Separation of indocyanine green boluses in the human brain and scalp based on time-resolved *in-vivo* fluorescence measurements

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Abstract. Non-invasive detection of fluorescence from the optical tracer indocyanine green is feasible in the adult human brain when employing a time-domain technique with picosecond resolution. A fluorescence-based assessment may offer higher signal-to-noise ratio when compared to bolus tracking relying on changes in time-resolved diffuse reflectance. The essential challenge is to discriminate the fluorescence originating from the brain from contamination by extracerebral fluorescence and hence to reconstruct the bolus kinetics; however, a method to reliably perform the necessary separation is missing. We present a novel approach for the decomposition of the fluorescence contributions from the two tissue compartments. The corresponding sensitivity functions pertaining to the brain and to the extracerebral compartment are directly derived from the in-vivo measurement. This is achieved by assuming that during the initial and the late phase of bolus transit the fluorescence signal originates largely from one of the compartments. Solving the system of linear equations allows one to approximate time courses of a bolus for each compartment. We applied this method to repetitive measurements on two healthy subjects with an overall 34 boluses. A reconstruction of the bolus kinetics was possible in 62% of all cases.

Keywords: biomedical; optics; bolus; fluorescence; indocyanine green.

1 Introduction

Optical contrast agents have been promoted to allow for an estimation of perfusion via dynamic measurements and may be used for imaging of pathology via specific molecular probes non-invasively as well. In animal studies, both options have been explored analyzing the fluorescence of different contrast agents, while in humans mostly the optical contrast agent indocyanine green (ICG) has been employed to assess brain perfusion by non-invasive optical approaches. While most studies in humans rely on the absorption properties of ICG, we have previously shown that also laser-induced fluorescence originating from an ICG bolus in the adult human brain can be detected non-invasively at the head surface. The results are promising since the fluorescence signal yielded a superior signal-to-noise ratio and higher sensitivity related to the inflow of ICG when compared to the diffuse reflectance signal. The major challenge of all non-invasive optical measurements, which is the differentiation between truly cerebral and extracerebral tissue contribution, has not been sufficiently solved for fluorescence measurements. The methods developed for determining depth-resolved absorption changes from changes in the time-of-flight distribution or from changes in their moments cannot be applied since fluorescence is generated in the presence of the dye only.

Here we propose a method to separate fluorescence signals from the scalp and the brain based on measured distributions of times of arrival (DTA) of fluorescence photons. We describe an approach based on a linear equation and sensitivity functions that can be derived directly from the measurement without prior knowledge of background optical properties of the head.

2 Experimental Setup

The measurements were performed with a prototype time-domain brain imager described in Ref. that was modified for fluorescence detection. Two detection channels were equipped with longpass filters (793RLP, TFI Technologies) to suppress excitation light and red-sensitive photomultiplier tubes (R7400-20, Hamamatsu Photonics). Continuous sequences of DTAs were recorded with a collection time of 0.2 sec per DTA. The instrumental response function (IRF) was measured according to Ref.

In-vivo measurements were performed on the forehead and over the left motor area of two healthy male volunteers resting in supine position (43 and 37 years). ICG was diluted in water and manually injected into the cubital vein within 2 sec. The dose of ICG was varied within the range from 1 to 6 mg. Up to six injections were applied per session. The experimental procedure was approved by the local ethical committee. The volunteers were two of the authors (Obrig and Steinbrink).

3 Algorithm and Data Analysis

Here we describe our method to decompose fluorescence DTAs into the contributions from the scalp and from the brain. Note
that in the following two time scales are considered, (i) microtime, i.e., a timescale of ps to ns denoted by $t$ relevant for the DTAs, (ii) macrotime, i.e., a timescale of $T$ of seconds to minutes on which the fluorochrome concentration changes during the bolus passage. Thus, the DTAs are functions of the microtime $t$ as well as the macrotime $T$, i.e., the measured fluorescence signal can be expressed as $M_F(t, T)$.

We assume a simple two-compartment model, i.e., the fluorochrome can be present in the brain (index $b$) and in the scalp (index $s$). It is assumed that the total time-resolved fluorescence arriving at the detector can be represented by the sum of the two contributions from both compartments as follows:

$$M_F(t, T) = a_b(T)S_b(t) + a_s(T)S_s(t).$$

(1)

The contributions from the brain and scalp are taken as products in which the microtime and macrotime dependencies are separated. The coefficients $a_s(T)$ and $a_b(T)$ are the macrotime courses of fluorescence from scalp and brain. $S_s(t)$ and $S_b(t)$ denote microtime-dependent sensitivity functions which describe the DTAs for fluorescence generated in the corresponding compartments. The factors $a(T)$ are assumed to be proportional to the concentration, whereas the $S(t)$ are assumed to be independent of fluorochrome concentration. Note that absolute magnitudes of the factors $a(T)$ are not known.

A similar approach is often used in continuous wave fluorescence tomography. In many publications (e.g., Ref. [5]) the sensitivities are determined via photon migration models. For our analysis we developed a novel approach, where the sensitivity functions are extracted directly from the in-vivo measurement assuming that the contribution of one of the two compartments is negligible during certain parts of the bolus transition through the head (macrotime $T$). This assumption, based on physiological knowledge, is sketched in Fig. 1. The rationale behind the assumption is the physiological difference in bolus arrival time in the brain compared to the extracerebral tissue. Hence, the inflow of the dye into the brain tissue [$T = T_b$], “brain-only” period 2 in Fig. 1 is dominant during the early bolus phase, and the contribution from the scalp $a_s$ can be neglected ($a_s = 0$); thus

$$S_b(t) = M_F(t, T_b)/a_b(T_b).$$

(2)

Immediately after the first bolus passage there is virtually no dye in the sampled brain tissue, and fluorescence is assumed to originate from the scalp only [Fig. 1a, period 3 where $a_b = 0$]; thus

$$S_s(t) = M_F(t, T_s)/a_s(T_s).$$

(3)

To find the corresponding periods in the macrotime we used the time courses of the integral $m_0(T) = \int M_F(t, T)dt$ and mean time of flight $m_1(T) = \int M_F(t, T)dt/m_0$ of DTAs. The latter depends on the depth at which fluorescence is generated. Consequently, we identified periods corresponding to $T_b$ and $T_s$ by searching for the local maximum and minimum of $m_1(T)$, respectively. The sensitivity functions were then taken as mean DTAs in these regions. To suppress the influence of ICG which remained in the tissue from previous injections we subtracted a background DTA calculated shortly before the bolus [period 1 in Fig. 1c].

For each time $T$, Eq. (1) represents an over-determined system of several hundred linear equations, corresponding to the number of time channels on the microtime scale, $t$. It is solved in the sense of a least-squares fit using the mrdivide operator of MatLab (Mathworks Inc.). As a result of such analysis for all times $T$, the time courses of the factors $a_s(T)$ and $a_b(T)$ are obtained. The partial contributions from scalp and brain to $I$-integrated fluorescence intensity can be defined as $m_1(T) = \int a_s(T)S_s(t)dt$ and $m_b(T) = \int a_b(T)S_b(t)dt$ with the total fluorescence $m_0(T) = m_b(T) + m_s(T)$.

4 Results and Discussion

Figure 1 shows the sensitivity functions (upper row) and the time courses of the total fluorescence intensity and the contributions from brain and scalp (lower row). The three cases I to III represent typical scenarios found within a total of 34 ICG injections performed in a number of sessions.

Scenario I: The extracted $S_b(t)$ and $S_s(t)$ differ clearly, showing a delayed sensitivity function of the brain. The decomposition results in physiologically reasonable time courses of the integrals $m_b(T)$ and $m_s(T)$, the latter exhibiting an early peak for the bolus arrival, typical for the brain. In brain perfusion studies this peak is the relevant signal, since small delays and broadening are the basis for cerebral perfusion assessment as in perfusion-weighted magnetic resonance imaging (MRI). Compared to the extracerebral compartment the first pass of the intracerebral bolus is clearly earlier and sharper than the passage through the extracerebral tissue.

Scenario II: The extracted $S(t)$ differ only slightly from each other. The time course of the separated integrals contains minor artifacts. The intracerebral contribution monotonously increases after the end of the bolus. This increase is unexpected.

Scenario III: The difference between both $S(t)$ is barely visible. The decomposition delivers substantial artifacts such as negative intensities and an implausible late increase of the intracerebral contribution. It should be noted though, that even in this scenario the first pass of the intracerebral bolus is clearly identifiable.

The time courses of all 34 measurements (28 on the forehead, 6 over the motor cortex) were classified according to the quality of the reconstructed fluorescence time courses for the brain, i.e., good (15 boluses), satisfactory (6), poor (11), and not separable (2). Since the true time course of the ICG bolus is unknown we applied three empirical criteria to judge the quality of the
decomposition results: i. Physiological plausibility of the time course of the reconstructed fluorescence contributions: For healthy subjects the intracerebral bolus should arrive earlier compared to the extracerebral bolus. Negative levels in the reconstructed contributions were regarded as artifacts and therefore reduced the quality level. ii. Difference between maximum and minimum in the course of mean time of flight \( m_1(T) \); a higher value means better quality. iii. Comparison of both sensitivity functions by visual inspection; a visible difference results in better quality.

Regarding the measurements performed on the forehead, in 75\% of all cases the decomposition provided physiologically reasonable results (i.e., classification as satisfactory or good). The six measurements over the motor cortex region were classified as poor (4) or not separable (2). The reproducibility of results obtained for multiple injections of the same dose was good, in particular for the well separable measurements. Within our series of subsequent injections we had seven datasets with two to four equal boluses from which reproducibility of the results could be estimated. The quality scores within these individual series did not differ by more than one grade.

A critical point in the analysis is the assumption of zero ICG concentration after the first bolus. The minimum in \( m_1(T) \) indicates that the concentration in the brain is minimal, but it is unlikely that it returns exactly to zero. A finite residual concentration at the minimum for the brain will introduce a small “parasitical” contribution to the sensitivity function determined on the contralateral, unaffected hemisphere. In this case we hypothesize that the corresponding sensitivity function determined on the contralateral, unaffected hemisphere might be used.

The practical application of the method sometimes showed problems due to only minor differences between the sensitivity functions derived for the cerebral and extracranial compartments. An optimization seems feasible. As could be demonstrated by the results of Monte-Carlo simulations (not shown here) the difference between the sensitivity functions could be increased by decreasing the source-detector separation, while the relative amplitude of the intracerebral contribution would drop.

The reconstruction was successful in 62\% of the measurements on two healthy male subjects. While this could be considered as a promising first step, further improvements are needed to allow for an integration of the proposed method
into a clinical diagnostic tool for monitoring of cerebral blood flow.

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