Collection efficiency of a single optical fiber in turbid media for reflectance spectroscopy

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Abstract: The effect of optical properties on the optical fiber collection efficiency in turbid media was studied experimentally and modeled by Monte Carlo simulations. An analytic expression was obtained to estimate the collection efficiency.

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1. Introduction

Single optical fibers have been commonly used as light delivering and collection tools for optical diagnosis. Authors have proposed their use to determine tissue optical properties [1], measure chromophore relative concentration [2-4] and monitor drug pharmacokinetics [5]. This paper explores the effect of the tissue optical properties on the optical fiber collection efficiency (f) when a single optical fiber is used as light source and detector. The optical fiber collection efficiency is defined as the total number of photons collected by the optical fiber that exit the tissue with an angle less than the fiber acceptance angle (numerical aperture – NA [6]) divided by all the photons collected by the fiber. In other words, f is the number of photons that couple to the fiber CORE divided by the number of photons that couple to the fiber CORE and CLAD as stated in eq. 1, and can be determined by Monte Carlo simulations.

\[ f = \frac{\text{CORE}}{\text{CORE} + \text{CLAD}} \]  

(1)

\[ f_\text{**} = \frac{\text{CORE}}{\text{CORE} + \text{CLAD} + \text{AIR}} \]  

(2)

It is difficult to determine f experimentally since the light lost in the fiber cladding is difficult to measure accurately. For that reason a different collection efficiency (f*, eq. 2; defined by Saidi [7]) was determined experimentally to compare experiments to theory. The term AIR is used for all the photons that exit the medium outside the optical fiber. The sum CORE + CLAD + AIR is equal to the total diffuse reflectance exiting the tissue and can be calculated by Monte Carlo simulations [8, 9] or Adding Doubling [9].

2. Materials and Methods

2.1. Acrylamide Gel Optical Phantoms Preparation

A 6x2 matrix of different optical property acrylamide gel phantoms was prepared using Intralipid as scattering element and India ink as absorber. Stock Intralipid-20% was calibrated with the added absorber technique. Stock Intralipid-20% (\(\mu_s'\approx 200\ \text{cm}^{-1}\) at 630nm) was mixed with water and stock India ink (\(\mu_a\approx 58\ \text{cm}^{-1}\) at 630nm) in appropriate proportions. Solutions had absorption coefficients (\(\mu_a\)) of 0.01, 0.1, 0.5, 1, 5 and 10 cm\(^{-1}\) and reduced scattering coefficients (\(\mu_s'\)) of 10 and 20 cm\(^{-1}\) at 630nm with a final volume of 40ml. Gels were prepared by adding 8g of acrylamide gel crystals, 0.16g of BIS-acrylamide, 0.100g of amonium persulphate and 0.2ml of TEMED to the solutions while stirring at 38° C. Samples gelled after approximate 2 minutes. Final sample volume was 48ml and samples were assumed to be a semi-infinite homogeneous medium for the purpose of modeling.

2.2. Monte Carlo Simulations

Monte Carlo simulations [8, 9] were performed for the same set of optical properties to establish f. Photons were randomly launched within the radius of the fiber forming a collimated beam into a homogenous semi-infinite medium. Henyey-Greenstein phase functions were used to randomly assign propagation angles (\(\Theta\)). The average cosine (or anisotropy, g) was fixed to a typical tissue value of 0.9. Photons were tracked until totally absorbed or until crossing the air/medium boundary. The roulette method [8, 9] was used to conserve energy. Photons that crossed the boundary were summed into three different bins according to the exit location at the surface and the exiting propagation angle (outside fiber = AIR, inside the fiber = CORE, for \(\Theta < \text{NA}\) and = CLAD for \(\Theta > \text{NA}\)).

2.3. Single fiber Reflectance Measurements

Samples were measured with a single 600\(\mu\)m optical fiber (Tefzel, Ceramoptec) coupled to a bifurcated optical bundle through a SMA adapter. The bifurcated bundle was composed of two 300\(\mu\)m optical fibers (Tefzel,
Ceramoptec) coupled to a single 600µm SMA connector at the distal end and two 300µm SMA connectors at the proximal ends. One fiber was connected to a tungsten-halogen white lamp (Ocean Optics) and the other to a spectrometer (Ocean Optics) controlled by a laptop computer. The setup of the system is shown in figure 1. Acquisition time was set to 30ms and a filter (2OD) was used with all samples and a stronger filter (3OD) was used with a mirror as a sample. The 600µm fiber was placed in contact with the samples with a water drop to avoid the fiber/air/gel refractive index mismatch.

![Experimental setup](image)

**Fig. 1.** Experimental setup. Acquisition time was set to 30ms and filters were placed before coupling to optical fiber.

\[ f^{*}\text{sample} = \frac{M_{\text{mirror}}}{S \cdot R_{\text{mirror}} \cdot f_{\text{mirror}} \cdot D} = f^{*\text{sample}} \cdot R_{\text{mirror}} \cdot 1 \]  
\[ f^{*\text{mirror}} = \frac{M_{\text{sample}} \cdot 0.90}{M_{\text{mirror}} \cdot R_{\text{sample}}} \]  

\( f^{*\text{mirror}} \) was assumed to be 1 since all the light that hits the mirror returns to the fiber with an angle smaller than the acceptance angle. \( R_{\text{sample}} \) (total diffuse reflectance) was obtained by a forward Adding Doubling calculation [8] with knowledge of the sample optical properties.

### 3. Results

Figure 2 shows the results for \( f^{*} \) determined by Monte Carlo (empty symbols) and experiments (filled symbols) for two \( \mu_{s}' \left( = 10\text{cm}^{-1}, \text{ and } \phi = 20\text{ cm}^{-1} \right) \) and six \( \mu_{a} \) (greater \( \mu_{a} \) to the left). \( f^{*} \) is plotted as a function of \( X[-] \) defined as \( X = \delta \cdot \text{mfp}'/d^2 \), where \( \delta = (3 \mu_{a}(\mu_{a}+\mu_{s}'))^{-1/2} \), \( \text{mfp}' = 1/(\mu_{a}+\mu_{s}') \) and \( d \) is the diameter of the optical fiber.

![Graph](image)

**Fig. 2.** \( f^{*} \) from Monte Carlo (empty symbols) and experiment (filled symbols). See text for details.

![Graph](image)

**Fig. 3.** Collection efficiency \( f \) per 5° bin. Integration of these values from 0 to the NA equals \( f \) for that NA.

Figure 3 demonstrates the dependence of the collection efficiency on the optical properties by plotting \( f \) as a function of collection angle bin (three \( \mu_{s}' \) 80, 20 and 1 cm\(^{-1}\), top to bottom). In the same figure plots of \( \text{COS(θ)SIN(θ)} \) (dashed lines; see eq. 5) show the similarities of the data to this simple expression for higher scattering and the differences for low scattering.

Monte Carlo simulations for a fiber diameter of 600µm and NA of 0.39 immersed in a medium with index of refraction of 1.35 (equivalent to a half angle of \( \sin^{-1}(0.39/1.35) = 16.79° \)) is showed in figure 4. A side view of the 3D plot shows that for \( \mu_{s}' \) above 20 cm\(^{-1}\) \( f \) approaches a value of 0.0835.
4. Discussion

Comparison between Monte Carlo (empty symbols) and experimental (filled symbols) $f^*$ in figure 2 show good agreement for physiologically relevant $\mu_s'$. Larger errors were observed for higher absorption (left-most points) probably due to the reduced signal/noise ratio for those measurements. Data tends to collapse to a monotonic sigmoidal curve when plotted against the non-dimensional parameter $X$ in a semi-log plot.

Equation 1 can also be interpreted as the total fraction of light that reaches the optical fiber with an angle smaller than that defined by the NA ($\theta_1$) divided by the total light that couples to the fiber at all angles, as stated in eq.5. Integration of eq.5 in spherical coordinates shows that $f$ should be equal to $\sin^2(\theta_1)$.

\[
    f = \frac{\int_0^{2\pi} \int_0^{\theta_1} \cos(\theta) \sin(\theta) d\theta d\phi}{\int_0^{2\pi} \int_0^{\pi} \cos(\theta) \sin(\theta) d\theta d\phi} = \frac{\pi \sin^2(\theta_1)}{\frac{2}{3} \pi} = \sin^2(\theta_1)
\]

(5)

For a NA of 0.39 and a medium with index of refraction of 1.35, $\theta_1$ is equal to 16.79°. Applying this angle in equation 5 gives $f = 0.0835$. The value of $f$ calculated from equation 5 is a good first approximation for most optical properties, as shown in figure 4. But it does not agree for small (physiologically relevant) values of $\mu_s'$ (figure 4 RIGHT). The cone of collection of an optical fiber (defined by the fiber NA) is dependent only on the indices of refraction of the fiber core/clad and the medium where the fiber is immersed [6]. Changes in the optical fiber collection efficiency for turbid media arises from differences in the angular distribution of the photons that reach the fiber for different optical properties, as observed in figure 3 for low scattering (bottom curve). These changes are not caused by an intrinsic parameter of the optical fiber but arise from its use in a turbid media, like biological tissues. The parameter $f$ can be used as a practical guide for choosing the NA of an optical fiber for biomedical applications.

5. References