

A Self-Calibrating Fiber-Optic Probe for Tissue Optical Spectroscopy

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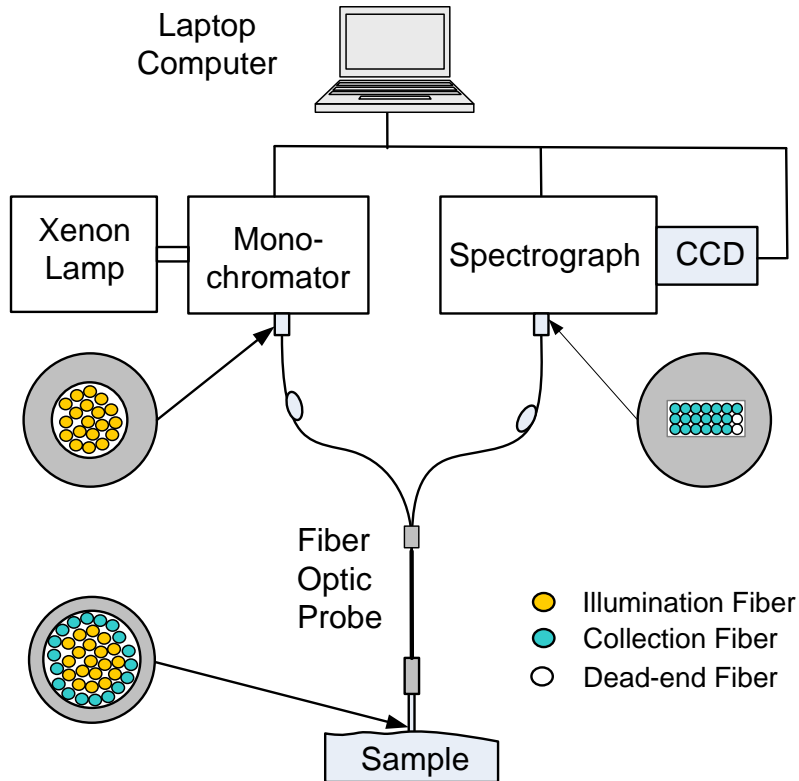
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Tissue Optical Spectroscopy



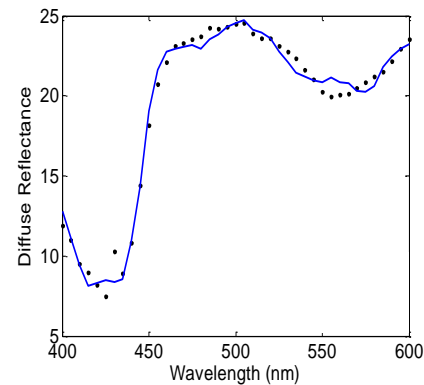
Optical spectroscopy is sensitive to the absorption, scattering and/or fluorescence properties of biological molecules in tissue.



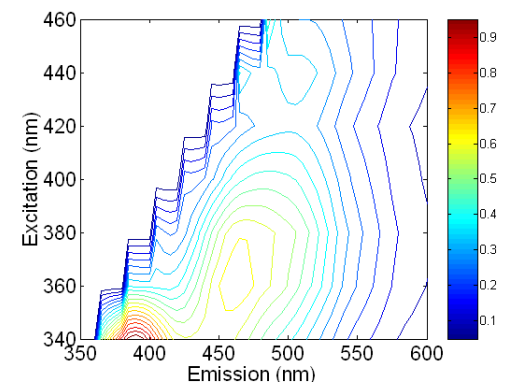
Tissue Optical
and Physiological Properties

Scalable Monte
Carlo Model

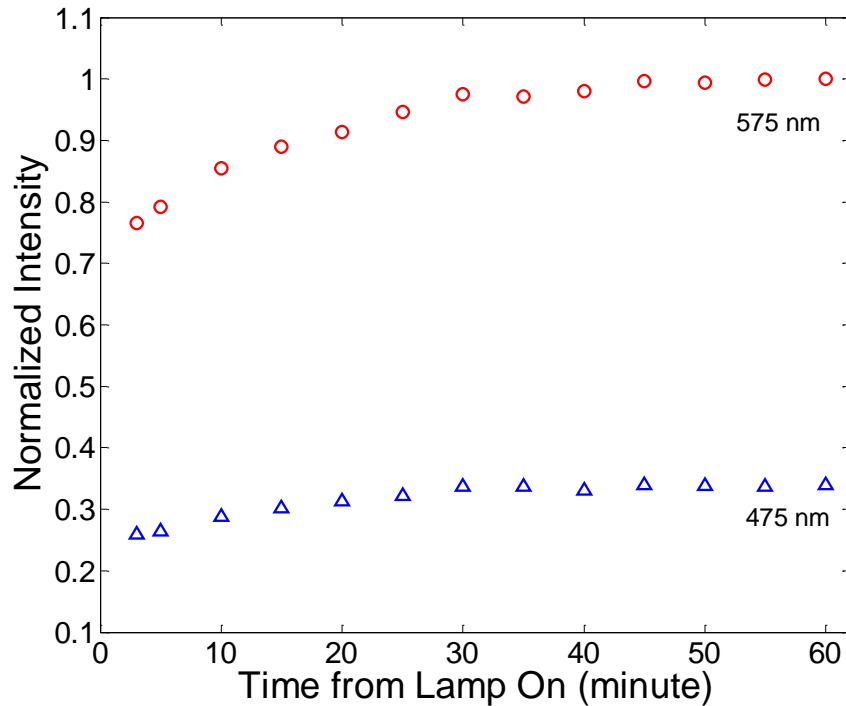
Diffuse Reflectance



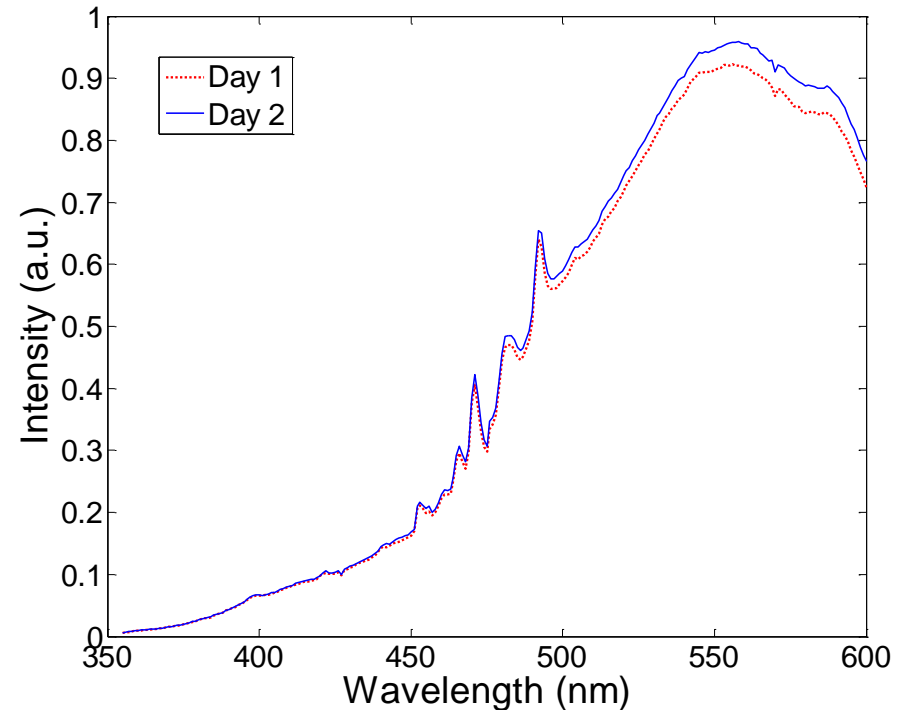
Fluorescence EEM



Sources of Intensity Fluctuations



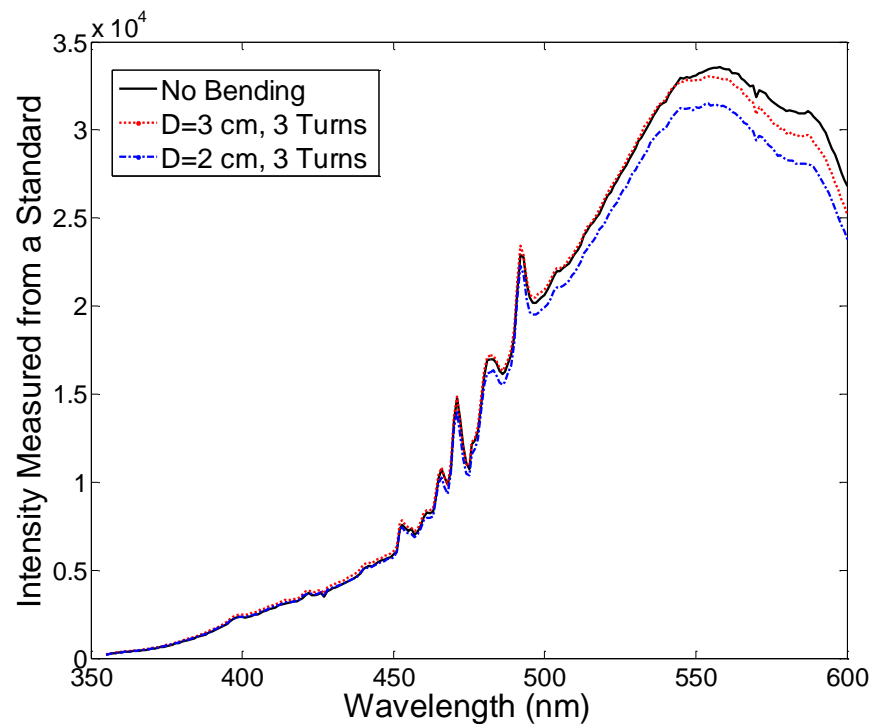
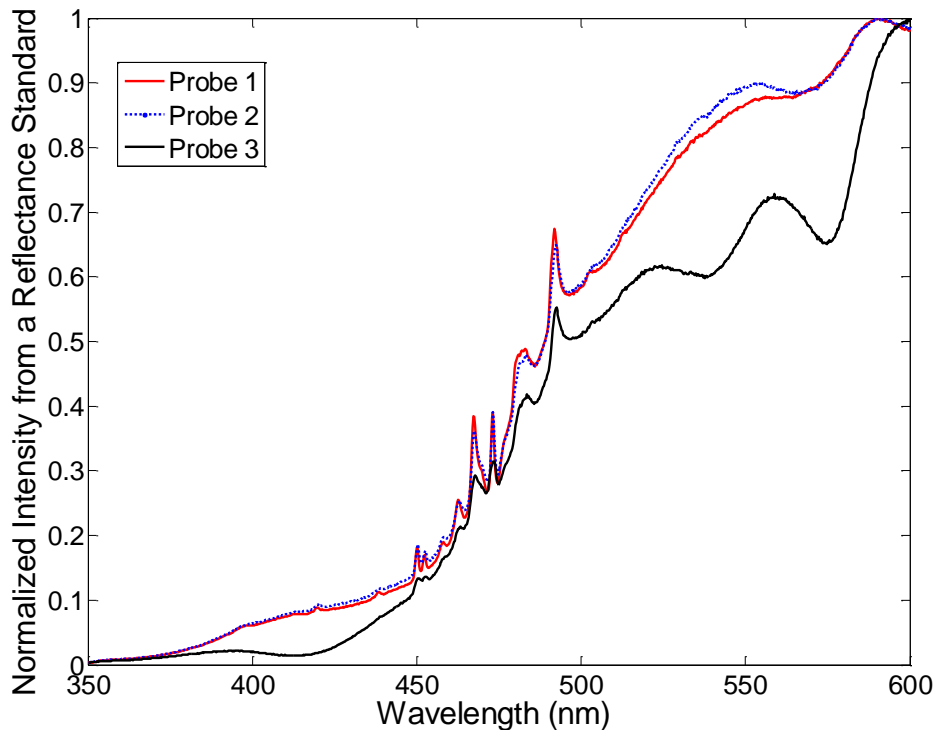
- Lamp intensity fluctuates 25% from 3-60 minutes and 3% after 30 minutes for 575 nm.
- The ratio I_{575}/I_{475} changes 4.4% after 30 minutes.



- The lamp intensity varies as much as 6.8% from day to day

5% change in intensity \rightarrow ~20% error in μ_a & μ_s'

Sources of Spectrum Variations



- **Instrument: wavelength-dependence**
- **Devices: degradation over time**

**5% change in intensity \rightarrow \sim 20% error
in μ_a & μ_s'**

- **Fiber bending loss during measurements**
- **$> 6\%$ at a diameter of 3cm**
- **$> 11\%$ at a diameter of 2cm**

Traditional Calibration Methods & Limitations



	Tissue Type	Calibration Method	Corrected for				Warm-up NOT Required	Cal. Meas. NOT Required in Clinic
			Through-put	Day-to-Day	Real-time			
					Lamp Drift	Fiber Bend		
Feld M.S. ^[1]	Colon polyps	Liquid phantom	Y	?	N	N	N	N
Richards, Kortum P. ^[2;3]	Ovarian, Cervix	Liquid phantom	Y	?	N	N	N	N
Thueler P. ^[4]	Stomach	Solid phantom	Y	Y	N	N	N	N
Foster T. ^[5]	Phantoms	Ref. fiber & liquid phantom	Y	Y	N	N	N	N
Our Group (Old) ^[6]	Breast, Cervix	Diffuse refl. Standard & Phantom	Y	Y	N	N	N	N

A real-time, faster, more robust and accurate calibration approach is needed!

Accuracy

Efficiency

¹ Zonios G, et al., *Appl. Opt.* 38(31): 6628-37, 1999.

² Utzinger U, et al., *Lasers Surg Med* 28(1): 56-66, 2001.

³ Mirabal YN, et al., *JBO* 7(4): 587-94, 2002.

⁴ Thueler P, et al., *JBO* 8(3): 495-503, 2003.

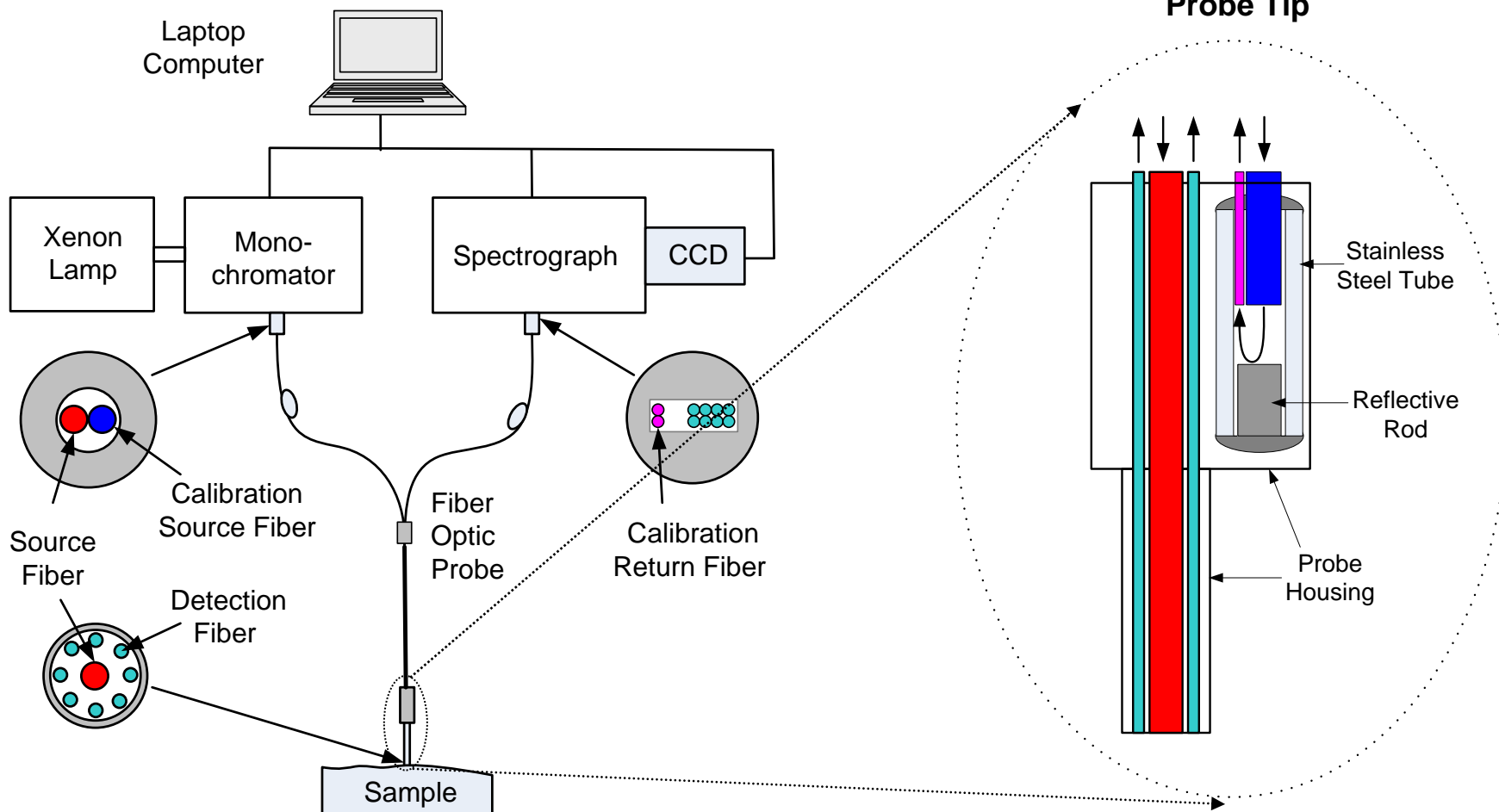
⁵ Nichols MG, et al., *Appl. Opt.* 36(1), 1997.

⁶ Zhu C, et al., *JBO* 10(2): 24-32, 2005.

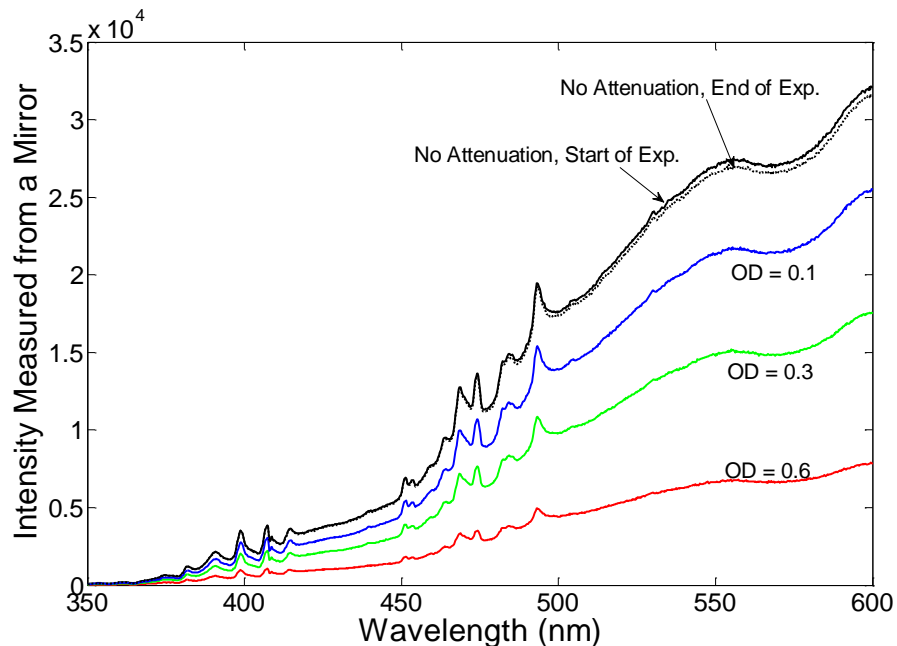
Experimental Setup



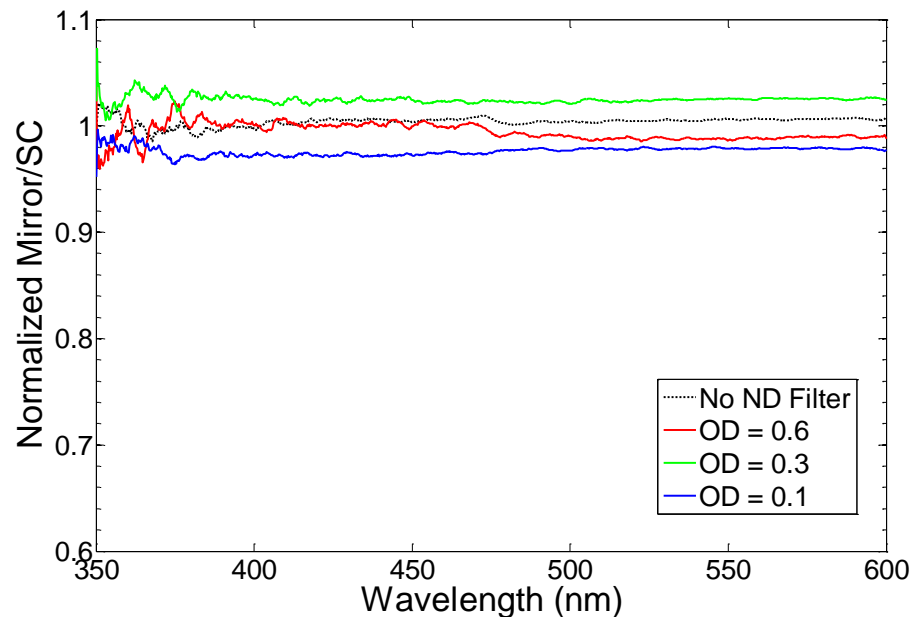
Schematic of the System



Correction for Intensity Variations



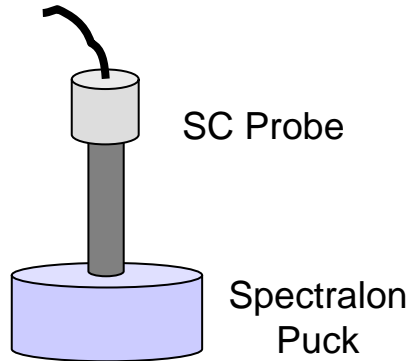
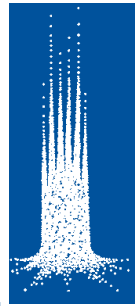
Raw spectra from a mirror at different levels of attenuation by an ND filter and no attenuation.



Ratio between mirror and self-calibration spectra normalized to the first scan (no filter).

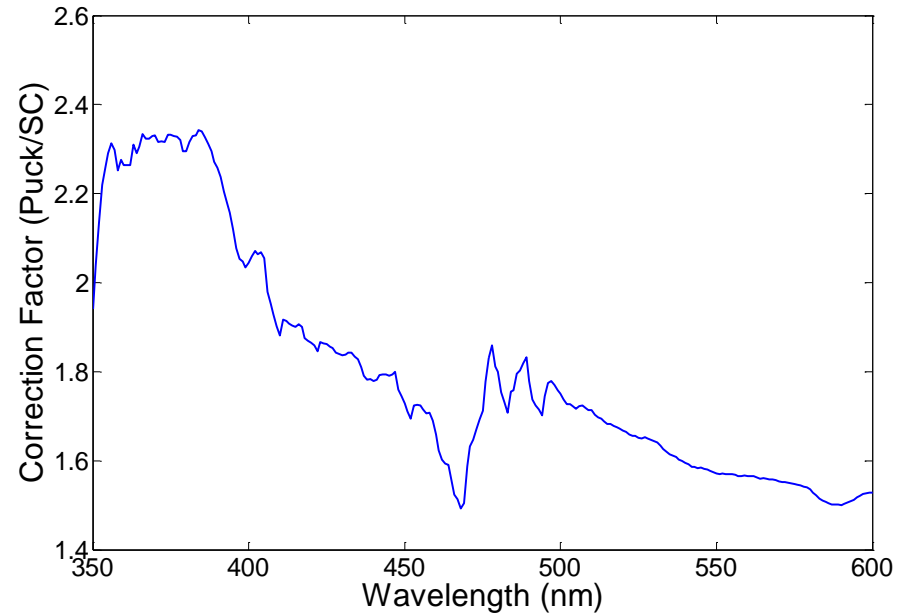
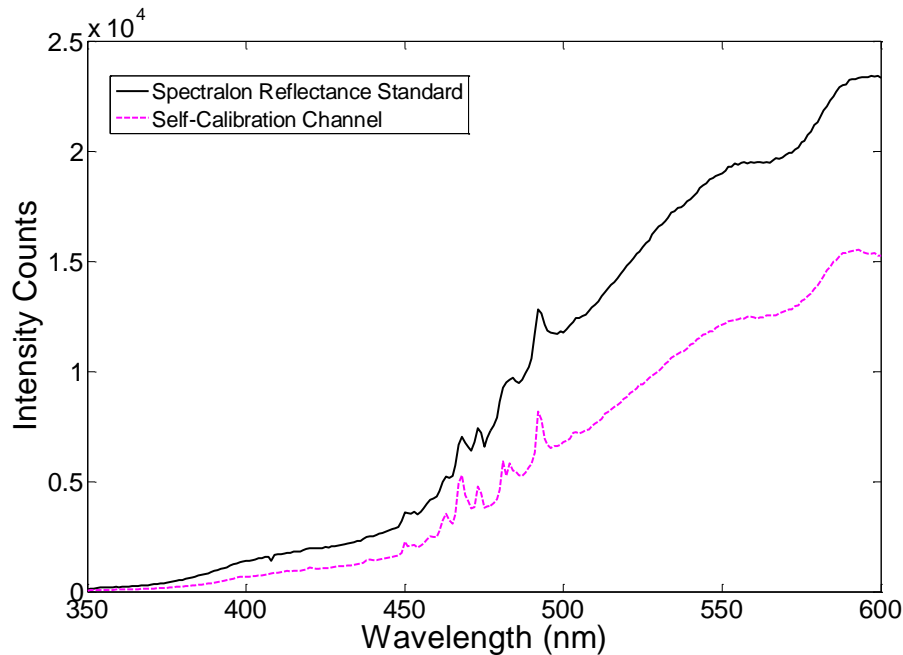
The variation in the illumination intensity due to the ND filters is minimized to less than $\pm 3\%$.

Correction for Wavelength-Dependence



Correction of the SC channel for wavelength response:

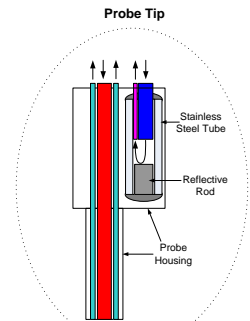
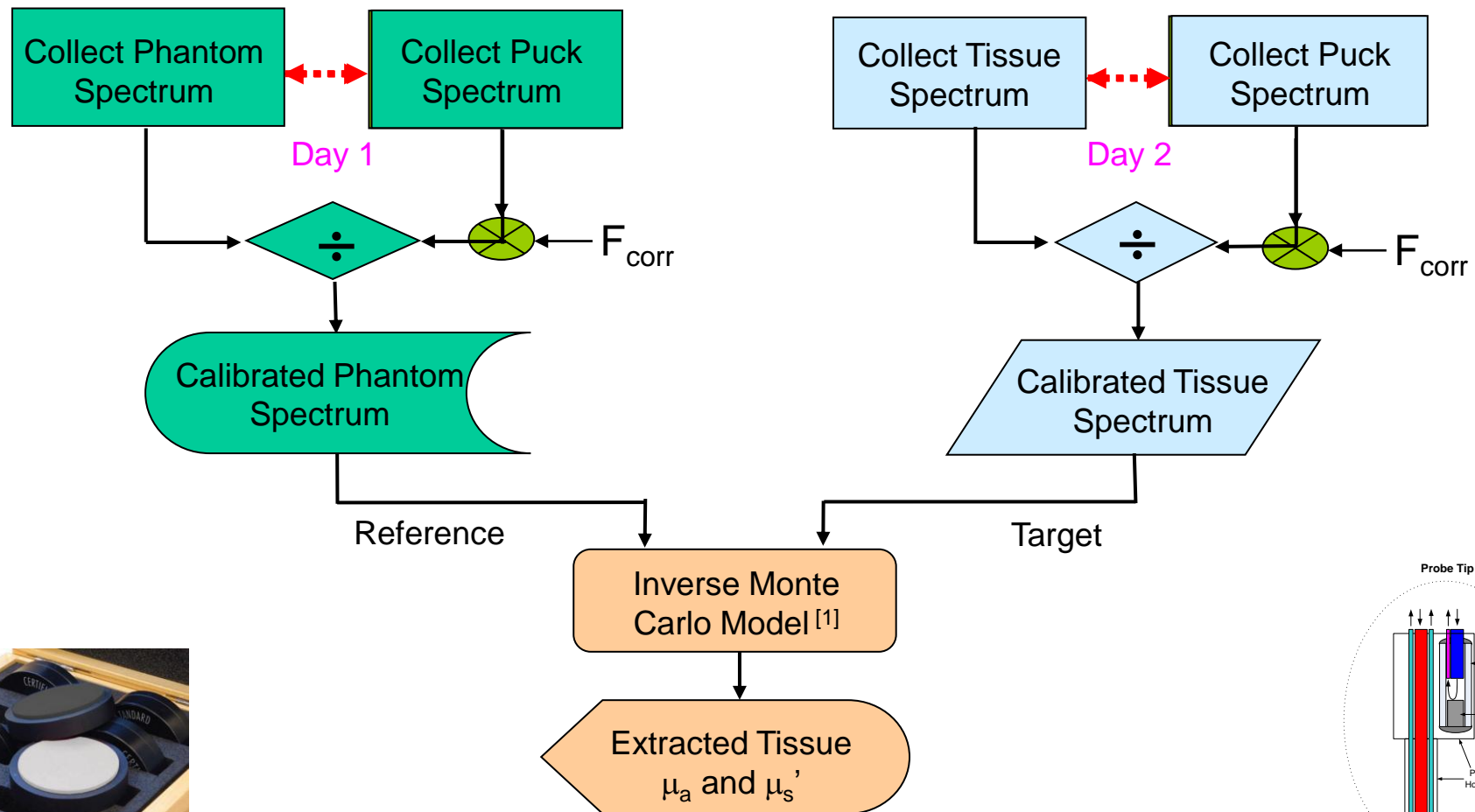
$$F_{corr}(\lambda) = \frac{R_{Puck}(\lambda)}{R_{SC}(\lambda)}$$



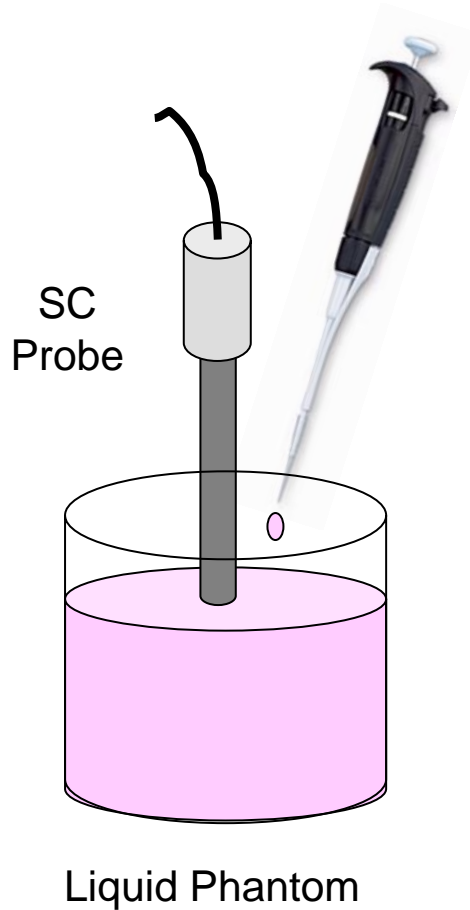


Data Analysis Procedures

Calibration with Spectral Reference Standard (Puck)



Phantom Experiments



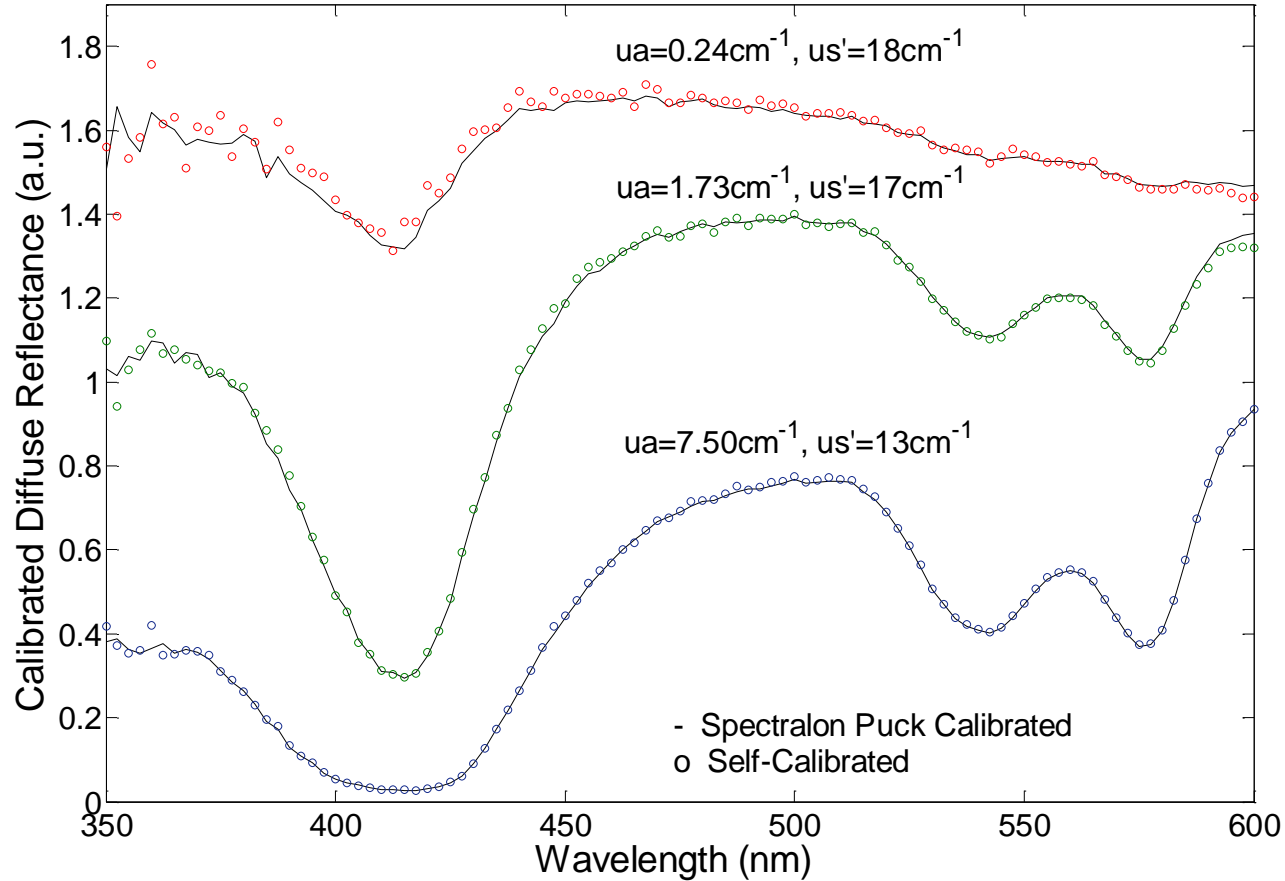
Tissue-mimicking phantoms:

- Absorber: hemoglobin
- Scatterer: 1 μm polystyrene microspheres
- [Hb]: 1 – 32 μM
- $\mu_a = 0.006 - 37 \text{ cm}^{-1}$
- $\mu_s' = 11.2 - 22.3 \text{ cm}^{-1}$
- Reference phantom: [Hb]=8.42 μM , $\mu_s' = 16.8 \text{ cm}^{-1}$

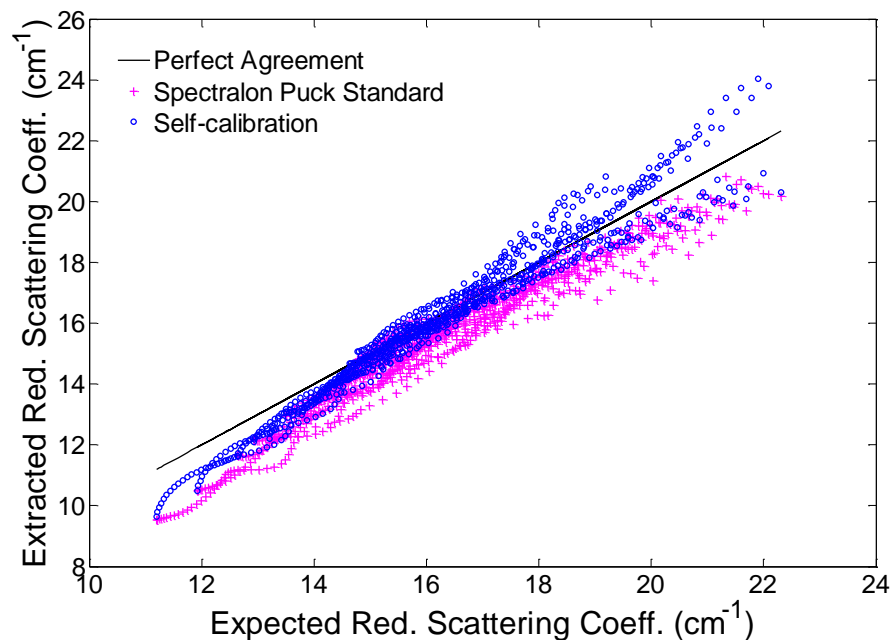
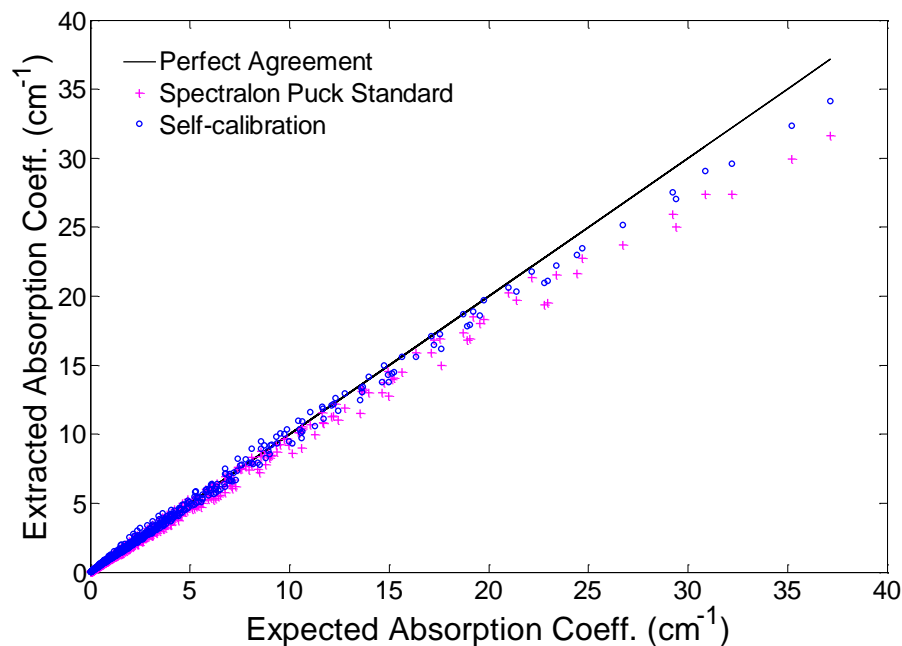
Calibrated Phantom Spectra



$$R_{puck_calibrated} = \frac{R_{tissue}}{R_{puck}} \quad v.s. \quad R_{sc_calibrated} = \frac{R_{tissue}}{R_{sc} \cdot F_{corr}(\lambda)}$$



Extracted vs Expected Phantom Optical Properties



	Spectralon Standard	Self-Calibration
Error in μ_a	$6.9 \pm 7.2\%$	$6.8 \pm 5.1\%$
Error in μ_s'	$3.5 \pm 1.5\%$	$5.6 \pm 2.9\%$



Conclusions

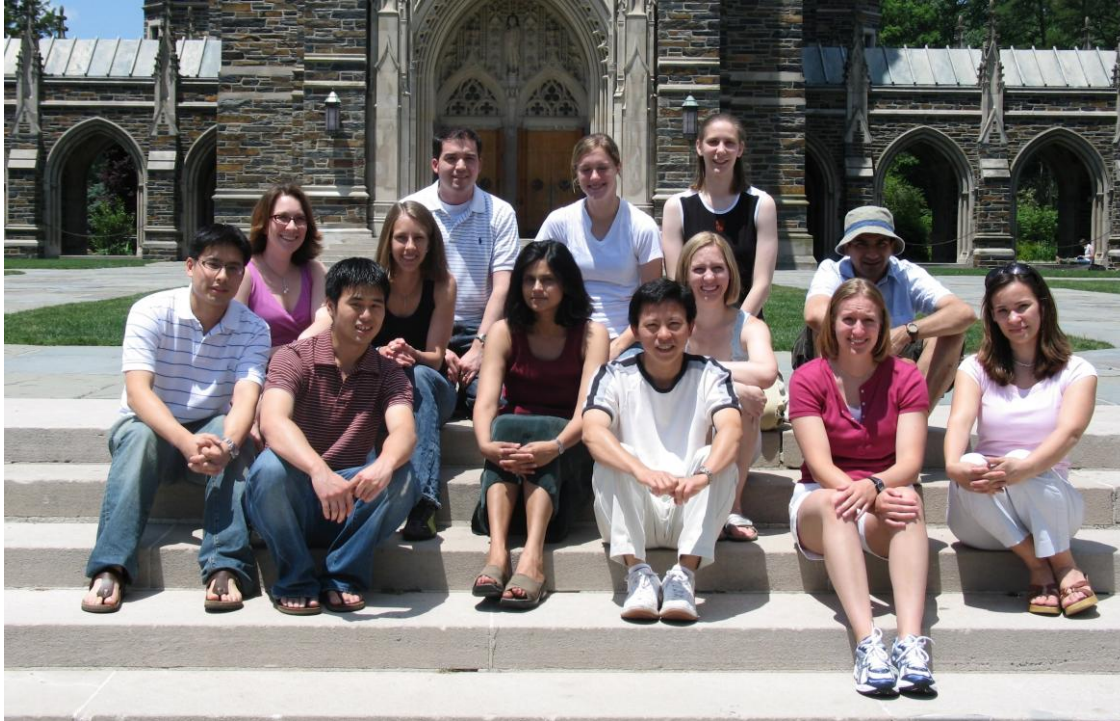
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Our Group (Old) ^[6]	Breast, Cervix	Diffuse refl. Standard & Phantom	Y	Y	N	N	N	N
Our Group (New)	Phantoms	Self-cal. & Phantom	Y	Y	Y	Y	Y	Y

Accuracy

Efficiency

Combined with a one time, single-reference phantom measurement, the self-calibrating probe can provide instrument-independent optical properties.

Acknowledgements



**Tissue Optical
Spectroscopy Lab
Dept. of Biomedical Eng.
Duke University**

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