## Measuring Reaction Rate Constants With Optical Biosensors

Ryan Evans<sup>1,\*</sup>, Tony Kearsley<sup>1</sup>, <u>David A. Edwards</u><sup>3</sup>, Arvind Balijepalli<sup>2</sup>

<sup>1</sup>Applied and Computational Mathematics Division, NIST, 100 Bureau Drive, Gaithersburg, MD 20899

<sup>2</sup>Engineering Physics Division, NIST, 100 Bureau Drive, Gaithersburg, MD 20899 <sup>3</sup>Department of Mathematical Sciences, University of Delaware, Newark, DE 19716 \*ryan.evans@nist.gov

## Abstract

Many biochemical reactions involve a stream of chemical reactants flowing through a fluid-filled volume, over a surface to which receptors are confined. Such surface-volume reactions occur during blood clotting, drug-protein interactions, and DNA-damage repair. Scientists measure reaction rate constants associated with these reactions using optical biosensors: an instrument in which reactants are convected through a flow-cell, over a surface to which other reactants are immobilized.

Scientists currently study biosensor experiments which involve multiple interacting components on the sensor surface. We discuss a partial differential equation model for multiple-component reactions in optical biosensors. Thanks to high Peclet number flow, this model reduces to a set of nonlinear integrodifferential equations for the reacting species concentrations, which in turn reduces to a set of ordinary differential equations which can be used to measure rate constants using biosensor data. We conclude by discussing recent developments on a related problem concerning instruments involved in creating personalized medicine.