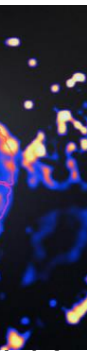
Exercise ECG	Cardiovascular mortality >3% per year according to Duke Treadmill Score
SPECT or PET perfusion imaging	Area of ischaemia ≥10% of the left ventricle myocardium
Stress echocardiography	≥3 of 16 segments with stress-induced hypokinesia or akinesia
<b>CMR</b>	≥2 of 16 segments with stress perfusion defects or ≥3 dobutamine-induced dysfunctional segments
<b>Coronary CTA or ICA</b>	Three-vessel disease with proximal stenoses, LM disease, or proximal anterior descending disease
Invasive functional testing	FFR ≤0.8, iwFR ≤0.89

89 (85-92)

87 (83-91)

7.10 (5.07-9.95)

0.13 (0.09-0.18)



Tabl  
func

Te

Anatr

Exi

Str

Co

SPI

PE

Str

Funct

Co

SPI

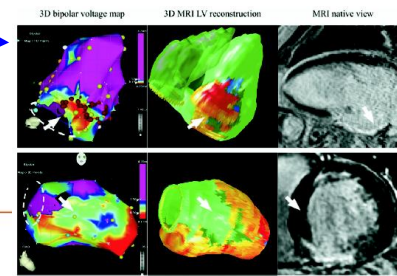
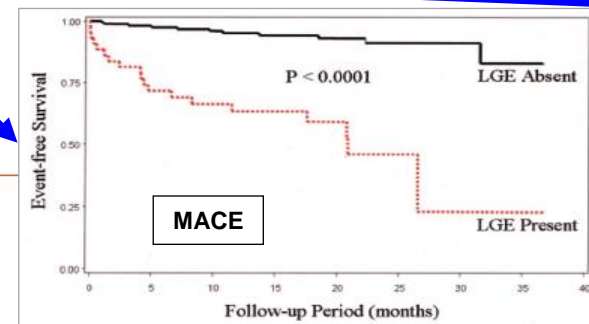
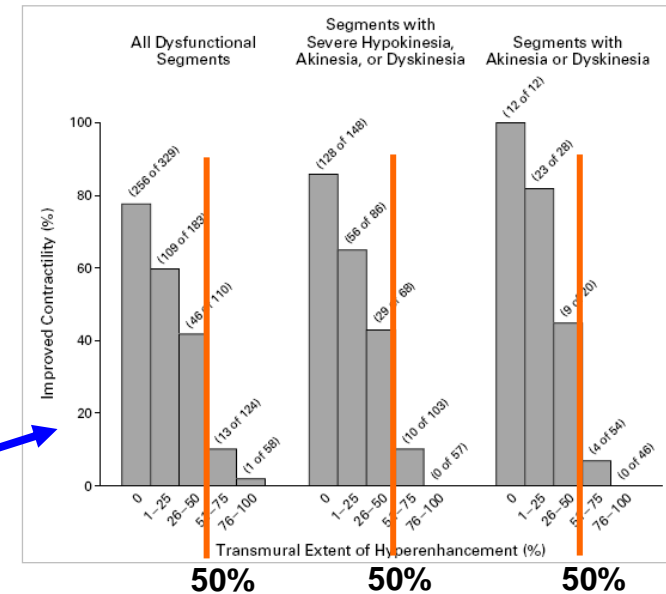
PE

Stress CMR

# CRM EN ESTUDIO DE VIABILIDAD

El miocardio viable recupera su función contráctil tras la revascularización  
**Criterio CRM: transmuralidad de la necrosis  $\leq 50\%$  del grosor parietal**

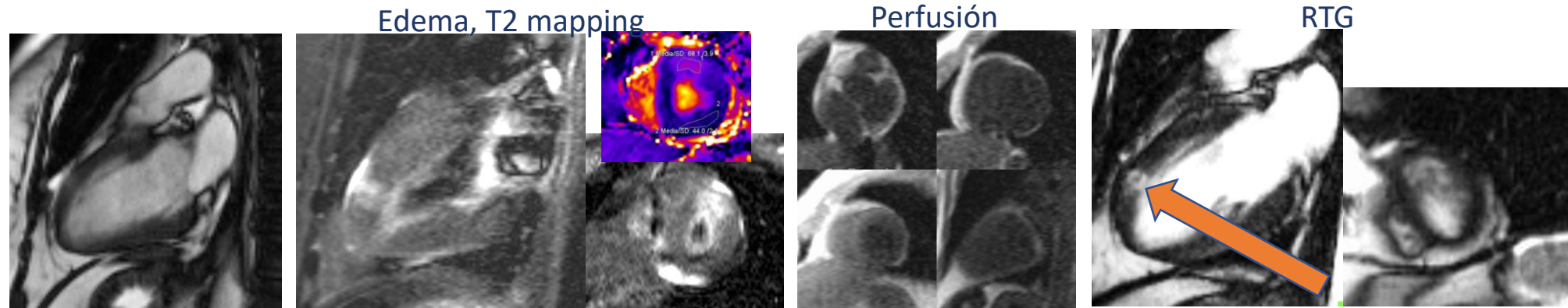
1. Morfología	grosor telediastólico
2. Cines	FE VI y VD alteraciones segmentarias de la contractilidad enr. reposo / dobutamina
3. STIR	detección de edema
4. Perfusión en reposo	obstrucción microvascular
5. Realce tardío de gadolinio	detección de necrosis transmuralidad predicción de eventos, arritmias
(Torsión, velocidad mioc.)	tagging reposo / dobutamina
((Espectroscopia	metabolismo))



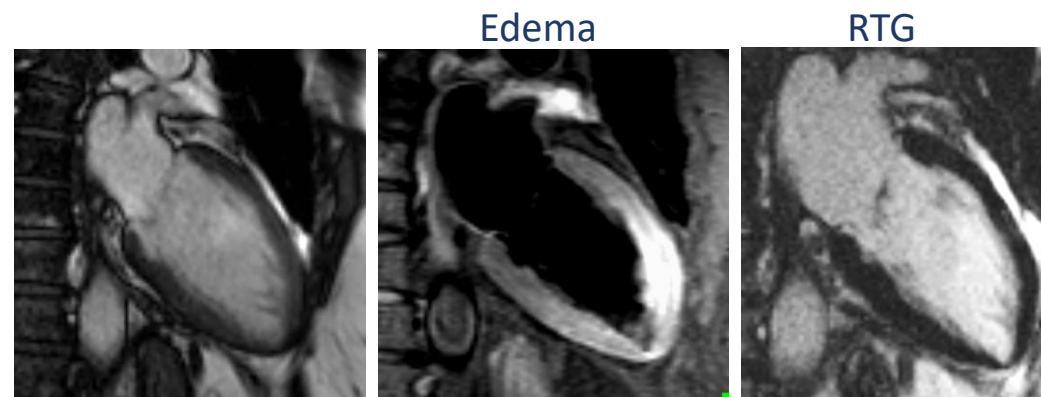


# CRM EN D.D. SCA

**A** L.F.G., Mujer 54 años, HTA, exfumadora. Ingreso por dolor torácico, elev. ST, elev. enzimática Coronarias normales  
Miocarditis vs IAM con coros normales



**B** P.R.P., Mujer 58a, HTA, exfumadora. Idéntica H<sup>a</sup>C<sup>a</sup>

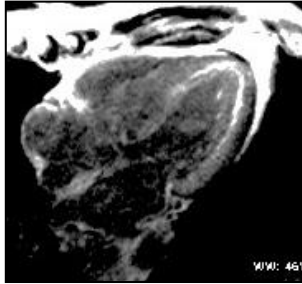




# 3. Miocardiopatías y miocarditis

Miocardiopatías, 14% de nuestras indicaciones

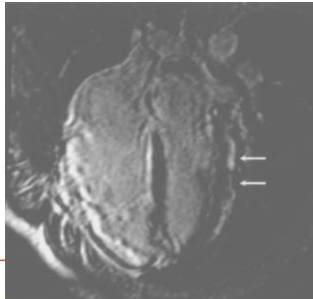
**Amiloidosis**



**MCH**



**DAVD**



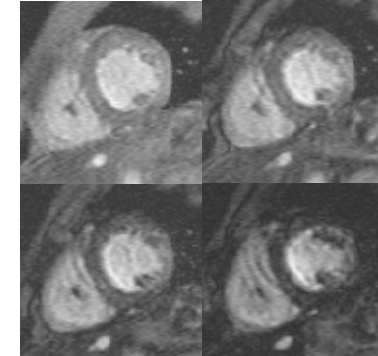
**Table 4 Indications for CMR of cardiomyopathies**

Indication	Class
1. Dilated cardiomyopathy	I
2. Myocarditis	I
3. Hypertrophic cardiomyopathy	I
4. Arrhythmogenic cardiomyopathy	I
5. Cardiac amyloidosis	I
6. Myocardial iron overload	I
7. Left-ventricular noncompaction	I
8. Fabry's disease	I
9. Cardiac sarcoidosis	I
10. Stress-induced (Takotsubo) cardiomyopathy	I
11. Endomyocardial fibrosis	I
12. Restrictive cardiomyopathy	II
13. Chemotherapy induced CMP	II
14. Athlete's heart	II

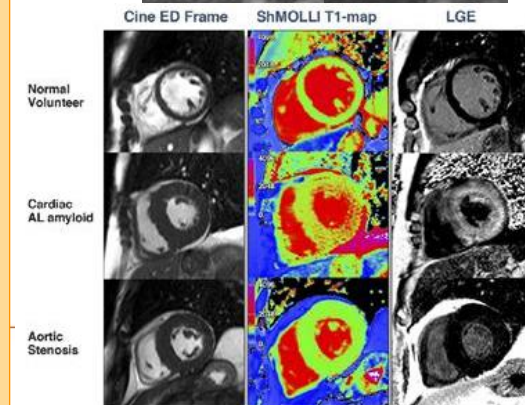
**MCPNC**



**MCP siderótica**



1. Estudio anatómico y de función ventricular
2. Medición de gradientes, velocidades subaórticas
3. Valoración de perfusión en reposo/estrés
4. Angiografía coronaria
5. Caracterización tisular:
  - Detección de edema miocárdico
  - Imagen multiparamétrica
  - Realce tardío de gadolinio



# MIOCARDITIS\_UPDATED LAKE LOUISE CRITERIA

JACC STATE-OF-THE-ART REVIEW

## Cardiovascular Magnetic Resonance in Nonischemic Myocardial Inflammation



Expert Recommendations

Vanessa M. Ferreira, MD, DPHIL,<sup>a</sup> Jeanette Schulz-Menger, MD,<sup>b</sup> Godtfred Holmvang, MD,<sup>c</sup> Christopher M. Kramer, MD,<sup>d</sup> Iacopo Carbone, MD,<sup>e</sup> Udo Sechtem, MD,<sup>f</sup> Ingrid Kindermann, MD,<sup>g</sup> Matthias Gutberlet, MD,<sup>h</sup> Leslie T. Cooper, MD,<sup>i</sup> Peter Liu, MD,<sup>j</sup> Matthias G. Friedrich, MD<sup>k,l,m</sup>

Más especificidad con combinación de edema y otros marcadores de inflamación:

**Al menos un criterio T2 + al menos un criterio T1**  
**T2map o T2-STIR + T1map o RTG**

**Un único marcador apoya el diagnóstico si la clínica acompaña, pero menos específico**

**CENTRAL ILLUSTRATION: Overview of the Updated Lake Louise Criteria**

	2018 Lake Louise Criteria	CMR Image Examples
Main Criteria	<b>Myocardial Edema</b> (T2-mapping or T2W images)	Regional or global increase of native T2 
	<b>Non-ischemic Myocardial Injury</b> (Abnormal T1, ECV, or LGE)	Regional or global increase of T2 signal intensity Regional or global increase of native T1 Regional or global increase of ECV Regional LGE signal increase 
Supportive Criteria	<b>Pericarditis</b> (Effusion in cine images or abnormal LGE, T2, or T1)	Pericardial effusion 
	<b>Systolic LV Dysfunction</b> (Regional or global wall motion abnormality)	Regional or global hypokinesis 

Ferreira, V.M. et al. J Am Coll Cardiol. 2018;72(24):3158-76.

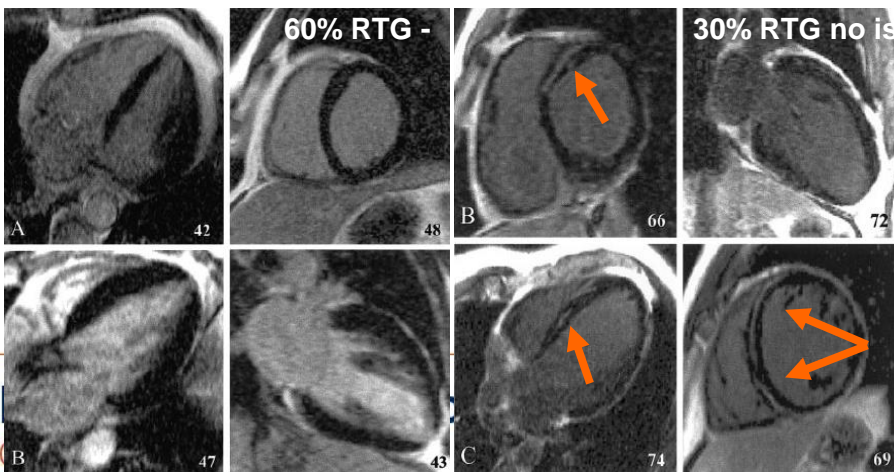
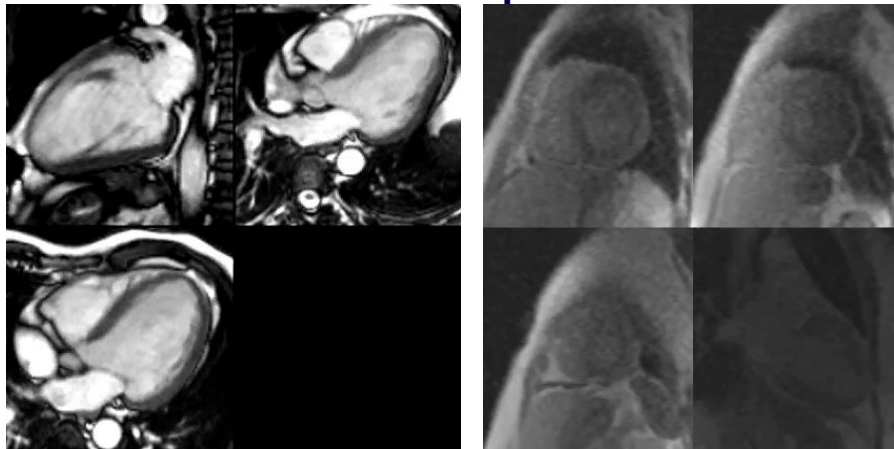
# CRM en MCD

## Valor diagnóstico y pronóstico

Excluir componente isquémico. Caracterización tisular: edema, cicatriz, fibrosis, infiltración

**Considerar (al menos una vez) en todos los pacientes con MCD**

### MCD no isquémica



### Diagnóstico de MCD no isquémica:

60% RTG negativo  
 25% RTG no isquémico  
 9% RTG isquémico

### Pronóstico:

Volúmenes ventriculares  
 FEVI  
 FEVD  
 Extensión de la fibrosis focal (RTG)

### A más RTG peor pronóstico:

Más arritmias  
 Menos recuperación funcional

### Valoración para resincronización

## Multimodality imaging in the diagnosis, risk stratification, and management of patients with dilated cardiomyopathies: an expert consensus document from the European Association of Cardiovascular Imaging

Erwan Donal<sup>1,2\*</sup>, Victoria Delgado<sup>3</sup>, Chiara Bucciarelli-Ducci<sup>4</sup>, Elena Galli<sup>1,2</sup>, Kristina H. Haugaa<sup>5</sup>, Philippe Charron<sup>6,7†</sup>, Jens-Uwe Voigt<sup>8</sup>, Nuno Cardim<sup>9</sup>, P.G. Masci<sup>10</sup>, Maurizio Galderisi<sup>11</sup>, Oliver Gaemperli<sup>12</sup>, Alessia Gimelli<sup>13</sup>, Yigal M. Pinto<sup>14†</sup>, Patrizio Lancellotti<sup>15</sup>, Gilbert Habib<sup>16,17</sup>, Perry Elliott<sup>18,19†</sup>, Thor Edvardsen<sup>5</sup>, Bernard Cosyns<sup>20†‡</sup>, and Bogdan A. Popescu<sup>21‡</sup>

**Table 4** Diagnostic criteria for relatives of familial DCM<sup>1</sup>

#### Major

1. Unexplained decrease of LVEF  $\leq 50\%$  but  $>45\%$

#### OR

2. Unexplained LVED dilatation (diameter or volume) according to nomograms (LVED diameter/volume 2 SD + 5% since this more specific echocardiographic criterion was used in studies that demonstrated the predictive impact of isolated dilatation in relatives).

#### Minor

1. Complete LBBB or AV block (PR  $\geq 200$  ms or higher degree of AV block).

2. Unexplained ventricular arrhythmia (100 ventricular premature beats per hour in 24 h or non-sustained ventricular tachycardia,  $\geq 3$  beats at a rate of  $\geq 120$  bpm).

3. Segmental wall motion abnormalities in the left ventricle in the absence of intraventricular conduction defect

4. Late enhancement (LGE) of non-ischaemic origin on cardiac magnetic resonance imaging.

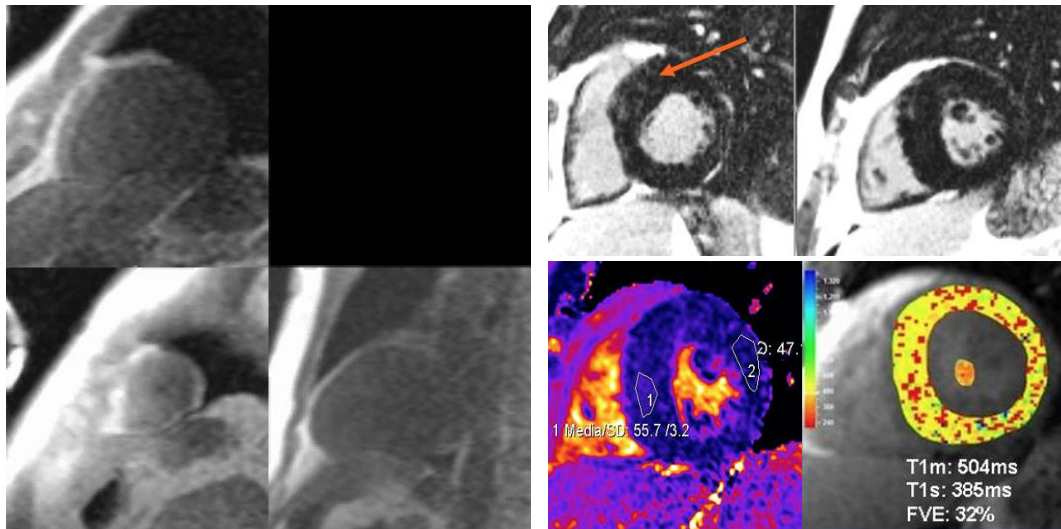
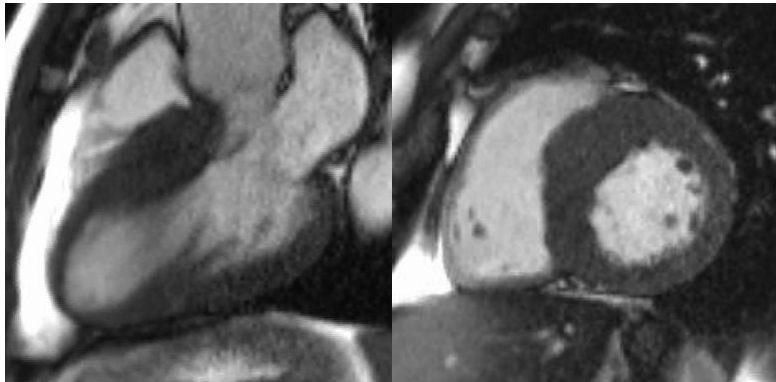
5. Evidence of non-ischaemic myocardial abnormalities (inflammation, necrosis, and/or fibrosis) on EMB.

6. Presence of serum organ-specific and disease-specific AHA by one or more autoantibody tests.

Note: Feature shown by two independent imaging modalities.<sup>1</sup>



# CRM en MCH



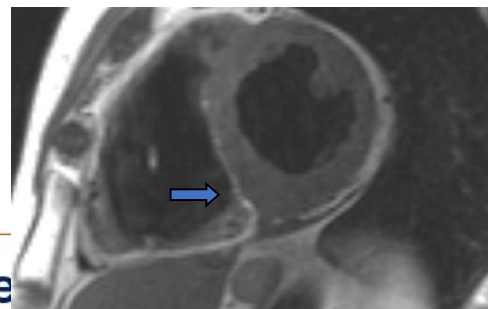
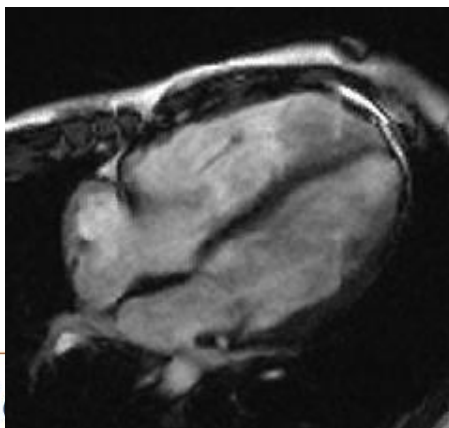
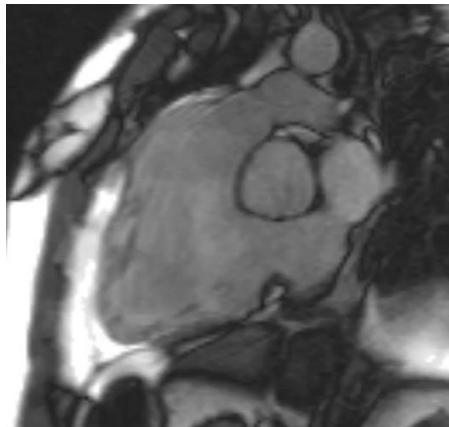
**Diagnóstico:** Detección de fenotipos, formas apicales  
 Diagnóstico diferencial con otras causas de hipertrofia  
 Gradiente subaórtico, grosor parietal, SAM, tamaño auricular

**Pronóstico:** Grosor parietal > 30mm  
 RTG (si existe: peor pronóstico, cuanto más haya: peor  
 Dilatación de AI, gradiente subaórtico....

Recommendations for CMR Imaging Referenced studies that support the recommendations are summarized in Online Data Supplement 4.		
COR	LOE	Recommendations
1	B-NR	1. For patients suspected to have HCM in whom echocardiography is inconclusive, CMR imaging is indicated for diagnostic clarification. <sup>74-80</sup>
1	B-NR	2. For patients with left ventricular hypertrophy in whom there is a suspicion of alternative diagnoses including infiltrative or storage disease as well as athlete's heart, CMR imaging is useful <sup>74-80</sup> (Figure 1).
1	B-NR	3. For patients with HCM who are not otherwise identified as high risk for sudden cardiac death (SCD), or in whom a decision to proceed with implantable cardioverter-defibrillator (ICD) remains uncertain after clinical assessment that includes personal/family history, echocardiography, and ambulatory electrocardiographic monitoring, CMR imaging is beneficial to assess for maximum left ventricular (LV) wall thickness, ejection fraction (EF), LV apical aneurysm, and extent of myocardial fibrosis with late gadolinium enhancement. <sup>38,74-87</sup>
1	B-NR	4. For patients with obstructive HCM in whom the anatomic mechanism of obstruction is inconclusive on echocardiography, CMR imaging is indicated to inform the selection and planning of SRT. <sup>88-92</sup>
2b	C-EO	5. For patients with HCM, repeat contrast-enhanced CMR imaging on a periodic basis (every 3 to 5 years) for the purpose of SCD risk stratification may be considered to evaluate changes in late gadolinium enhancement and other morphologic changes, including EF, development of apical aneurysm, or LV wall thickness (Figure 1, Table 7).

# CRM EN MCA

**Diagnóstico:** Disfunción sistólica de VD  
 Alt. segmentarias de la contractilidad VD  
 Fibrosis (RTG) VI y VD  
 Imagen paramétrica  
**Pronóstico** (si hay RTG: peor)



Categoría	VD (criterios ITF 2010 ACTUALIZADOS)	VI (nuevos criterios)
-----------	--------------------------------------	-----------------------

I. Morpho-functional ventricular abnormalities

II. Structural myocardial abnormalities

**By echocardiography, CMR or angiography:**  
**Major**

- Regional RV akinesia, dyskinesia, or bulging plus one of the following:
  - global RV dilatation (increase of RV EDV according to the imaging test specific nomograms)
  - global RV systolic dysfunction (reduction of RV EF according to the imaging test specific nomograms)

**Minor**

- Regional RV akinesia, dyskinesia or aneurysm of RV free wall

**By CE-CMR: Major**

- Transmural LGE (stria pattern) of  $\geq 1$  RV region(s) (inlet, outlet, and apex in 2 orthogonal views)

**By EMB (limited indications): Major**

- Fibrous replacement of the myocardium in  $\geq 1$  sample, with or without fatty tissue

**By echocardiography, CMR or angiography: Minor**

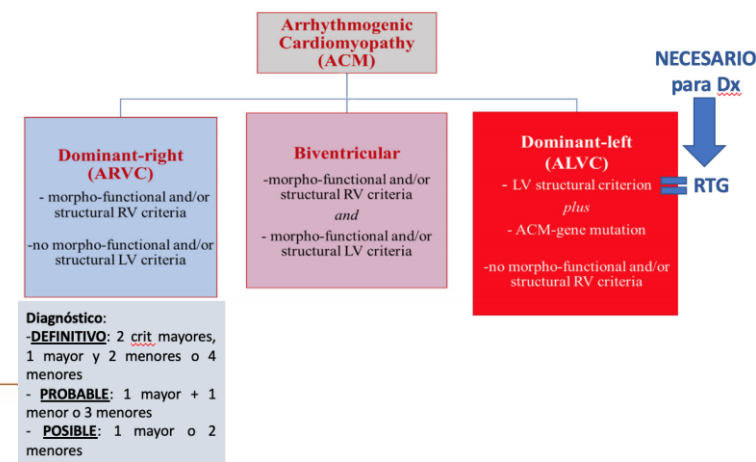
- Global LV systolic dysfunction (depression of LV EF or reduction of echocardiographic global longitudinal strain), with or without LV dilatation (increase of LV EDV according to the imaging test specific nomograms for age, sex, and BSA)

**Minor**

- Regional LV hypokinesia or akinesia of LV free wall, septum, or both

**By CE-CMR: Major**

- LV LGE (stria pattern) of  $\geq 1$  Bull's Eye segment(s) (in 2 orthogonal views) of the free wall (subepicardial or midmyocardial), septum, or both (excluding septal junctional LGE)





# CRM EN MCR y MC INFLAMATORIA

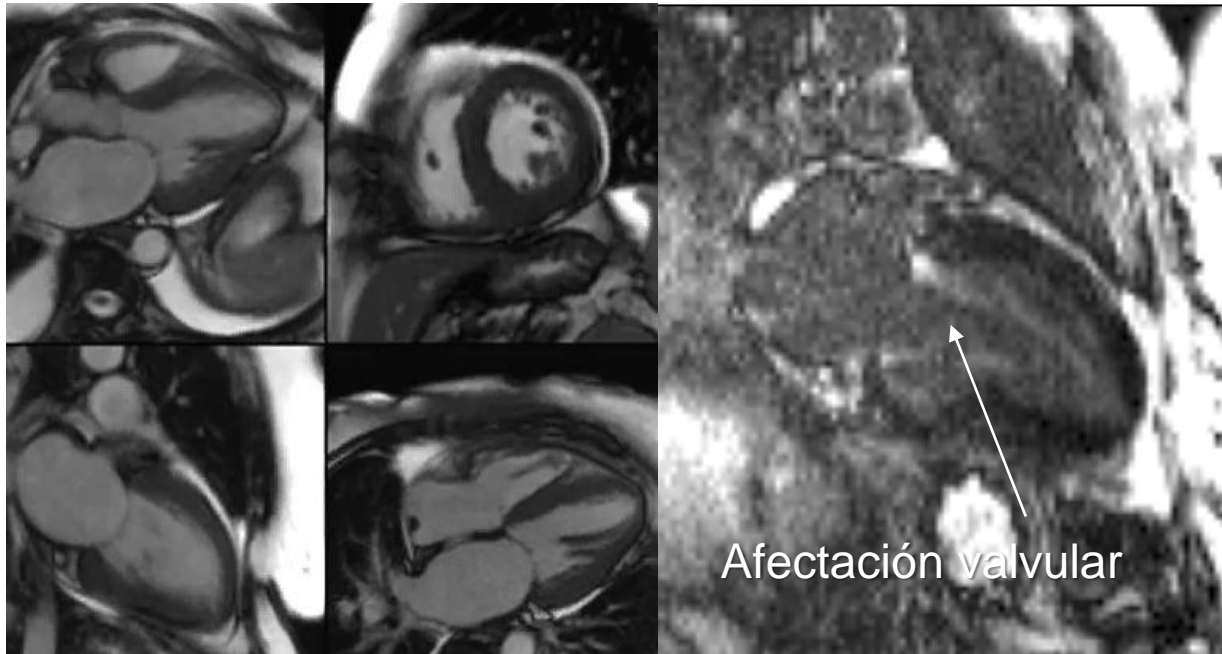
Detección de restricción, hemodinámica restrictiva

**Estudio etiológico** (sarcoidosis, amiloidosis, fibrosis endomiocárdica,...)

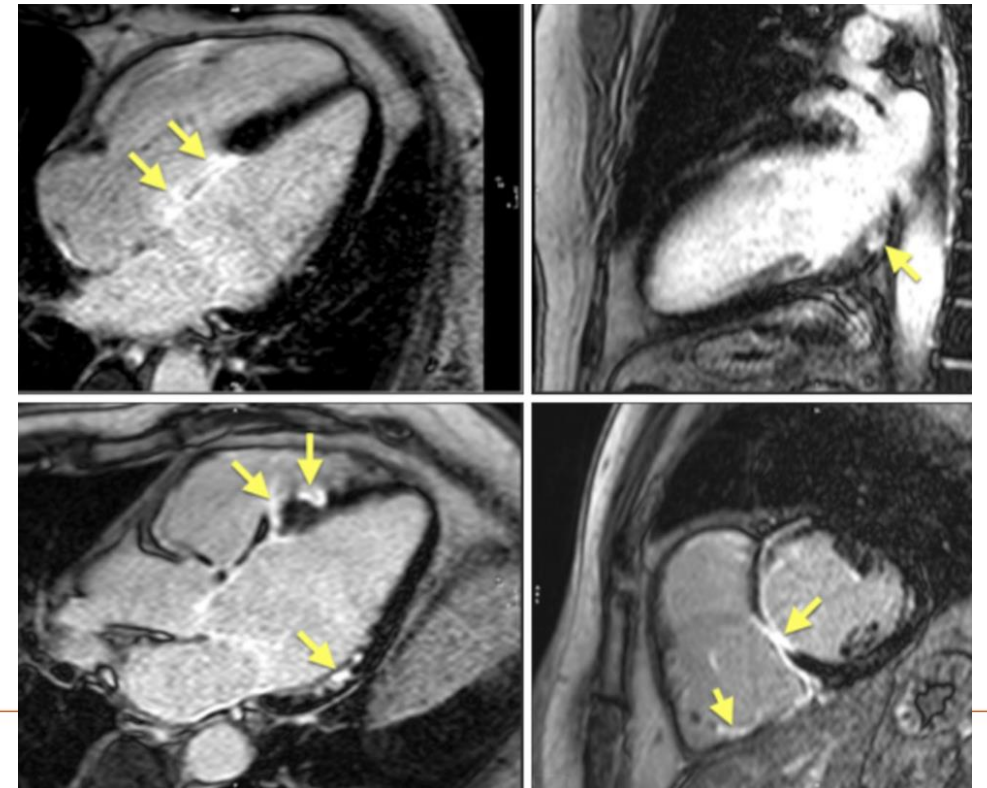
D.D. con pericarditis constrictiva

**Pronóstico:** edema, RTG, imagen multiparamétrica

Amiloidosis



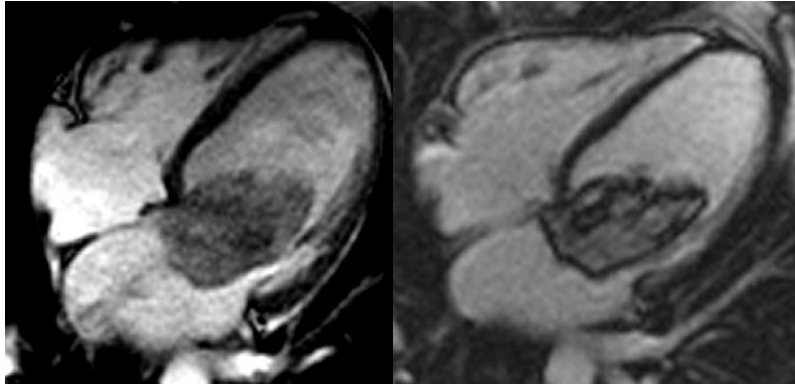
Sarcoidosis



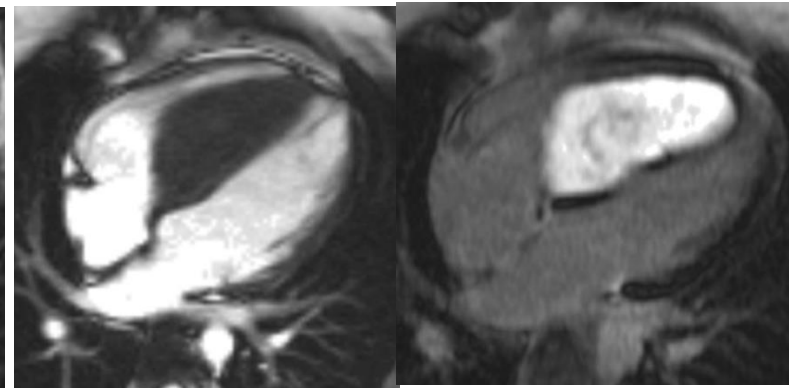


# 4. Pericardiopatías, tumores, trombos

Mixoma



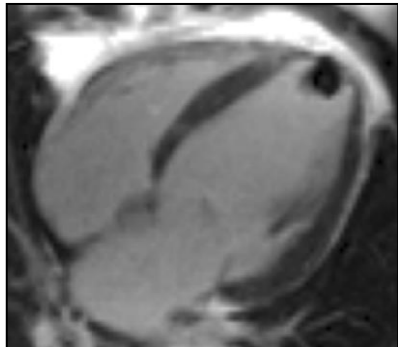
Fibroma



**Table 5 Indications for CMR in pericardial disease**

Indication	Class
1. Pericardial effusions	III
2. Pericardial inflammation	I
3. Pericardial constriction	I
4. Congenital anomalies of the pericardium	I

Trombo VI



Trombo AD



**Table 6 Indications for CMR of cardiac masses**

Indication	Class
1. Suspected cardiac mass	I
2. Differentiation between benign, malignant and non-tumourous masses	I
3. Guide surgery and/or biopsy if this is deemed appropriate	I
4. Follow-up of benign cardiac tumours that do not require urgent intervention for changes over time	I
5. Evaluation of tumour resection/debulking, monitoring recurrence after surgery and regression or progression after chemotherapy or radiotherapy	I
6. Extra-cardiac extension of cardiac tumours or cardiac extension of tumours originating from surrounding structures	I
7. Impact of cardiac masses on hemodynamics	I

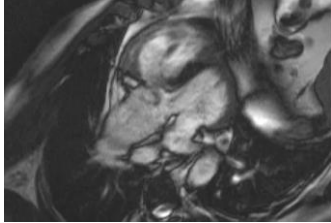
# 5. Cardiopatías congénitas

6% de indicaciones

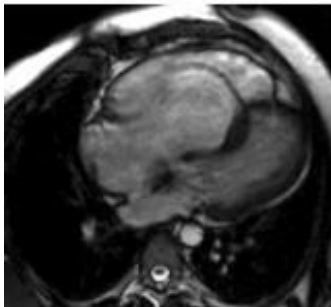
## Valoración inicial y seguimiento en GUCH

- 1. Shunt I-D:** Intracardíacos: CIV, CIA, DVPA parcial, Canal AV, Extracardíacos: PDA, Fístula de seno de Valsalva
- 2. Malformaciones izquierdas sin shunt/ shunt I-D / bidireccional:**  
Coartación de aorta  
Estenosis aórtica (supra, valvular, sub), válvula bicúspide  
Regurgitación mitral
- 3. Malformaciones derechas sin shunt / shunt D-I / bidireccional:**  
Estenosis pulmonar con septo íntegro, T.Fallot, Ebstein
- 4. Malformaciones de las conexiones venosas pulmonares:** DVPAT
- 5. Malposiciones:** Levocardia, Dextrocardia, Mesocardia  
D-TGA, VD de doble salida, L-TGA, V. Único
- 6. Malformaciones coronarias:** Fístula AV coronaria, ALCAPA

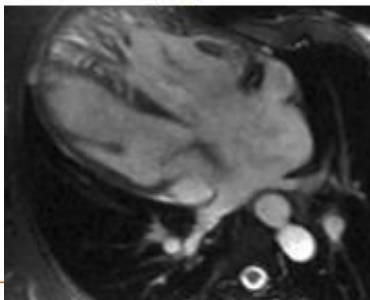
TF



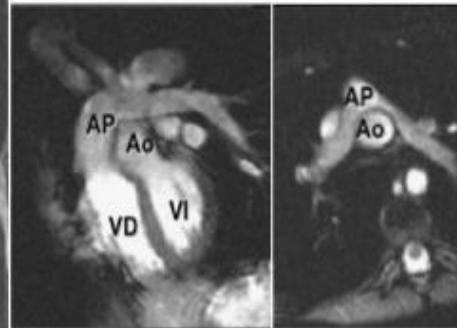
Ebstein



CIV



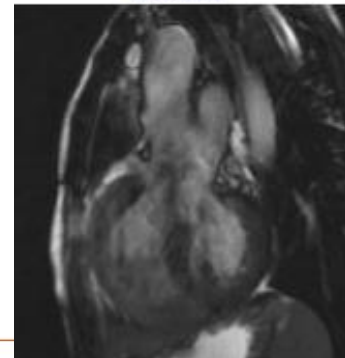
TGA



DVPAP



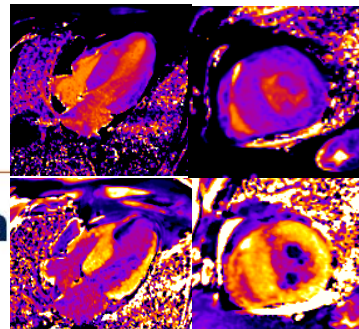
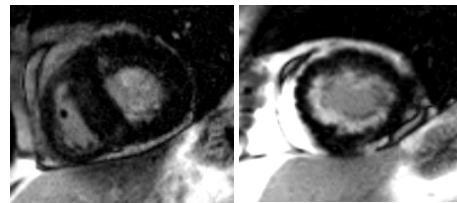
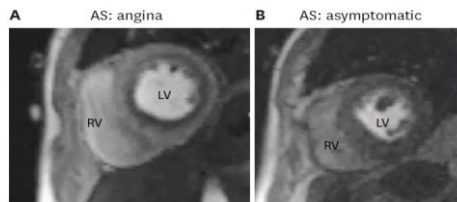
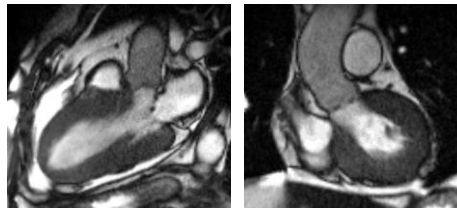
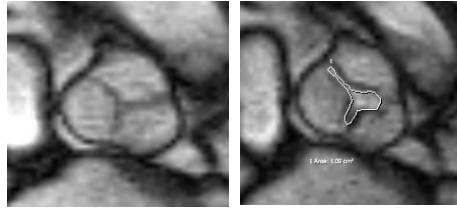
Truncus



# 6. Valvulopatías

CRM EN ESTENOSIS AORTICA

- Masa y función VI
- Anatomía valvular. AVAo →
- Detección de estenosis subvalvular y supravalvular
- Raíz aórtica y Ao ascendente →
- Hipoperfusión en estrés →
- Realce tardío de gadolinio: fibrosis focal, necrosis →
- Imagen multiparamétrica →



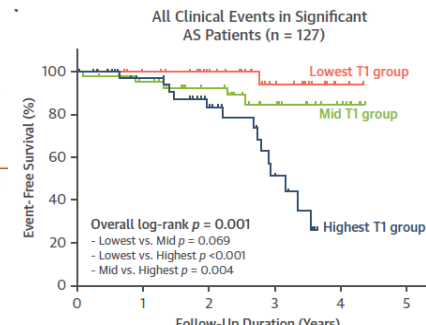
T1 nativo (1020 ms)

Post-Gd (ECV 33%)

En > 50% de ptes  
↑ mortalidad x 2  
(total y CV)

**Table 7 Indications for CMR in valvular heart disease**

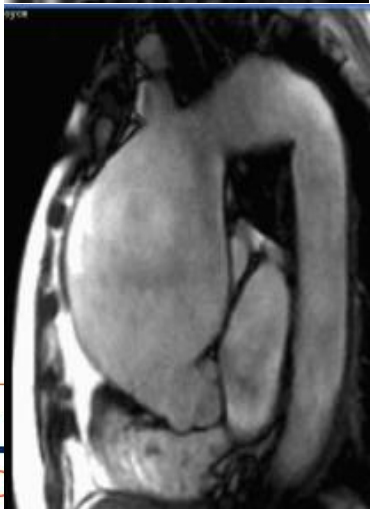
Indication	Class
1. Aortic stenosis	II
2. Identification of sub- and supravalvular stenosis	I
3. Aortic regurgitation	II
4. Ascending aortic flow patterns in aortic stenosis	Inv
5. Mitral stenosis	III
6. Mitral regurgitation	II
7. Pulmonary stenosis	I
8. Pulmonary regurgitation	I
9. Tricuspid stenosis	III
10. Tricuspid regurgitation	II
11. Prosthetic valve disease	II





# 8. Patología de grandes vasos

Aneurisma de aorta

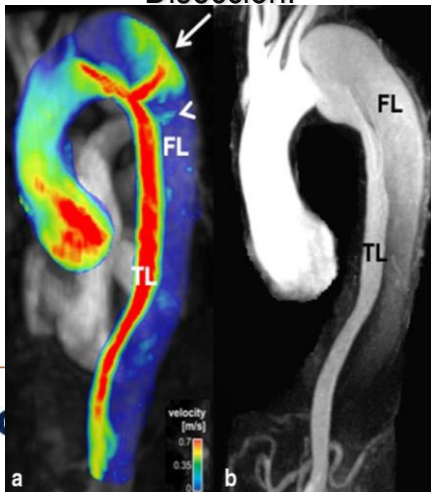


4% de indicaciones

**Estudio anatómico y funcional**  
**Medición de velocidad de flujo, cuantificación p. valvular**  
**AngioRM**  
**¿Flujo 4D?**

**Aneurisma: localización, extensión, seguimiento**  
**Disección, HIM, UPA**

Disección.



H.I.M.



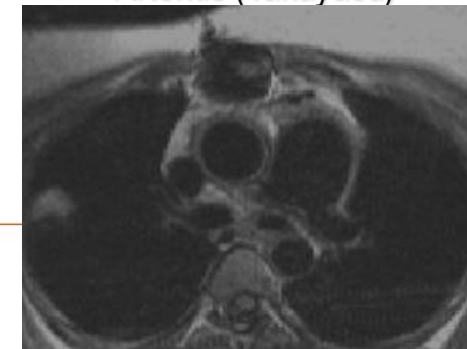
U.P.A



**Table 2 Indications for CMR in acquired diseases of the vasculature**

Indication	Class
1. Diagnosis and follow-up of thoracic aortic aneurysm including connective tissue diseases	I
2. Diagnosis and planning of stent treatment for abdominal aortic aneurysm	II
3. Follow-up of stented abdominal aortic aneurysm	III
4. Aortic dissection	
Diagnosis of acute aortic dissection	II
Diagnosis and follow-up of chronic aortic dissection	I
5. Diagnosis of aortic intramural hematoma	I
6. Diagnosis of penetrating aortic ulcers	I
7. Pulmonary artery anatomy and flow	I
8. Pulmonary emboli	
Diagnosis of central pulmonary emboli	III
Diagnosis of peripheral pulmonary emboli	III
Assessment of chronic pulmonary embolic disease	III
9. Assessment of aortic arch arteries	I
10. Assessment of aortic branch arteries including the Adamkiewicz artery	II
11. Assessment of carotid, vertebral and circle of Willis arteries	I
12. Assessment of upper extremity arteries	I
13. Assessment of hand arteries	II
14. Assessment of renal arteries	I
15. Assessment of mesenteric arteries	I
16. Assessment of pelvic and lower extremity arteries	I
17. Assessment of pulmonary veins	I
18. Assessment of thoracic, abdominal and pelvic veins	I
19. Assessment of lower extremity veins	I
20. Assessment of atherosclerotic plaque in the carotid artery	II
21. Assessment of atherosclerotic plaque in the aorta	II
22. Assessment of vascular wall inflammation in large and medium sized arteries	II
23. Assessment of aortic pulse wave velocity	Inv
24. Endothelial function	Inv

Arteritis (Takayasu)



CRM puede proporcionar información exacta y reproducible para diagnóstico y pronóstico, y para decidir el manejo en un amplio espectro de patologías cardiovasculares

Los rápidos avances tecnológicos han permitido una expansión de la capacidad diagnóstica e indicaciones clínicas de la técnica