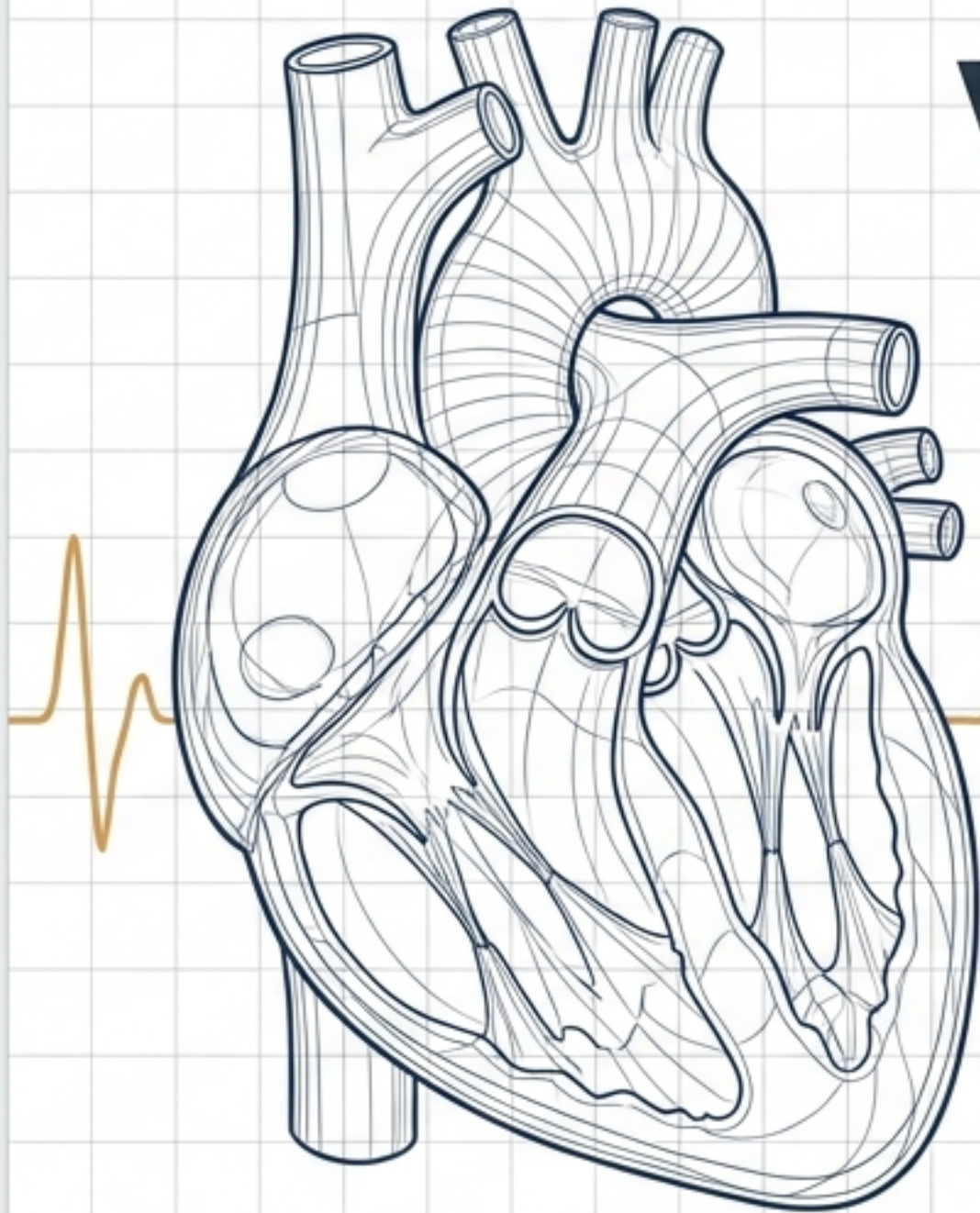


# **Ventricular Tachyarrhythmias in the Australian Context**

**An Evidence-Based Clinical Playbook  
for Acute Crisis Management and  
Sudden Cardiac Death Prevention**



# The National Burden of Sudden Cardiac Death Demands Immediate System Responses

## The Mortality Toll



**20,000–30,000**

**Annual Sudden Cardiac Deaths** in Australia.

Cardiovascular disease accounts for **25%** of all national deaths (AIHW).

## The Substrate

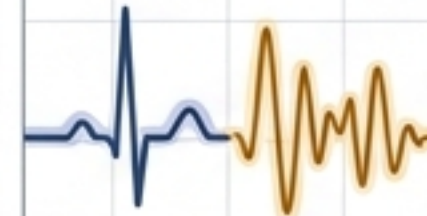


**60–70%**

of VT/VF events occur in patients with prior myocardial infarction or ischaemic cardiomyopathy.

**Ischaemia** is the undisputed primary enemy.

## Out-of-Hospital Cardiac Arrest (OHCA)



**30,000**

annual OHCA. Overall survival-to-discharge is only **10–12%**.

VF/pulseless VT occurs in **20–25%** of cases (Aus-ROC registry).



Aboriginal and Torres Strait Islander peoples experience sudden cardiac death at **2–3 times the rate** of non-Indigenous Australians, driven by rheumatic heart disease and remote geographic delays.



# Identifying and Triaging Sustained Monomorphic Ventricular Tachycardia

**Sustained Monomorphic VT (SMVT):**  
Uniform QRS lasting  $\geq 30$ s or causing haemodynamic compromise.

## Unstable

Systolic BP  $< 90$  mmHg, altered consciousness, acute pulmonary oedema



**ACTION:**  
Immediate synchronised cardioversion. Do not delay for pharmacology.

## Stable

Systematic structural assessment required.



**TTE (Echo):**  
Assess LVEF, wall motion, valves (Nationwide availability).



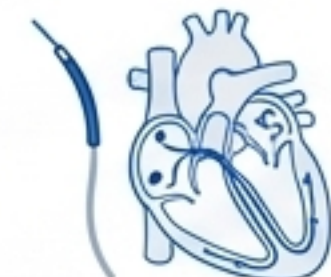
**CMR (MRI with LGE):**  
Localise myocardial scar/fibrosis.



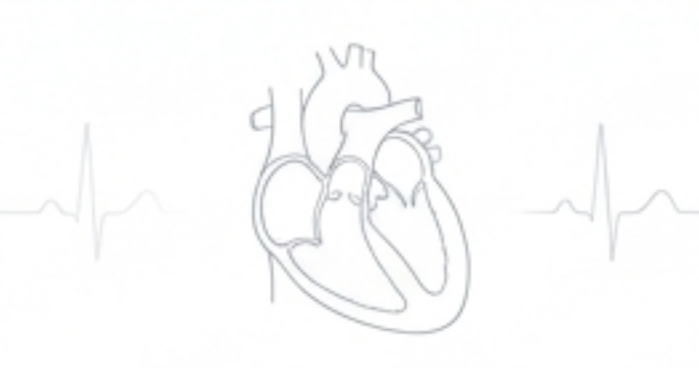
**Angiography (CT or Invasive):**  
Exclude ischaemic aetiology.



**EP Study:**  
Map circuits and identify ablation targets.




# The Pharmacological Dashboard for Ventricular Arrhythmia Management




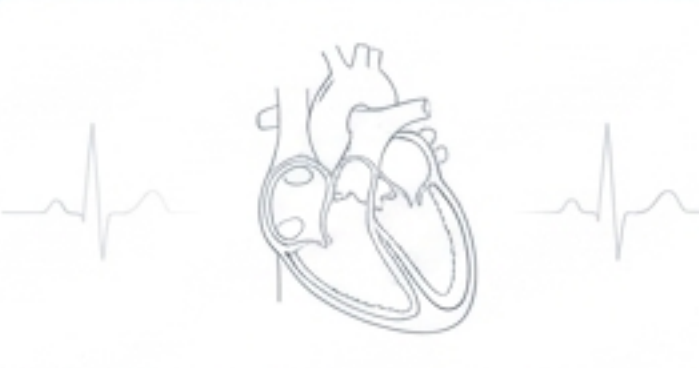
**Amiodarone (Class III)**

**Loading:** 300 mg IV over 20–30 min, then 900 mg/24h.

**Maintenance:** PO titrating to 200 mg daily.


**Toxicity:** TFTs, LFTs, PFTs,  slit-lamp exam.

**Status:** PBS General Benefit. 




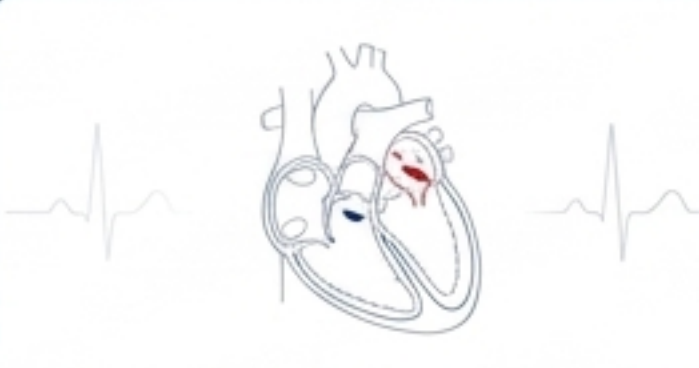
**Sotalol (Class III)**

**Dose:** 80–160 mg PO BD.

**Caution:** Contraindicated  if QTc >500 ms. Initiate in hospital.


**Renal:** Extend interval if eGFR 30–60.


**Status:** PBS General Benefit. 



**Flecainide (Class IC)**

**Dose:** 50–200 mg PO BD.

**CRITICAL WARNING:**  Contraindicated in structural heart disease (increased mortality per CAST trial).

**Status:** PBS General Benefit. 



**Mexiletine (Class IB)**

**Dose:** 200 mg PO TDS.

**Use:** Adjunctive therapy for refractory VT or channelopathies. 

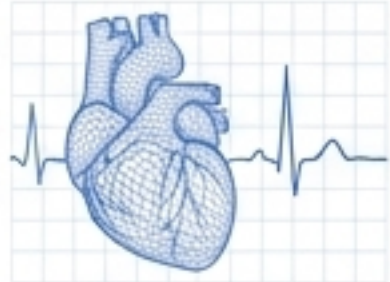
**Status:** Not PBS-listed (Requires TGA Special Access Scheme).  

# The Polymorphic VT Bifurcation Requires Immediate QT Interval Assessment

## Assess Bazett QTc

### Normal QT

QTc <440ms (M) / <460ms (F)



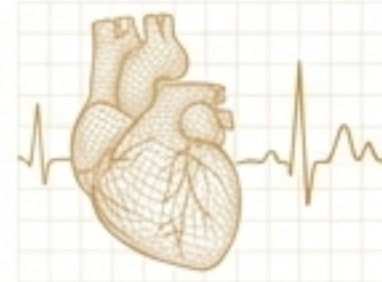
**Mechanism:** Primarily acute myocardial ischaemia.

#### Action:

- Anti-ischaemic therapy
- Emergency coronary angiography
- Beta-blockers
- Amiodarone or lidocaine (1–1.5 mg/kg IV)

### Prolonged QT

QTc ≥500ms



**Mechanism:** Torsades de Pointes (TdP). High Risk.

#### Action:

- Cease all offending drugs immediately (check CredibleMeds database for macrolides, psychotropics, antiemetics).
- Correct K<sup>+</sup> (>4.5) and Mg<sup>2+</sup> (>1.0).
- Continuous telemetry mandatory.

# Reversing Torsades de Pointes Demands Rapid Membrane Stabilisation

## The TdP Twist



### 1. Cease Agents



Immediate halt of all suspected agents (sotalol, haloperidol, escitalopram).

### 2. IV Magnesium Sulphate



First-line therapy. 2g (8 mmol) IV over 5–10 min, regardless of serum Mg level. Mechanism is membrane stabilisation, not replacement.

### 3. Increase Heart Rate



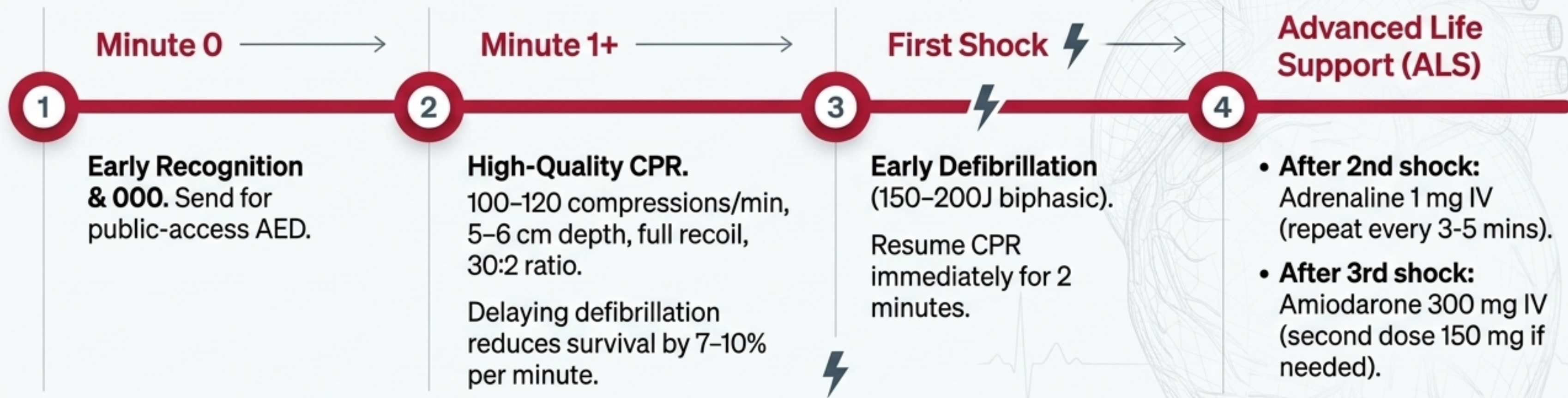
Isoprenaline (2–10 mcg/min) or transvenous pacing (90–110 bpm) to physically shorten the QT interval.

### 4. Defibrillate



If degenerating to VF, deliver immediate unsynchronised shock.

# Execution of the Out-of-Hospital Cardiac Arrest Defibrillation Protocol



## Identify Reversible Causes

### The 4Hs:

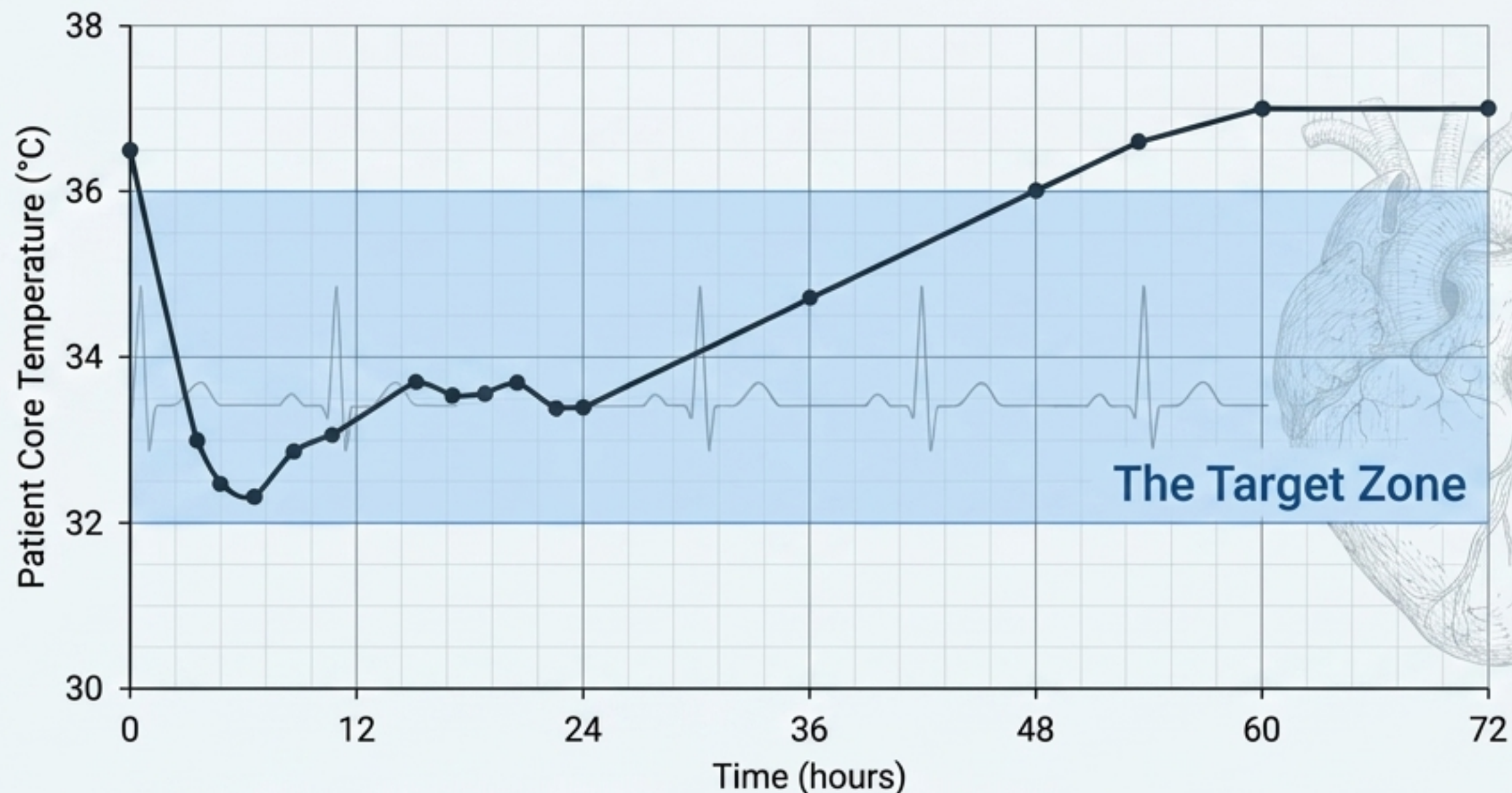
- Hypoxia
- Hypovolaemia

- Hypo/Hyperkalaemia
- Hypothermia

### The 4Ts:

- Tension pneumothorax
- Tamponade
- Toxins
- Thrombosis

# Post-Resuscitation Targeted Temperature Management Stabilizes the Neurological Substrate



## The Target Zone

Maintain 32–36°C for ≥24 hours post-ROSC for comatose patients (VF/pVT initial rhythm).



## The Rewarming Phase

Controlled rewarming at 0.25–0.5°C/hour.  
Maintain strict normothermia (<37.7°C) for 72 hours post-arrest.



## Monitoring Methods

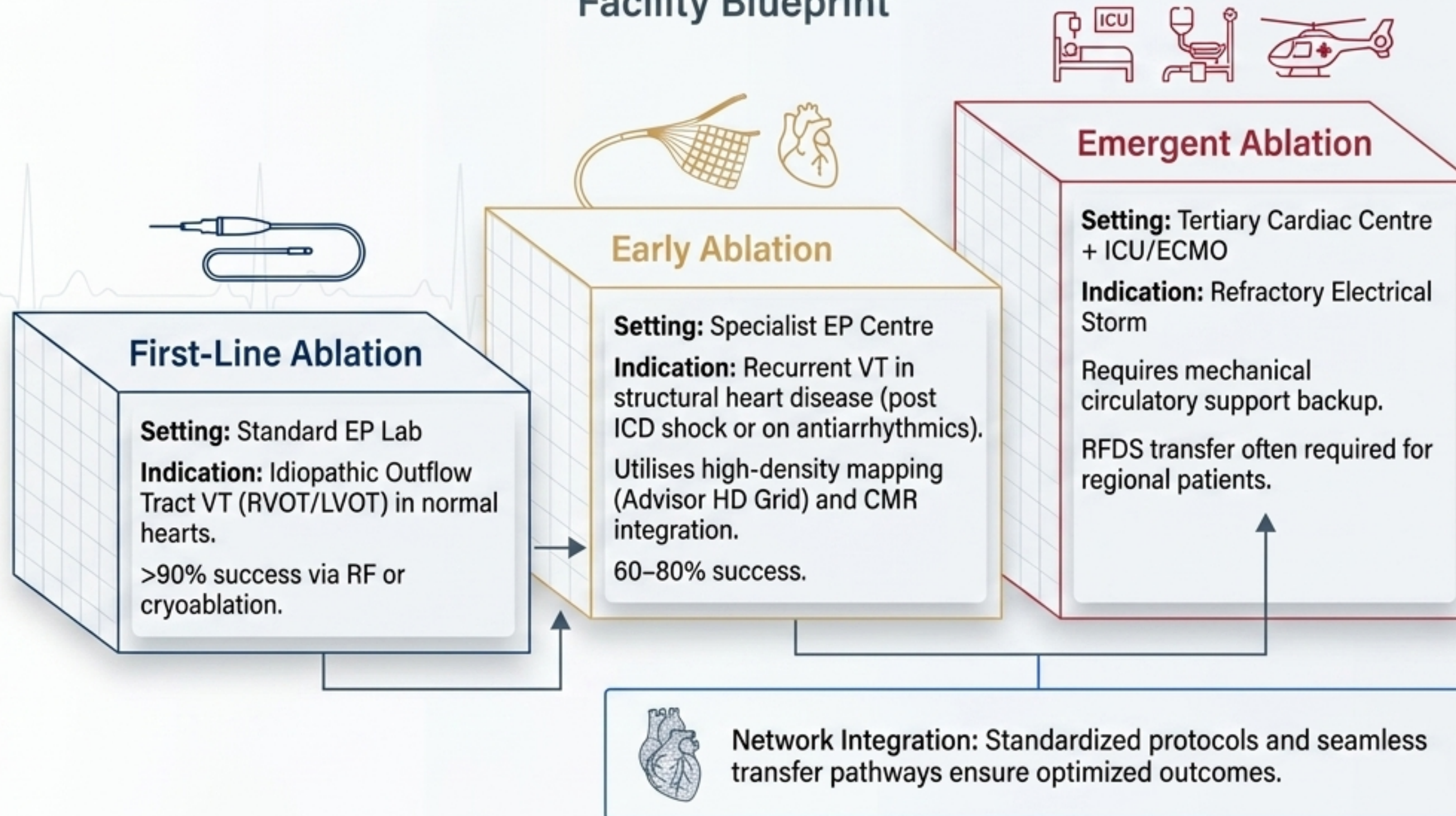
Surface cooling (Arctic Sun) or endovascular cooling catheter.  
Continuous core temp via oesophageal, bladder, or intravascular lines.

### Clinical Update (The TAME Trial 2023)

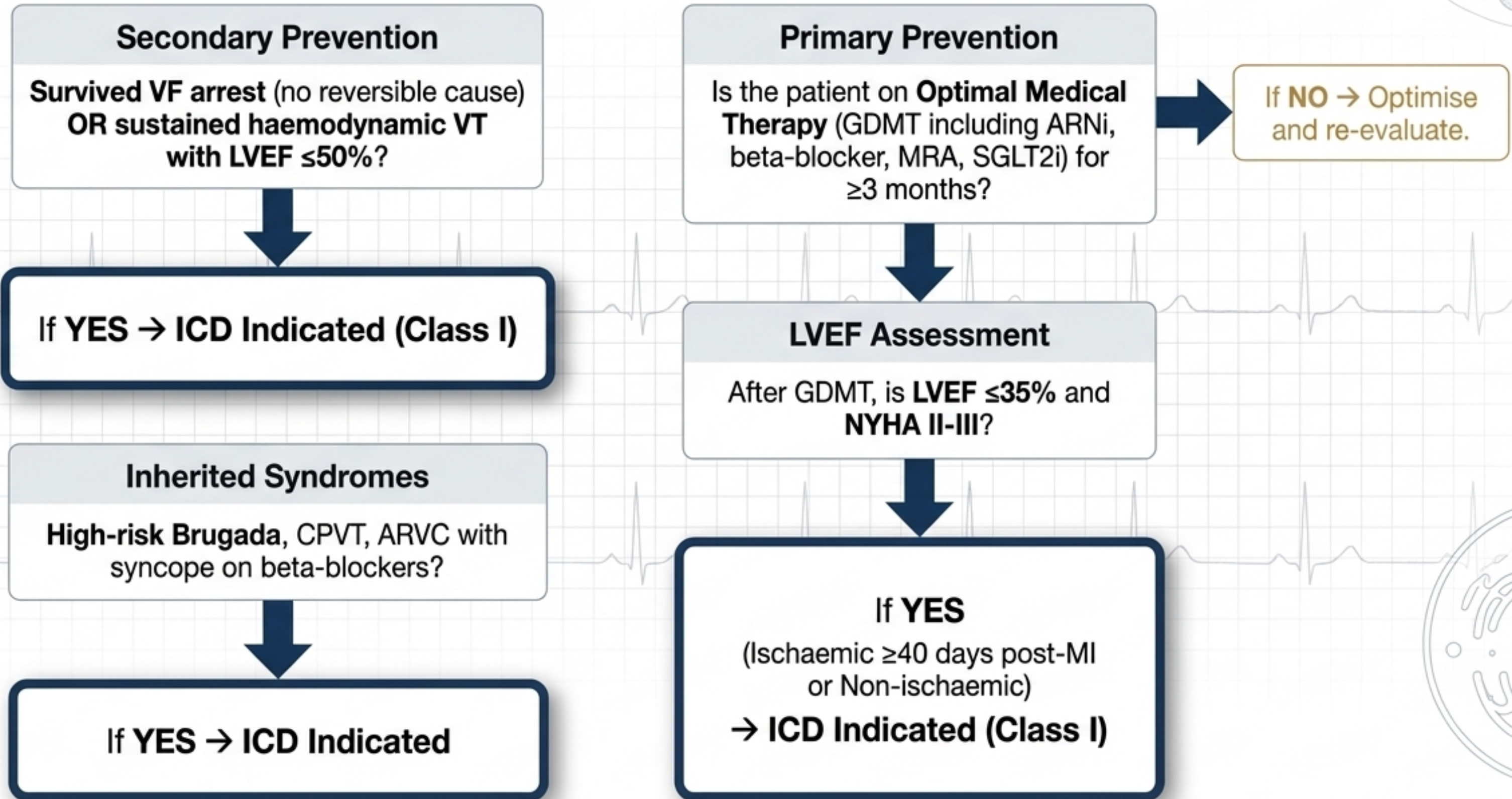
Routine pre-hospital rapid infusion of cold fluids is no longer recommended.  
Strict normothermia (avoiding active fever >37.7°C) may be as effective as deep hypothermia (33°C).

# Deploying Catheter Ablation Across the Health System Network

## Facility Blueprint



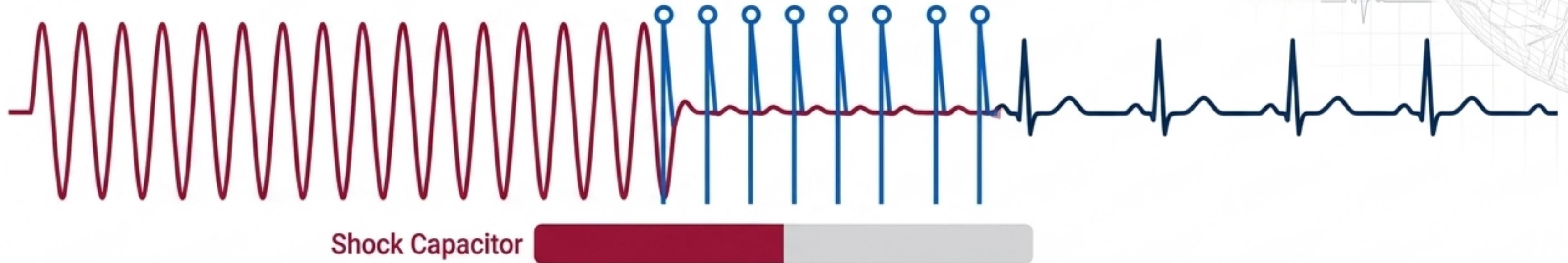
# Primary and Secondary Prevention ICD Implantation Logic



# Modern ICD Programming Minimizes Inappropriate Therapies Through Interception

## ATP Interception Model

Burst ATP – Anti-Tachycardia Pacing at 88% cycle length



ATP intercepts arrhythmia before shock delivery.

### Single-Zone Programming

Monitor zone with ATP before shock at  $\geq 200$  bpm.

No therapy  $< 200$  bpm.



### Detection Duration

Prolonged detection (30/40 intervals or  $\geq 6-12$ s) allows self-termination, avoiding painful, unnecessary shocks.



### Subcutaneous ICD (S-ICD)

Uses a conditional zone (200-250 bpm with morphology discrimination). Cannot deliver ATP, but avoids transvenous lead complications.



# Troubleshooting the Discharging ICD

Alert Type	Aetiology	Management
Appropriate Shock	True VT/VF detected.	Optimise <b>antiarrhythmic therapy</b> , consider <b>catheter ablation</b> , reassess device programming.
Inappropriate SVT Shock	AF, flutter, AVNRT crossing rate zones.	<b>Rate control, SVT ablation</b> , reprogram <b>discriminators</b> (sudden onset/morphology).
T-Wave Oversensing	Tall T-waves double-counted as R-waves; <b>hyperkalaemia</b> .	Reprogram <b>sensitivity vector</b> ; <b>correct electrolytes</b> .
<b>Lead Fracture / Noise</b>	Insulation breach or physical artifact.	<b>Urgent lead revision</b> ; evident via <b>impedance monitoring</b> on remote follow-up.

# Multidisciplinary Management of the Electrical Storm

**Definition:**  $\geq 3$  episodes of VT/VF within 24 hours requiring intervention.  
**15–25% in-hospital mortality.**

## 1. Haemodynamic Support

ICU admission. Consider *Impella/ECMO* for refractory hypotension. Correct ischaemia/electrolytes.



## 2. Intravenous Amiodarone

300 mg over 20-30 min, then **900 mg** over 24 hours.



## 3. Sympathetic Blockade

IV *metoprolol*, *propranolol*, or *esmolol* (ultra-short acting for haemodynamic concern).



## 4. Deep Sedation

*Propofol/ketamine* intubation to forcefully blunt the catastrophic catecholamine surge.



## 5. Urgent EP Rescue

*Catheter ablation* under circulatory support. Consider stellate ganglion block.

# Decoding Inherited Arrhythmia Syndromes

Brugada Syndrome	CPVT	ARVC	Congenital LQTS
<ul style="list-style-type: none"><li>• <b>Mutation:</b> SCN5A (20-30%).</li><li>• <b>Hallmark:</b> Type 1 coved ST elevation V1-V3.</li><li>• <b>Trigger:</b> Fever, large meals, sodium channel blockers.</li><li>• <b>Therapy:</b> ICD, avoid fever, Quinidine.</li></ul>	<ul style="list-style-type: none"><li>• <b>Mutation:</b> RYR2 (~60%).</li><li>• <b>Hallmark:</b> Normal resting ECG; Bidirectional VT on exercise.</li><li>• <b>Trigger:</b> Exercise, emotion.</li><li>• <b>Therapy:</b> Nadolol + Flecainide. Strict sport avoidance.</li></ul>	<ul style="list-style-type: none"><li>• <b>Mutation:</b> Desmosomal (PKP2).</li><li>• <b>Hallmark:</b> Fibro-fatty RV replacement, epsilon waves.</li><li>• <b>Trigger:</b> High-intensity endurance exercise.</li><li>• <b>Therapy:</b> Beta-blockers, Sotalol, Epicardial ablation.</li></ul>	<ul style="list-style-type: none"><li>• <b>Mutation:</b> LQT1 (exercise), LQT2 (alarms), LQT3 (sleep).</li><li>• <b>Hallmark:</b> QTc &gt;500ms.</li><li>• <b>Therapy:</b> Beta-blockers, Left cardiac sympathetic denervation (LCSD).</li></ul>

# The Structural and Electrical Investigation Toolkit



## The Baseline (ECG & Telemetry)

QTc measurement, bundle branch morphology, Holter/loop recording to document arrhythmia burden and correlate with symptoms.



## The Anatomy (Echocardiography)

Nationwide MBS access. Defines LVEF, RV structure (RVOT dimensions for ARVC), and valvular integrity.



## The Substrate (Cardiac MRI)

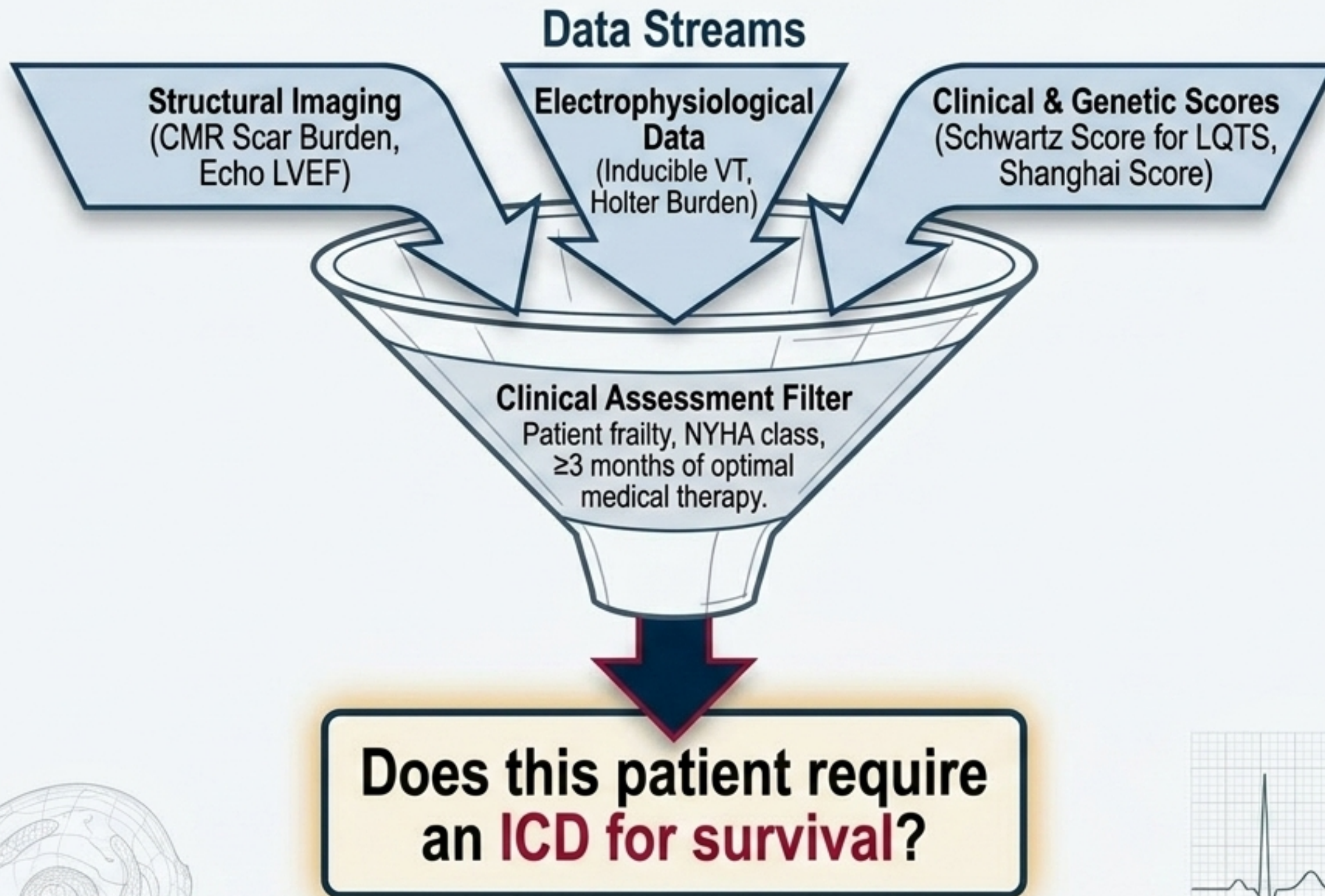
Essential for tissue characterisation. Late gadolinium enhancement (LGE) precisely maps myocardial scar and fibro-fatty infiltration.



## The Circuit (EP Study & Provocation)

Inducibility testing. Ajmaline/Flecainide provocation for Brugada (via SAS). Exercise testing specifically for CPVT diagnosis.

# The SCD Risk Stratification Funnel Unifies Diagnostic Pathways



# Modifying Treatment Parameters for Special Populations

## Pregnancy

- Synchronised cardioversion is safe.
- Metoprolol/labetalol preferred (Category C). Avoid atenolol.
- ICD implantation feasible with foetal shielding.
- LQT2 at high risk postpartum.

## Renal Impairment

- High risk of pro-arrhythmic electrolyte shifts ( $K^+$ ,  $Mg^{2+}$ ).
- Sotalol requires mandatory dose reduction/avoidance if eGFR <30.
- High TdP risk.
- Dialysis alters ICD infection risk.

## Paediatrics

- Outflow tract VT highly amenable to ablation (>85% success).
- Subcutaneous ICDs preferred to avoid transvenous lead growth issues.
- Weight-based dosing critical.

## Hepatic & Elderly

- Amiodarone heavily hepatically metabolised; avoid in Child-Pugh C.
- Elderly face high amiodarone toxicity and high risk of falls causing inappropriate ICD shocks.

# Geographical Disparities and First Nations Health Considerations

## First Nations Burden

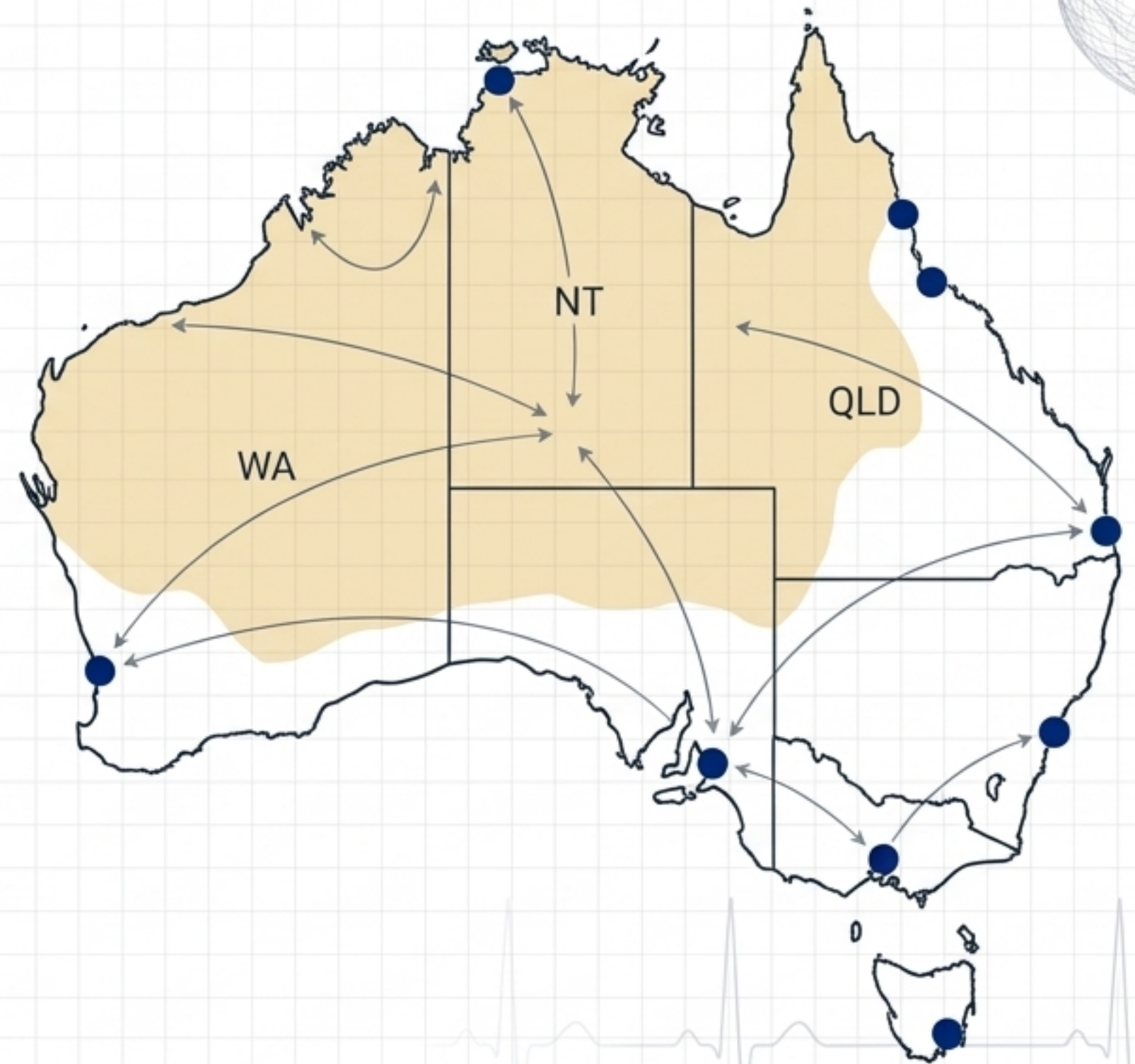
2–3x higher rate of SCD. Driven heavily by Rheumatic Heart Disease (RHD) substrate and high rates of non-ischaemic cardiomyopathy.

## The Access Gap

Catheter ablation and ICD implantation are geographically constrained. Remote communities rely on public-access AED expansion, RFDS retrieval, and the Closing the Gap (CTG) PBS co-payment measure.


## Cultural Safety

Coordinate with Aboriginal Medical Services (AMS). Accommodate Sorry Business in follow-up scheduling. Use family-based approaches for genetic cascade screening.




# Long-Term Device Surveillance and Psychosocial Integration


## Remote Monitoring Networks

- Daily transmissions via Medtronic CareLink, Abbott Merlin, etc. 
- Ensures early detection of VF, lead alerts, and heart failure metrics.
- Vital for remote Australian continuity of care.

## Driving Restrictions (Austroads)

- Primary prevention ICD: No restriction. 
- Secondary prevention ICD: Cease driving  $\geq 2$  weeks post-implant.
- **Post-Shock:** Cease driving for  $\geq 6$  months. Commercial licenses generally disqualified.

## The Psychological Impact

**ICD shocks induce severe anxiety, PTSD, and depression.** 

Routine PHQ-9/GAD-7 screening is mandatory. Refer to cardiac psychology and patient networks like SADS Australia.

Physical survival must be matched by psychological recovery.