

Oral presentation

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## 12-lead ECG in a 1.5 Tesla MRI: Separation of real ECG and MHD voltages with adaptive filtering for gating and non-invasive cardiac output

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### Introduction

The Magneto-Hydro-Dynamic (MHD) effect arises when conductive blood flows in the MRI magnetic field ( $B_0$ ). MHD generates a voltage which distorts the real electrocardiogram ( $ECG_{real}$ ), especially during the S-T segment where flow from the left ventricle (LV) into the aorta contributes to a large MHD voltage [1]. A dominant QRS and undistorted S-T segment are important for MRI gating and physiological monitoring for ischemia during cardiac imaging/interventions [2].

### Purpose

We hypothesized that adaptive filtering could separate between MHD and  $ECG_{real}$ , and that the MHD signal could non-invasively estimate cardiac output.

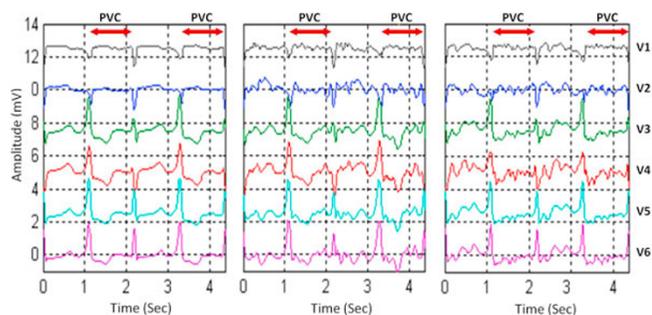
### Methods

MRI-compatible 12-lead ECGs were acquired with a modified ECG-recording system [3] from three healthy volunteers and one patient with idiopathic outflow tract Premature Ventricle Contractions (PVCs) (Ejection Fraction 20-25%, LV wall thickening, mitral regurgitation). Three sets of 20-sec breath-held ECGs (Fig. 1), were measured in a 1.5 scanner with subjects placed (i) outside the scanner with their head-in ( $ECG = ECG_{real}$ ), (ii) at iso-centre

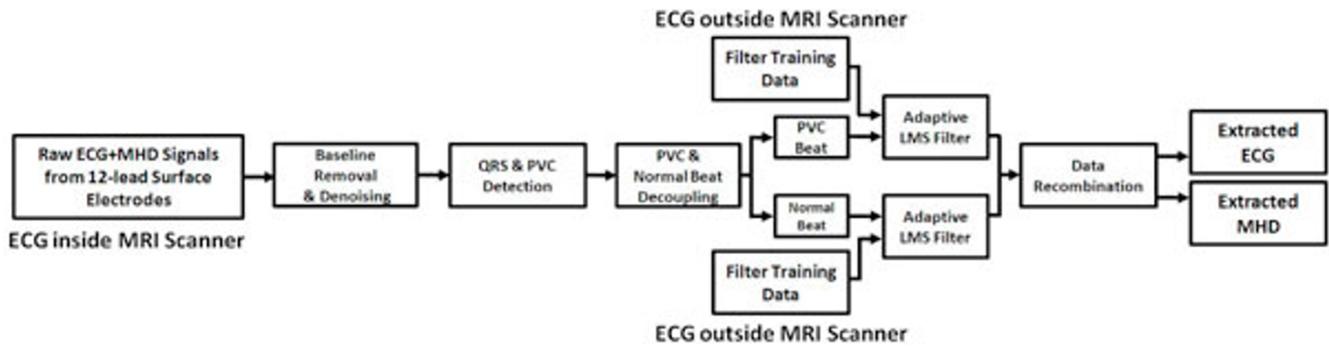
with their head-in ( $ECG = ECG_{real} + MHD_{head-in}$ ), and (iii) at iso-centre with their feet-in ( $ECG = ECG_{real} + MHD_{feet-in}$ ), which reverses  $B_0$  polarity ( $MHD_{feet-in} \sim -1 \times MHD_{head-in}$ ). Data processing (Fig. 2) involved application of an adaptive Least-Mean-Square filter to (ii) and (iii), whilst (i) was used to train the filter to decouple the MHD signal from  $ECG_{real}$ .

### Results

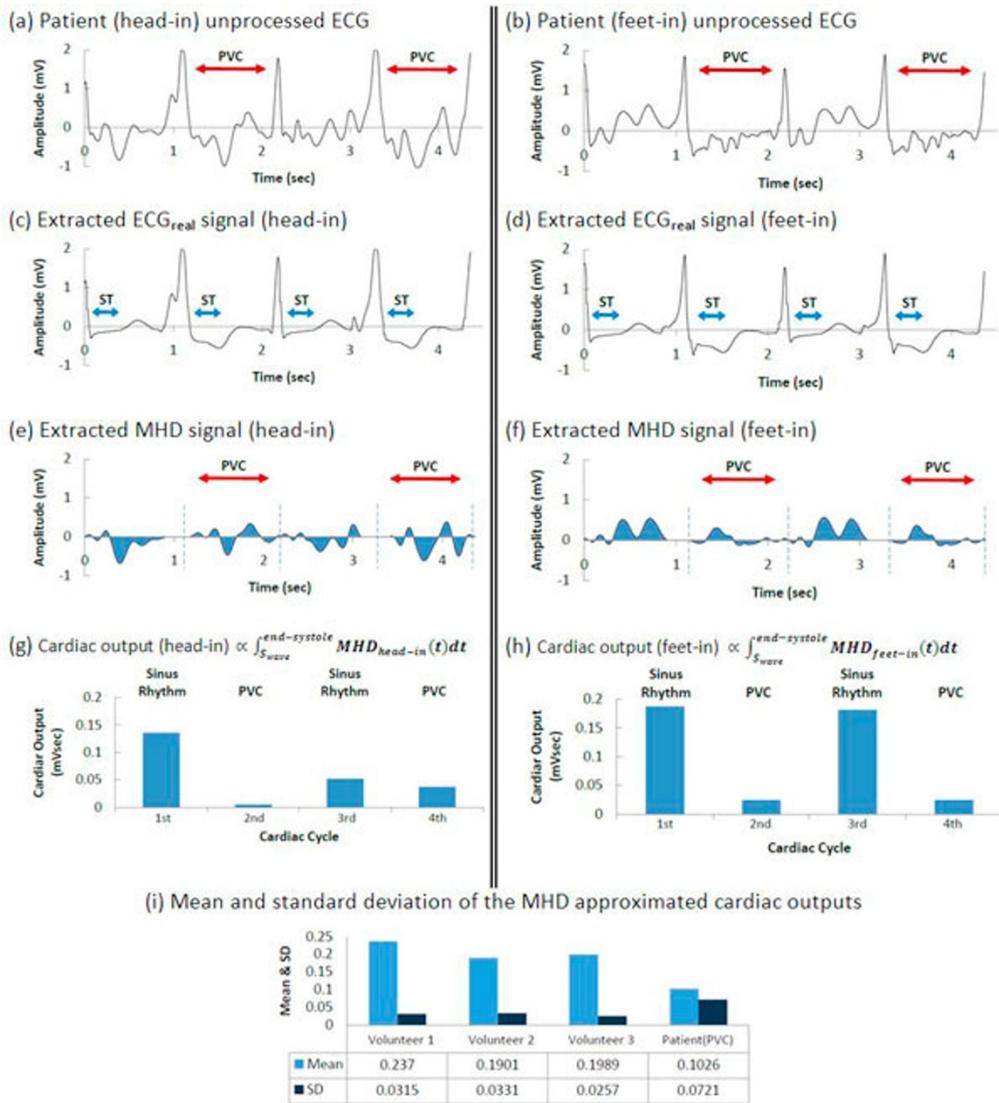
Fig. 3(a-d) show processing of the patient's V6 ECGs in positions (ii) and (iii). MHD signals are effectively



**Figure 1**  
Unprocessed PVC patient ECGs.



**Figure 2**  
Adaptive filtering procedure.



**Figure 3**  
(a-h) PVC patient ECGs at iso-centre. (i) Cardiac outputs of healthy volunteers versus the patient.

ment preserved. The MHD signals, Fig. 3(e-f), are maximal during the S-T segment. Oscillating positive and negative MHD voltages during systole in each PVC cycle can be explained by flow eddies, consistent with the patient's mitral regurgitation. Fig. 3(g-h) show the cardiac output, calculated from the systolic time-integrated MHD. Cardiac output during PVC cycles is much smaller than during normal beats. Fig. 3(i) indicates that the PVC patient's average cardiac output is 44-54% of the healthy volunteers', due to less effective PVC beats.

### Conclusion

The filtering procedure separates the ECG<sub>real</sub> and MHD signals in 12-lead ECGs acquired within the MRI. The QRS complex becomes dominant, as required for good MRI gating, while preserving S-T segment fidelity for physiological monitoring during imaging/interventions. MHD signals allow for non-invasive monitoring of beat-to-beat cardiac output.

### References

1. Gupta : *IEEE Trans BioMed Eng* 2008.
2. Haberl : *ECG pocket*, Borm Bruckmeier Publishing; 2006.
3. Dukkupati : *Circulation* 2008.

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