

# Numerical Identification of Motor Units using an Optimal Control Approach

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**Abstract:** A numerical approach to locate motor units in human muscles by high density surface EMG measurements is presented. For this purpose a mathematical model has been derived which can be evaluated by finite element computations. On that basis an optimal control problem is specified that can be solved by a function space oriented optimization method. Numerical results are reported for a test problem.

*Keywords:* Optimal control, Finite elements, Electromyography, Medical applications, Biomedical modelling

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## 1. INTRODUCTION

High density surface Electromyography (sEMG), is a non-invasive method of measuring the activity of muscle whereby an array of electrodes is placed above the skin and a spatially and temporally resolved measurement of the electric potential on the skin is obtained. Recent advances in high-density sEMG measurement have opened the possibility of extracting information about single motor units (groups of muscle fibers controlled by the same motor neuron) from the sEMG signal.

While significant advancements have been made in identifying the activity of individual motor units from the surface EMG signal through EMG decomposition methods (cf. e.g. Kleine et al. (2007)), a reliable and accurate method to determine where the motor units are located and where the trajectory of the muscle fibers run from the sEMG signal is not yet available. Previous works consider spatial data only [van den Doel et al. (2008, 2011); Liu et al. (2015)] or use simple parametric models within a least squares approach [Mesin (2015)].

In this work we describe an approach to automate the identification of motor units using techniques from numerical simulation and non-linear optimization.

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## 2. CONDUCTION OF ACTION POTENTIALS

The basis of our approach is a mathematical model for the physiology and for the physical situation.

### 2.1 Propagation of electric fields in the human body

Consider some part of the body, e.g. a limb or a part of the head, represented by a Lipschitz domain  $\Omega \subset \mathbb{R}^3$  with sufficient smooth boundary  $\Gamma$ . We denote the spatial variable by  $x \in \Omega$  and the temporal variable by  $t \in [0, T]$ . We are interested in the electric potential  $\Phi(x, t)$  in  $\Omega$  caused by a given distribution of electric charge  $\rho(x, t)$ .

The human body consists of various types of tissue with different physical properties. We assume that the different tissues behave like a volume conductor [Stegeman et al. (2000) and Lowery (2016)]. This means Ohm's law is applicable and thus  $\Phi(x, t)$  can spread out in the domain and can therefore be measured at the surface. Like the majority of previous sEMG simulation studies we assume that at the frequencies of interest the tissue is purely resistive and its conductivity  $\sigma(x)$  is independent of  $\Phi$  [Stegeman et al. (2000), Lowery (2016) and Gootzen et al. (1991)]. For skin and subcutaneous fat tissue we assume that  $\sigma$  is isotropic [Stegeman et al. (2000), Lowery (2016), Andreassen and Rosenfalk (1981) and Gootzen et al. (1991)]. For muscle tissue however one assumes that the conductivity is anisotropic, i.e. that the conductivity is higher along the fiber axis.

On  $\Gamma$  we impose different kinds of boundary conditions. In the part of the boundary  $\Gamma_0$  where skin is adjacent to air we note that in general the electrical potential in air is zero. Further we assume that there are no sources outside of the domain and thus we get the following Robin-boundary condition

$$\partial_\nu \Phi(s, t) = -\mu \Phi(s, t) \quad \text{at } \Gamma_0 \quad (1)$$

with skin conductivity  $\mu$ . In addition, since we only model a part of the body, there are artificial boundaries  $\Gamma \setminus \Gamma_0$  that result from removing the rest of the body. We assume that the potential is zero for all points which are far away from the inner source and thus we get homogeneous Neumann boundary conditions.

As usual we can write now the electric potential equation and the boundary conditions in the weak form with a

solution  $\Phi(\cdot, t) \in H^1(\Omega)$

$$\int_{\Omega} \rho(x, t) v(x) dx = \int_{\Omega} \sigma(x) \nabla \Phi(x, t) \nabla v(x) dx + \int_{\Gamma_0} \mu \Phi(s, t) v(s) ds \quad \forall v \in H^1(\Omega) \quad (2)$$

We remark that the left hand side is well defined, for  $\rho(\cdot, t) \in L_2(\Omega)$ . However, this equation can still be formulated rigorously for concentrated charges, i.e., if  $\rho(\cdot, t)$  is a measure on  $\Omega$ , if  $v$  is chosen in  $W^{1,p}(\Omega)$  with  $p > 3$ . Then it is known (cf. e.g. Haller-Dintelmann et al. (2009); Alibert and Raymond (1997)) that  $\Phi(\cdot, t) \in W^{1,p'}(\Omega)$  for  $1/p + 1/p' = 1$ .

In our model the charge distribution  $\rho(\cdot, t)$  in (2) is caused by ionic activity, the so called action potential that propagates along the muscle fiber of a motor unit and activates the contraction of the muscle. Its definition will be explained in the following two subsections.

## 2.2 Motor units

A motor unit is a bundle of muscle fibers which are innervated by the same motor neuron. The motor unit is the smallest controllable unit of muscle. If now a muscle fiber is activated, two action potentials propagate in opposite directions from the neuro-muscular junction to the ends of the muscle fiber. The velocity  $v$  with which the action potential propagates is almost constant. In general the neuro-muscular junction lies approximately in the middle of the fiber. In Figure 1 one can see a schematic view of such a motor unit. As all of the fibers within a

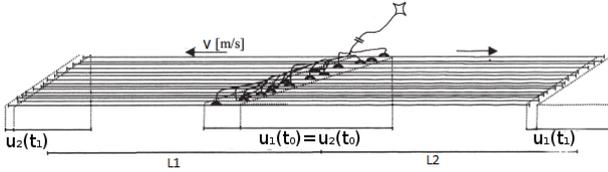


Fig. 1. Sketch of a motor unit [Stegeman et al. (2000)]

motor unit are activated simultaneously, we can treat a bundle of muscle fibers in a motor unit as a single fiber. For such a fiber we assume that the radius (a few  $\mu m$ ) is much smaller than its length (several  $cm$ ) [Andreassen and Rosenfalck (1981) and Gootzen et al. (1991)] and thus we can represent the trajectory of the moving action potential along the fibers by a pair of regular curves

$$u_1, u_2 : [t_0, t_1] \mapsto \Omega_m \\ u_1, u_2 \in V := H^1([t_0, t_1])^3.$$

The point  $u_1(t_0) = u_2(t_0)$  represents the neuro-muscular junction and  $u_1(t_1)$  and  $u_2(t_1)$  represent the fiber ends. With  $\dot{u}_k(\tau) := \frac{d}{d\tau} u_k(\tau)$  we denote the tangent vector of the curves and with  $s_k(\tau) := |\dot{u}_k(\tau)|$  the speed of the curves.

## 2.3 Propagating action potentials

Before we define the action potential and thus also the source density we make some assumption. First we assume

that only one motor unit is active and thus the support of the source density  $\rho$  is included in the curves, i.e.

$$\rho(x, t) = 0 \quad \forall t \in [0, T], x \notin u_k([t_0, t_1]), k = 1, 2. \quad (3)$$

Therefore  $\rho$  is a measure with support on  $u_k$ . Furthermore we assume that the source density fulfills the condition

$$\int_{\Omega} \rho(x, \cdot) dz = 0 \quad (4)$$

which expresses the conservation of charge in the human body.

We now consider that the source density is a spatially distributed signal, the so called action potential, which propagates along a trajectory of a motor unit. Similar to Rosenfalck (1969) we define the action potential in terms of an artificial real variable  $z$  as follows:

$$i_m(z - z_0(t)) := \begin{cases} \sigma_{in} \pi r^2 \frac{d^2 V_m(z - z_0(t))}{dz} & \text{if } z \leq z_0 \\ 0 & \text{if } z > z_0 \end{cases} \quad (5)$$

where  $\frac{d^2 V_m(z)}{dz} = -96 \exp(z)(6z + 6z^2 + z^3)$  is the second spatial derivative of the transmembran potential,  $\sigma_{in}$  is the intracellular conductivity and  $r$  is the radius of the motor unit. Here we changed the orientation of the action potential, by replacing  $z$  through  $-z$  in  $V_m$ , and added the origin of the signal  $z_0(t) := v(t - t_0)$ . With this modification we generate a signal which propagates in time from left to right along the artificial axis as the time  $t$  increases. This moving action potential can be seen in Figure 2 for three different times.

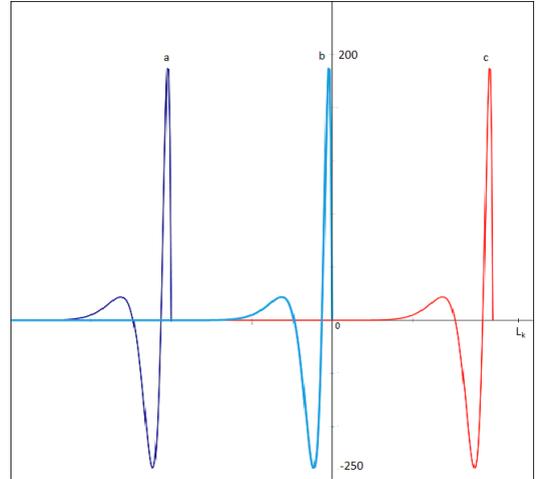


Fig. 2. Moving action potential at different times  $t$

For fixed  $t$  we can now uniquely identify each point on the trajectory  $u_k$  with a point  $\tilde{z} \in [0, L_k]$  by the arc length of the corresponding curve segment. Here  $L_k$  is the length of the trajectory  $u_k$ . Thus we can define the diffeomorphism

$$z(u_k(\tau)) := \int_{t_0}^{\tau} |\dot{u}_k(\xi)| d\xi. \quad (6)$$

If we combine now (3), (5) and (6) we get for the source density

$$\rho(x, t) = \begin{cases} i_m(z_k(\tau) - z_0(t)) & \text{if } x = u_k(\tau) \\ 0 & \text{else} \end{cases}. \quad (7)$$

One can now easily proof that the source density fulfills the condition (4).

*End-effects.* By the simulation of sEMG it is well known that if the fiber length is finite so called end-effects can appear [Gootzen et al. (1991)] and thus we have to correct our model of the source density. Before we can correct the model of the source density we have to explain when and why those end-effects appear. To this end we simulated an sEMG measurement with our current model. In Figure 3 one can see the result of a simulation with the current model (red graph). Here we simulated a straight fiber and the electrode was positioned in the middle between the neuro-muscular junction and one end of the fiber. Comparing the simulation with real measurements one can easily see that there appear some unphysical peaks at the left and right end of the measurement. We have noted before in the modeling that the integral over the full support of the action potential is zero (compare (4)), but since the signal is represented through a moving stationary source it can happen that not the full support of the signal lays on the curve which leads to an imbalance of charges in the tissue. There are two possible situations when such an imbalance can appear, namely when the signal arrives at the neuro-muscular junction ( $t \in I_0$ ) and when the signal vanishes at the fiber end ( $t \in I_1$ ). One possibility for correcting this imbalance is given in [Gootzen et al. (1991)] by adding a term  $g$  that represents stationary sources at the fiber-ends. These term has then to be chosen such that

$$\int_{u_k} \rho(x, t) dx + g(u_k(t_0), t) + g(u_k(t_1), t) dx \stackrel{!}{=} 0.$$

Furthermore we assume that

$$\begin{aligned} g(u_k(t_0), t) &= 0 & \text{if } t \in I_1 \\ g(u_k(t_1), t) &= 0 & \text{if } t \in I_0 \end{aligned}$$

holds, which means that only one correction term is not equal to zero at the same time. This assumption is only sufficient if the lengths of the curve is longer than the support of the action potential. In our model problems we will assume that this is always true. With this assumption we get for the correction terms

$$g(u_k(t_0), t) = \begin{cases} -\int_0^1 |\dot{u}_k(\tau)| i_m(z(u_k(\tau)) - z_0(t)) d\tau & \text{if } t \in I_0 \\ 0 & \text{else} \end{cases}$$

$$g(u_k(t_1), t) = \begin{cases} -\int_0^1 |\dot{u}_k(\tau)| i_m(z(u_k(\tau)) - z_0(t)) d\tau & \text{if } t \in I_1 \\ 0 & \text{else} \end{cases}$$

If we now add these correction terms to the source density and simulate again the measurement, one can see that the end effects has almost vanished (see green graph in Figure 3).

### 3. ADJOINT APPROACH

In the previous section we have described a mapping  $(u_1, u_2) \rightarrow \Phi$ , where  $\Phi$  is defined on the space-time cylinder  $\Omega \times [0, T]$ . Although our problem is merely quasi-static, the evaluation of this mapping for reasonably high temporal and spatial resolution is computationally expensive,

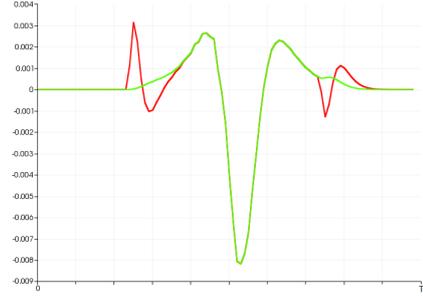


Fig. 3. Comparing the corrected simulated measurement (green) with the uncorrected one (red)

in particular if this evaluation has to take place multiple times within an optimization algorithm.

However, by sEMG measurements, only part of the information that is present in  $\Phi$  is actually used. Measurements are taken only at finitely many electrodes  $D_i \subset \Gamma_0$  on the skin, given by

$$\tilde{y}_i(t) = \int_{D_i} \Phi(s, t) ds \quad \text{for } t \in [0, T].$$

For fixed  $t$  each measurement is thus a linear functional  $l_i : H^1(\Omega) \rightarrow \mathbb{R}$  with argument  $\Phi(\cdot, t)$ .

To reduce the numerical effort of evaluating  $\tilde{y}_i$  we now introduce an alternative way to compute the measurements. The scalar quantity

$$\tilde{y}_i(t) = l_i(\Phi(\cdot, t)) \quad \text{for } t \in [0, T]$$

can be evaluated efficiently by the following formula

$$\tilde{y}_i(t) = \int_{\Omega} w(x) \rho(x, t) dx \quad (8)$$

where  $w$  is the solution of the adjoint problem

$$\int_{\Omega} \sigma(x) \nabla w(x) \nabla v(x) dx + \int_{\Gamma_0} \mu w(s) v(s) - \mathbb{1}_D v(s) ds = 0. \quad (9)$$

This formula follows from the following simple abstract computation: Let  $\Phi \in V$  satisfy:

$$a(\Phi, v) = r(v) \quad \forall v \in W,$$

where  $a : V \times W \rightarrow \mathbb{R}$  is bilinear and  $r \in W^*$  is linear. Let further  $w$  satisfy:

$$a(\phi, w) = l(\phi) \quad \forall \phi \in V,$$

then

$$l(\Phi) = a(\Phi, w) = r(w).$$

### 4. OPTIMAL CONTROL PROBLEM

Now we assume that we have a measurement array with  $J$  electrodes. For each of these electrodes we can compute a weight function  $w_j$  by solving the adjoint problem (9). If we now define the vector  $w = (w_0, \dots, w_J)$  we can, by using the model (8), compute the vector valued potential with the vector valued integral

$$y(t, u) = \sum_{k=1}^2 \left[ \int_0^1 w(u_k(\tau)) |\dot{u}_k(\tau)| i_m(z_k(\Theta_k(\tau), t)) d\tau + w(u_k(0))g(u_k(0), t) + w(u_k(1))g(u_k(1), t) \right] \quad (10)$$

such that the potential at the electrode  $j$  is the  $j$ -th component of  $y$ . Furthermore with  $y_m(t)$  some measured potential at the electrodes is given. For our optimal control problem we then want to minimize the  $L^2$ -norm of the distance between measurement and simulation. Furthermore we add a penalty term which shall ensure that the speed of the curve is nearly constant and equal in magnitude to a given reference velocity  $v_r$ . We get then the following optimization problem

$$\min J(u) = \|y(u, t) - y_m(t)\|_{L^2([0, T])}^2 + \frac{\alpha}{2} c(u) \quad (11)$$

with

$$c(u) = \int_{t_0}^{t_1} (|\dot{u}(\tau)| - v_r)^2 d\tau$$

## 5. NUMERICAL IMPLEMENTATION

In this section we want to take a closer look on how to solve the above stated optimization problem. Therefore we first specify the geometric setting. As domain  $\Omega$  we choose a cuboid with size  $1\text{cm} \times 10\text{cm} \times 1\text{cm}$ . This cuboid shall represent an idealized piece of some limb. We divide this cuboid into two horizontal layers where the lower layer has a thickness of  $8\text{mm}$  and represent the muscle tissue. The second layer is  $2\text{mm}$  thick and represent a fat layer under the skin. Furthermore we define the upper boundary as  $\Gamma_0$  where the domain is bounded by skin. At the other boundaries we assume that the domain would continue. One can see a schematic view of this cuboid in Figure 4.

Furthermore on the time interval  $[0\text{s}, 0.015\text{s}]$  gener-

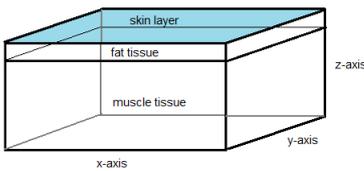


Fig. 4. Schematic view of the domain

ate artificial measurements by forward simulation. The action potential is active in the subinterval  $[t_0, t_1] = [0.0025\text{s}, 0.0125\text{s}]$ , which is the domain of the trajectories. Finite element discretization of the spaces  $H^1(\Omega)$  and  $H^1([t_0, t_1])^3$  is performed as follows. We choose a triangulation  $\mathfrak{T}$  of  $\Omega$  and a triangulation  $\mathfrak{I}$  of the interval  $[t_0, t_1]$ . The domain  $\Omega$  is thereby divided into tetrahedra, whereas the  $I$  is simply divided into  $n$  intervals. Using this triangulation we can then define the space of linear ansatz functions for  $H^1([t_0, t_1])^3$  and the space of quadratic ansatz functions for  $H^1(\Omega)$  by

$$V_n := \{v : [t_0, t_1] \mapsto \mathbb{R}^3 \mid v|_K \in P_1(K) \forall K \in \mathfrak{I}\} \subset V$$

$$W_n := \{w : \Omega \mapsto \mathbb{R} \mid w|_K \in P_2(K) \forall K \in \mathfrak{T}\} \subset H^1(\Omega)$$

The corresponding finite element method is implemented with the help of the finite element Toolbox Kaskade7 [Götschel et al. (2012)].

### 5.1 Solving the adjoint problem

For the optimization it is essential to compute the weight functions  $w$  by solving the adjoint problem and be able to evaluate them at each point in the domain  $\Omega$ . Since we have to compute, store and evaluate the weight functions efficiently, we decided to use a hierarchically and adaptively built triangulation of  $\Omega$ . Therefore we first generate a coarse grid and refine it first globally to a certain mesh size. After that we refine then the area where the electrodes are placed and the the area where the optimal solution is expected to be. One possible triangulation can be seen in Figure 5. We then uses a Galerkin-Method to solve

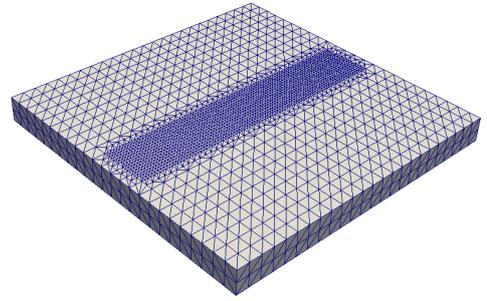


Fig. 5. Mesh with about 750000 tetrahedra. In the critical region the mesh has been refined, a-priori.

the adjoint problem (8), which as usual this leads to the discrete replacement problem

$$\text{find } w_n \in W_n \text{ s.t.} \\ a(w_n, \eta) = r(\eta) \quad \forall \eta \in W_n \quad (12)$$

With the usual techniques one can show that the bilinear  $a(\cdot, \cdot)$  is V-elliptic and since  $r$  is a bounded linear functional we know from the Lemma of Lax-Milgram that the problem (12) has then a unique solution.

After finite element discretization we end up with a large sparse linear system of equation that is solved by the preconditioned conjugate gradient method. We use a BPX-preconditioner [Bramble et al. (1990)], which takes advantage of our hierarchic grid.

### 5.2 Evaluation of weight functions

As we have seen in (10) and (11), the objective function of our optimal control problem depends on line integrals that involve the weight functions  $w_j$ , evaluated along given trajectories. For optimization purposes we also have to evaluate spatial derivatives of  $w$ , i.e.,  $w_x$  and  $w_{xx}$  to compute the derivative and the hessian of the objective, since a small perturbation of  $u_k$  leads to a perturbation of the points, where  $w$  is evaluated.

Since  $w$  is only available as a finite element function we cannot expect  $w_x$  to be continuous. Even more,  $w_{xx}$  is only defined in the interior of the tetrahedra, and may not properly reflect the global curvature of the solution. For

example  $w_{xx} = 0$  for linear finite elements. Nevertheless, there are theoretical results available (cf. Ovall (2007)) that indicate that second derivatives of finite element functions of order higher than one asymptotically approximate second derivatives of regular solutions of elliptic PDEs. In our numerical experiments, see Figure 8 below. The convergence behaviour of our optimization algorithm depends on the resolution of the finite element discretization. We observe fast linear convergence, the finer the grid, the faster the rate. The deeper understanding of this interesting phenomenon is subject to current investigations.

The evaluation of the line integrals is performed by numerical quadrature along  $u_k$ . The necessary evaluation of the finite element function  $w$  at a quadrature point  $x$  requires a search for the tetrahedron where  $x$  is located. To do this efficiently we exploit that the quadrature points are ordered along the trajectory and thus use a neighborhood search. If this fails, we fall back to a hierarchic search over the whole grid.

### 5.3 Numerical solution of the optimization problem

To solve the minimization problem we use a simple SQP line search method. This means in each step we compute a direction of descent  $\delta u$  and a sufficient step size  $\beta$ . For the step computation we establish a quadratic model of the functional by

$$J(u + \delta u) \approx m_u(\delta u) := J(u) + J'(u)\delta u + \frac{1}{2}q_u(\delta u, \delta u).$$

The bilinear form

$$q_u(\delta u, \delta u) = J''(u)(\delta u, \delta u) + \gamma \|\delta u\|_{V \times V}^2.$$

employs second order information of  $J$ . The term  $\gamma \|\delta u\|_{V \times V}^2$  is added to overcome possible indefiniteness of  $J''(u)$ . So  $\gamma$  is chosen adaptively to make  $q_u$  positive definite, if  $J''(u)$  isn't. We thus have a modified Hessian-method which differs from classical modified Hessian methods by the choice of regularization term.

To get the direction of descent from this model one has then to minimize  $m_u(\delta u)$  over  $\delta t$  which is equivalent to solving the variational problem

$$\begin{aligned} &\text{find } \delta u \in V_n \times V_n \text{ s.t.} \\ &q(\delta u, \eta) = J'(u)\eta \quad \forall \eta \in V_n \times V_n \end{aligned} \quad (13)$$

Since  $u$  is defined on a one dimensional domain this problem is of moderate size after discretization and can thus be solved by a Cholesky factorization.

For the so computed direction it remains to compute a sufficient step-size. We do this by using a simple backtracking algorithm with Armijo acceptance criterion as one can find it e.g. in [Nocedal and Wright (2006)]. Finally we get the following simple optimization algorithm 1:

*Algorithm 1.* (line search).

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choose  $u_0$ 
while  $J'(u)\delta u > \epsilon$  do
    solve  $q(\delta u, \eta) = -J'(u)\eta$ 
    compute step length  $\beta$ 
     $u_{k+1} = u_k + \beta\delta u$ 
end while

```

## 6. NUMERICAL EXAMPLE

To test the above described algorithm we first simulate a measurement for a reference trajectory  $\tilde{u}$ . Therefore we choose a measurement array of 63 electrodes, which are placed in three rows of 21 electrodes above the reference trajectory. The electrodes have the shape of circles with diameters of 2 mm and the distance between the centers of two neighbouring electrodes is 4 mm. In Figure 6 we illustrate the setting, by plotting the the position of the electrodes at the skin (black circles) and the reference trajectory (green). We divide then our time interval  $[0.0025s, 0.0175s]$  into 150 time steps and compute for each electrode the potential  $y_i(t_k)$  at each time step  $t_k$ . From this measured potential one can then make an initial guess for the starting trajectory by placing it in the regions where the highest potential is measured. This is also a good option in practical applications. In Figure 6 one can also see our choice for the starting trajectory (red).

From this measurement we identify the reference trajectory by our optimization algorithm. For this example we assume that we know the interval  $[t_0, t_1]$ , the velocity  $v$  of the signal, which is  $4 \frac{m}{s}$ , and the position of the neuromuscular junction  $u_1(t_0) = u_2(t_0)$ . The position of the fiber ends  $u_1(t_1)$  and  $u_2(t_2)$  and the depth of the trajectory are unknown and shall be identified during the optimization.

We stop the algorithm when the energy norm of the gradient is sufficiently small, i.e.  $J'(u)\delta u \leq 10^{-9}$ . In Figure 6 one can see the computed solution (dashed blue) compared to the reference trajectory (green). One can see that the reference trajectory is identified very well by the solution of our optimization problem, the two graphs coincide.

To assess the influence of the discretization of the weight functions we performed the optimization with two different grids for the computation of  $w$ . In the first run we used approximately 750000 tetrahedra. In a second run we reduced the mesh of 142000 tetrahedra. It can be observed that both solutions are quite similar in accuracy, but the rate of convergence differs significantly. For the coarse solution, about 14 iterations are needed, while the fine solution requires only 8 steps. We attribute this behaviour to the fact that more accurate second order information is available for the fine solution. To illustrate this we compare in Figure 8 the energy norm of the gradient for the different mesh sizes.

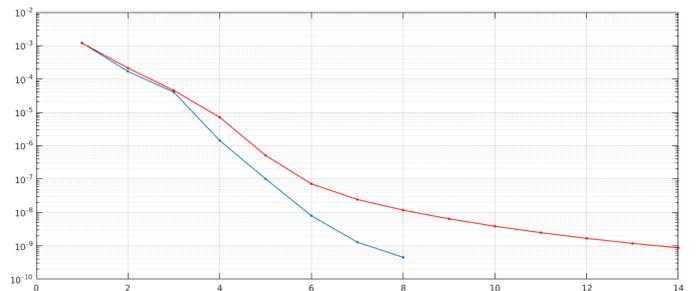


Fig. 8. Comparison of the energy norm of the gradient for 750000 tetrahedra (blue) and 142000 tetrahedra (red)

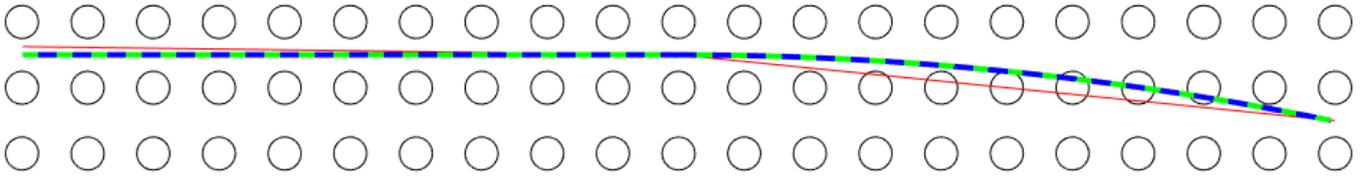


Fig. 6. Position of the electrodes and comparison of the reference trajectory (green), the start trajectory (red) and the computed solution (dashed blue) in the  $xy$ -plane.

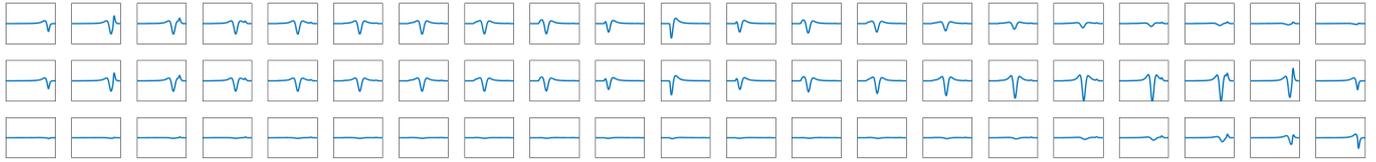


Fig. 7. Simulated measurement data of the array of electrodes. In each square the temporal signal of the corresponding electrode is displayed.

## 7. CONCLUSION

We have constructed a numerical algorithm that can take into account the full spatio-temporal information, gained by high-density sEMG measurements in order to locate motor units in human muscles. It is based on an accurate finite element model of the physiological situation. An adoint approach makes the problem tractable numerically. For a test problem our optimization method converges in a few iterations and yields accurate results.

In future research our method has to be applied to real measurement data to assess the accuracy of our forward model and the influence of modelling errors and noisy data on the identified solution. From a numerical point of view, adaptive solution techniques for the computation of the weight functions as well as for the trajectories will be explored. Finally, a deeper understanding of the accuracy of the spatial derivatives of  $w$  is desirable, giving rise to further theoretical investigations.

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