

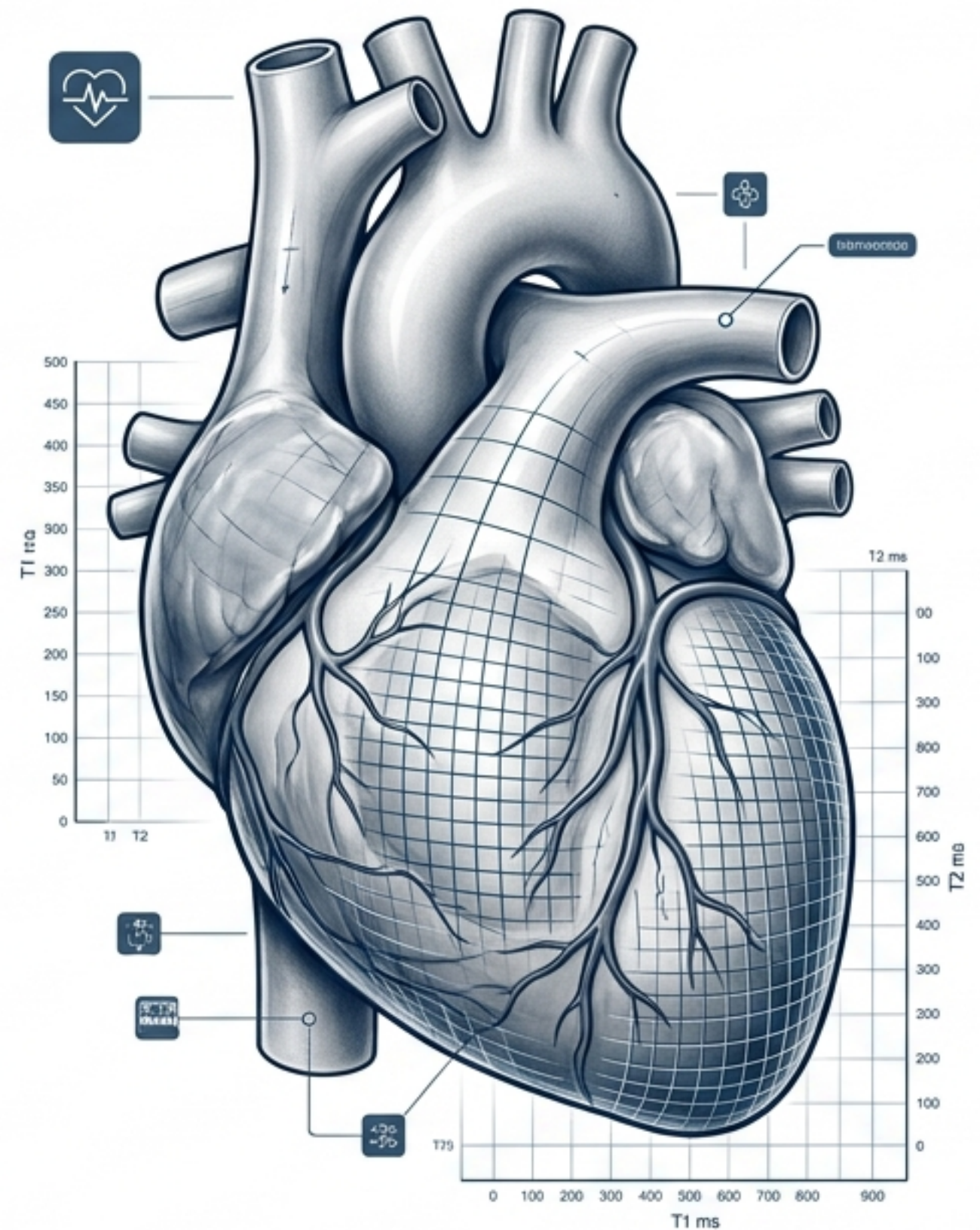
# Diagnosing and Managing Acute Myocarditis

## A Clinical Dashboard for the Australian Practitioner

Source: Med2Date Clinical Guidelines

Framework: 2018 Lake Louise & 2020 ESC Criteria

Last Updated: 15 May 2026





## THE BASELINE BURDEN

**4-14 cases** per **100,000** persons annually in Australia.

**Underestimated** due to limited regional CMR access.

**Median age:** 30-40 years.  
Male:Female ratio 1.5-2:1.



## THE LETHAL TAIL

**5-12%** of **Sudden Cardiac Death** in young adults.

Accounts for up to **9%** of unexplained dilated cardiomyopathy presentations nationally.



## THE COVID-19 ERA

NCIRS data shows increased incidence from both **SARS-CoV-2 infection** (20-30% of hospitalized patients show myocardial injury) and **mRNA vaccination** (active TGA surveillance).



**CLINICAL RED FLAG:** Maintain a low diagnostic threshold for young patients presenting with chest pain, troponin elevation, and non-ischaemic ECG changes.

# THE ETIOLOGY MATRIX



## VIRAL (Most Common)

**Pathogenesis:** Direct cytopathic effects followed by immune-mediated response.

### Key Agents:

- Parvovirus B19 (30-50% by PCR, endothelial tropism)
- Coxsackievirus B (15-25%, direct myocyte lysis, summer/autumn)
- HHV-6
- Adenovirus (paediatric)
- SARS-CoV-2



## IMMUNE-MEDIATED

**Pathogenesis:** Aberrant immune activation requiring targeted therapy.

### Key Subtypes:

- Giant-Cell Myocarditis (Fulminant, young adults, poor prognosis without transplant)
- Eosinophilic (Drug reactions, EGPA)
- Cardiac Sarcoidosis
- Autoimmune (SLE, rheumatoid arthritis)



## DRUG-INDUCED

**Pathogenesis:** Toxic or hypersensitivity reactions. Key emerging causes.

### Key Agents:

- Immune Checkpoint Inhibitors (Nivolumab, Pembrolizumab - 0.3-1.5% risk, fulminant onset <6 weeks)
- Clozapine (Requires baseline, 4wk, 12wk troponin per TGA)
- Antimicrobials
- Recreational (Methamphetamine)

# The Clinical Spectrum Matrix



## Mild / Acute Subclinical

- **Setting:** Ward / Outpatient
- **Presentation:** Pleuritic/atypical chest pain, viral prodrome 1-4 weeks prior.
- **Parameters:** Mild troponin elevation, normal/mildly reduced LVEF (>50%), non-specific ST-T changes.



## Moderate / Acute Symptomatic

- **Setting:** Cardiac monitoring ward / CCU
- **Presentation:** Significant chest pain mimicking ACS, dyspnoea, palpitations.
- **Parameters:** Troponin >5x ULN, elevated BNP, LVEF 30-50%, regional wall motion abnormalities, pericardial effusion (~30%).



## Severe / Fulminant

- **Setting:** ICU / Transplant-capable centre
- **Presentation:** Rapid-onset cardiogenic shock, multi-organ dysfunction.
- **Parameters:** LVEF <30%, biventricular failure, high-degree AV block, ventricular arrhythmias. Mortality 50-70% without Mechanical Circulatory Support.



# DIFFERENTIAL DIAGNOSIS MATRIX

DIAGNOSIS	CLINICAL PARAMETERS	
<b>Myocarditis</b>	<ul style="list-style-type: none"> <li>⚡ Atypical/variable pain</li> <li>⚡ Diffuse concave-up ST elevation (non-coronary)</li> </ul>	<ul style="list-style-type: none"> <li>⚡ Troponin elevated beyond expected ACS curve-RWMA in non-coronary distribution on Echo</li> </ul>
<b>Acute Coronary Syndrome (ACS)</b>	<ul style="list-style-type: none"> <li>✗ Exclude via coronary angiography if patient &gt;40 or has cardiovascular risk factors.</li> <li>✗ Variable presentation</li> <li>⚡ Typical coronary ST-T changes</li> </ul>	<ul style="list-style-type: none"> <li>⚡ Expected troponin curve</li> </ul>
<b>Acute Pericarditis</b>	<ul style="list-style-type: none"> <li>✓ Predominant pleuritic pain, pericardial rub</li> <li>⚡ PR depression</li> <li>✓ Normal troponin and normal ventricular function</li> </ul>	<ul style="list-style-type: none"> <li>⚡ Diffuse ST elevation</li> </ul>
<b>Takotsubo Cardiomyopathy</b>	<ul style="list-style-type: none"> <li>✓ Apical ballooning pattern on Echo/CMR</li> <li>⚡ Triggered by emotional/physical stress</li> <li>✓ Reversible ventricular dysfunction</li> </ul>	<ul style="list-style-type: none"> <li>✓ Triggersible ventricular dysfunction</li> <li>⚡ Modest troponin elevation</li> </ul>
<b>Pulmonary Embolism</b>	<ul style="list-style-type: none"> <li>✓ Acute right heart strain (Echo/ECG)</li> <li>⚡ D-dimer elevation</li> <li>✓ CTPA diagnostic</li> </ul>	<ul style="list-style-type: none"> <li>✓ D-dimer elevation</li> <li>⚡ Sinus tachycardia</li> </ul>

# Initial Non-Invasive Workup

## 1. High-sensitivity Troponin

- **Elevated** in >90%. Serial measurements at 0, 3, 6 hours.
- Abnormal rise/fall trajectory is key. (MBS 66504).



## 2. 12-lead ECG

- **Abnormal** in 90%. Look for diffuse ST elevation, T-wave inversions, low voltage QRS, or new bundle branch blocks.
- **Note:** High-degree AV block suggests Giant-Cell or Sarcoidosis.



## Suspected Myocarditis

## 3. Transthoracic Echo (TTE)

- Assess LVEF, regional wall motion abnormalities, myocardial wall thickness (transient oedema hypertrophy), and pericardial effusion.
- Excludes valvular disease.



## 4. Bloods & Serology

- BNP/NT-proBNP for prognostication.
- Autoimmune markers (ESR, CRP, ANA) and Viral PCR/Serology (Coxsackie, Parvovirus, HIV, SARS-CoV-2) where available.

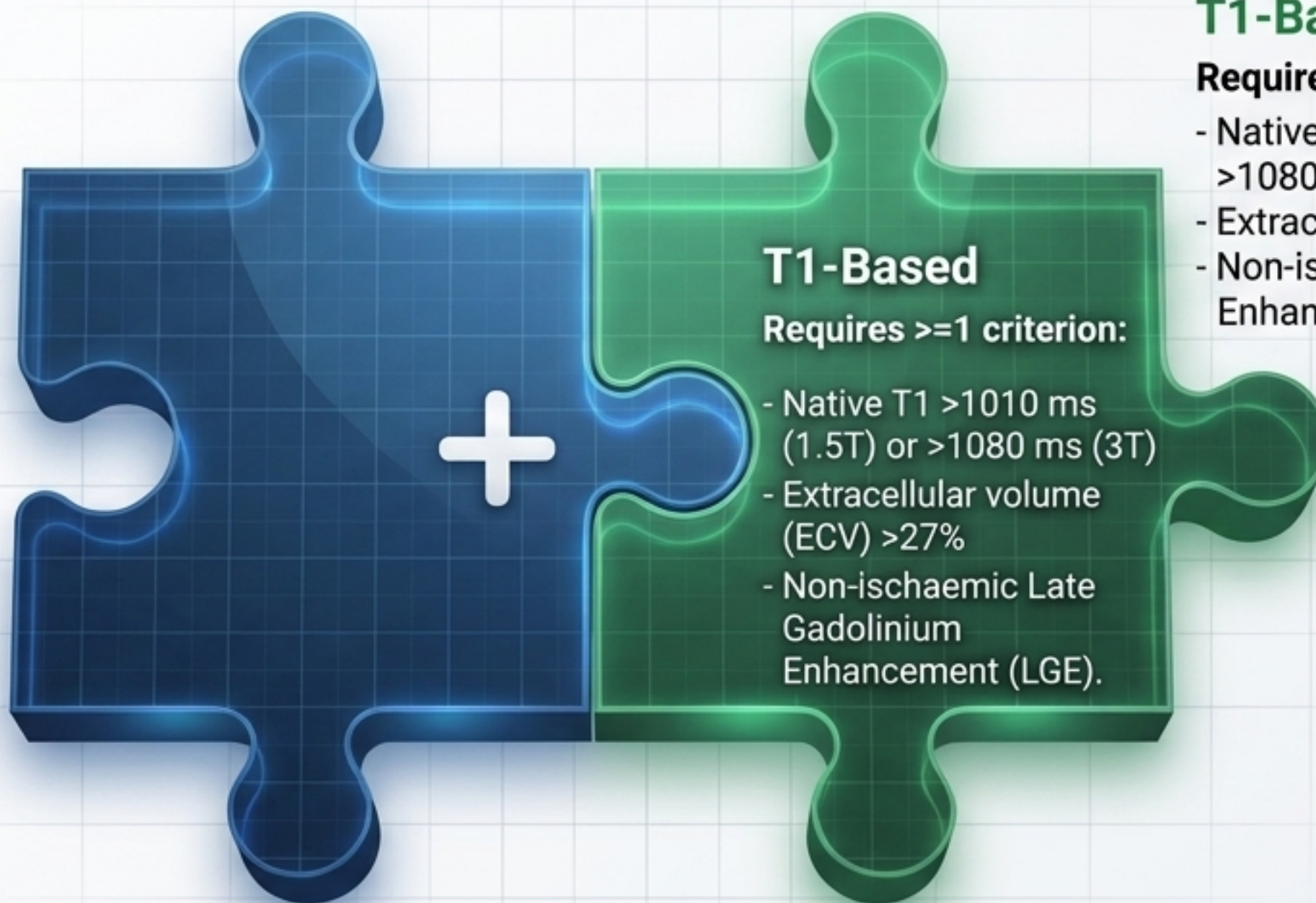


# The 2018 Lake Louise Intersection Diagram

## T2-Based (Oedema)

Requires  $\geq 1$  criterion:

- T2 ratio  $>1.9$  (STIR)
- Native T2  $>52$  ms (1.5T) or  $>54$  ms (3T)
- Elevated T2 signal  $\geq 2$  SD above normal.



## T1-Based

Requires  $\geq 1$  criterion:

- Native T1  $>1010$  ms (1.5T) or  $>1080$  ms (3T)
- Extracellular volume (ECV)  $>27\%$
- Non-ischaemic Late Gadolinium Enhancement (LGE).

## T1-Based (Injury/Fibrosis)

Requires  $\geq 1$  criterion:

- Native T1  $>1010$  ms (1.5T) or  $>1080$  ms (3T)
- Extracellular volume (ECV)  $>27\%$
- Non-ischaemic Late Gadolinium Enhancement (LGE).

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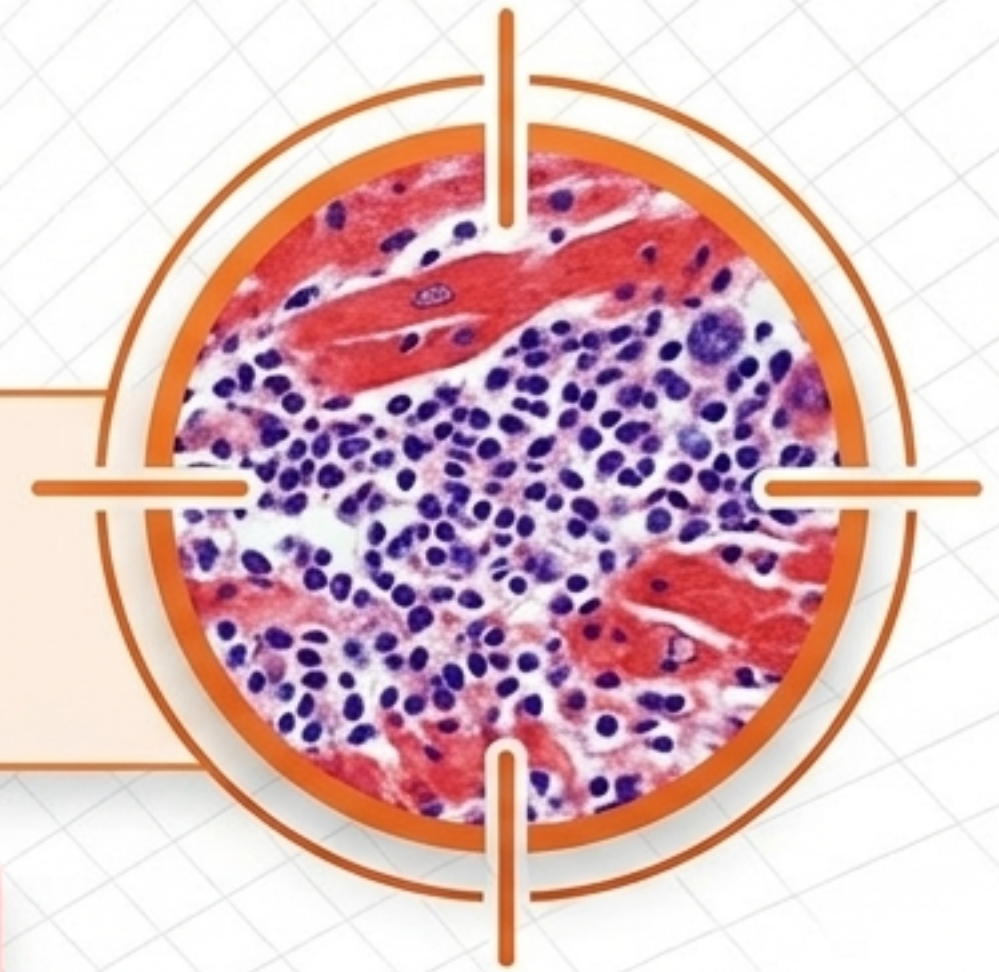


**Positive CMR  
Diagnosis**

**Note:** A CMR study is negative for acute myocarditis if neither T2 oedema nor T1 injury criteria are present.

# Endomyocardial Biopsy (EMB): The Histological Gold Standard

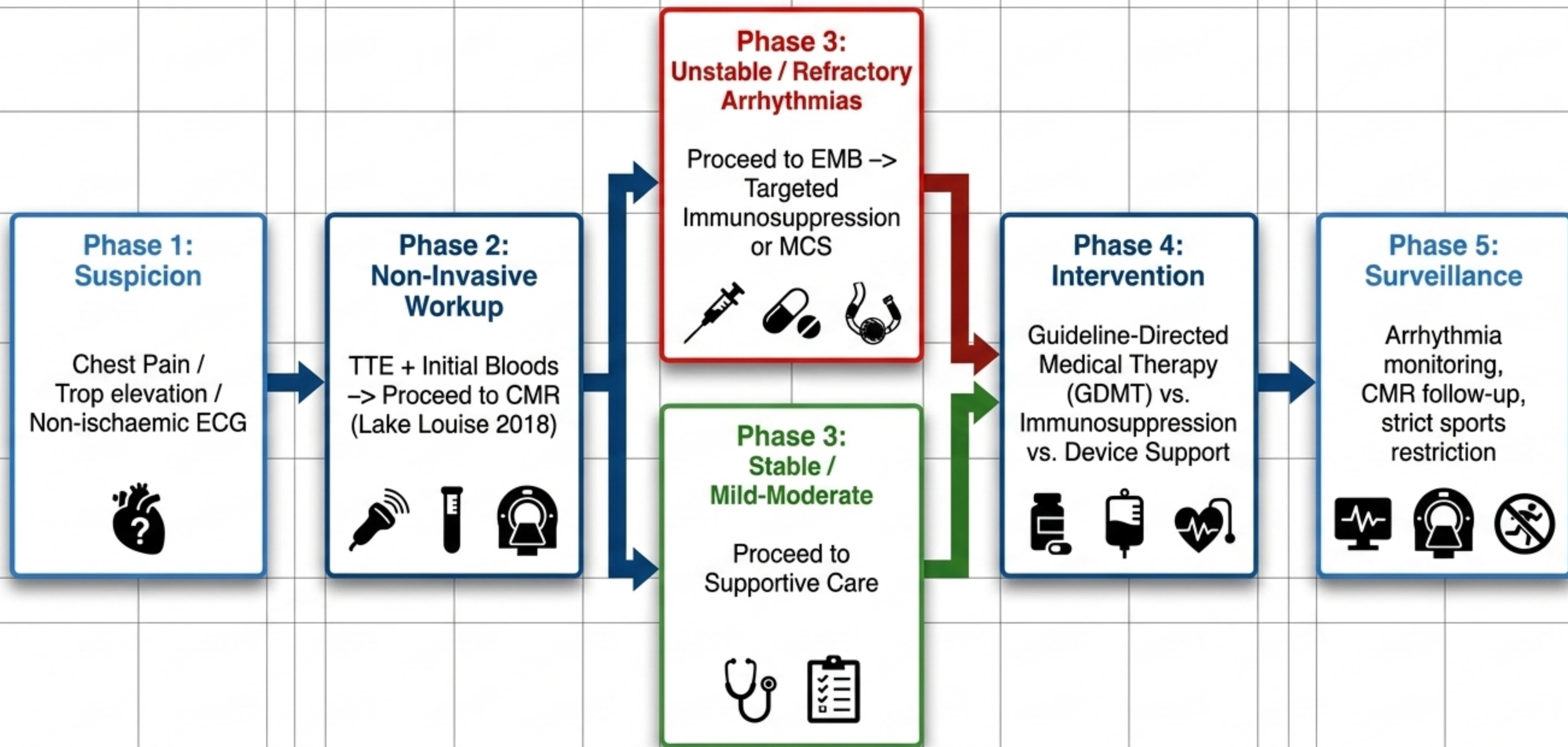
**Dallas criteria** requires lymphocytic infiltrate  $\geq 14$  cells/mm<sup>2</sup> with myocyte necrosis. Yield improved via immunohistochemistry (CD3, CD68) and viral PCR.



## Hard Indications (CSANZ/AHA/ESC)

- ✓ 1. **Fulminant heart failure** with haemodynamic instability (suspect Giant-Cell or Eosinophilic).
- ✓ 2. **Progressive LV dysfunction** refractory to standard therapy for  $\geq 2$  weeks.
- ✓ 3. Suspected **cardiac sarcoidosis** or **eosinophilic myocarditis** where CMR is non-diagnostic.
- ✓ 4. **New-onset HF** (<2 weeks) with LV dilation and new ventricular arrhythmias or AV block.
- ✓ 5. **HF** associated with **cutaneous drug rash**, eosinophilia, or immune checkpoint inhibitors.

# The Master Care Pathway



# Medication Dashboard

## Perindopril (ACEi)

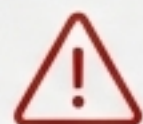
**Adult:** 2mg PO daily, titrate to 5-10mg.

**Renal:** Start 2mg every other day if eGFR <30.

## Carvedilol (Beta-Blocker)

Initiate only when stable.

**Adult:** 3.125mg PO BD, titrate to 25-50mg BD based on weight.



**Avoid NSAIDs**  
(increases injury/fluid retention).  
**Complete alcohol cessation.**  
**Strict exercise restriction.**

## Spironolactone (MRA)

**Adult:** 12.5-25mg PO daily.

Contraindicated if eGFR <30 or K<sup>+</sup> >5.0.

## Furosemide (Diuretic)

**Adult:** 20-80mg IV/PO daily for congestion.

**Note:** Anticoagulation reserved for LVEF <30%, thrombus, or AF.

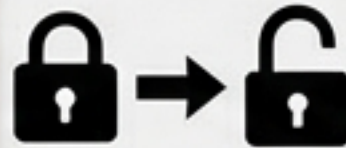
# Critical Safety Principle: Immunosuppression

Immunosuppression is **CONTRAINDICATED** in **active viral myocarditis (PCR-positive on EMB)**. It impairs viral clearance and worsens outcomes.



## The "Lock" (Do Not Give)



Empiric treatment for standard lymphocytic/viral myocarditis.



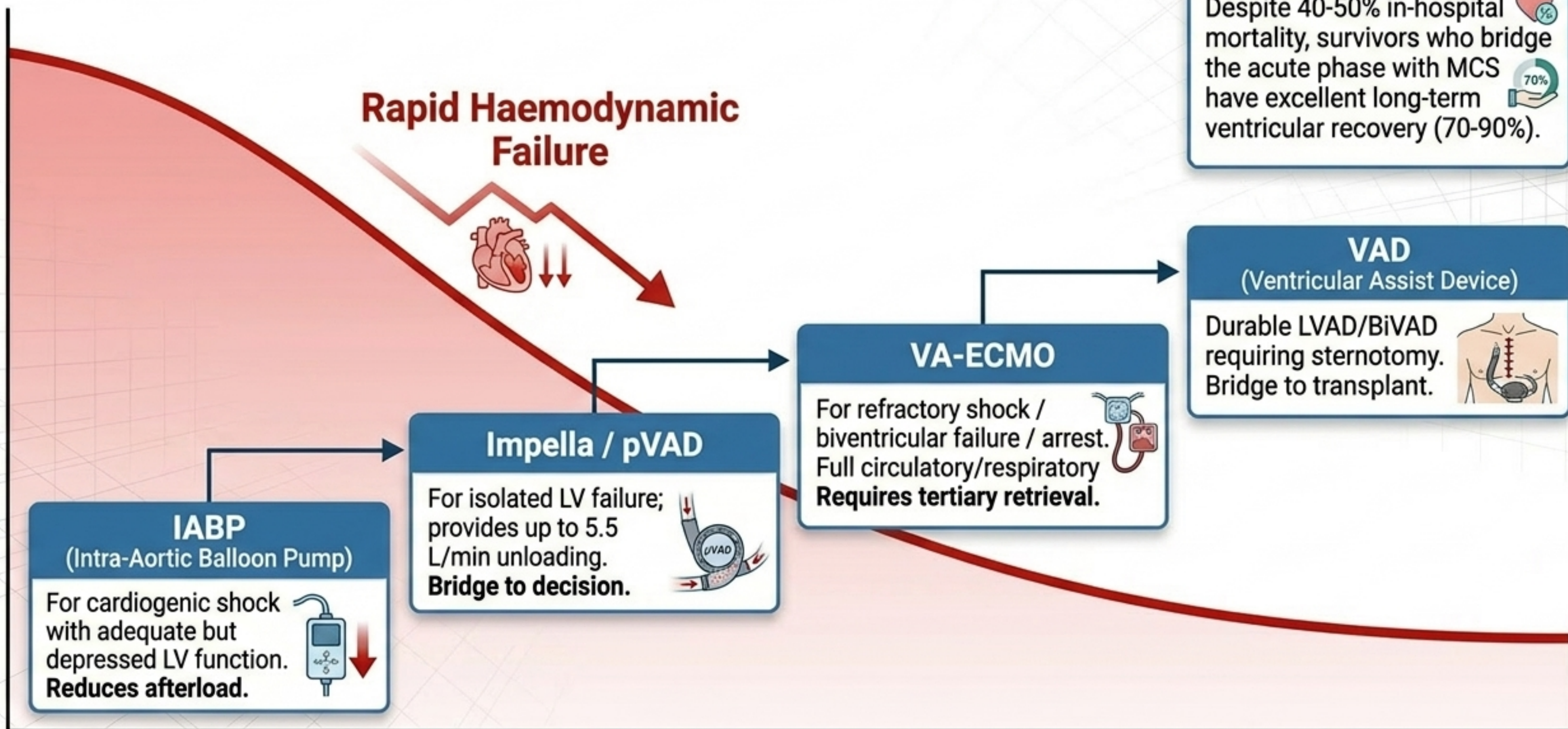
## The "Unlock" (Mandatory Indication)

Biopsy-proven, PCR-negative immune-mediated subtypes. Requires distinct histologic proof (Giant cells, Eosinophils, non-caseating granulomas) before initiating high-dose steroids.

# Targeted Immunosuppressive Regimens

<b>Giant-Cell Myocarditis</b>	<b>Methylprednisolone 1g IV x3 days → Prednisolone 1mg/kg taper. 2nd line: Ciclosporin or Azathioprine + ATG. Requires lifelong immunosuppression and early transplant evaluation.</b>
<b>Eosinophilic Myocarditis</b>	<b>Methylprednisolone → Prednisolone taper over 2-3 months. 2nd line: Mepolizumab  300mg SC (for EGPA). Minimum 6-12 months therapy.</b>
<b>Cardiac Sarcoidosis</b>	<b>Prednisolone for 4-8 weeks, taper over 6-12 months. 2nd line: Methotrexate, Mycophenolate. Minimum 12-24 months therapy.</b>
<b>ICI-Related Myocarditis</b>	<b>Stop ICI immediately. Methylprednisolone  1-2g IV x3-5 days. 2nd line: Abatacept for fulminant cases.</b>

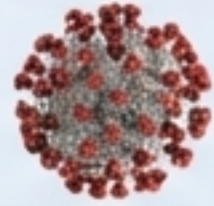
# The Fulminant Escalation Staircase



# COVID-19: Infection vs. Vaccination



## COVID-19 Infection



- **Direct invasion via ACE2, cytokine storm.**
  - Troponin elevated in 20-30% of hospitalized cases.
  - Clinical myocarditis in 2-5%.
  - CMR shows oedema/fibrosis in 30-60% of post-COVID patients.

## mRNA Vaccines



- **Highest risk in males aged 12-30 after dose 2 (BNT162b2 or mRNA-1273).**
  - Onset: 1-7 days post-vax.
  - TGA data: ~30/100k for males 12-17.
  - Prognosis: Excellent, typical LVEF >50%, resolves in 1-7 days.

### Clinical Guidance:

Avoid further doses of the same mRNA platform; consult cardiology for alternatives (Novavax). Echo follow-up at 3-6 months.

# The Australian Context: Remote Practice & ATSI Health

## Rheumatic Heart Disease Overlap

ARF/RHD rates are significantly higher and can mimic or coexist with viral myocarditis, complicating diagnosis in remote communities.



## Geographical Diagnostics

CMR is limited to major centers (Perth, Brisbane, Adelaide).

Remote diagnosis relies heavily on Point-of-care Echo by remote area nurses/FIFO cardiologists.

Early retrieval is critical.



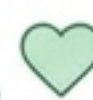
## Cultural Safety & Access

Emphasize yarning-based approaches for explaining rest requirements. Utilize Aboriginal Health Workers (AHWs) in care planning.



## Medication Adherence

Ensure PBS Close the Gap (CTG) utilization for zero/low-cost long-term heart failure medication access.



# Special Populations Management



## Pregnancy

**Overlap:** Difficult to differentiate from Peripartum Cardiomyopathy.

**Med Safety:** ACEi/ARBs are strictly **CONTRAINDICATED** (fetopathy).  
Use Hydralazine + Nitrates.  
Spironolactone is anti-androgenic (**avoid**).  
Metoprolol preferred.

**Delivery:** MDT required if LVEF <40%;  
vaginal delivery with epidural preferred  
over C-section unless unstable.



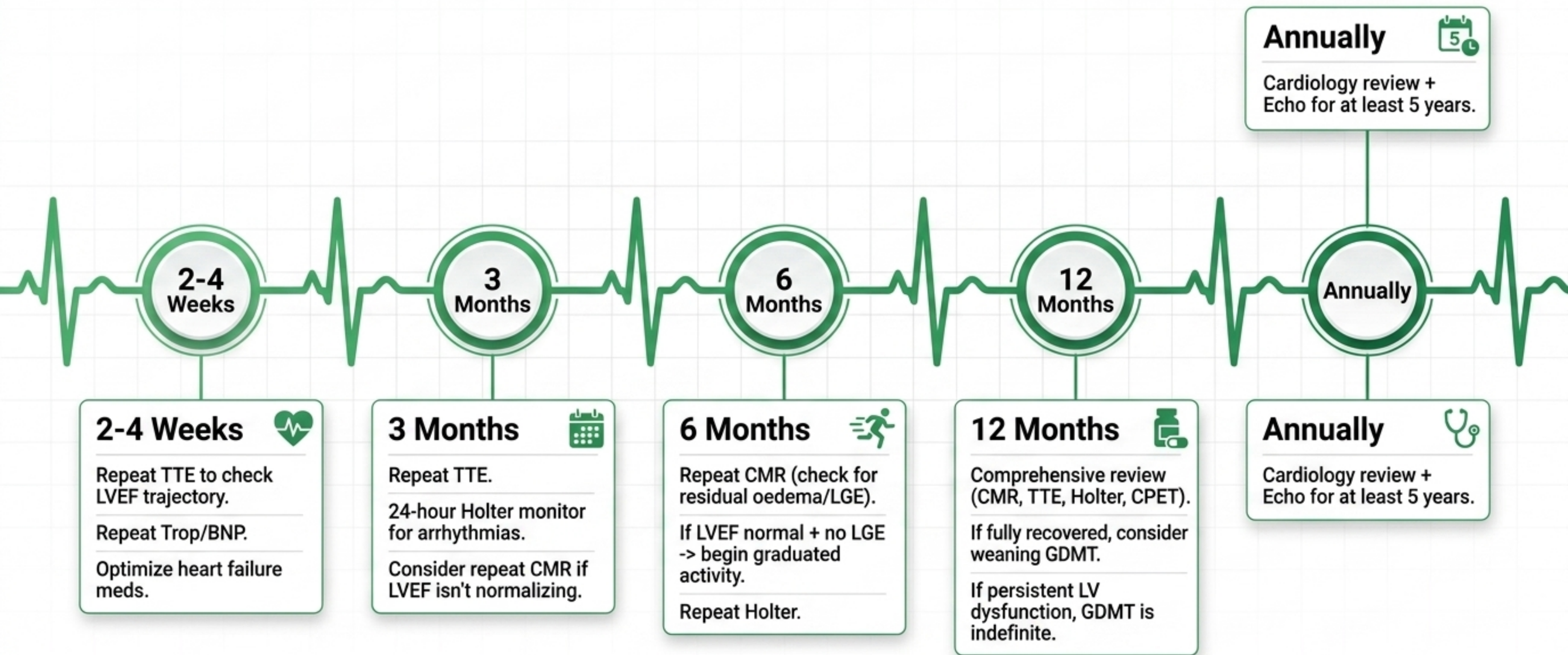
## Paediatrics

**Neonates:** High **fulminant risk, 50-70% mortality** in shock.

**Intervention:** IVIG (1-2 g/kg) is commonly used empirically despite limited RCTs. ECMO is the primary MCS modality.

**Sports:** Minimum 6 months **strict restriction**.

# The Recovery Roadmap Timeline



# Exercise & Return to Sport

## The Pathophysiology Note:

Myocardial inflammation renders the heart electrically unstable. Catecholamines from exercise trigger lethal arrhythmias.



## Red Zone (0-3 Months)

Complete avoidance of competitive sport and moderate/vigorous exercise. Light walking only.

## Yellow Zone (3-6 Months)

Re-assessment. If LVEF normalized and Holter clear, start graduated recreational exercise under guidance.

## Green Zone (6-12 Months)

Return to competitive sport **ONLY** if: LVEF  $\geq 55\%$ , **NO** residual oedema/LGE on CMR, **NO** arrhythmias on Holter, normal BNP, and normal CPET.

# Chronic Sequelae & Surveillance Dashboard

## Dilated Cardiomyopathy (DCM)

20-30% of biopsy-proven cases progress to DCM over 5-10 years. Genetic testing recommended.



## Arrhythmia Surveillance

ILR for unexplained syncope. ICD indicated if LVEF  $\leq 35\%$  persists beyond 3-6 months or secondary prevention for VF/VT.



## Recurrence

Relapse rate is 10-15% for lymphocytic, >50% for Giant-Cell without maintenance immunosuppression.



## Psychological Impact

High rates of anxiety/depression. Routine screening with PHQ-9/GAD-7 and referral to cardiac rehab recommended.



Recovery is a continuum. Lifelong surveillance for structural and electrical stability is the final cornerstone of care.