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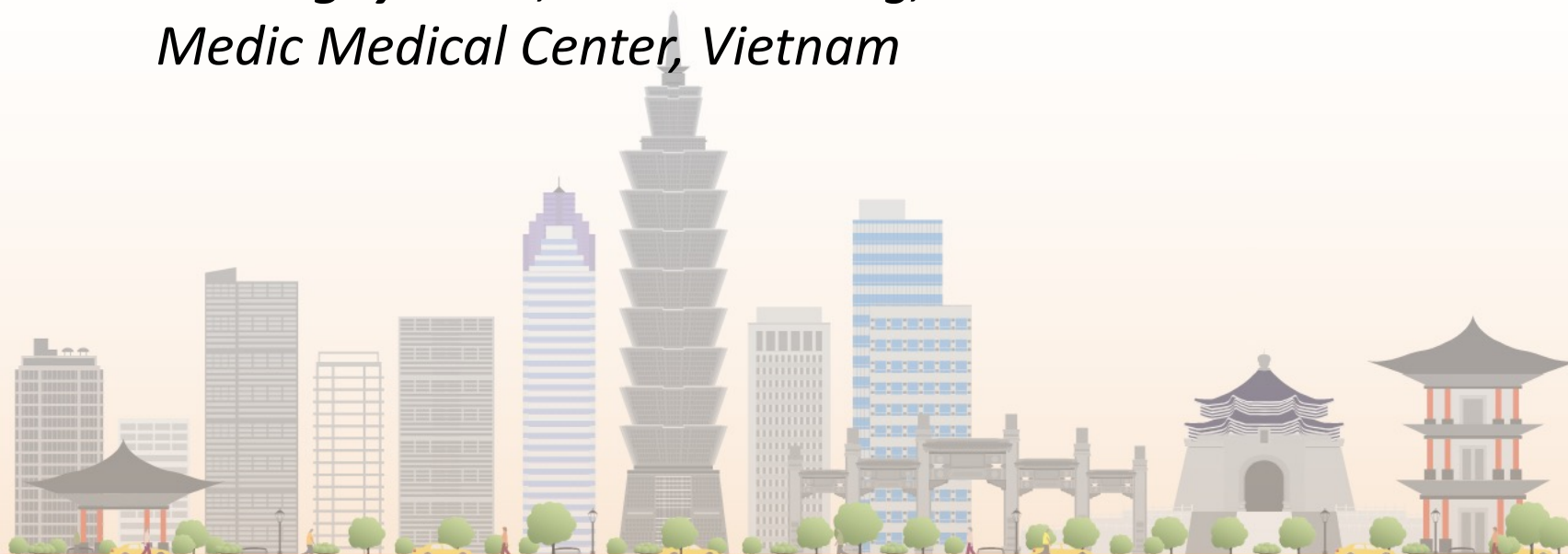
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Acute Myocarditis with Ring-Like Late Gadolinium Enhancement: A Case of Suspected Genetic Cardiomyopathy

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CLINICAL TIMELINE

A 28 year-old man. No prior cardiac history

Day-7 to admission

- Fever
- Abdominal fullness
- Intermittent chest pain
- Progressive dyspnea

Admission

- Troponin I: 164 -> 162 pg/ml
- NT-proBNP: 2255 pg/ml
- LVEF 22%
- AF

Early work-up

CMR: extensive non-ischemic injury

At first, everything fit acute myocarditis

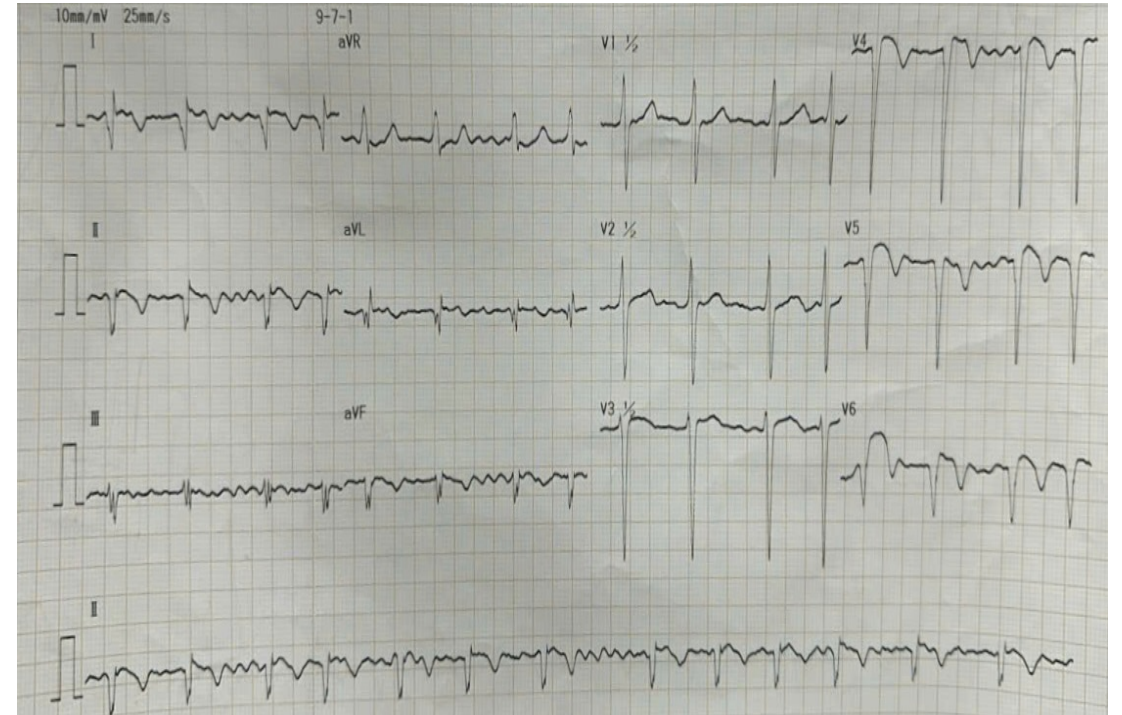
- Fever
- Chest pain
- Troponin elevation
- New-onset heart failure



But the ventricle was already too sick

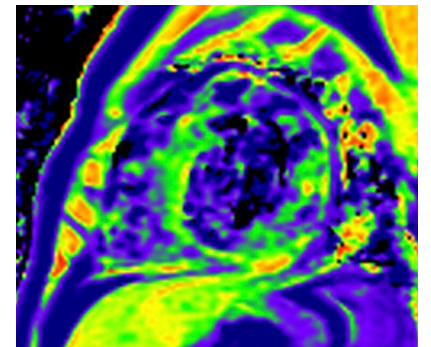
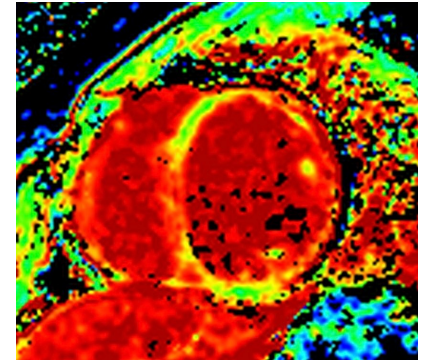
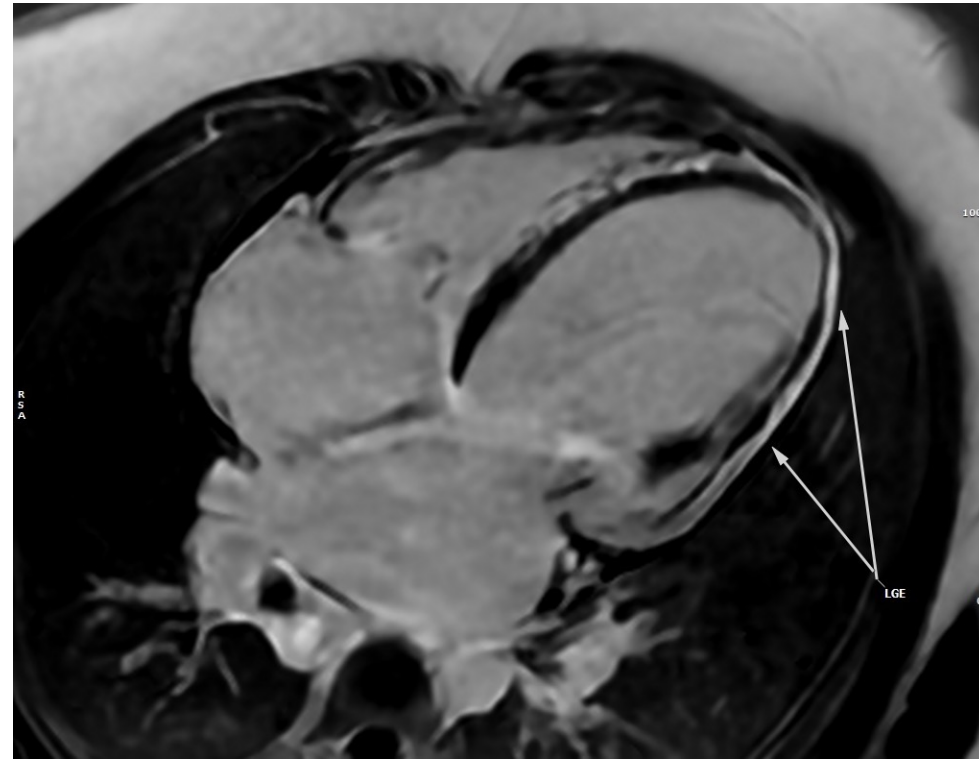
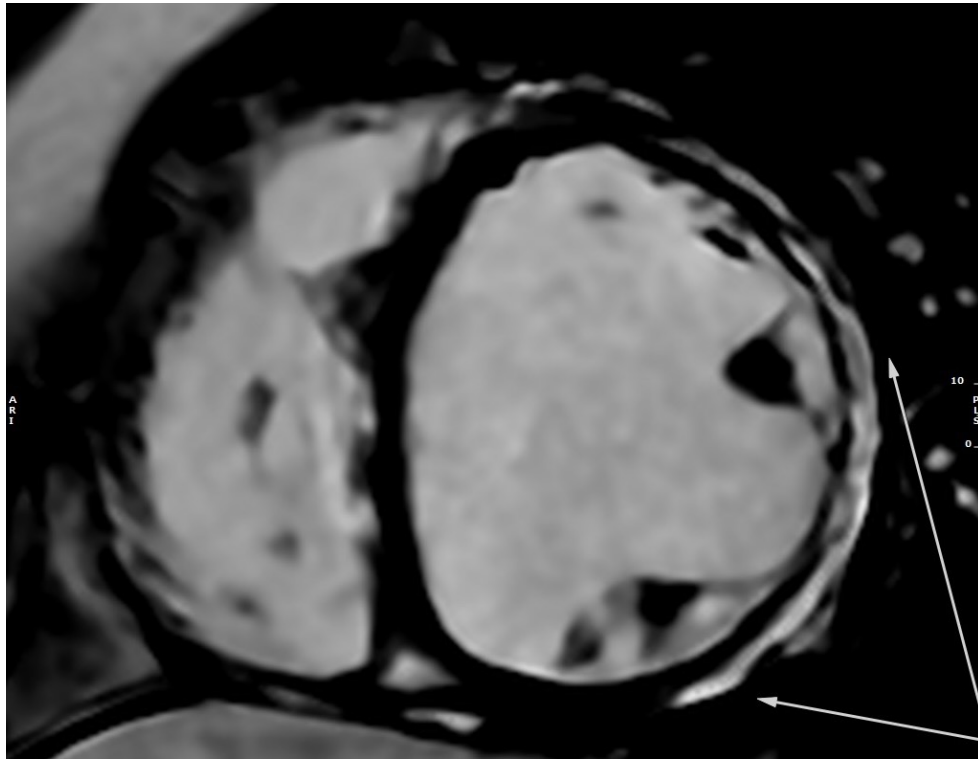


LV dilation and LVEF 22%



AF with controlled ventricular response

CMR made myocarditis alone uncomfortable



- Ring-like non-ischemic LGE. The total LGE mass was **21 g**, corresponding to approximately **21%** of the left ventricular mass.

→ **Suspected genetic cardiomyopathy**

- ECV 46%
- Native T1 elevated: 1160ms (Native T1 base: 1000ms)
- T2 mapping elevated: 63ms (T2 mapping base: 58ms)



What still fit — and what did not

Fit myocarditis

- Fever
- Chest pain
- Troponin rise
- Inflammatory CMR features

Did not fit

- EF 22%
- LV dilation
- Persistent AF
- Disproportionate fibrosis



Symptoms improved, but the substrate did not

- Chest pain improved
- Dyspnea improved
- EF did not significantly recover
- Holter ECG:
 - Atrial fibrillation throughout the entire recording period, with a ventricular response ranging from **35 to 128 bpm**, average **66 bpm**.
 - Atrial fibrillation with rapid ventricular response (>100 bpm) accounted for **<1%**; atrial fibrillation with slow ventricular response (<50 bpm) accounted for **5%**.

Why think of genetic DCM?

- Young patient
- Severe non-ischemic cardiomyopathy with ring-like LGE
- Persistent arrhythmia
- Incomplete recovery



The final clue: NGS result

- **Gene:** DMD
- **Variant:** NM_004006.3:c.31+1G>T
- **Zygoty:** Hemizygous
- **Type:** Canonical splice-site variant
- **Classification:** Pathogenic
- **Associated phenotypes:** DCM 3B / dystrophinopathy spectrum

Why this changed management

- Long-term HF follow-up
- Rhythm surveillance
- Maternal / family evaluation
- Cascade screening after confirmation



Learning point

- Do not stop at “myocarditis”
- Ring-like scar + poor recovery should trigger re-evaluation
- CMR did not end the diagnosis — it redirected it

