## Monte Carlo Analysis of Near Infrared Wavelengths Towards an Optical NIR Blood Analyte Sensor

3D Monte Carlo (MC) light-tissue interaction simulations was conducted for seven near-infrared (NIR) wavelength LEDs, 1050, 1200, 1300, 1450, 1460, 1550, and 1650 nm, to assess the penetration depth, pathlength, and detected power to develop noninvasive sensing methods for blood analytes.

Methods: The MC simulations were performed using an adapted version of "MCmatlab: an open-source, user-friendly, MATLAB-integrated three-dimensional Monte Carlo light transport solver with heat diffusion and tissue damage". Figure 1 presents the emitter-detector geometry, and 7-layer tissue composition used.



Fluence rate

1050 nm

Absorption

1050 nm

**Results:** Each wavelength was simulated for 12 hours where 9e9 - 9e10



photon packets were launched. Figure 2 presents the fluence rate and absorption [W/m^2/W-incident] for each wavelength on the centre of the emitter-detector plane.

The mean photon pathlength was between 2.330 and 2.548 mm (Figure 3A). The mean maximum photon depth was between 0.986 and 1.141 mm (Figure 3B). The percentage of photons collected by the detector were all less than 0.1% (Figure 3C).



Figure 3: (a) Average pathlength, (b) depth, and (c) percentage of photons collected for each wavelength [nm].

The maximum sum normalised fluence rate was between 87.86 and 56.10 W/m2/W-incident located in the stratum corneum layer for all wavelengths (Figure 4A). The maximum normalised absorption was between 10.14, and 37.03 W/m2/W-incident located in the upper dermis (Figure 4B).



Figure 2: Fluence rate and absorption [W/m^2/W-incident] for each wavelength

Figure 4: Normalised fluence rate and normalised absorption [W/m^2/W-incident] through each layer for each wavelength

**Conclusions:** The penetration depth increases with wavelength as suggested by the increase in penetration depth from visible to SWIR. The percentage of photons detected is less than 0.1%, highlighting the importance for highly sensitive photodetectors with these wavelengths. MC simulations were carried out on NIR wavelengths to better understand the light-tissue interaction. The pathlength and penetration depths found will be used to develop sensor emitter-detector separation and device selection, and post processing algorithms for a non-invasive blood analyte concentration sensor.



<u>Josephine A. Dixon</u>, University of Canterbury, Christchurch, New Zealand, josie.dixon@pg.canterbury.ac.nz Jordan F. Hill, University of Canterbury, Christchurch, New Zealand, jordan.hill@pg.canterbury.ac.nz J. Geoffrey Chase, University of Canterbury, Christchurch, New Zealand, geoff.chase@canterbruy.ac.nz Christopher Pretty, University of Canterbury, Christchurch, New Zealand, chris.pretty@canterbruy.ac.nz

