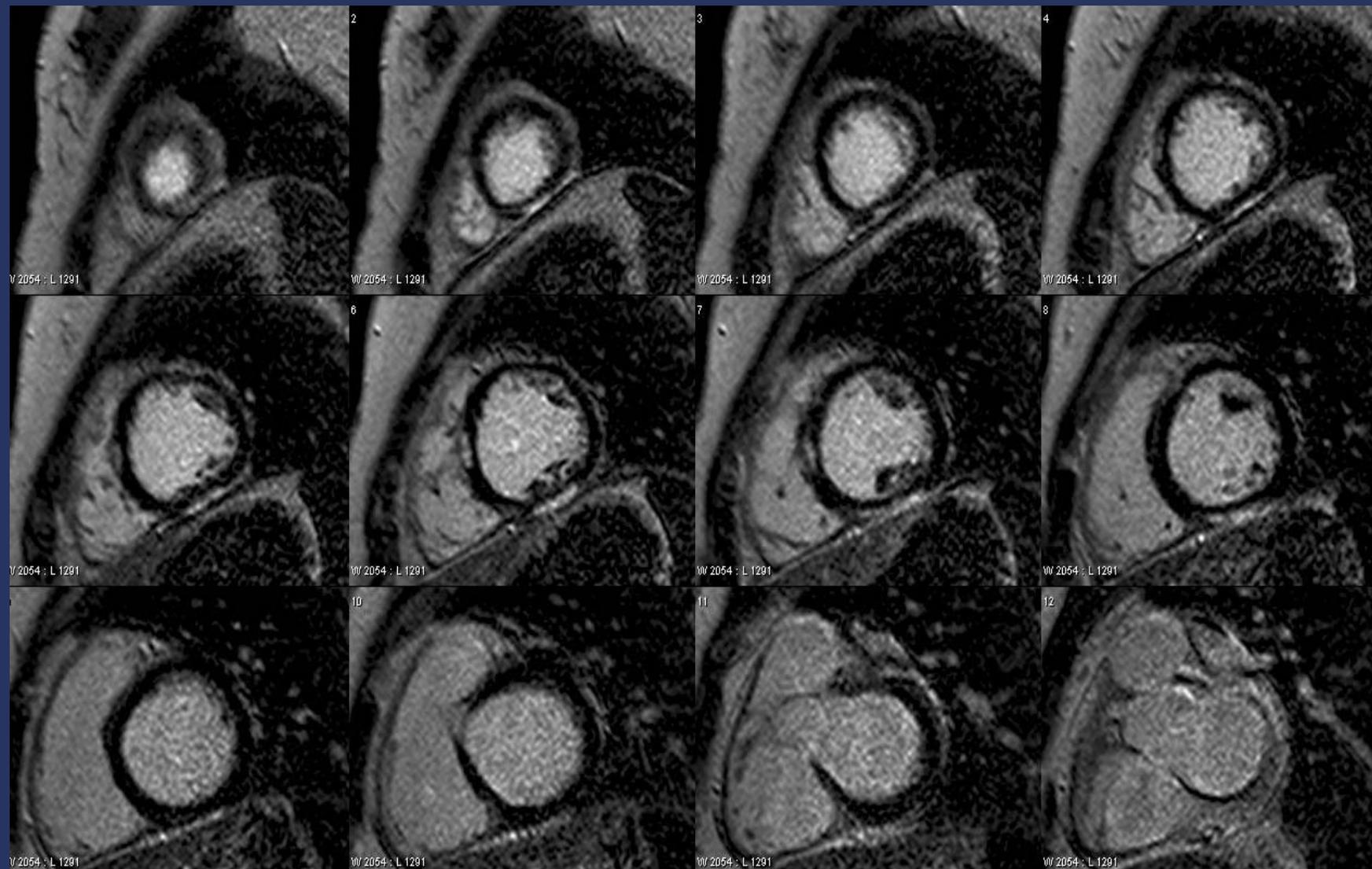


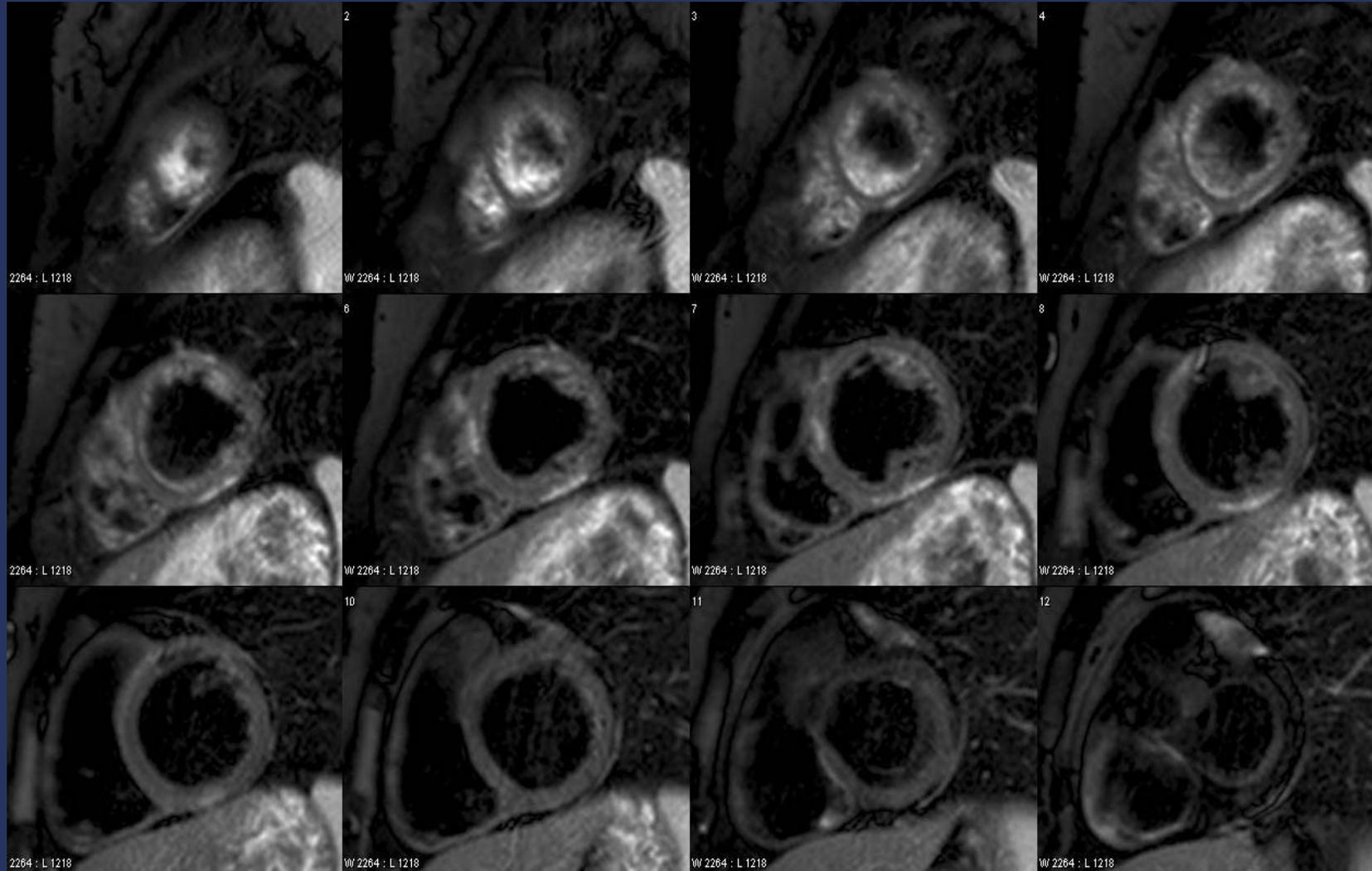
Case

- F/27
- C/C: acute chest pain (known vivax malaria))
- Cardiac enzyme: elevated
- Outside CAG: Normal

DE-MRI



T2-MRI





Myocarditis

- Occasionally sudden death or chronic DCM (5 - 10%)
- The onset of myocarditis is difficult to recognize clinically.
- Even when the diagnosis is considered, currently used diagnostic procedures suffer from limitations.
- (1) history of flu-like symptoms within 8 weeks before admission
- (2) one of the following symptoms:
fatigue/malaise, chest pain, dyspnea, or tachycardia
- (3) ECG signs: AV block, ST depression, or V. Tachycardia.

Mahrholdt H et al, Circulation. 2004;109:1250-1258

Myocarditis

- Contrast enhancement:
 - 28/32 patients (88%)
 - usually one or several foci in the myocardium.
(most frequently located in the lateral free wall)
- Initial vs F/U 3 month MRI
 - the area of contrast enhancement:
decreased from $9 \pm 11\%$ to $3 \pm 4\%$ of left ventricular mass
 - the left ventricular ejection fraction:
improved from $47 \pm 19\%$ to $60 \pm 10\%$.

Mahrholdt H et al, Circulation. 2004;109:1250-1258

Myocarditis

- Biopsy

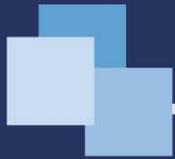
- 21 patients in whom biopsy was obtained from the region of contrast enhancement:
active myocarditis in 19 patients
(parvovirus B19, n=12; human herpes virus type 6, n=5).
- 11 patients in whom biopsy could not be taken from the region of contrast enhancement:
active myocarditis in one case only (HHV6).

Mahrholdt H et al, Circulation. 2004;109:1250-1258

Myocarditis

- Combined approach using T2-WI, CE-T1WI (before 4 min after contrast injection) and DE-MRI
- Global T2 SI and CE-T1WI were higher in patients than in controls.
 - T2 WI: 2.3 ± 0.4 vs 1.7 ± 0.4 ,
 - CE-T1WI : 6.8 ± 4.0 vs 3.7 ± 2.3

Myocarditis



	Sensitivity	Specificity	Diagnostic accuracy
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T2 WI	84%	74%	79%
CE-T1WI	80%	68%	74.5%
DE-MRI	44%	100%	71%
“Any-two”	76%	95.5%	85%

Abdel-Atyet al, JACC 2005;45:1815-1822



AMI vs Myocarditis

- Acute myocarditis may mimic acute myocardial infarction (AMI) when the patient has various combinations of chest pain, hemodynamic instability, ischemia-like electrocardiographic (ECG) changes, biochemical marker (troponin I and T and/or creatine kinase) changes, and segmental wall motion abnormalities at presentation.