

## Ventricular Tachycardia

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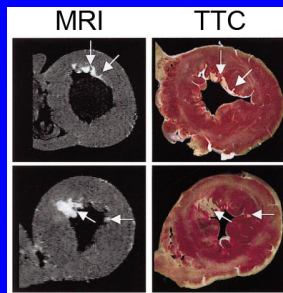
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## Presenter Disclosure Information

- There are no financial relationships relevant to this presentation.
- Off label use: MRI of patients with implanted devices; Gadolinium for cardiac studies

## Delayed Enhancement MRI

- Detailed imaging of scar and viable myocardium
- Extensively validated
- High spatial resolution



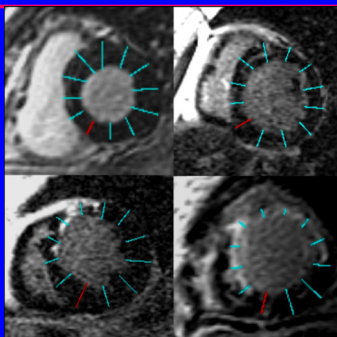
Lima, J.A., et al. *Circulation*, 1995; 92(5): p. 1117-25.  
Kim, R.J., et al. *NEJM*, 2000; 343(20): p. 1445-53.  
Kim RJ, et al: *Circulation* 1999;100:1992-2002  
Ashikaga et al. *Circ Res*, 2007; 101: p. 939-47

## Non-Ischemic Cardiomyopathy

Scar can be present and may be the substrate for VT

## Scar Patterns: Different Degrees of Transmurality

No Scar



1-25%

26-75%

76-100%

Nazarian, et al. *Circulation*. 2005;112:2821-2825

## Scar Pattern predicts VT Inducibility

- 7 different morphologies of sustained monomorphic ventricular tachycardia were inducible in 5 patients
- Median cycle length of 300 ms (interquartile range 240-345 ms)

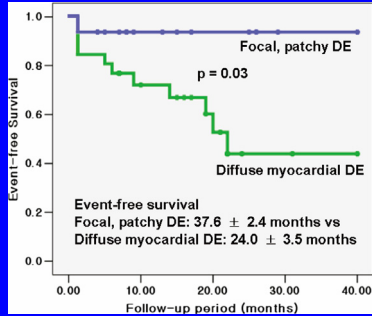
TABLE 2. MRI Parameters and Comparison After Stratification by Inducibility at Electrophysiological Study\*

	All Patients (n=26)	Electrophysiological Study		P
		Inducible (n=5)	Noninducible (n=21)	
Left ventricular ejection fraction, %	27 (17-43)	16 (13-29)	29 (19-44)	0.18
Left ventricular end-diastolic volume, mL	201 (129-273)	159 (154-273)	205 (129-259)	0.72
Left ventricular mass, g	153 (113-192)	184 (160-197)	144 (113-189)	0.38
Patients with scar, n (%)	17 (65)	5 (100)	12 (57)	0.13
Scar volume, %	4.6 (0.8-7.2)	6.0 (5.7-7.2)	1.5 (0.7-6.3)	0.09
Predominant scar distribution involving 26%-75% of wall thickness, n (%)	6 (23)	5 (100)	1 (5)	<0.001

\*Data are expressed as median (interquartile range) or absolute numbers (column percentage).

Nazarian, et al. *Circulation*. 2005;112:2821-2825

## Scar Pattern Predicts Event-Free Survival



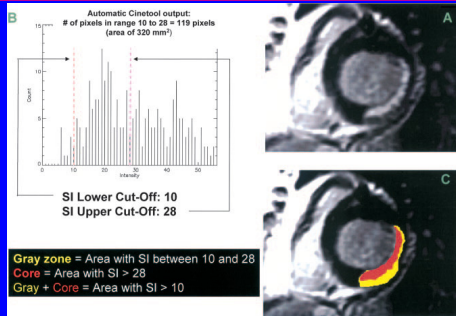
- 79 patients
- No ICDs
- EF 25-28%

Cho, et al. *Circ J.* 2010; 74: 476 – 483

## Ischemic Cardiomyopathy

Hyperenhancement may not have uniform intensity

## Gray Zone: Intermediate level of hyperenhancement



Schmidt et al. *Circulation.* 2007;115:2006-2014

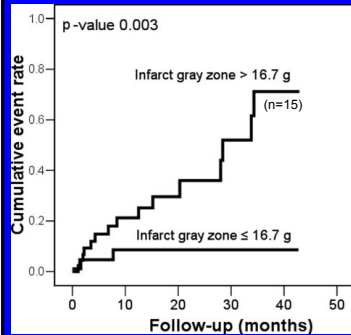
## Extent of Gray Zone Predicts Inducibility

Schmidt et al. *Circulation.* 2007;115:2006-2014

TABLE 2. MRI Indices According to Inducibility Status at Electrophysiology Study

Variable	Noninducible (n=27)	Inducible (n=20)	P
MRI LVEF	0.30±0.10	0.29±0.07	0.79
LV end-diastolic volume, mL	220±70	228±57	0.68
LV end-systolic volume, mL	156±61	162±44	0.71
LV end-diastolic mass, g	146±46	132±30	0.23
Infarct location, n (%)			0.23
Anterior±other territory	15 (56)	15 (75)	
Inferior and/or lateral only	12 (44)	5 (25)	
Transmurality infarct extent: % of sectors grouped by quartiles of transmural extent			
No infarct	51±15	45±9	0.11
1% to 25% infarct transmural extent	8±4	7±2	0.61
26% to 50% infarct transmural extent	8±3	8±5	0.88
51% to 75% infarct transmural extent	11±5	12±5	0.39
76% to 100% infarct transmural extent	23±14	28±11	0.17
Extent of hyperenhancement, g			
Total (core+gray)	34±17	40±11	0.17
Infarct core	21±10	21±5	0.95
Gray zone	13±9	19±8	0.015

## Extent of Gray Zone Predicts ICD Firings



- 91 patients with MI
- 18 patients with ICD firing

Roes S, Bax J et al. *Circ Cardio Imaging.* 2009;2:183-190.

## Extent of Gray Zone Predicts ICD Firings

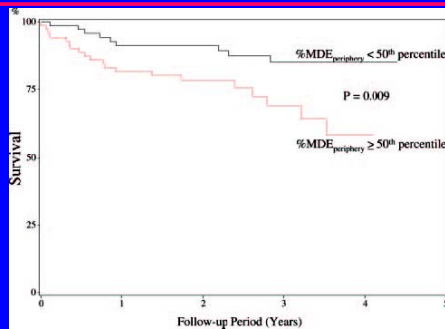
Table 2. Baseline MRI Variables

Variable	Total Population (n=91)	No Appropriate ICD Therapy (n=73)	Appropriate ICD Therapy (n=18)	P Value
LVEF, %	28±9	29±9	25±7	0.06
LV EDV, mL	333±112	331±117	339±95	0.8
LV ESV, mL	245±107	241±110	259±94	0.5
LV mass, g	148±40	148±41	149±38	0.9
Total infarct (infarct core+gray zone), g	46±25	43±23	58±29	0.02
Infarct core, g	26±17	25±16	30±17	0.2
Infarct gray zone, g	20±13	18±11	28±16	0.002

Data are expressed as mean±SD. LV indicates left ventricular; LV EDV, LV end-diastolic volume; LV ESV, LV end-systolic volume; LVEF, LV ejection fraction.

Roes S, Bax J et al. *Circ Cardio Imaging.* 2009;2:183-190.

## Extent of Gray Zone Predicts Survival



- 144 patients with CAD
- No ICD's
- Mean EF 42-45%

Yan A, Stevenson W, et al. *Circulation*. 2006;114:32-39

## Extent of Gray Zone Predicts Survival

TABLE 2. Univariable and Multivariable Associations With All-Cause Mortality

Variable	Univariable Analysis		Multivariable Analysis 1		Multivariable Analysis 2	
	Unadjusted HR (95% CI)	P	Adjusted HR (95% CI)	P	Adjusted HR (95% CI)	P
Age*	1.37 (0.96-1.97)	0.09	...	...	1.45 (0.98-2.14)	0.06
Diabetes	2.21 (1.05-4.66)	0.04	...	...	...	...
Previous percutaneous coronary intervention	0.37 (0.16-0.88)	0.02	...	...	...	...
Previous coronary bypass surgery	2.39 (1.08-5.30)	0.03	...	...	...	...
QRS duration >120 ms	2.29 (0.98-5.38)	0.06	...	...	...	...
Left bundle-branch block	3.21 (1.22-8.44)	0.02	...	...	...	...
Corrected QT >440 ms	2.37 (1.14-4.93)	0.02	...	...	...	...
LV end-diastolic volume index†	1.13 (1.04-1.22)	0.004	...	...	...	...
LV end-systolic volume index†	1.13 (1.05-1.23)	0.002	1.16 (1.07-1.26)	<0.001	...	...
LVEF‡	1.36 (1.08-1.72)	0.01	...	...	1.52 (1.19-1.94)	<0.001
%MDE <sub>periphery</sub> §	1.31 (1.06-1.63)	0.01	1.45 (1.15-1.84)	0.002	1.42 (1.11-1.81)	0.005
MDE <sub>periphery</sub> ¶	1.06 (1.00-1.11)	0.035	...	...	...	...

For model 1, predictors on univariable analysis ( $P < 0.10$ ) were evaluated in this multivariable model by forward stepwise selection ( $P < 0.10$  for entry and  $P > 0.05$  for removal) criteria. For model 2, age, LVEF, and %MDE<sub>periphery</sub> were all entered into this multivariable model.

\*HR per decade increase.

†HR per 10-mL/m<sup>2</sup> increase.

‡HR per 10% decrease.

§HR per 10% increase.

Yan A, Stevenson W, et al. *Circulation*. 2006;114:32-39

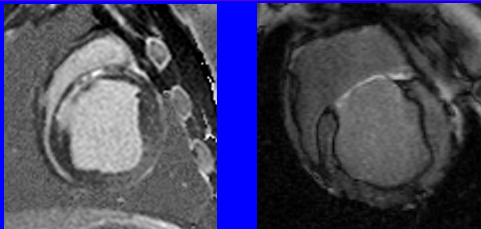
## Extent of Scar can Predict Mortality

- 349 patients (76% men) with EF of 24%.
- 56 events (51 deaths and 5 CTx)
- Mean scar percentage higher in patients with events: ( $39 \pm 22$  vs.  $30 \pm 20$ ,  $p = 0.003$ )
- Transmurality score higher in patients with events ( $9.7 \pm 5$  vs.  $7.8 \pm 5$ ,  $p = 0.004$ ).
- Effect of Gray Zone ???

Kwon et al. *JACC: Cardiovascular Imaging* 2009; 2(1): 34-44.

## What is the Gray Zone?

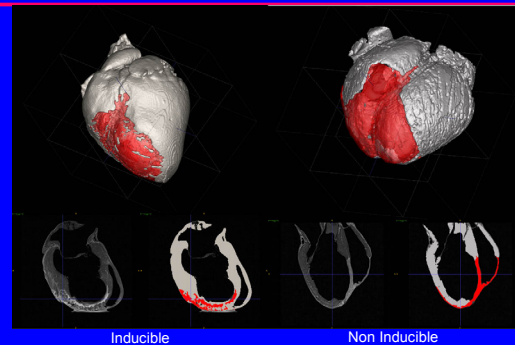
## 10 Week Swine Infarct Model: Inducibility Correlates with Gray Zone



- Inducible VTs in 8 of 17 pigs
- Gray Zone:  $21 \pm 8$  % infarct
- Non-Inducible in 9 of 17 pigs
- Gray Zone:  $12 \pm 3$  % infarct

$p < 0.009$

## Gray Zone = Tissue Heterogeneity High Resolution MRI (0.4x0.4x0.4 mm voxels)

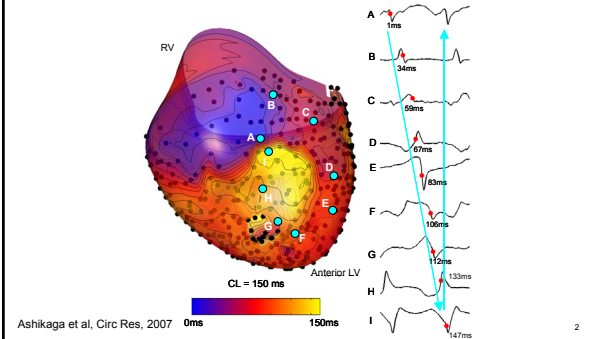


What do the images tell us about the mechanisms of VT?

Electroanatomic Correlations

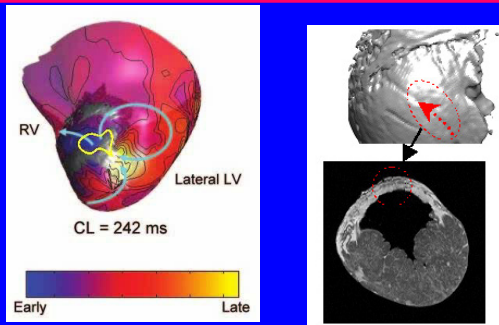
### Epicardial Electrograms during VT

Recordings from Multielectrode Epicardial Sock



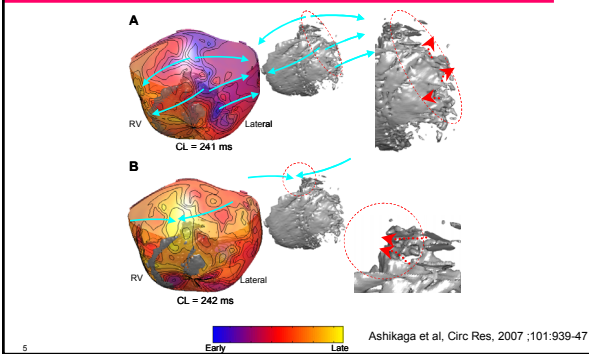
### Epicardial Reentry Registered with Scar:

Reentry Through Viable Tissue in Scar



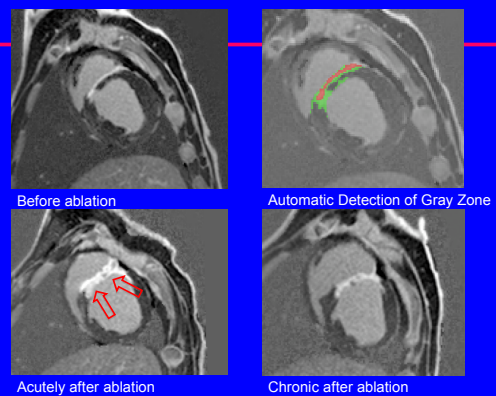
### Epicardial Centrifugal Pattern VT

Reentry Through and Around Scar

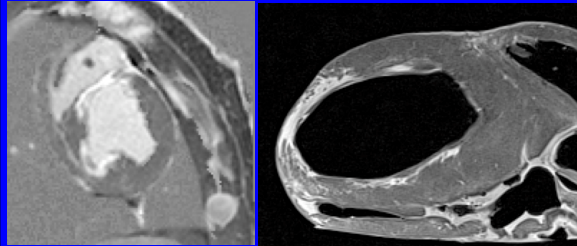


Is the Gray (Heterogeneous) Zone identified on MRI the critical pathway for VT generation?

### Ablation in Gray Zone Can Eliminate VTs Chronically



## Incomplete ablation of the Gray Zone: Inducible VT



Standard resolution MRI

High resolution MRI

## Sarcoidosis

### Sarcoidosis

DE-MRI

Patient	DE-CMR	
	Baseline	Follow up
<b>Patient A</b> Subendocardial HE in LV inferolateral wall - CAD-type		
<b>Patient B</b> Transmural HE in LV anterior wall - CAD-type RV side of septum HE - non-CAD-type		 None because of cardiothoracic implantation
<b>Patient C</b> Midwall HE in LV inferolateral wall - non-CAD-type		
<b>Patient D</b> Episcleral HE in LV inferior wall - non-CAD-type RV side of septum HE - non-CAD-type RV free wall HE - non-CAD-type		
<b>Patient E</b> Multiple, patchy, primarily mural HE - non-CAD-type		

Jaroudi, Kim, et al: Circulation. 2009;120:1969-1977.

### Outcomes

Any Event	DE-CMR		JMH		Cardiac Death	DE-CMR		JMH	
	(n=21)	(n=80)	(n=10)	(n=71)		(n=21)	(n=60)	(n=10)	(n=71)
No	15 (71%)	58 (97%)	7 (70%)	66 (93%)	No	17 (81%)	59 (98%)	8 (80%)	68 (96%)
Yes	6 (29%)	2 (3%)	3 (30%)	5 (7%)	Yes	4 (19%)	1 (2%)	2 (20%)	3 (4%)

**A**

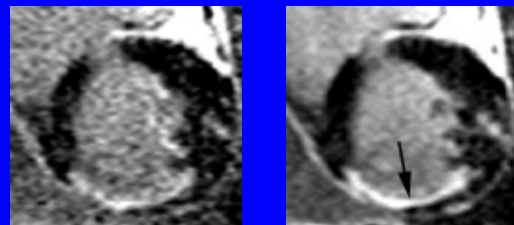
**B**

Jaroudi, Kim, et al: Circulation. 2009;120:1969-1977.

## Limitations of MRI Scar Imaging

- Motion Effects
- Partial Volume Effects
- Artifacts from ICD
- Nephrogenic Systemic Fibrosis

## Motion can Reduce Resolution



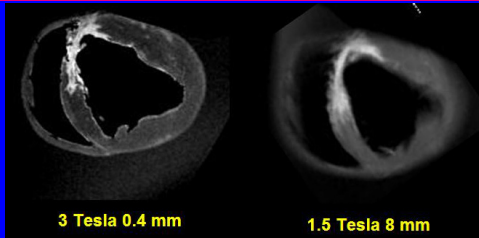
No Motion Correction

Motion Corrected

Kellman P, McVeigh ER, et al. 2002;47(2):372-383. Magn Reson Med.

## Partial Volume Effects can Reduce Resolution

Intensity Averaging with thick slices



3 Tesla 0.4 mm

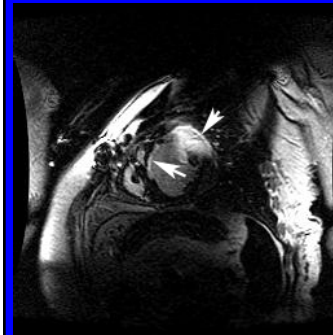
1.5 Tesla 8 mm



Lardo Lab

## Artifacts from ICD

Can Mimic HyperEnhancement



Imaging safe with ICD

- 555 MRI examinations performed in 460 patients
- Special protocol used
- No adverse events
- No clinically significant changes in pacing parameters
- Adequate scar imaging in 60-80%

Fixes are being developed

## Nephrogenic Systemic Fibrosis

- Scleroderma like syndrome
- Develops in patients with severe renal disease
  - Generally on dialysis
- Rare, not reversible
- First reported in 2000
- Linked to Gadolinium use in 2006
- Mechanism(s) unknown
- Does not occur in patients with GFR > 30 ml/min

## Summary

- Delayed enhancement MRI can accurately identify detailed anatomy of scar
- Multiple VT circuits can exist through and around areas of scar
- Scar transmural distribution appears to be related to arrhythmogenic potential in Non-Ischemic Cardiomyopathy
- The critical zones for occurrence of VT in ICM are likely areas of tissue heterogeneity (gray zone) that can be imaged with MRI
- The extent of scar heterogeneity is related to the risk of sudden death in ischemic cardiomyopathy
- Delayed enhancement MRI may be very useful in diagnosing cardiac sarcoidosis
- Limitations include motion effects, partial volume effects, and the presence of an ICD

## Clinical Implications

- Delayed enhancement MRI may help predict risk of arrhythmias, and probably sudden death
- Delayed enhancement MRI has added predictive power over that of EF alone in ICM
- Delayed enhancement MRI will likely lead to more accurate indications for ICD placement, and may help guide VT ablation
- Delayed enhancement MRI may be useful in guiding therapy of possible cardiac sarcoidosis

## Investigators

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