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When studying gas channels in cells, surface and intracellular pH are determined in an oocyte using electrodes which measure the concentration of hydrogen ions ( $H^+$ ). It has been hypothesized that the presence of the electrode creates microenvironmental effects on pH measurements near the cell membrane. We develop a mathematical reaction-diffusion model of the electrode tip at the cell membrane which is restricted into a subdomain with artificial boundary conditions to keep the computational burden manageable. We consider an oocyte immersed in a liquid in which the pH is modulated by controlling the contents of carbon dioxide,  $CO_2$ , which reacts with water forming carbonic acid,  $H_2CO_3$ , which further dissociates to bicarbonate  $HCO_3^-$ , thus releasing a proton  $H^+$ . The local concentrations change due to diffusion, reaction, and gas transport through the cell membrane. To implement the solution numerically, we first discretize the solution in the spatial direction using a finite element scheme, reducing the problem into a system of ODEs. Through a series of numerical experiments, we show that the electrode does, in fact, create a microenvironment impacting the extracellular pH measurements. (Received September 26, 2017)