

# Comparison of two equivalent dipole layer based inverse electrocardiography techniques for the non-invasive estimation of His-Purkinje mediated ventricular activation

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**Background:** Inverse electrocardiography (*i*ECG) is used to non-invasively estimate the cardiac activation sequence. This becomes increasingly difficult during His-Purkinje mediated ventricular activation when the myocardium is activated through multiple activation wavefronts. Equivalent dipole layer based *i*ECG requires an initial estimate of the ventricular activation sequence, wherein cardiac physiology can be incorporated. A physiologically realistic initial estimation should be based on a combination of waveforms initiated at His-Purkinje associated endocardial locations, instead of an iterative multi-focal search. In this study, performance of multi-focal *i*ECG and multi-wave *i*ECG was compared.

**Methods:** Six subject specific CT-based geometries of ventricles, lungs and thorax were created and supplemented with electrode positions. In ventricular geometries, papillary muscles and moderator band were incorporated. For each subject, His-Purkinje mediated activation with (RBBB/LBBB) and without (normal) conduction defect were simulated. Corresponding 67-lead body surface potentials (BSP) were computed and used as input for both *i*ECG techniques. Estimated activation sequences of both *i*ECG techniques were compared to simulated activation sequences. In one subject, 67-lead BSP measurement was performed during His-Purkinje mediated activation and used as input for both *i*ECG techniques. For this subject, *i*ECG estimated maps were compared to invasive mapping.

**Results:** Figure 1 displays an example of the estimation of normal activation. Mean inter-map correlation between simulated and estimated activation sequences for multi-focal *i*ECG vs. multi-wave *i*ECG was  $59\pm 36\%$  vs.  $86\pm 7\%$  for normal activation,  $79\pm 8\%$  vs.  $72\pm 17\%$  for RBBB activation and  $41\pm 31\%$  vs.  $62\pm 25\%$  for LBBB activation. In the subject with measured BSP and invasive mapping, epicardial inter-map correlation between invasive and *i*ECG maps was  $-4\pm 8\%$  for multi-focal *i*ECG and  $65\pm 1\%$  for multi-wave *i*ECG.

**Conclusion:** This study shows improved performance of multi-wave *i*ECG for the estimation of His-Purkinje mediated activation sequences with and without conduction defect. Future research focusses on further validation of multi-wave *i*ECG using invasive mapping.

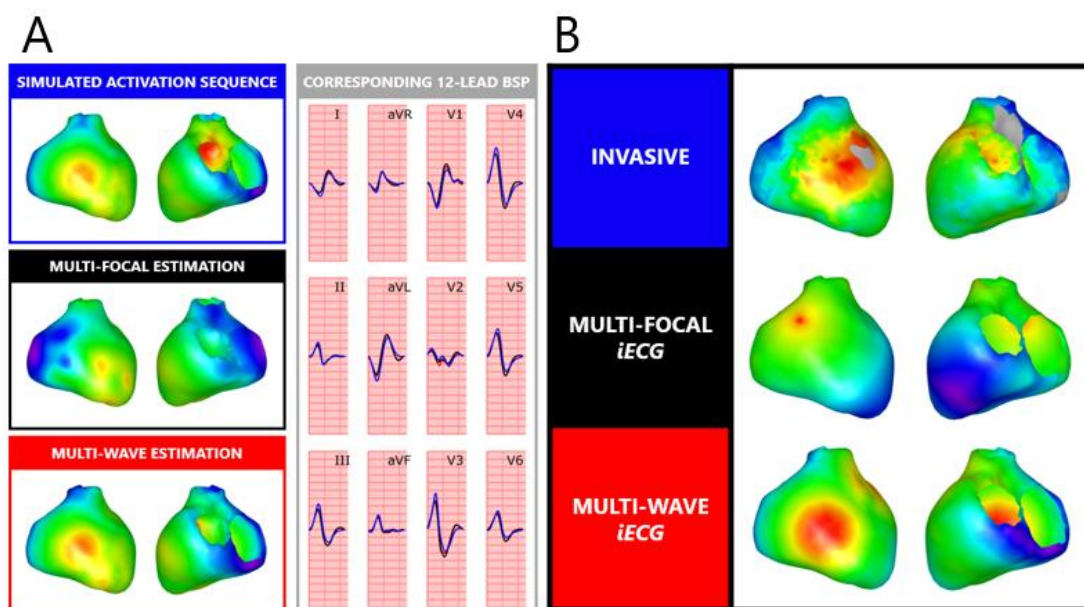


Figure 1: The estimation of the activation sequence using multi-focal *i*ECG (black) and multi-wave *i*ECG (red) of normal His-Purkinje mediated activation from early (red) to late (purple) activation. **A:** Representative results obtained from the simulation study. Activation sequences and corresponding 12-lead BSP are displayed. **B:** Epicardial invasive map, multi-focal *i*ECG and multi-wave *i*ECG estimation of one subject are displayed.