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ESTIMATION OF MONOPOLAR EMG SIGNALS FROM SPHINCTER MUSCLES

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1. INTRODUCTION



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Surface electromyogram (EMG) is usually recorded by means of spatial filters with vanishing sum of weights. More information could be extracted from monopolar signals measured with respect to a reference electrode away from the muscle. Under some assumptions, surface EMG detected along a line parallel to the fiber path has zero mean value in space at any time. This property is a constraint which can be used to estimate monopolar signals from single differential (SD) EMG signals and is satisfied in the case of a circumferential electrode array surrounded by a circular muscle.

2. METHODS

The problem of estimating monopolar signals $\vec{m}(t)$ from SD $\vec{s}(t)$ is not well posed. Indeed, there are infinite solutions, as an arbitrary function of time f(t) can be added to the monopolar signals without affecting the SD signals $s_{i}(t) = m_{i}(t) + f(t) - (m_{i+1}(t) + f(t)) = m_{i}(t) - m_{i+1}(t)$

SD signals from an array of N electrodes can be expressed in terms of monopolar signals as follows

Matrix \underline{A} cannot be inverted as it has a vanishing eigenvalue, associated to an eigenvector with constant entries.

Nevertheless, the pseudoinverse of matrix \underline{A} can be evaluated and monopolar signals can be estimated as $\vec{m}_{est}(t) = \underline{A}^{\#}\vec{s}(t)$



A) B) Simulated SD signals Simulated monopolar signals SD signals with noise (SNR 15 dB)

Under the assumption that the volume conductor is space invariant, the monopolar surface EMG detected along a curve parallel to the fiber path has zero mean value in space at any time.

B) Section of the impulse response

C) Surface potential as a sum of 3 impulse responses, corresponding to the three impulses of the tripole approximation of the source



Figure 1 A) Example of simulation of monopolar SFAP, using a model of sphincter. B) SD signals obtained from the monopolar signals. C) SD signals with 15 dB white noise. D) Estimate of monopolar signals from perturbed SD signals.

3. RESULTS





Figure 2 A) Sketchy representation of an impulse response. B) Section of the impulse response at the location of the detection system in the direction of muscle fibres and transmembrane current (source of the volume conductor problem). C) Surface potential along the direction of the muscle fibres expressed as a convolution of the source (tripole approximation) with the impulse response.

Application to simulated noise free MUAPs and interference signals



Figure 3 Performance of the method. Mean and STD of the RMS error in estimating monopolar from SD SFAP corresponding to 88 positions of simulated fibers (depth between 2 and 8 mm within the muscle with 1 mm step, distance of the fibres from the detection electrodes in the axial direction between 0 and 10 mm with 1 mm step)

Figure 4 A) Application of the method to a MUAP (MU constituted by 154 fibers) and B) to an interference signal (60 MUs, force level 40% MVC). SD signals were obtained from the simulated monopolar signals, and then monopolar signals were reconstructed.

Considerations on the estimation error



Application to signals simulated by a planar model



considering for the estimation all the 55 channels detection as in A).

Importance of covering the entire potential distribution

4. CONCLUSIONS

Under the hypothesis of space invariance of the volume conductor, monopolar signals detected along the direction of the muscle fibres with an array covering the entire spatial support of the potential distribution have vanishing spatial mean at any time. This provides a constraint for estimating monopolar from SD signals from sphincter muscles.

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Different angles between the detection array and the fibre direction

Figure 7 A) Square "noise error" for realisations of Gaussian noise (mean 0, STD 1) compared to the theoretical expected value as a function of the number of channels N. B) RMS "approximation error", "noise error" and estimation error normalised with respect to the square root of the energy of the monopolar signals of a representative SFAP (fibre located under the detection array, 1 mm deep in the muscle).