

## Barts and The London Cardiovascular Biomedical Research Unit

# Bipolar vs Unipolar mapping for VT

Richard Schilling

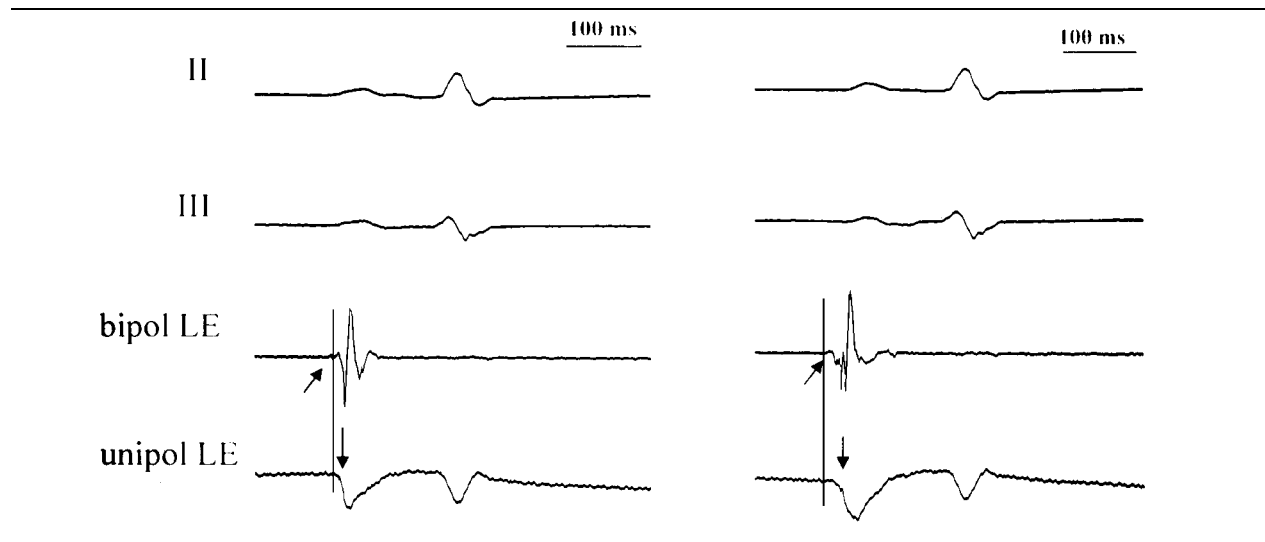


# Disclosures

- Research agreements with:
  - Biosense Webster
  - St Jude
  - Medtronic
  - Hansen medical
  - Boston Scientific

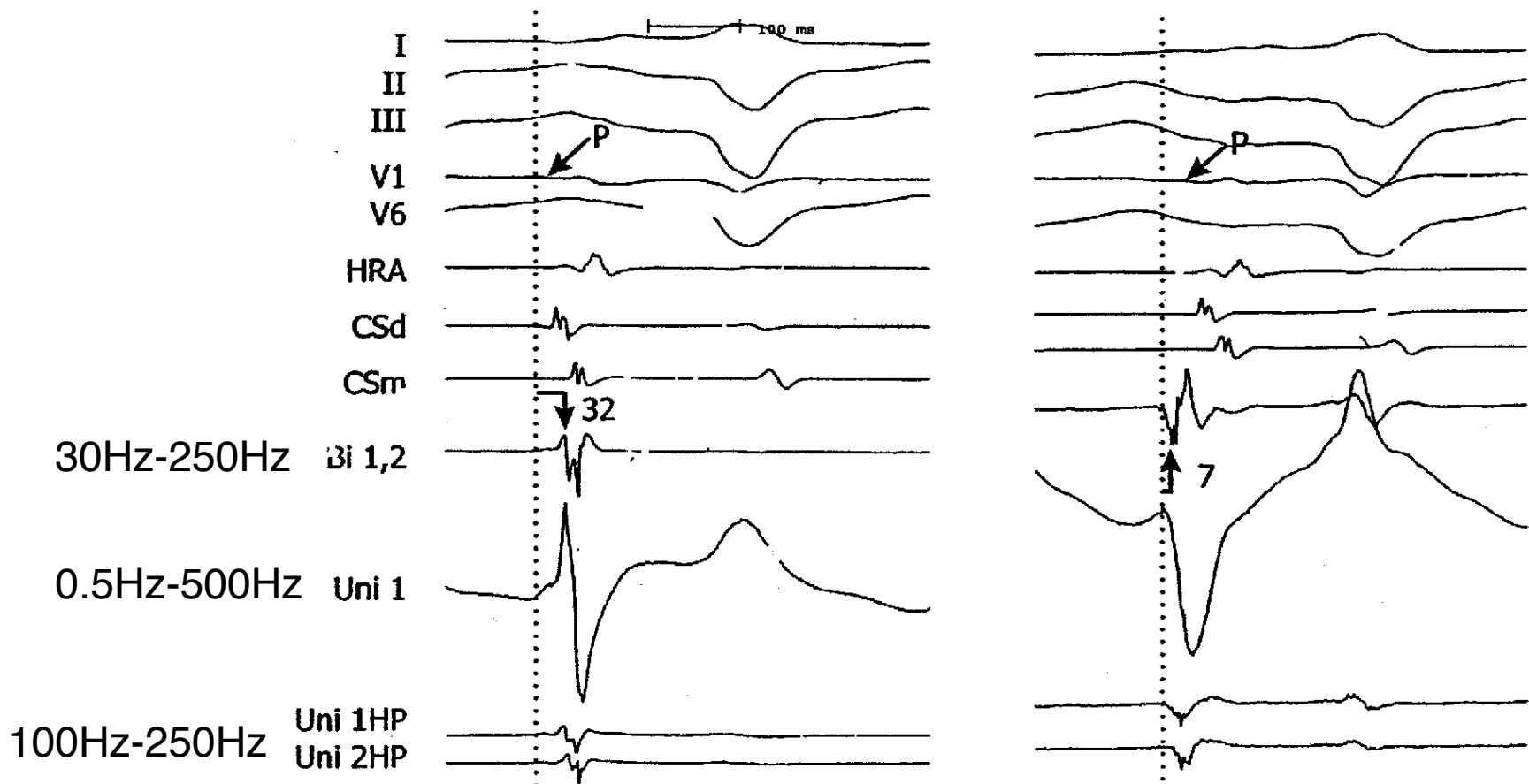
# Bipolar vs unipolar

- Bipolar easier to time than unipolar for LAT



# Single catheter mapping of AT

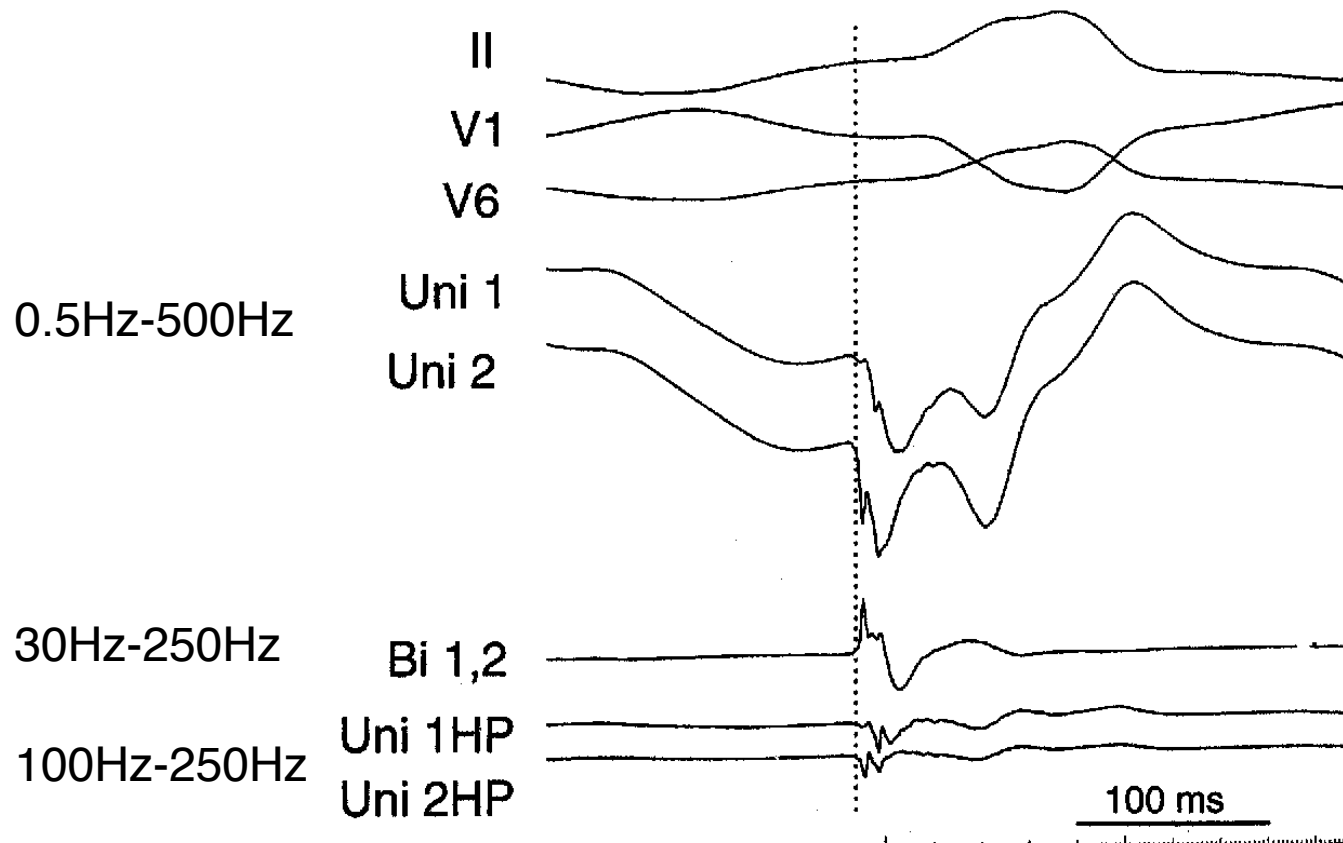
- Unipole marks p wave onset





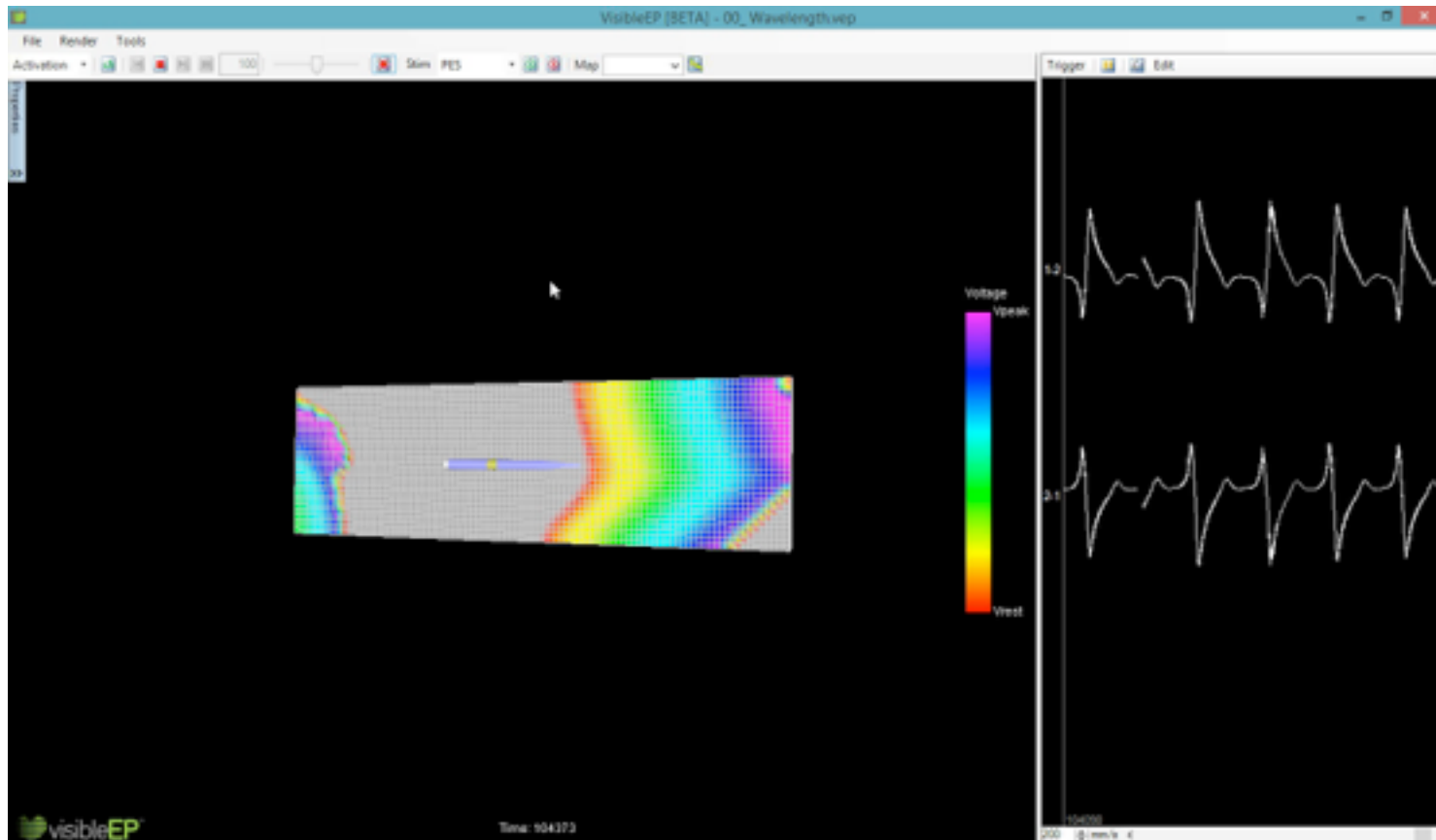
# Single catheter mapping of AT

- Potential errors



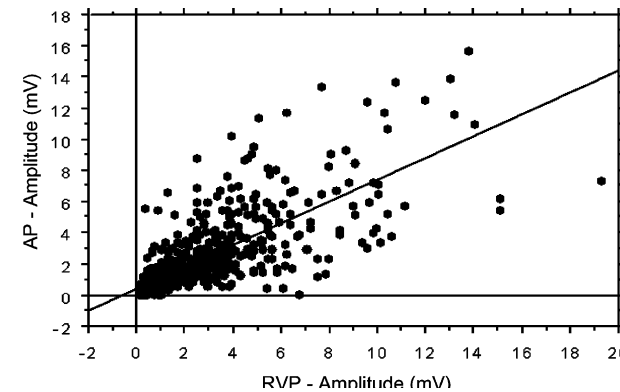
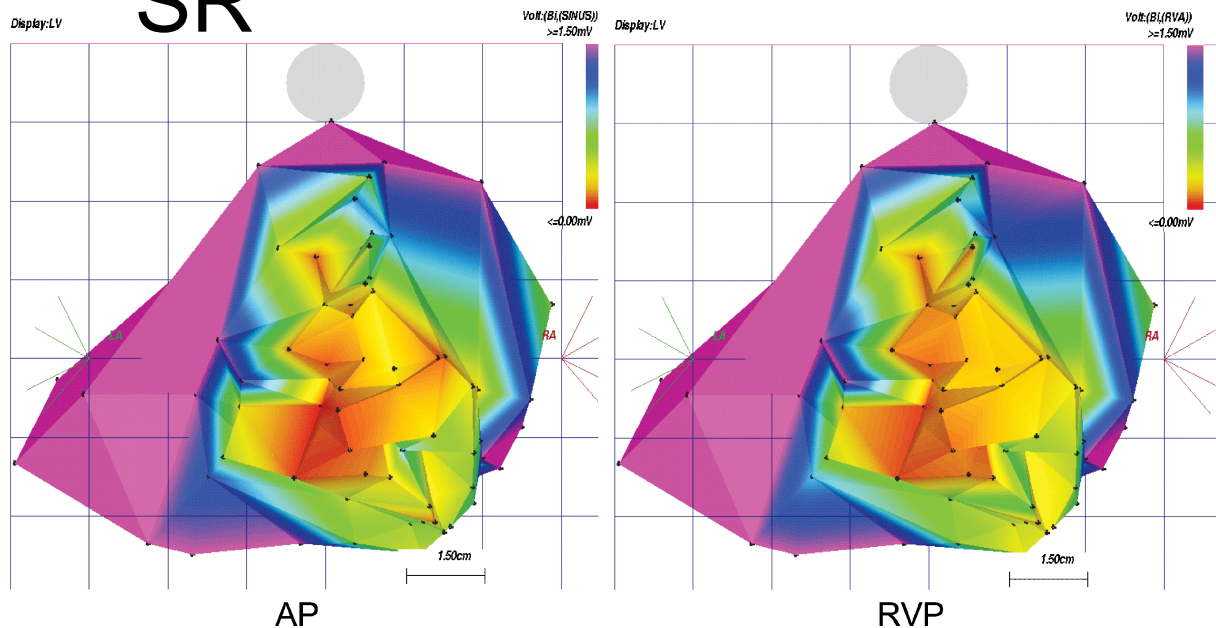
# Bipolar voltage mapping

- Voltage dependent on catheter orientation and electrode spacing



# Bipolar voltage mapping

- Wavefront changed by V or A pacing during SR



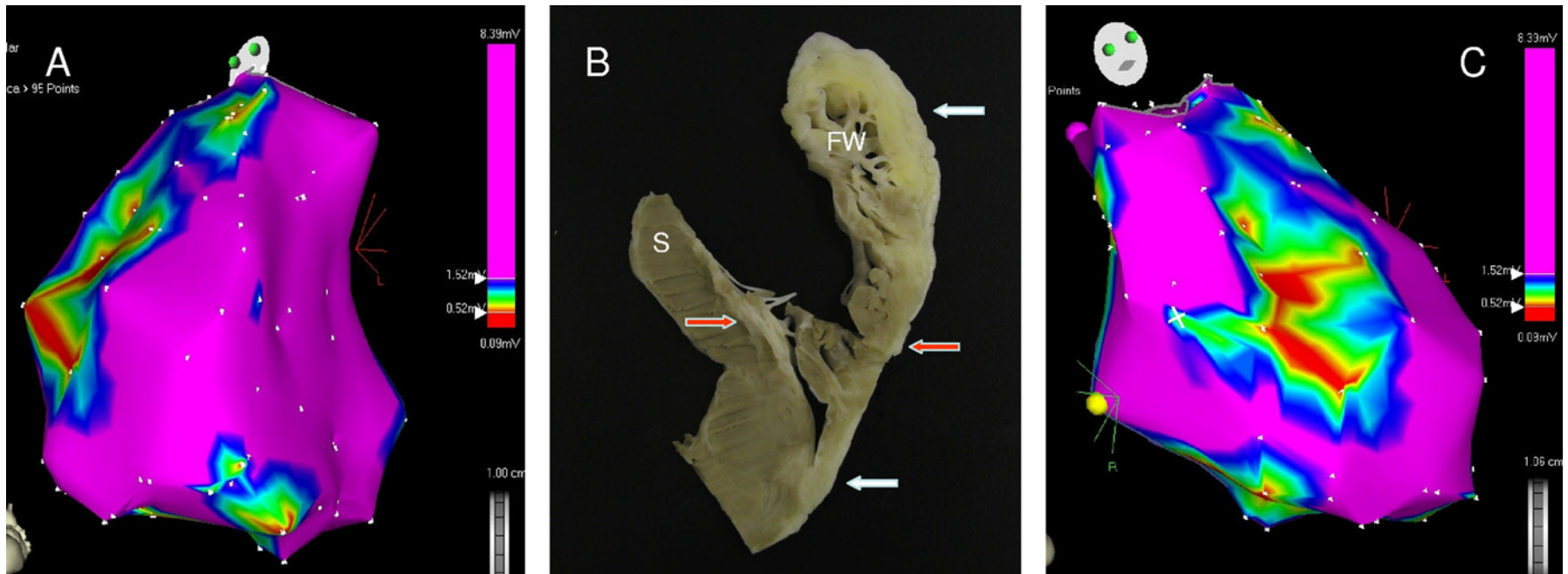
- $\Delta$  wavefront  $\Rightarrow$   $\Delta$  voltage but primarily in normal voltage areas

# Identification of epicardial ventricular scar

- Useful if:
  - failed endocardial ablation/  
substrate map
  - potential epicardial source e.g.  
non-ischaemic cardiomyopathy

# Identification of non-endocardial scar

- Pt undergoing transplant for ARVC

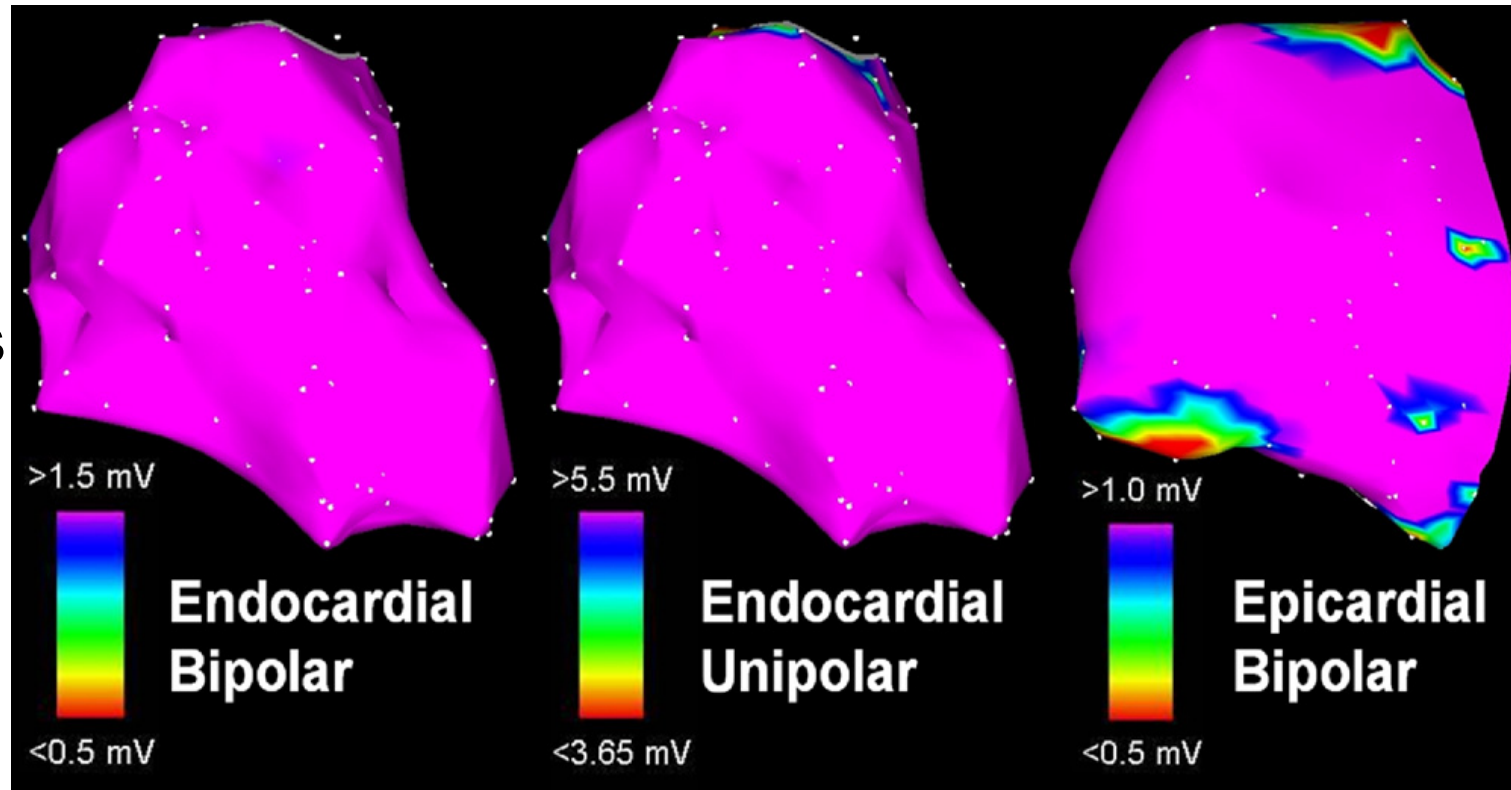


- Scar missed if  $>40\%$  of myocardial wall preserved

# Arrhythmogenic cardiomyopathy

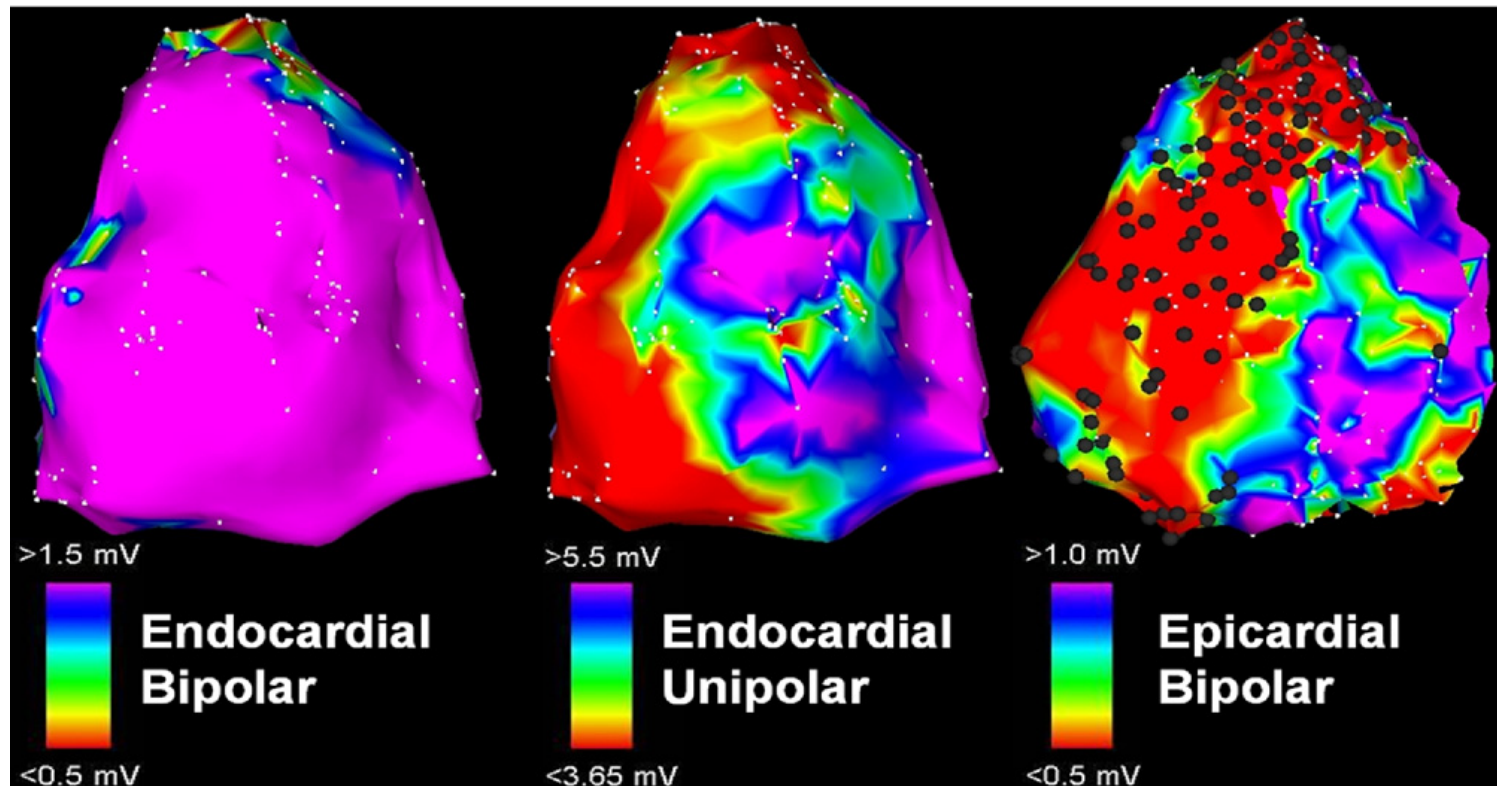
- Endocardial and epicardial bipolar maps cf unipolar endocardial

95% Uni  
>5.5mV in  
normal hearts



# Arrhythmogenic cardiomyopathy

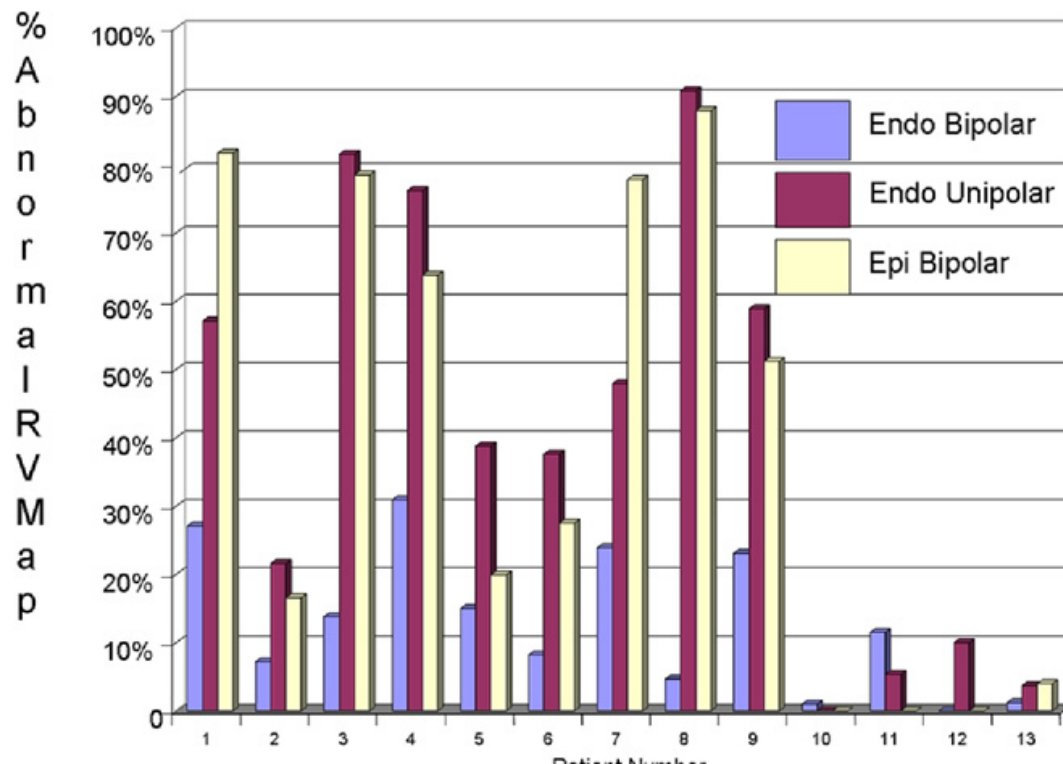
- Endocardial and epicardial bipolar maps cf unipolar endocardial





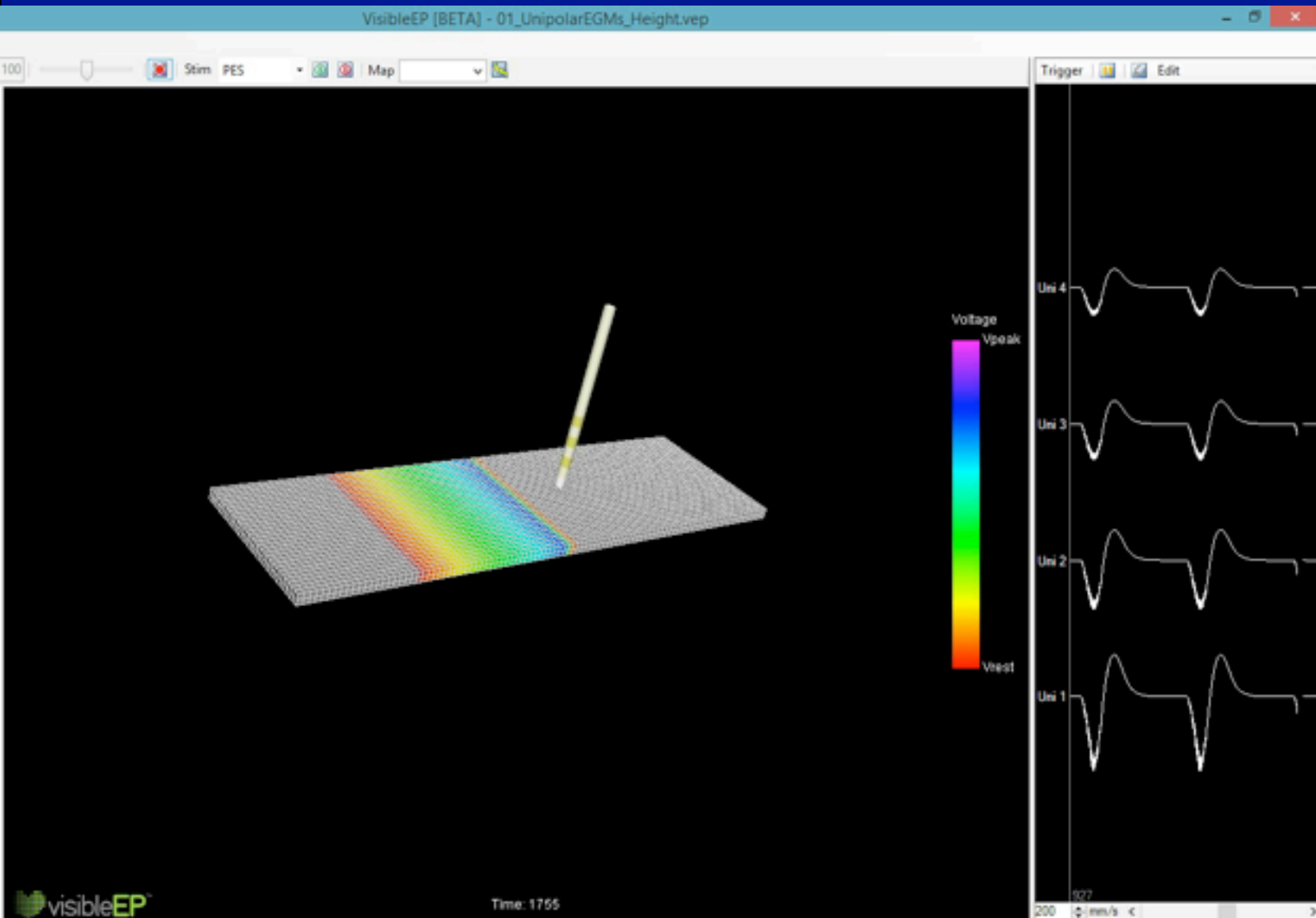
# Both bipolar and unipolar may overestimate scar

- Bipolar - epicardial fat
- Unipolar - contact etc





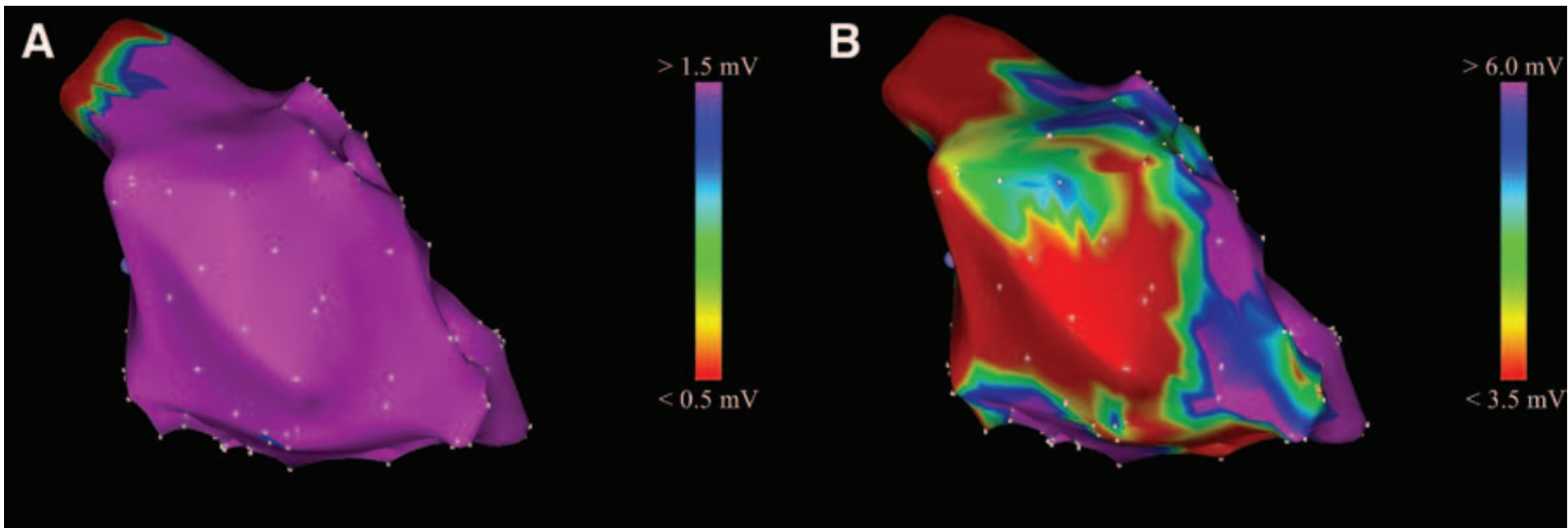
# Unipolar contact affects voltage



Egram  
“sharpness” or  
frequency  
a better  
predictor

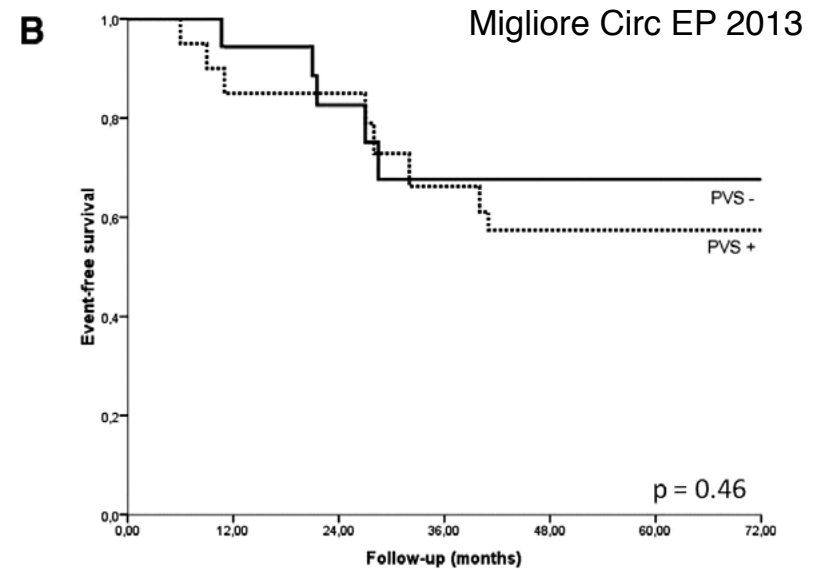
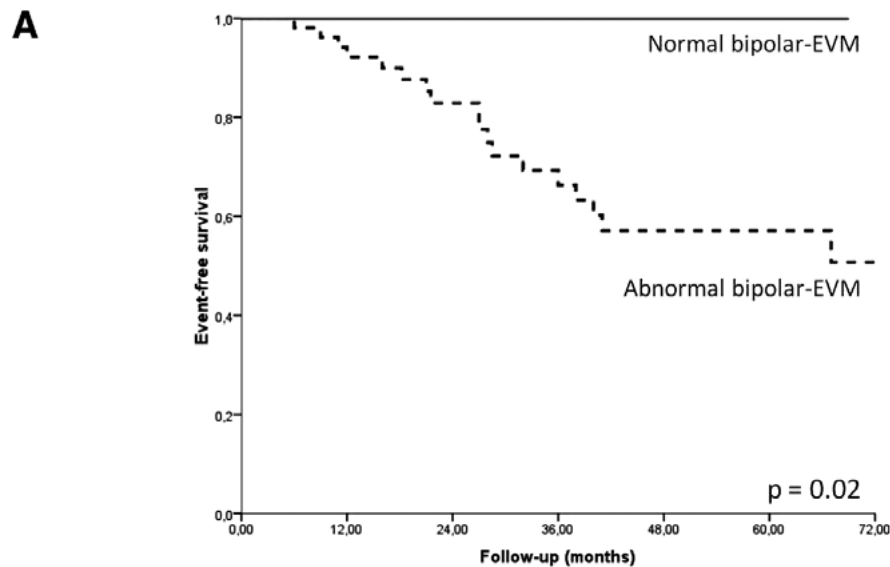
# Can scar predict risk in ARVC?

- n=69 ARVC:
  - abnormal EVM  $>1\text{cm}^2$  bipolar  $<1.5\text{mV}$  or unipole  $<6\text{mV}$



# Risk prediction with voltage mapping in ARVC

- Event rate (sudden death or VT/F)



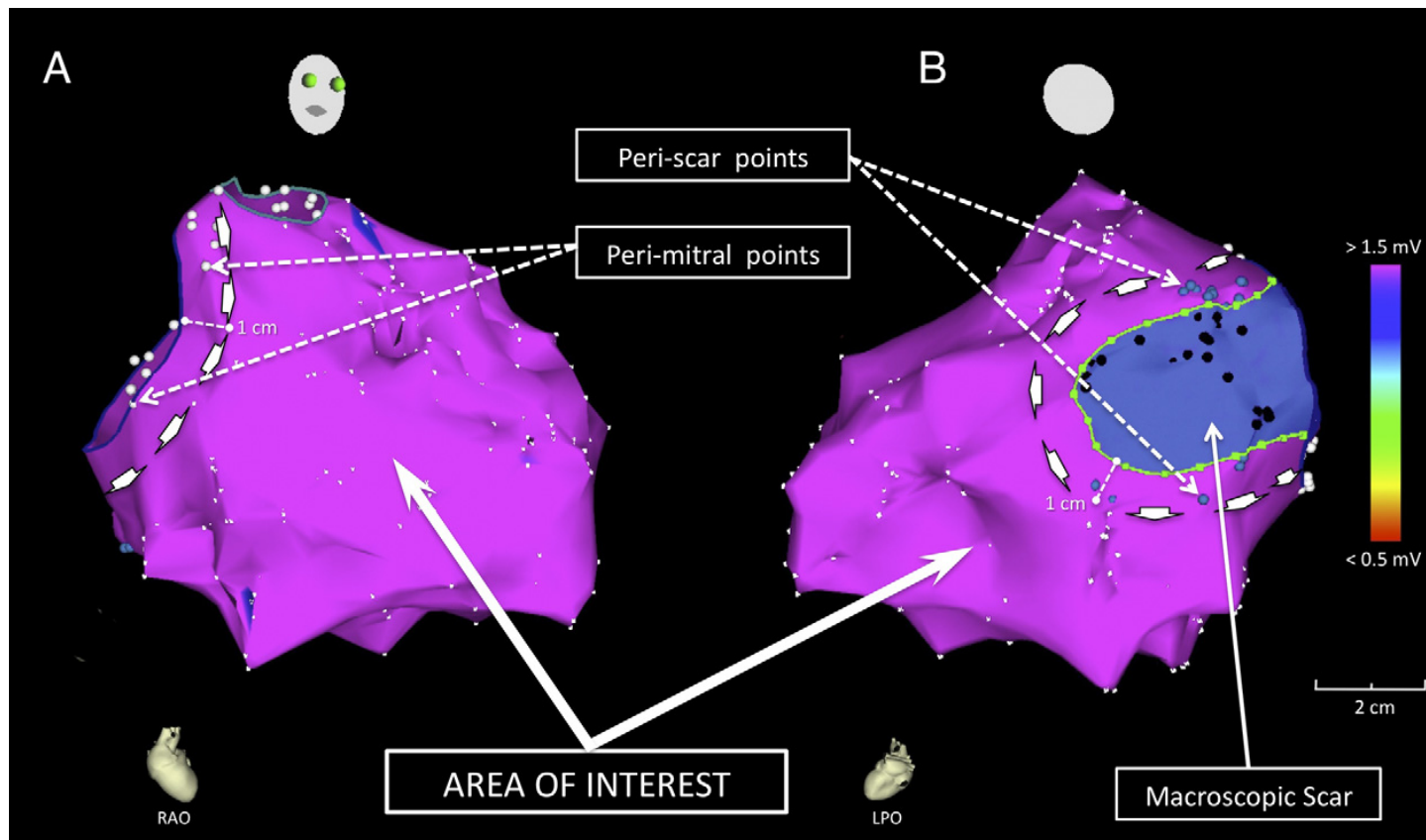
Abnormal map	HR	CI	P	HR	CI	P
Bipole	1.7	1.5-2.0	<0.001	1.6	1.2-1.9	<0.001
Unipole	1.3	0.6-2.3	0.3			

# Risk prediction with ARVC

- Unipolar mapping too sensitive
- Risk likely to be related to volume rather than transmural
- Bipole may be too crude a method for defining risk

# Unipolar mapping to identify irreversible myocardial damage

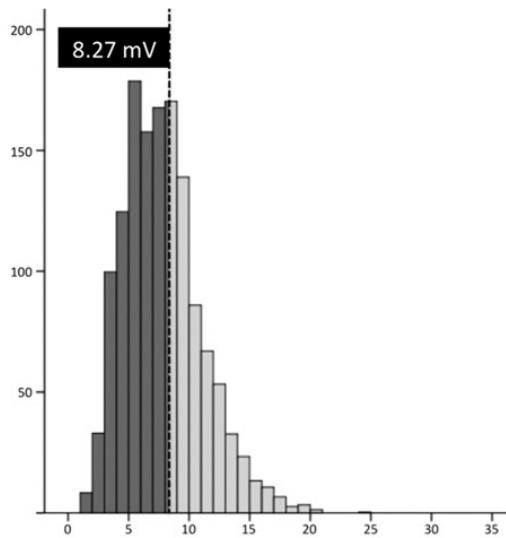
- Normal/Ectopy induced/Idiopathic cardiomyopathy
- Scar excluded (MRI, bipolar endo/epi defined)
- Unipolar voltages mapped



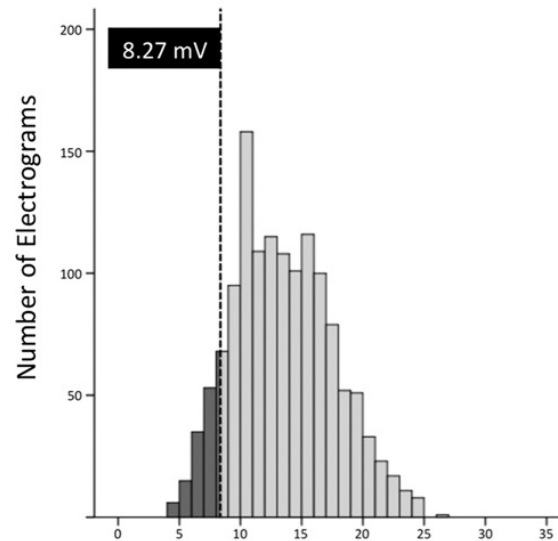
Campos et al  
JACC 2012

# Unipolar voltages of “normal” myocardium

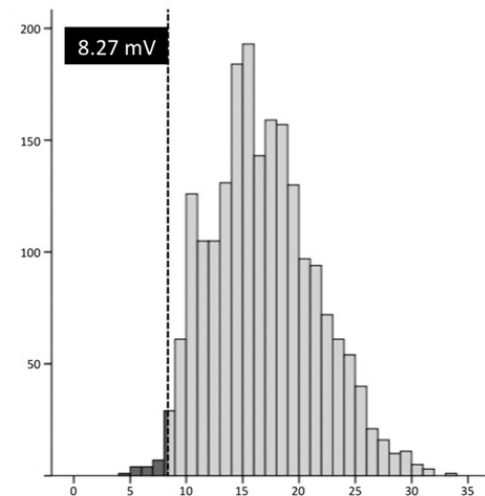
Irreversible



Reversible



Normal

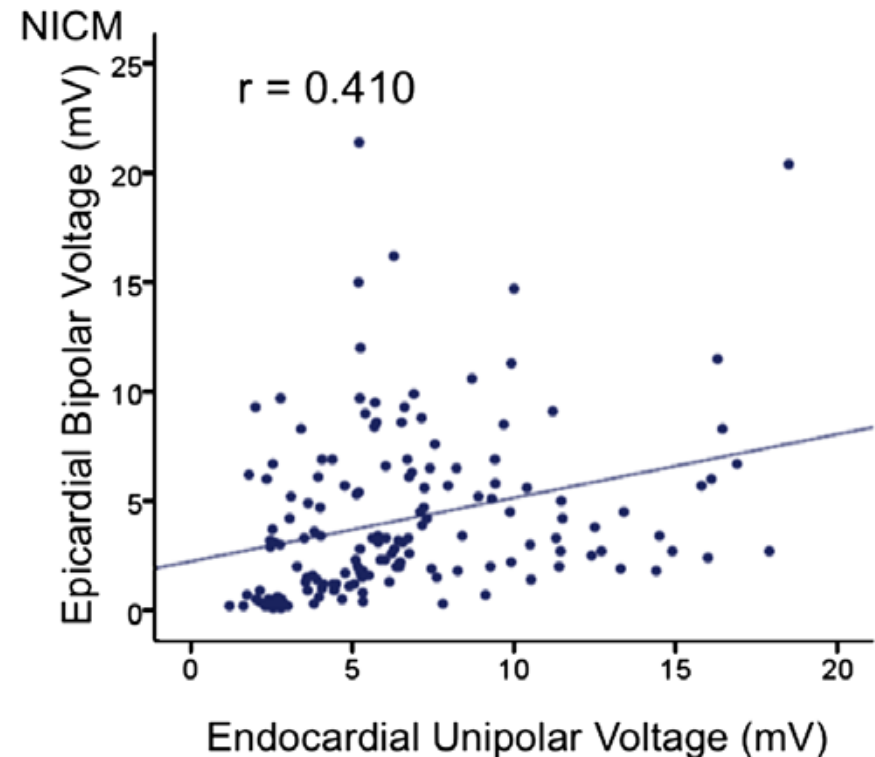
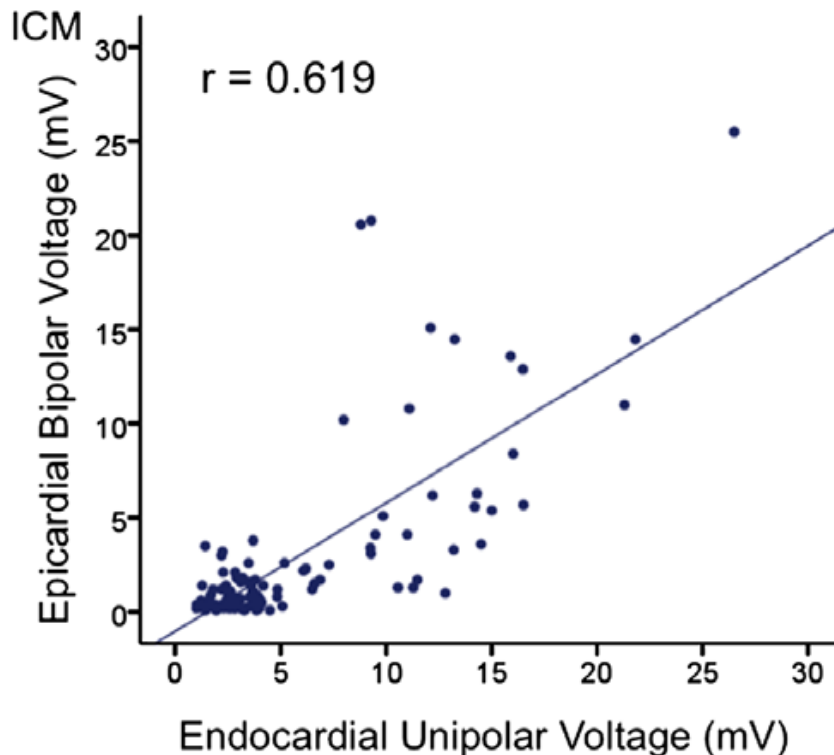


Unipolar voltage

Campos et al  
JACC 2012

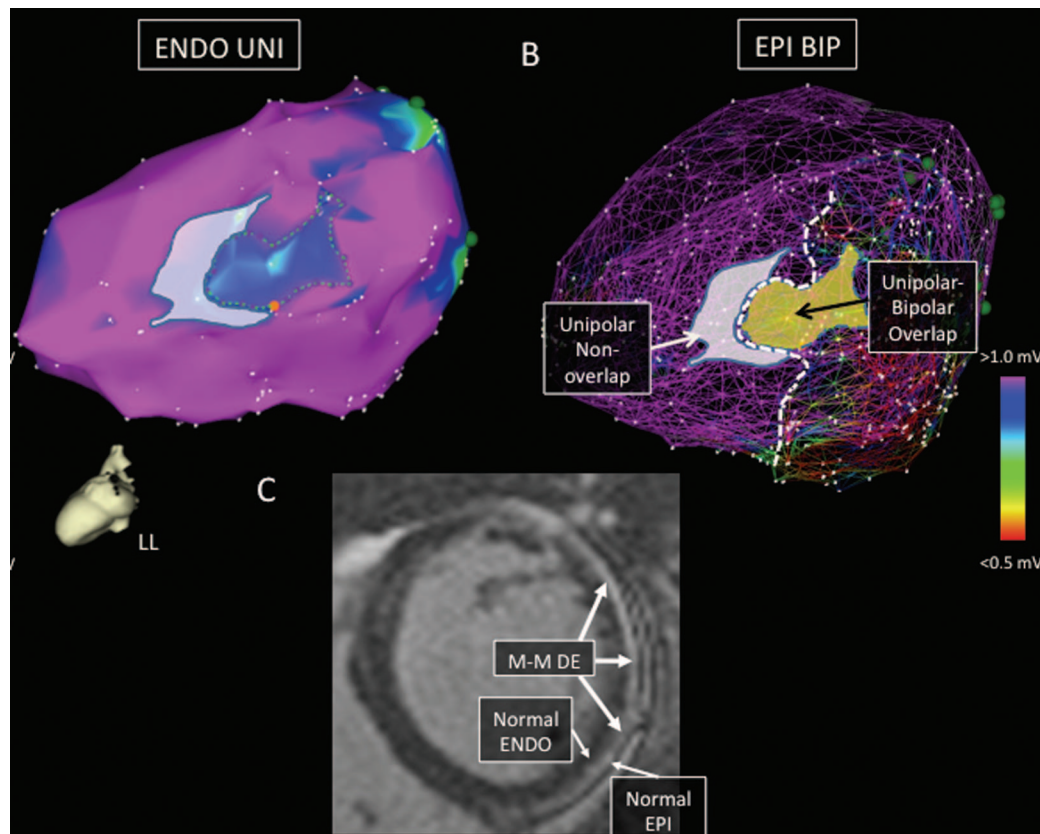
# Ischaemic vs non-ischaemic voltage mapping

- Unipolar endo correlated to epicardial scar



# Identification of mid-myocardial scar

- Abnormal endo uni with normal epi bi

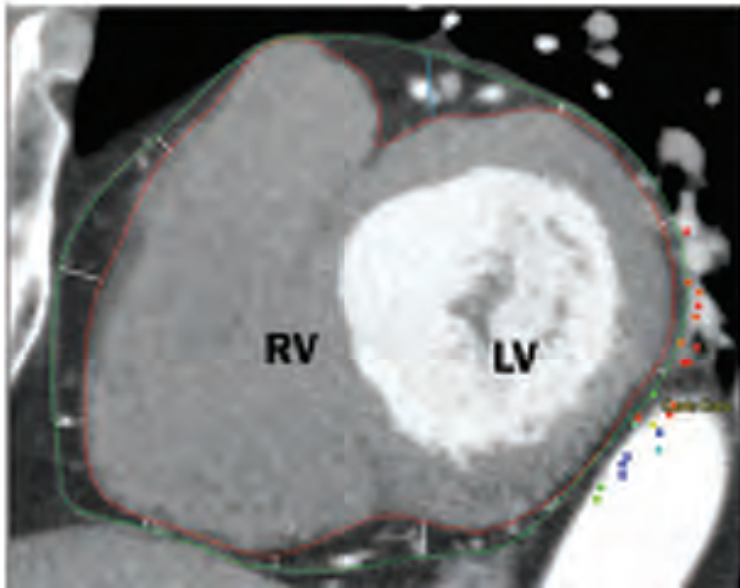




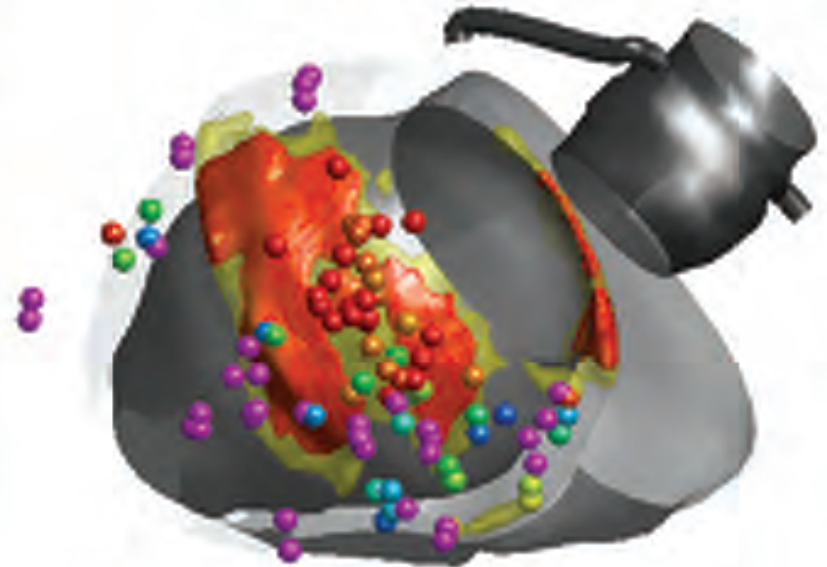
# Distinction of scar/fat/viable myocardium

- Epicardial bipolar/unipolar EAM on 3D imaging

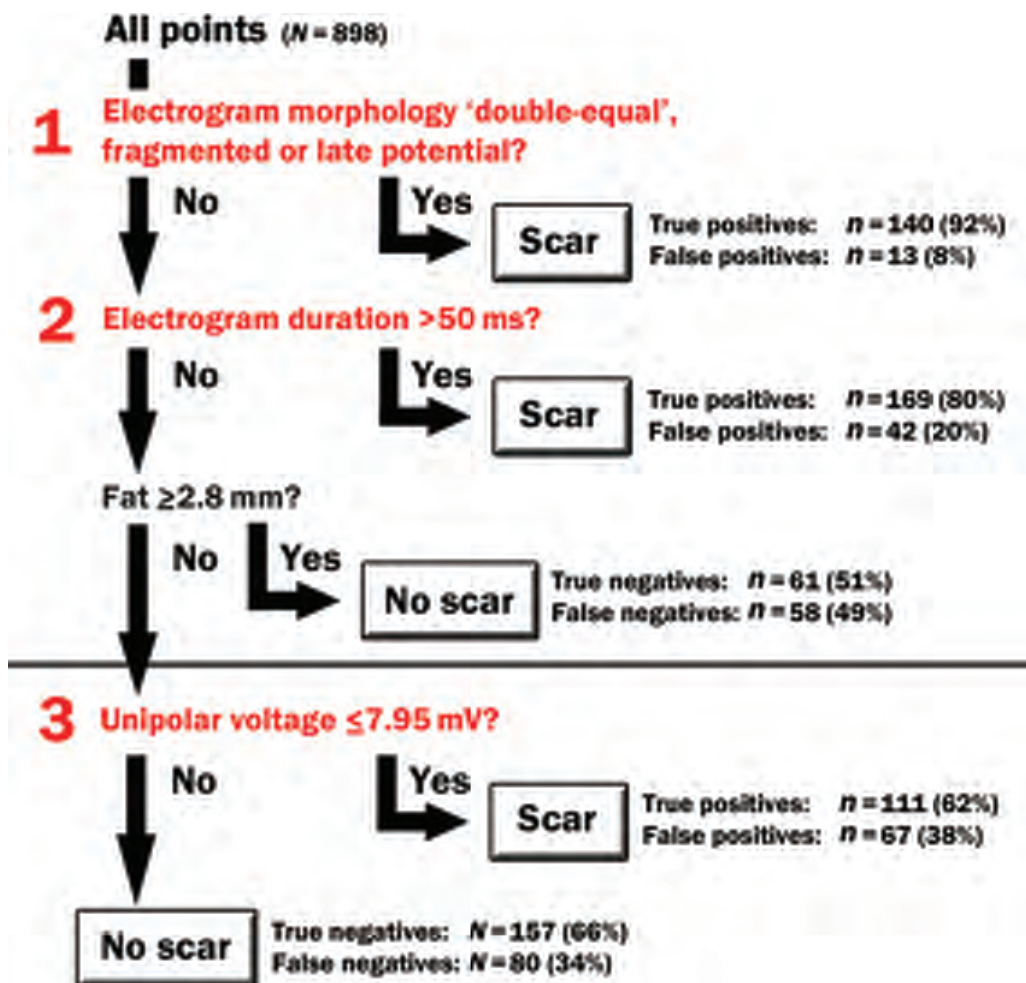
CT



CE-MRI



# Algorithm for identification



# Other considerations

- Voltage definitions vary
- Contact often difficult to achieve epicardially
- Contact detection may offer new insights

# Conclusions

- Bipolar easier to achieve clean signals and easier to measure
- Unipolar may offer insights into 3 dimensionality of tissue
- True clinical advantages yet to be proven