

Nanoscale

Accepted Manuscript



This article can be cited before page numbers have been issued, to do this please use: B. M. Lowe, C. Skylaris, N. G. Green, Y. Shibuta and T. Sakata, *Nanoscale*, 2018, DOI: 10.1039/C8NR00776D.



This is an Accepted Manuscript, which has been through the Royal Society of Chemistry peer review process and has been accepted for publication.

Accepted Manuscripts are published online shortly after acceptance, before technical editing, formatting and proof reading. Using this free service, authors can make their results available to the community, in citable form, before we publish the edited article. We will replace this Accepted Manuscript with the edited and formatted Advance Article as soon as it is available.

You can find more information about Accepted Manuscripts in the [author guidelines](#).

Please note that technical editing may introduce minor changes to the text and/or graphics, which may alter content. The journal's standard [Terms & Conditions](#) and the ethical guidelines, outlined in our [author and reviewer resource centre](#), still apply. In no event shall the Royal Society of Chemistry be held responsible for any errors or omissions in this Accepted Manuscript or any consequences arising from the use of any information it contains.

Cite this: DOI: 10.1039/xxxxxxxxxx

Molecular Dynamics Simulation of Potentiometric Sensor Response: Effect of Biomolecules, Surface Morphology and Surface Charge[†]

B. M. Lowe,^{a*} C-K. Skylaris,^b N. G. Green,^c Y. Shibuta^a and T. Sakata^{a*}

Received Date

Accepted Date

DOI: 10.1039/xxxxxxxxxx

www.rsc.org/journalname

The silica-water interface is critical to many modern technologies in chemical engineering and biosensing. One technology used commonly in biosensors, the potentiometric sensor, operates by measuring changes in electric potential due to changes in the interfacial electric field. Predictive modelling of this response caused by surface binding of biomolecules remains highly challenging. In this work, through the most extensive molecular dynamics simulation of the silica-water interfacial potential and electric field to-date, we report novel prediction and explanation of the effects of nano-morphology on sensor response. Amorphous silica demonstrated a larger potentiometric response than an equivalent crystalline silica model due to increased sodium adsorption, in agreement with experiments showing improved sensor response with nano-texturing. We provide proof-of-concept that molecular dynamics can be used as a complementary tool for potentiometric biosensor response prediction. Effects that are conventionally neglected, such as surface morphology, water polarisation, biomolecule dynamics and finite-size effects are explicitly modelled.

1 Introduction

Over the last century, significant progress in the field of study of electrified charged interfaces has enabled development of a range of new technologies, such as electrochemical fuel cells¹, the ability to engineer colloids² and materials for water purification^{3,4}. Despite this progress, improved understanding is required for reliable design and modelling of these technologies. In particular, the complex interaction between surface charge, the electrostatics of water polarisation, charged analyte molecules and electrolyte ions remains a topic of active research.

An example of one such technology are potentiometric sensors. The response of potentiometric sensors originates from a change in electric potential within the system resulting from interfacial electrostatics. A popular class of potentiometric sensors are ion-sensitive field-effect transistor (IS-FET) sensors. IS-FETs were initially popularised in the 1970s by Bergveld^{5,6} and can detect changes in pH due to surface charging reactions and ion-adsorption at the oxide-electrolyte interface. Starting with the work of Cui et al.⁷ in 2001, field-effect sensors have been func-

tionised with receptors specific to biomolecular analyte for the task of sensing biomolecular analyte, often referred to as 'BioFET' sensors. This functionalisation facilitates a type of biosensor which offers many advantages over standard immunological detection methodologies such as the capability for label-free, low-cost electrostatic analyte detection⁸. In the present work, field-effect sensors are focused upon as an example application system, however, the interfacial physics described is relevant to all kinds of potentiometric biosensors^{9,10}.

The silica-water interface in particular was investigated for the present work as it is a popular choice of oxide material for potentiometric sensors given its low-cost and simple production by thermal treatment of silicon wafers^{11–13}. Silica-water interfaces are also one of the most abundant on the planet, and therefore the relevance of understanding their interfacial electrostatics extends beyond sensor response, into understanding geochemical processes such as dissolution kinetics^{14–16} and prebiotic biochemistry¹⁷.

Reliable and accurate quantitative predictions of the response of field-effect sensors due to interaction of analyte molecules is currently unavailable using even the most sophisticated models, and even qualitative predictions of the response of field-effect sensors remains challenging^{8,13}. The majority of field-effect sensor models incorporate a model of the electrical double layer primarily based on either the Poisson-Boltzmann equation^{18–28} or the linearised Poisson-Boltzmann equation^{13,29–31}, which were first developed in the early 20th century³². This approach neg-

^a Department of Materials Engineering School of Engineering, The University of Tokyo 7-3-1 Hongo, Bunkyo, Tokyo 113-8656, Japan. E-mail: sakata@biofet.t.u-tokyo.ac.jp

^b School of Chemistry, University of Southampton, UK.

^c Department of Electronics and Computer Science, Nano Research Group, University of Southampton, UK. E-mail: ng2@ecs.soton.ac.uk

[†] Electronic Supplementary Information (ESI) available: [details of any supplementary information available should be included here]. See DOI: 10.1039/b000000x/

* Corresponding Authors



lects effects that are expected to be important to the generation of a potentiometric signal, examples of which would include, but are not limited to; the effect of Van der Waals interactions, water polarisation, electrolyte-protein ion dynamics and the finite size of biomolecules. Furthermore, difficulty in controlling experimental conditions necessary for reproducible response, such as the analyte-receptor density at the sensor surface⁸, has contributed to difficulties in validating and improving the predictive power of field-effect response models.

There are several research questions which are particularly problematic to understand via Poisson-Boltzmann-based models. For example, the experimental observation of potentiometric response due to binding of electrically neutral non-polar organic molecules^{33,34} and the anomalously large signal of hexanol compared to butanol on bare silica³³. Other examples include the enhancement of potentiometric biosensor response at high ionic strength by addition of a biomolecule-permeable polyethylene glycol surface layer³⁵ and the changes in high frequency signal induced by biomolecule-electrolyte dynamics^{8,35–37}.

The aim of the present work is to use molecular dynamics simulations as a novel tool for prediction of potentiometric biosensor response and for improving understanding of the interfacial electrostatics. The simulated surface potential is calculated as a function of surface charge for a simple well-experimentally characterised system - the silica-water interface. The results are validated against analytical models and experimental pH sensing data. Using the relationship between response and surface charge as a baseline, the magnitude of response due to addition of a model charged biomolecule-DNA, is investigated. For biomolecular systems, a molecular dynamics-based approach has several advantages over conventional mean-field models due to its explicit treatment of biomolecule dynamics, water polarisation and finite-size effects.

Various durations of molecular dynamics simulation have been used in the literature to calculate the surface potential at the silica-water interface: ~ 1 ns³⁸ (reactive forcefield), 3 ns^{39–42}, 20 ns⁴³ and ~ 30 ns⁴⁴. Whilst sufficient for obtaining qualitative changes in potential, this duration is too short to reliably quantitatively distinguish the millivolt changes in surface potential expected from biomolecular signal and pH changes. For example DNA is known to take at least 300 ns for its ionic atmosphere to equilibrate⁴⁵ and at least 90 ns is required to obtain a silica-water surface potential with a mean potential stable to within several millivolts⁴⁶. In the present work, we perform 24 simulations of the silica-water interface for 320 ns and several simulations including DNA molecules for 480–680 ns totalling an extensive total simulation duration of at least 8 μ s between all simulations. Furthermore, compared to our previous work^{39–42}, we use a forcefield which has increased reliability for describing the silica-water-bio interface and a methodology for evaluating the electrostatics with increased accuracy^{43,46,47}. Combined, these considerations ensure the present work is amongst the most rigorous molecular dynamics simulation of the silica-water interfacial potential and electric field to-date. This work is of particular novelty because molecular dynamics-based oxide-water surface potential calculations remain rare, likely due to historical diffi-

culties in obtaining accurate forcefields which integrate both organic and inorganic molecules.

The surface potential calculation methodology used has been detailed and discussed in our most recent work for a simple crystalline silica-water system⁴⁶. In the present work, we focus on the application of understanding potentiometric sensor response and nanoscale electrostatics. We report a novel prediction and explanation of the effects of nano-morphology on surface potential changes, and provide the first proof-of-concept that molecular dynamics can be used as a complementary tool for potentiometric biosensor response prediction.

The background section is structured as follows: Section 2.1 presents commonly used models for the electrical double layer as these will be directly compared to the molecular dynamics simulations of the present work. Then in Section 2.2, the commonly-used site-binding model for prediction of response of potentiometric sensors to changes in pH and its limitations, are introduced. In Section 2.3, a brief outline of the surface chemistry of silica is provided because accurate modelling of the surface chemistry is crucial to an accurate molecular dynamics model of the interface. Finally, literature experimental data is presented for the effects: surface morphology on sensor response (Section 2.4), and the addition of DNA on sensor response (Section 2.5).

2 Background

2.1 Mean-Field Electrical Double Layer Models

In this section, a brief overview of analytical mean-field electrical double layer models relevant to potentiometric sensor response modelling is presented. A more detailed background for these models can be found in Supplementary Information 1

The simplest and most common model of field-effect sensor response treats the response due to biomolecules as charge on a parallel plate capacitor model of the surface (also called the ‘constant capacitance’ model or Helmholtz-Perrin model⁴⁸). In such models, the measured change in potential is assumed to be equal to the change in surface charge from biomolecule binding, divided by a capacitance value which describes the coupling of the analyte to the FET semiconducting channel^{49,50}. Even though this method may be useful for estimating the density of bound molecules from an experiment, it provides no generally transferable insight into the precise relationship between the contents of the electrical double layer (i.e. biomolecule binding, electrolyte ionic strength etc.) and sensor response, and therefore is not considered in the present work.

Aside from the Helmholtz-Perrin model, field-effect sensor response models almost exclusively utilise the Poisson-Boltzmann equation (or the more general Poisson-Nernst-Planck equation) to describe the electrical double layer. Such models therefore explicitly incorporate the effect of ionic strength on the response. Primary differences between field-effect biosensor models occur in the way that the biomolecular component is treated in a mean-field manner. For example, whether the biomolecular charge is treated as a ‘smeared out’ surface charge over an infinitely thin surface charge layer^{27,51} or as an ion-permeable membrane of finite thickness^{52,53}. Some recent attempts at modelling field-



effect sensor response model the charges on the biomolecule discretely in a continuum solvent^{13,30,54,55}. Such an approach provides a good compromise between atomistic and continuum approaches, but neglects the important effects of biomolecule dynamics, surface morphology and water polarisation.

In the present work, the surface charge to surface potential relationship for the Poisson-Boltzmann equation, Debye-Hückel theory and modified Poisson-Boltzmann equation are presented for comparison with the molecular dynamics results; expressions will be labelled using subscripts *pb*, *dh* and *mpb*, respectively. The surface potential ϕ_s refers to the potential only at the surface. The equations relating to the position dependent potential shown in Supplementary Information 1. These models are chosen in the present work due to their simplicity and wide-spread usage. In addition they require no or minimal system-specific empirical parameters, and therefore provide predictions that are transferable across oxide systems without overfitting to a particular oxide-electrolyte system.

From the Poisson-Boltzmann equation, an analytical expression for the surface charge density (σ) to surface potential (ϕ_s) relationship can be obtained, often termed the Grahame equation (Supplementary Information 1). Rearranging the Grahame equation for surface potential provides the following expression, shown assuming a symmetric valency electrolyte⁵⁶:

$$\phi_{s,pb} = \frac{2k_b T}{qz} \operatorname{arcsinh} \left(\frac{\sigma}{\sqrt{8c_\infty \epsilon_r \epsilon_0 k_b T}} \right), \quad (1)$$

where ϕ_s is the electric potential at the surface, q is the elementary charge, ϵ_0 is the permittivity of free space, ϵ_r is the relative permittivity of the medium, z is the valence of the electrolyte ion i , c_∞ is the bulk concentration of electrolyte ion i (expressed as a number density, units of m^{-3}), k_b is the Boltzmann constant and T is the temperature.

Assuming ϕ is small ($|\frac{qz\phi(r)}{k_b T}| \ll 1$), the Poisson-Boltzmann equation can be linearised and rearranged resulting in Equation 2, the Debye-Hückel model⁵⁷⁻⁶⁰ surface-charge potential relationship⁵⁶:

$$\phi_{s,dh} = \frac{\sigma}{\epsilon_0 \epsilon_r \kappa}, \quad (2)$$

in which the reciprocal of κ is the Debye length which, for a symmetric 1:1 monovalent electrolyte is equal to:

$$\frac{1}{\kappa} = \sqrt{\frac{\epsilon_0 \epsilon_r k_b T}{2q^2 c_\infty}}. \quad (3)$$

This approximation of assuming ϕ is small is termed the Debye-Hückel approximation. Despite this approximation not being strictly valid in most situations of interest in colloid science and electrochemistry⁵⁷, it is commonly used. One reason for this is its convenience, whereby electrostatic screening by electrolyte can be described in a simple parameter - the Debye length, which is inversely proportional to the square root of the ionic strength.

The Poisson-Boltzmann equation neglects finite size effects, and thus provides inaccurate results for highly concentrated systems. One way of dealing with this is the incorporation of a

parallel plate-like layer at the surface which represents accumulated charge, called the ‘Stern’ layer^{48,61}. In reality, the region modelled by the Stern layer contains both highly polarised water molecules and accumulated electrolyte ions⁶¹. An alternative method which does not require empirical fitting of a Stern layer capacitance is the ‘modified’ Poisson-Boltzmann equation, as shown in Equation 4 for a $z:z$ symmetric electrolyte⁶². The modified Poisson-Boltzmann equation simply constrains the maximum density of counterions permissible at the surface to an (semi-)empirical finite value based on packing constraints^{62,63}. The corresponding surface charge to surface potential relationship, which can be solved numerically for $\phi_{s,mpb}$, is:

$$\sigma = \frac{\operatorname{sgn}(\phi_s) 2zqc_\infty}{\kappa} \sqrt{\frac{2}{v} \ln \left[1 + 2v \sinh^2 \left(\frac{zq\phi_{s,mpb}}{2k_b T} \right) \right]} \quad (4)$$

where $v = 2a^3 c_\infty$ is a dimensionless measure of the non-diluteness by which a represents the mean spacing of ions at their maximum possible concentration ($a = c_{\max}^{-1/3}$). The phenomenological parameter a could be taken to be the radius of an ion (e.g. 1.84 Å hydrodynamic radius of a sodium ion⁶⁴) but considering at high electric fields (i.e. high surface charges) there is a well-known decrease in permittivity of the liquid at the interface⁶⁰, ion-ion correlation effects could extend the value to as high as 10 nm^{62,63}.

2.1.1 Mean-Field Models: Biomolecular Response

As a result of electrostatic screening, potentiometric sensors show a decreased signal with both increased distance of the biomolecule⁶⁵ from the surface, and increased ionic strength^{8,66}. To quantify this, it is common in the field of biomolecular potentiometric sensing to use the Debye-Hückel model (Equation 2) to estimate the characteristic length after which electrostatic interactions are significantly weakened, termed the ‘Debye length’. At a distance of several Debye lengths from the biomolecular charge the charge from the biomolecule is expected to be highly screened and thus have little effect on the sensor. The simplicity of an analytical expression provides a useful and rapid tool for evaluation of the importance of ionic screening, however, the justifiability of using the Debye-Hückel model for highly charged polyelectrolytes such as DNA is questionable, due to the assumption of low surface potential and neglect of steric interactions.

An alternative which is more justifiable at high surface potentials is the Poisson-Boltzmann equation without linearisation (Grahame equation, Equation 1). The Grahame equation has commonly been used to calculate the surface potential due to binding of charged biomolecular analyte such as DNA or, conversely, to calculate the charge due to bound biomolecules, given a measured surface potential^{65,67-69}. For example, the change in surface potential can be calculated before and after analyte binding, assuming that the intrinsic (i.e. silanolate, in the present work) surface charge (σ_{surf}) is additive with the charge contribution from DNA (σ_{DNA}). In other words, the DNA is assumed to be an infinitely thin smeared layer of uniform charge density on the surface which does not change the intrinsic charge, i.e. the biomolecule does not affect the charging of silanolate groups:

$$\Delta\phi = \phi(\sigma_{\text{surf}} + \sigma_{\text{DNA}}) - \phi(\sigma_{\text{surf}}) \quad (5)$$



The calculated shift in surface potential is strongly influenced by the choice of intrinsic surface charge density, and often this information is not measured but estimated, providing a large source of error in such calculations. Such calculations also often result in signals too weak compared to experimental response and thus the theoretical basis for the observed biomolecular signals remains unclear⁷⁰. This has provided motivation for the creation of improved models⁶⁸ and the present work.

2.2 Site Binding Models

The Nikolsky-Eisenman equation and ion exchange theory is widely used to describe glass-membrane electrodes and ion-sensitive electrodes⁷¹, however this approach is limited to estimating the selectivity of the sensor against competing ions⁷². In the field of IS-FET sensing research, site-binding models are instead commonly used to describe the surface potential as a function of pH^{73–75}, in which the surface charge is calculated based on several chemical equations for reactions of hydroxyl groups at the surface combined with a model of the electrical double layer. Potentiometric ion sensors modelling has been extensively reviewed by Bermejo⁷⁶ and Bobacka et al.⁷¹.

In the site-binding model^{73,77}, for an oxide with a 'sensitivity factor' of 1, the change in potential due to change in pH is the same as that predicted from the Nernst equation for a semi-permeable membrane with a change in proton activity (i.e. ~ 59 mV/pH at room temperature). For many oxides, including silica, a potentiometric response weaker than 59 mV/pH is often measured and in the site-binding model this is explained via a reduced value of the sensitivity factor. The sensitivity factor is a function of both the ability of the oxide surface to deliver or take up protons ('surface buffer capacity') and the differential double-layer capacitance, with this capacitance being primarily determined by the ion concentration of the bulk solution^{73,77}.

Even though site-binding models have proved useful in IS-FET design and will likely continue to do so, they present several fundamental limitations. Firstly, they do not provide any information on the effect of surface morphology upon response and, secondly, they rely upon empirical parametrisation of many properties. These properties can vary greatly even for a single material, for example, there is debate over the correct acid-base dissociation constant value for silica⁷⁸ and the density of surface sites is preparation dependent⁷⁹ so there is a risk of the model being overfit, and used descriptively rather than predictively. Finally, site-binding models are not suitable for prediction of sensor response to binding of biomolecular analyte. In the present work, we propose molecular dynamics as a complementary approach to existing models to explore response to surface morphology and biomolecular binding.

2.3 Silica Surface Chemistry

Silanol groups (Si–OH) at the silica surface react with H^+/OH^- in water to form a charged surface layer⁸⁰. These chemical reactions are evidenced to be the primary surface charging mechanism for hydrated silica⁵⁸, with electrolyte effects having a measurable but less significant effect⁸¹. Ionic strength can be import-

ant, for example, for ultrapure water (< 0.001 M), negligible surface ionisation can occur due to the lack of availability of charge stabilising cations⁸². In the present work, the ionic strength of the simulated system was 300 mM. This ionic strength is relevant to physiological conditions (e.g. high ionic strength biosensing experiments), biomimetic synthesis⁸³, or applications such as ocean geochemistry (salt water = ~ 0.7 M⁸⁴).

The precise extent of surface ionisation depends on the pH, silanol density, type of silanol group (e.g. isolated, vicinal or occluded in pores), the ionic strength of the solution and the type of cations and anions⁸⁵. At pH values relevant to most biosensing conditions (pH 6–9), silica is negatively charged and only gathers a positive charge at extremely acidic conditions ($< \text{pH } 2$)^{86,87}, as evidenced by the measured point-of-zero charge (i.e. the pH for which the net charge of the surface is zero) of between pH 2–4⁸⁸.

2.4 Nano-morphology and Potentiometric Sensor Response

When comparing two systems with identical hydroxyl density and composition, commonly-used site-binding models predict no effect on sensor response from changes in surface morphology. Nonetheless, some experiments suggest amorphous surfaces are beneficial to pH sensor response: for example, sputtered silica was shown to have a higher pH sensitivity than thermally grown silica⁸⁹, addition of nanowires to a planar Al_2O_3/SiO_2 surface increased pH sensitivity by 5 mV/pH (55 mV/pH to 60 mV/pH)⁹⁰ and texturing a silanised silica surface with silanised silica nanoparticles increased pH sensitivity by 11 mV/pH (43 mV/pH to 54 mV/pH)^{91,92}. However, other experiments have shown no effect of increased surface porosity on the pH sensitivity, but did show an increase in capacitance^{93,94}. It is clear that further work is needed to clarify the precise relation between surface morphology and pH sensitivity. Molecular dynamics simulations provide the unique ability to directly investigate the effect of nanoscale changes in morphology on the electrostatics at the interface. In the present work, novel molecular dynamics simulations are used to investigate this relationship.

2.5 Experimental Potentiometric Sensing of DNA

Experimental DNA sensing experiments often involve first immobilisation of single stranded ('probe') DNA followed by hybridisation with a complementary strand. In the present work, the absence of DNA versus the presence of DNA is simulated, which is analogous to the sum of hybridisation and immobilisation signals. Experimental field-effect biosensor DNA hybridisation signals were reviewed by Poghossian et al., ranging from no significant response to ~ 120 mV response⁷⁰. Nishio et al. reviewed immobilisation signals to be 32–100 mV with hybridisation signals generally being weaker, between 11–14 mV⁵⁵. Such large variation between experiments is due to variation in factors such as probe density, buffers composition, reference electrode setup, DNA sequence and surface chemistry. This variation makes rational design difficult and is part of the motivation for the present work which can systematically explore changes in the electric double layer due to biomolecules.

At high ionic strength, the signal from the DNA is commonly



expected to be reduced due to screening. However, DNA has been demonstrated to produce signals with a magnitude of several millivolts even in high ionic strength solutions of 0.5 M or 1 M^{95,96}. This suggests that the 0.3 M ionic strength used in the present work should generate a detectable signal providing there is sufficient bound DNA. The density of surface bound DNA molecules used in the present work (8.58×10^{12} molecules cm^{-2}) is likely higher than typical densities but not unreasonable given that density of surface bound DNA molecules ('probe density') as high as 1×10^{13} molecules cm^{-2} has been measured by Surface Plasmon Resonance measurement⁹⁷.

3 Computational Methods

3.1 Model Setup

This section is divided into the setup of the silica surface models (Section 3.1.1), the surface charge and electrolyte setup (Section 3.1.2) and finally the DNA molecule setup (Section 3.1.3). Analytical model results were calculated at 298.15 K assuming a relative permittivity of 78.5, an ionic strength of 0.3 M and a 1:1 monovalent electrolyte.

3.1.1 Silica Model Setup

A crystalline silica model was used with Q^3 isolated silanol terminated surface, derived from the (10 $\bar{1}$) cleavage plane of α -cristobalite, with a silanol density of 4.804 OH nm^{-2} in its fully protonated state. The silanol density of $\sim 5 \text{ OH nm}^{-2}$ has been used by other authors successfully to reproduce experimental IS-FET data⁹⁸, is often used in IS-FET modelling⁷³ and is expected to be the density of fully hydrated silica as discussed in Supplementary Information 2.

A different silica model was also used to emulate an amorphous silica surface. The model consists of a periodic unit cell of silica with little internal ordering and a similar silanol density to the crystalline model, of 5.09 OH nm^{-2} . This model has increased surface area compared to the crystalline surface and has silanol groups distributed over a greater distance from the surface. A side-by-side comparison of the two models can be seen in Figures 1(a) and (b).

Both silica models originated from the INTERFACE model database⁸⁵. Details such as cell dimensions are summarised in Table 1. In both models, atoms within $\sim 11 \text{ \AA}$ of the base were rigidly constrained for all simulations to emulate a silica bulk. Comprehensive details of the silica model setup can be found in Supplementary Information 3.

3.1.2 Surface Charge and Solvent Box Setup

With increasing pH, surface charge density increases and the surface potential becomes more negative. Potentiometric titration experiments can be used to obtain the relationship between surface charge density and surface potential. This relationship is non-linear, but to a first order approximation is treated as a linear relationship, and in the present work, the approximate empirical relationship presented by Emami et al.^{85,99} was used in order to compare simulated systems of a given surface charge density to the corresponding 'effective pH'. Specifically, $0.024 \text{ Cm}^{-2} \text{ pH}^{-1}$ was used, which corresponds to measurements for silica sur-

faces terminated by Q^3 isolated silanol groups, with a density of $\sim 4.7 \text{ OH nm}^{-2}$ at 0.1 – 0.3 M ionic strength conditions^{85,99}.

Various silica models with differing extents of surface charge were prepared by deprotonating the top-surface silanol groups. For each system, neutralising Na^+ counterions were added to maintain charge neutrality. This procedure is analogous to addition of NaOH to a neutral silica surface in an experimental setup, where negatively charged silanolate groups would form at the surface due to reaction with hydroxide ions and there would be a simultaneous increase in bulk Na^+ concentration.

Once electroneutrality was obtained, NaCl was added to set the bulk ionic strength. An approximately 300 mM ionic strength of NaCl was used based on the initial volume of the water box, with the specific number of ions used and cell size shown in Table 1. The details of the solvent preparation are presented in Supplementary Information 3. In brief, two different initial configurations of counterions were considered in order to test convergence to thermodynamic equilibrium. A TIP3P solvent box with an initial height of 73 \AA was used with ions placed randomly. A control system at low ionic strength was also investigated as detailed in Supplementary Information 4.

3.1.3 DNA Model Setup

The initial canonical DNA dodecamer (5'-GGGGGGGGGGG-3') structure was generated and placed in a solvent box as described in Supplementary Information 3. A water box with an initial height of 100 \AA was used, and initialised with its helical axis normal to the surface of the crystalline silica model with surface charge density -0.083 Cm^{-2} . CHARMM36 forcefield parameters¹⁰⁰ were used with phosphate groups added to the terminal groups resulting in a DNA molecule with a net charge of $-24q$ which was neutralised with sodium ions. Lavery et al. analysed the amount of time needed for equilibration of the ionic atmosphere around DNA in bulk electrolyte and concluded at least 300 ns needed⁴⁵. Consequently, a 320 ns equilibration was performed, followed by 180 ns with harmonic constraints applied to the DNA atoms. Harmonic constraints ($0.5 \text{ kcal mol}^{-1} \text{ \AA}^{-2}$) were applied so the DNA remained surface-bound, and to remove any significant effect of biomolecule dynamics. These constraints were then removed, allowing free diffusion of the DNA, and simulated for a further 180 ns of dynamics. Snapshots of the constrained and unconstrained systems are shown in Supplementary Information 5. As described for the bare silica-electrolyte systems, two repeats were performed with different initial electrolyte configurations.

3.2 Simulation Parameters

Mean-field (analytical model) calculations in the present work were performed at 298.15 K assuming a relative permittivity for the medium of 78.5.

For molecular dynamics simulations, the INTERFACE forcefield was used. The forcefield provides CHARMM forcefield compatible parameters for silica and sodium ions¹⁰¹. Unlike most forcefields of silica-water interfaces, the INTERFACE forcefield has parameters to treat charged silica-water interface and had been shown capable of accurately reproducing a broad range of ex-



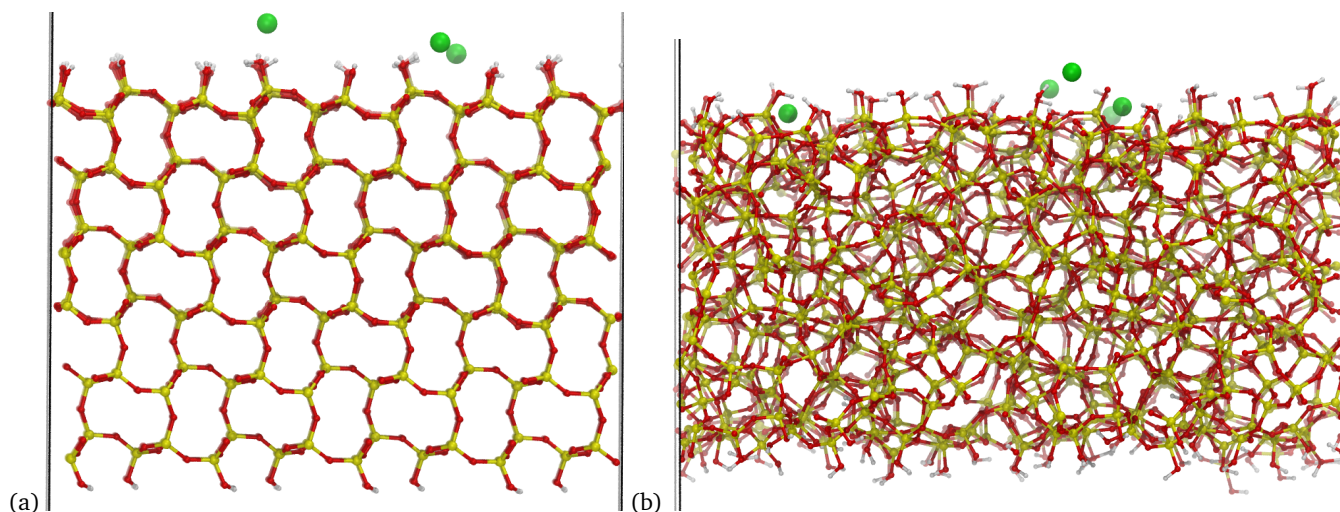


Fig. 1 Periodic slabs used to model the silica surface in the present work. Green, red, yellow and grey atoms are sodium, oxygen, silicon and hydrogen respectively: (a) side-on view of crystalline silica surface, (b) side-on view of amorphous silica surface. Both surfaces are analogous in that they contain approximately the same surface silanol density and surface charge density and are only terminated by Q^3 silanol groups. These images were taken from trajectory of the -0.083 C m^{-2} models, with the sodium ions introduced to maintain charge neutrality. For the crystalline system, the negative silanolate charges are all located randomly within the first atomic layer, whereas for the amorphous system, charges were distributed randomly between the first atomic layer and 5 \AA below.

Model	ρ (pH_{eff})	SiO^-	Cl^-	Na^+	Model	ρ (pH_{eff})	SiO^-	Cl^-	Na^+
	0 (~3.0)	0	15	0+15		0 (~3.0)	0	22	0+22
Crystalline	-0.041 (~4.7)	3	15	3+15	Amorphous	-0.048 (~5.0)	5	22	5+22
4.81 OH nm^{-2}	-0.083 (~6.4)	6	15	6+15	5.09 OH nm^{-2}	-0.086 (~6.6)	9	22	9+22
$33.5 \text{ \AA} \times 34.9 \text{ \AA}$	-0.12 (~8.1)	9	15	9+15	$40.3 \text{ \AA} \times 41.4 \text{ \AA}$	-0.12 (~8.2)	13	22	13+22
~300 mM NaCl	-0.17 (~9.9)	12	15	12+15	~300 mM NaCl	-0.16 (~9.8)	17	22	17+22
	-0.39 (~19)	28	15	28+15		-0.38 (~19)	40	22	40+22

Table 1 Simulated System Summary showing the structure (crystalline/amorphous), surface charge density (ρ) in units of C m^{-2} and number of silanolate, chloride and sodium ions respectively. pH_{eff} is shown in parenthesis, and is the approximate empirical pH for this surface charge density as presented in the methods Section 3.1.2. A separate simulation was also performed in which DNA was added to the crystalline 4.81 OH nm^{-2} simulation at 300 mM NaCl, with 24 additional Na^+ to neutralise the negatively charged DNA molecule. The crystalline system also had a repeat simulation at 0 mM bulk ionic strength as presented in Supplementary Information 4.



perimental observables such as water contact angle, adsorption energy of peptides (at a charged surface), water adsorption isotherm, immersion energy in water and the cell parameters of quartz^{85,101,102}. The forcefield likely represents the state-of-the-art for accurate classical molecular dynamics simulation of the electrified silica-water(-biomolecule) interface. A discussion of the forcefield parameters and validation is found in the Supplementary Information 6.

Details of the molecular dynamics software parameters can be found in our previous work⁴⁶ and in Supplementary Information 7. In brief, a 2 fs timestep was used with NVT Langevin dynamics at 298.15 K with the SETTLE algorithm for all hydrogen atoms¹⁰³. A minimum of 320 ns of dynamics was performed for each simulation, with analysis over the last 180 ns. A simple harmonic restraint was applied to water molecules which evaporated to return them to the bulk. The electrostatics were evaluated using particle-mesh Ewald (PME) method, using the EW3DC correction which provides enhanced accuracy for polarised systems^{43,46,47}.

3.3 Analysis Methodology

The analysis methodology is detailed in Supplementary Information 7 and discussed in our previous work⁴⁶. In summary, the charge density was calculated by averaging the charge density into xy slabs along the z -axis, thus reducing the system to a one-dimensional grid. The electric potential is calculated by double integration of the charge density from the Poisson's Equation^{43,47}, with integration performed using the trapezium rule and the electric potential and field set to zero at $z = 0$ Å^{43,104}. In order to calculate a surface potential the surface was defined as the z position of minimum electric potential in the region where there is a large potential drop due to silanolate charge; the surface location is indicated as a vertical solid black line in Figure 2 with the dashed lines indicating the maximum and minimum z -position. The surface potential was then calculated as the difference between the mean potential in the bulk water (between $z = 80 - 85$ Å) and at the surface. For potential calculations the analysis was performed over the last 180 ns of the trajectory in three 60 ns parts; the standard error of the mean (ddof=1) was calculated based on the mean of these three parts. For the plot showing electric field, the field was calculated every 100 ps over the last 180 ns (1800 frames) for both repeats of each simulation. The resulting data for the two repeat simulations was combined (3600 frames) and the mean electric field plotted, with the 95% confidence intervals shown, using 1000 bootstrap intervals. For the water polarisation plot, the z -component of the water polarisation was calculated as a function of water molecule orientation multiplied by water number density.

4 Results and Discussion

The results are split into five subsections. Firstly, in Section 4.1 the charge structure of the electric double layer is analysed. In the following sections, metrics directly related to the response of many potentiometric sensors are calculated and discussed. Specifically, Section 4.2 presents the electric field below the silica

substrate, with the aim of emulating the 'field effect' which controls the response of field-effect sensors. The change in surface potential due to changes in surface charge density/'effective pH' is presented in Section 4.3, and in Section 4.4 the change in surface potential due to addition of DNA is presented. Finally, in Section 4.5, the effect of molecular dynamics simulation duration on the accuracy and precision of calculated electrostatic properties for biosensing applications is presented.

4.1 Charge Density Profile

In Figure 2, the distribution of charge as a function of z is shown. The crystalline system, shown in subfigures (a), (b) and (c), is compared side-by-side with the amorphous system, shown in subfigures (d), (e) and (f). The z position defined as the surface for surface potential calculations is indicated by a vertical black line, and the coloured curves are drawn with increasing darkness corresponding to increased surface charge density, as shown in the figure legend. The charge is shown as integrated charge density of the system from $z = 0$ and therefore shows the total charge density in the unit cell up to a given z position (i.e. in the bulk the value is zero as the system is net electroneutral). In subfigures (a) and (d), the total charge density is presented, but in subfigures (b) and (e) only the contribution from sodium ions is presented. In subfigures (c) and (f), only the contribution from water charges is presented (i.e. the negative oxygen and positive hydrogen atoms of the TIP3P water model).

The position of the dashed vertical black lines in Figure 2 show that the silanols were distributed over a broader range of z positions in the amorphous structure than in the crystalline structure. An important result of this, combined with the effect of irregular morphology of the amorphous system, is that the amorphous model charge distribution, shown in Figure 2(d), becomes less structured compared to the charge distribution of the crystalline system, as shown in Figure 2(a). The sodium ion distribution was similar between crystalline and amorphous systems (Figure 2(b) versus (e)). Comparison of Figure 2(c) and (f) reveals that the water in the crystalline model had three discrete layers, in contrast to the amorphous system in which the layers were less distinguishable, showing broader water density peaks, with less structure. The formation of three discrete layers on crystalline glass surfaces is supported by *ab initio* molecular dynamics studies¹⁰⁵.

4.1.1 Stern Layer

The Stern layer is important in determining the surface potential for high charge density systems, and so, even though the Stern layer is only strictly defined in the Gouy-Chapmann-Stern model, in this paper, the interfacial region up to the second minima in the water density profile (2 layers of water) was used to define a Stern-like layer and analysed in more detail.

In Figure 3(a) the density of sodium ions within the Stern-like layer increased approximately linearly with increased surface charge density for both amorphous and crystalline silica models. This is in agreement with the molecular dynamics simulation of Lee et al. which showed similar behaviour at high surface charge densities, upon an ideal structureless surface¹⁰⁶ and is a result of increased electrostatic attraction between the negatively



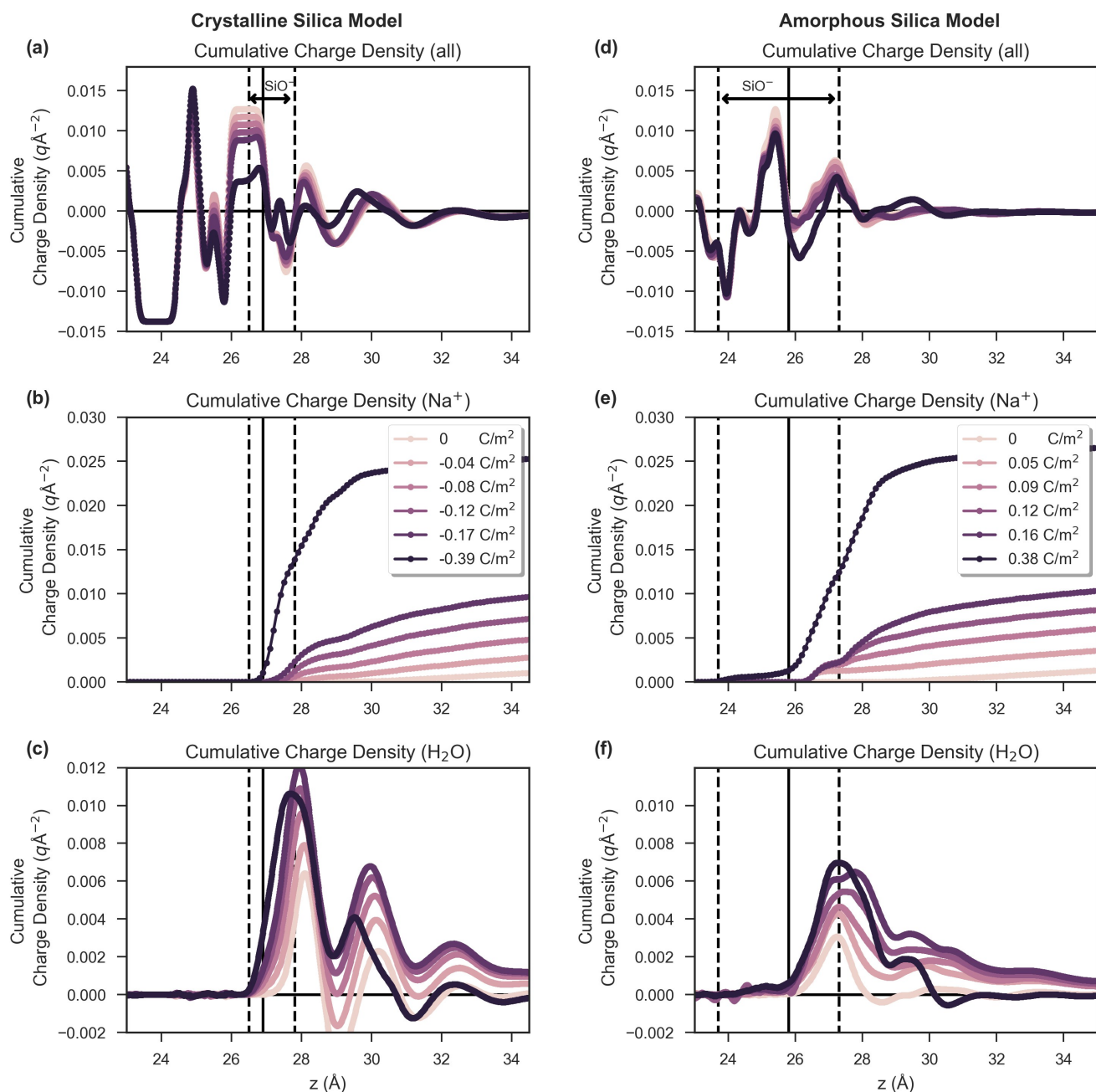


Fig. 2 The integrated charge density of the system as a function of z position for increasing surface charge density (from light to dark lines) with the crystalline silica model shown on the left (a,b,c) and the amorphous silica model shown on the right (d,e,f). Subfigures are shown for the combined charge density of all atoms (a, d), only the sodium ions only (b,e) and only the oxygen and hydrogen atoms in water molecules (c, f). For each plot, the integration was performed from the silica (at $z = 0 \text{ \AA}$) to the bulk water. For the crystalline model, the silanol density showed two peaks, the largest at $27.0 \pm 0.1 \text{ \AA}$ and a smaller peak at $27.5 \pm 0.1 \text{ \AA}$. For the amorphous system the silanol density showed silanols distributed between $22.0 \pm 0.1 \text{ \AA}$ and $28.4 \pm 0.1 \text{ \AA}$ with the highest density at $25.6 \pm 0.1 \text{ \AA}$. Shown on the figure by solid black vertical lines is the surface definition described in the methods section. Dashed vertical black lines indicate the minimum and maximum z -position at which charged silanolate groups were present. Both (b) and (e) show that with increased surface charge density there was an increase in sodium ion accumulation at the surface. Comparing (a) to (d), it can be seen that the amorphous structure resulted in a more diffuse charge distribution than the crystalline system primarily due to less structured water layers, evident by comparing (c) to (f).



charged surface and positively charged sodium ions. The sodium ion density for the amorphous system was higher than the crystalline system, which is explained by the increased surface area and amorphous morphology providing cavities with improved favourability for sodium binding. The surface accessible surface area (1.4 Å solvent probe radius) was 53% larger for the amorphous model compared to the crystalline model (details in Supplementary Information 3). Comparison between amorphous and crystalline charge density and water polarisation profiles will later be discussed.

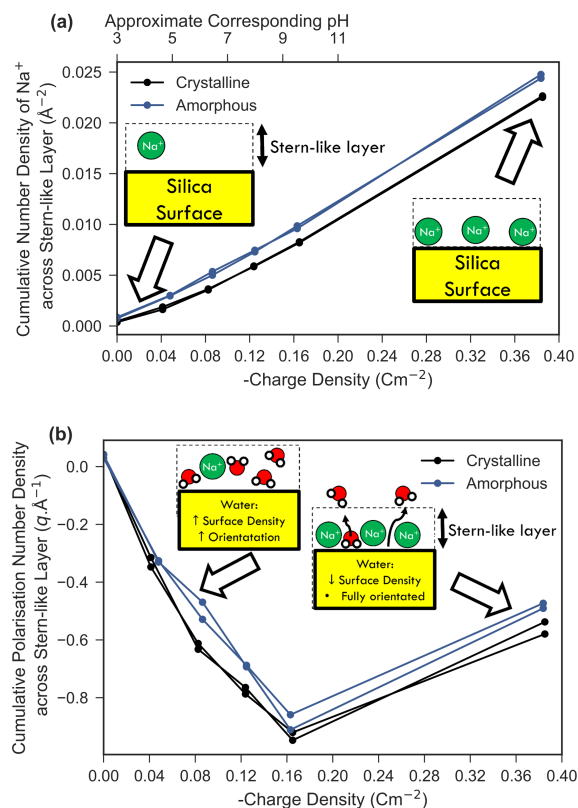


Fig. 3 Comparison of both the cumulative water polarisation (a) and sodium ion number density (b) across the Stern-like layer near the surface, as a function of the surface charge density. Schematic insets are included to show qualitative difference between low and high surface charge systems. Both the crystalline system (black) and the amorphous system (blue) are shown, for each surface charge density the result of the repeat simulation is also shown. The cumulative number density was calculated as the integral of the number density across the Stern-like layer similar to the work of Lee et al.¹⁰⁶. The approximate 'effective pH' (see methods) is shown on the top-most x-axis of subfigure (a). The cumulative polarisation density was calculated by integrating the polarisation across the Stern-like layer as described in the methods section.

In Figure 3(b) it can be seen that net polarisation of water within the Stern-like layer increased approximately linearly with charge density up to -0.17 Cm^{-2} for both the amorphous and crystalline model systems. The polarisation is a measure of both the net orientation of waters in the z direction but also their density, and thus is proportional to the electric-field imparted by the molecules and their effect on surface potential.

Figure 3 includes one result at very high surface charge density

which is not relevant to typical experimental conditions of silica-water interfaces as a surface charge density above -0.17 Cm^{-2} would require a highly alkaline medium ($> \text{pH } 11$)^{85,99} to form. Furthermore, experiments show that highly alkaline conditions ($> \text{pH } 9$) result in significant dissolution of silica¹⁴ which remains a challenging task to simulate even using 'reactive' molecular dynamics forcefields³⁸. Nonetheless, simulation of high surface charge densities is of interest both for exploring the limiting case and because it is attainable by other oxide systems. For example, δMnO_2 demonstrates a surface charge of -1 Cm^{-2} at pH 8 despite having similar point-of-zero charge to silica¹⁰⁷. At a surface charge density of -0.38 Cm^{-2} the linear trend in Figure 3(b) ceased and there was instead a relative decrease in water polarisation. This decrease was not a result of decreased orientational ordering (as evidenced in Supplementary Information 8), but rather due to a decrease in the density of bound water due to displacement by the particularly high density of sodium ions bound at these extreme surface charges as depicted in the schematic inset of Figure 3(b).

4.2 Electric Field and the Field Effect

Field-effect sensors measure the surface potential change (or, more precisely, a threshold voltage shift) due to binding of analyte molecules or changes in pH. The underlying physical mechanism is that the long-range electric field ('field effect') from the analyte-bound surface extends through the oxide, and thus modulates the carrier concentration in the semiconducting layer below the analyte-binding surface, resulting in the change in threshold voltage. To emulate this property via simulation, the electric field at a distance from the oxide interface was calculated via the force on a positive test charge centred below the oxide slab.

Prediction of biomolecular potentiometric response is a key motivation for the present work and thus the electric-field response due to DNA specifically was investigated. DNA sensing experiments often use an oxide surface which has been chemically functionalised with a layer of a material such as (3-Aminopropyl)triethoxysilane (APTES). This layer is chemically bonded to single stranded DNA to provide a highly selective template for binding of specific single stranded DNA via a hybridisation reaction. In the present work, we are primarily interested in evaluating whether the simulation methodology can provide potentiometric-response prediction for charged biomolecules in general and so we modelled DNA on a bare silica surface to eliminate additional complexity and to provide a more general model system given that many different surface functionalisation layers are used in the literature. The CHARMM force field used in this work is well-suited to incorporation of organic, solvated molecules like APTES, and has been used to model APTES in the literature¹⁰⁸.

DNA was added to the crystalline silica system with an intrinsic silanolate surface charge density of -0.083 Cm^{-2} and constrained near the surface and the electric field response for this DNA system is shown in the blue bar of Figure 4. A shift in the electric field due to addition of DNA and due to increased negative surface charge density was observed. The direction of the shift is consist-



ent with an increase in negative charge at the surface, as expected for negatively charged DNA. The graph shows a response due to changes in effective pH of $0.007 \text{ Vnm}^{-1} \text{ pH}^{-1}$ (linear regression of data from green bars of Figure 4). A shift of 0.007 Vnm^{-1} was also observed for addition of DNA (blue bar), and given that a typical pH sensor can detect changes of at least 1 pH unit, the simulated change in electric field of $0.007 \text{ Vnm}^{-1} \text{ pH}^{-1}$ due to DNA is significant with regard to expected limit-of-detection of a sensor.

This response to DNA provides a proof-of-concept for this molecular dynamics simulation methodology as a novel way of investigating the field effect for biosensing applications at a molecular scale and can provide a platform for systemically exploring molecular scale effects which may be relevant to optimising FET-sensor response such as surface morphology, buffer composition, biomolecule structure and biomolecule dynamics.

4.2.1 Improvement on Previous Work

Our previous work⁴⁶ had some limitations compared to the present work, such as shorter timescales, a less accurate force-field and a less accurate method of evaluating electrostatics but in that work we also investigated the effect of DNA on the electric field. In that work, the DNA was added to crystalline silica surface model with -0.2 Cm^{-2} surface charge density with electrolyte of similar ionic strength (0.2 M) to the present work (but with mixed valency electrolyte: NaCl:MgCl_2) and a statistically significant change in electric field due to DNA addition was calculated. It was not possible to determine whether the signal was significant with regards to the resolution of detection of experimental measurement. As improvement over the previous work, in the present work the surface charge density/electric field relation was used to provide model calibration method and evidence that this DNA induced change in electric field would be large enough to be experimentally measurable.

The effect of distance of the DNA from the surface was also observed in the present work. In the red bar of Figure 4, the result from the system after the DNA was permitted to freely diffuse from its initial surface position (unconstrained) is shown as a red bar. The DNA moved greater than 1 nm away from the surface and the electric field returned to that which was indistinguishable from the control shown in green. The distance-dependent reduction in electric field was a result of screening from polarised water and ions in solution. Experiments also show a strong distance-dependent reduction of DNA signal⁶⁵. At 300 mM NaCl, the Debye length is 0.55 nm and therefore, based on the Debye length screening arguments (Section 2.1.1), the influence on the interface is expected to be negligible after the DNA reaches 1–3 nanometres distance, which is in agreement with the present molecular dynamics simulation result.

While the electric field was used for this analysis, there was a strong correlation between the total electrostatic energy of the system (double integration of the entire charge density) and the electric field below the silica-water interface, as shown in Supplementary Information 9. Given this correlation, it is possible that the total stored electrostatic energy may also be a viable metric of estimating sensor response from molecular dynamics simulations.

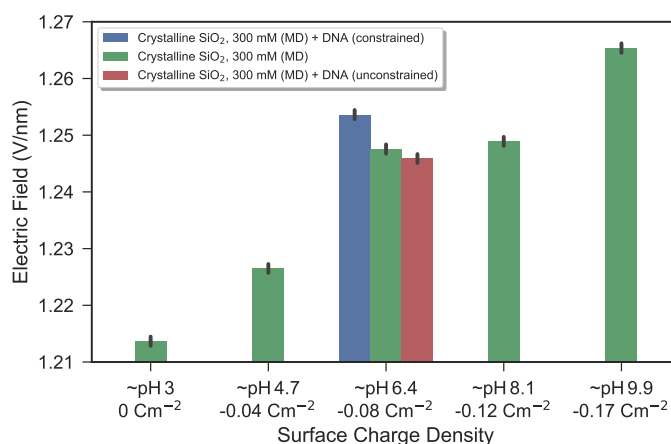


Fig. 4 Bar chart of the electric field as a function of surface charge density for the crystalline silica model. The electric field is calculated from the force on a test charge below the silica substrate with an increasingly positive value expected from a more negatively charged surface. The green bars show the electrolyte-only system, where each bar represents the combined data from two repeat simulations. The black bars show the bootstrapped 95% confidence interval as described in the methods. The blue and red bar show systems including DNA constrained near the surface (blue) and free to diffuse (red). From the blue bar, it can be seen that there is change in electric field due to the introduction of DNA which was constrained near the surface (within 1 nm of the surface). After the constraints were removed, the DNA diffused slightly away from the surface (lowest position of the DNA fluctuating between approximately 1–3 nm, from the surface), resulting in no significant change in the electric field compared to the control. To illustrate the difference between constrained and unconstrained systems, snapshots of the DNA system are shown in Supplementary information 5.



4.3 Surface Potential: Effect of pH

4.3.1 Mean-Field Models

The surface potential shift as a function of surface charge and effective pH are shown in Figure 5 for three mean-field models—the Grahame equation (Equation 1, solid line), the Debye-Hückel model (Equation 2, short dashed line), and the modified Poisson-Boltzmann model (Equation 4, circles for $a = 0.189$ nm and long dashed line for $a = 1$ nm). Monovalent 1:1 electrolyte is assumed (e.g. NaCl) with two ionic strengths shown: 300 mM (blue) corresponding to the same ionic strength as the molecular dynamics simulations performed, and a more dilute 100 mM (green) for comparison.

Figure 5 shows that the surface potential became increasingly negative with an increased surface charge density in all cases. A linear surface charge/surface potential relation occurs for the Debye-Hückel model due to the constant capacitance of this model whereby the Debye length is constant as a function of surface charge⁴⁸. In contrast, both the Poisson-Boltzmann and modified Poisson-Boltzmann models have a non-linear surface charge/surface potential relation^{48,106}. At low surface potentials, they provide similar predictions, however, at high surface potentials they differ because the modified Poisson-Boltzmann model prevents unphysically dense accumulation of ions at the surface, and thus has a lower differential capacitance⁶² or, equivalently, a steeper slope in Figure 5.

The a parameter for the modified Poisson-Boltzmann represents the maximum density of ions possible at the surface, with smaller values meaning higher maximal density. $a = 0.189$ nm is the hydrodynamic radius of a sodium ion⁶⁴ and $a = 1$ nm a plausible higher value considering contributions from solvent and ion-correlation effects resulting in more dispersed ions at the surface⁶². The parameter is being used here to provide a range of predictions from the modified Poisson-Boltzmann model, without resorting to empirical parametrisation. It can be seen that at these charge densities, if $a = 0.189$ nm then the results are equivalent to the Poisson-Boltzmann equation but that if $a = 1$ nm the modified Poisson-Boltzmann equation predicts a larger potential response. In all cases, the Debye-Hückel model predicts the largest response, but is technically invalid at potentials much greater than k_bT/q (~ 26 mV).

All three mean-field models show a strong dependence upon ionic strength, which is contradicted by experimental evidence.

4.3.2 Experimental Data

Figure 6 shows the change in surface potential as a function of surface charge density (effective pH) for experimental data (dashed lines). The amorphous silica molecular dynamics model (blue solid lines) and the crystalline silica molecular dynamics model (black solid lines) are also shown. Only surface charge densities below -0.12 Cm^{-2} are shown in the figure because anomalous surface potential behaviour at high surface charge densities has already been discussed in our previous work⁴⁶ and because such high surface charge densities are beyond the reach of most experiments^{16,85,99}. The surface potential simulated by molecular dynamics can only be approximate, due to the difficulty of defining a precise surface layer in atomistic simulations^{46,106}.

A comparison to experimental pH response data was discussed in our previous work⁴⁶. In summary, the experimental data is highly variable due to variation between silica surface preparation and electrolyte composition, however, molecular dynamics simulations and mean-field models show a qualitative agreement with some experimental data up to high effective pH⁴⁶. Silica often shows a sub-Nernstian response¹¹⁴ in the 30–40 mV/pH range^{8,46,112,114}, but ideal Nernst response (~ 59 mV/pH) has been reported⁸⁹ as shown in Figure 6 (pink circles). This untypically high pH response was likely due to differences in silica structure as the silica was sputtered as opposed to being thermally grown or naturally formed.

4.3.3 Molecular Dynamics Models

In the pH 3–8 range, the crystalline molecular dynamics silica simulation, shown in Figure 6, demonstrated approximately 25–45 mV per effective pH and the amorphous simulation showed approximately 59 mV per effective pH, this increased surface potential change with effective pH or surface charge can be explained by the amorphous system showing higher sodium ion accumulation, as shown in Figure 3(a) for a given surface charge. As discussed, this is likely due to increased surface area with the cavities of the amorphous surface providing more favourable surface adsorption sites. If sputtered silica demonstrates a higher nanoscale surface area, the results of the present work provide a potential theoretical explanation for why sputtered silica has evidenced enhanced pH response compared to thermally grown silica⁸⁹. As presented in the background, some experiments support the concept of nano-texturing resulting in increased pH response^{89–92}, but the relationship remains unclear with further work being required to clarify the precise relationship.

The amorphous system showed a decrease in the polarisation of water compared to the crystalline system (Figure 3(b)), due to the amorphous structure reducing the ability of water to form a highly ordered and oriented monolayer. This decrease in polarisation would be expected to cause a smaller magnitude surface potential, however, it is counteracted by increased sodium ion binding (Figure 3(a)), resulting in the observed larger magnitude of surface potential for the amorphous system versus the crystalline system.

4.3.4 Comparison between Mean-Field Models, Experiments and Molecular Dynamics

The modified Poisson Boltzmann equation predicts accumulation of cations in multilayers (i.e. extending away from the surface) due to increasing surface charge density, resulting in quadratic surface charge/surface potential dependence¹⁰⁶. In contrast, in molecular dynamics simulations, water polarisation can compensate a large component of the surface charge such that larger surface charge densities are required for multilayer cation formation. As a result, in the molecular dynamics simulation, the cations accumulated primarily within the first molecular layer (Stern-like layer), and the resulting potential response (Figure 6) showed a more linear surface charge/surface potential dependence than the modified Poisson-Boltzmann equation in Figure 5 in this charge density regime.



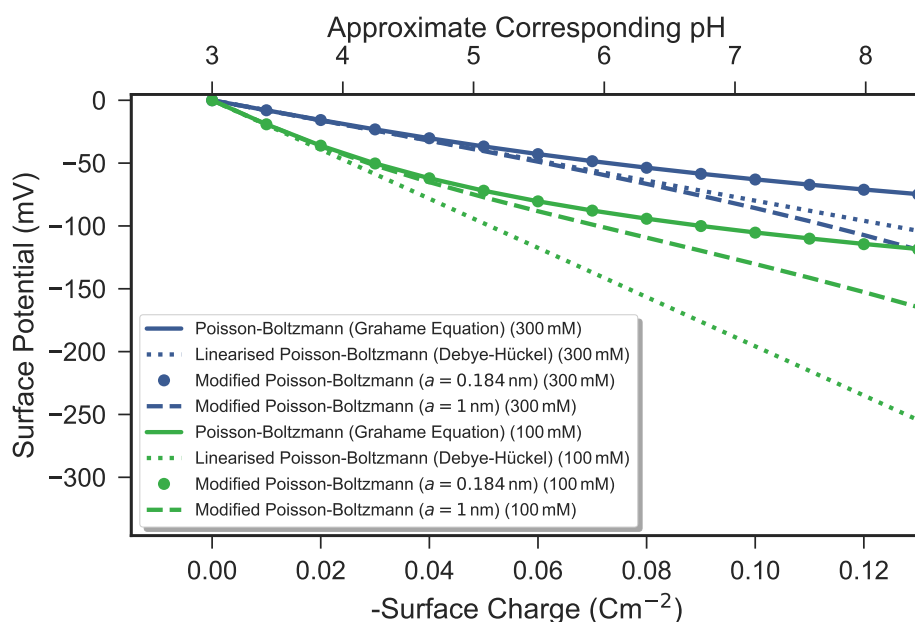


Fig. 5 Calculated surface potential as a function of surface charge from three mean-field models—the Grahame equation (Equation 1, solid line), the Debye-Hückel model (Equation 2, short dashed line), and the modified Poisson-Boltzmann model (Equation 4, circles for $a = 0.189$ nm and long dashed line for $a = 1$ nm). Two ionic strengths of monovalent 1:1 electrolyte are shown: 300 mM (blue) and 100 mM (green). Simulated data is plotted as a function of surface charge density, with the approximate linear relationship between surface charge density and pH shown as described in the methods section.

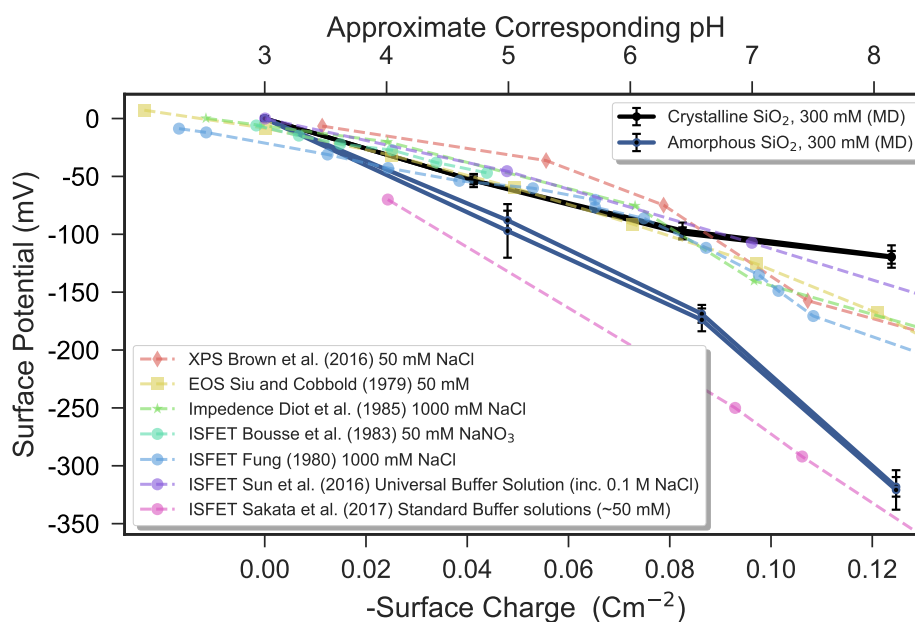


Fig. 6 Comparison of the surface potential simulated via molecular dynamics (solid lines) with experimental data (dashed lines), as a function of surface charge. The simulated surface potentials are presented relative to the system with no surface charge by subtraction of the surface potential for the zero surface-charge system. Literature data is displayed with the electrolyte used in the legend, and were measured by methods indicated by marker: \blacklozenge X-Ray Photoelectron Spectroscopy (XPS)¹⁰⁹, \blacksquare Electrolyte-on-insulator (EOS)⁸⁸, \blackstar Impedance¹¹⁰, \bullet IS-FET^{89,111–113}. Experimental data is plotted as a function of measured pH, whereas simulated data is plotted as a function of surface charge density, with both axis aligned using the approximate linear empirical relationship between surface charge density and pH described in the methods section. The high pH sensitivity of the data of Sakata et al. was obtained using a sputtered silica sample, as opposed to conventional thermally grown or native oxide, and for this data, the point-of-zero charge was not identified so only potential differences can be shown, with the data manually aligned to a point-of-zero charge of ~ 3 in agreement with typical experiments. Molecular dynamics (MD) simulation of the crystalline silica-electrolyte system (black lines) and amorphous silica-electrolyte system (blue lines) are shown. The error bars indicate the 95% confidence interval over three 60 ns repeats and quantify uncertainty in the mean due to temporal fluctuations, with two repeat simulations for each surface charge performed to quantify thermodynamic convergence.



Comparison of Figures 5 and 6 shows that the mean-field models at 300 mM ionic strength significantly underestimate the potential response of the two experiments shown at 1 M ionic strength^{110,111} and that predicted by the molecular dynamics models. The reason for this underestimation is in part due to ignoring the effect of water polarisation. The mean-field models use a constant permittivity for the liquid throughout the system, which is in contrast to the present molecular dynamics simulations which simulate the spatially-dependent polarisation of water. If a lower permittivity is used in the mean-field model, a larger potential results.

Another reason for the underestimation is due to the strong ionic-strength dependence observed in the mean-field models. Experiments show ionic strength to have weaker effect, with pH being the dominant determinant of response⁸¹. The strong ionic strength dependence compared to that observed experimentally is due to neglect of the acid-base chemical equilibria; for example, in ultrapure water (ionic strength less than 0.001 M), negligible surface ionisation can occur due to the lack of availability of charge stabilising cations⁸². Modelling such equilibria is often performed empirically via site-binding models as it remains too computationally intensive to simulate the effects of ionic strength of acid-base equilibria from first-principles⁸⁰. While site-binding models can provide accurate prediction of potentiometric pH response with sufficient parametrisation, they cannot provide molecular-scale insight such as the discussed effects of surface morphology.

4.4 Surface Potential: Effect of DNA

As presented in the background (Section 2.5) potentiometric sensor response to DNA binding is highly variable due to difficulties in controlling experimental conditions, and the theoretical basis for the magnitude of experimentally observed response remains an open research question. Thus in this section the effect of DNA on potentiometric response are investigated.

The shift in surface potential due to addition of DNA (i.e. combined immobilisation and hybridisation signal) was calculated to be -5 ± 12 mV in one simulation and -24 ± 8 in a repeat simulation (95% confidence intervals calculated as per Figure 6), with only the latter simulation being statistically significant (unpaired t-test $p = 0.48$ and 0.011 respectively). The high uncertainty of the potential calculations can be contrasted with the electric field results for the same simulations, the data for which was previously shown in Figure 4. The electric field calculations showed strong statistical significance between the absence and presence of DNA, with $p = 0.0005$ and $p = 0.0001$ respectively. As discussed in our previous work⁴⁶, surface potential calculations via molecular dynamics are highly sensitive to changes at the position selected as the 'surface' layer, whereas the long-range electric field calculations performed in this work are more reliable. The electric field calculations showed a response equivalent to one effective pH unit, which is consistent in magnitude with the approximately 25 mV surface potential shift observed due to DNA in one of the two DNA simulations.

For the present work, the simulated DNA-crystalline silica

model system has a DNA density of 8.58×10^{12} molecules cm^{-2} and 12 base pairs (24 negative charges per DNA) and thus a surface charge density of $\sigma_{\text{DNA}} = -0.33 \text{ Cm}^{-2}$. The silanolate surface charge density was $\sigma_{\text{surf}} = -0.083 \text{ Cm}^{-2}$. Equation 5 can be used with the Grahame equation (Equation 1) to predict a shift in surface potential of 77 mV. Thus the Grahame equation predicts a shift in surface potential greater than that predicted by the molecular dynamics simulation of 5 – 24 mV, however, given the variation in experimental DNA sensing data, the quantitative accuracy of the molecular dynamics and mean-field models are currently difficult to evaluate.

Further complicating the issue, the predictions between different mean-field models are highly variable. For example, while response calculated using Equation 5 is often presented using the Grahame equation in the literature, it is instead possible to use Equation 5 with either the Debye-Hückel model or modified Poisson-Boltzmann equation. In such cases, different predictions are obtained; specifically -79 mV and -756 mV for the modified Poisson-Boltzmann equation with $a = 0.184$ nm and 1 nm respectively; and -264 mV for the Debye-Hückel model. Supplementary Information 1 presents visualisations of the differences in predictions of each model. As these equations have broad utility to the biosensing community, easy-to-use open-source code is also provided to facilitate members of the biosensing community to easily investigate the effects of changing surface charge (intrinsic or biomolecule), ionic strength and electrolyte composition for this set of three equations.

Future work will further validate the model by investigation of biomolecular systems which show more reproducible experimentally measured surface potential shifts such as polystyrenesulfonate (PAS) and polyallylamine hydrochloride (PAH) which form a polyelectrolyte multilayer^{67,69}.

4.5 Effect of Simulation Duration and Convergence

Large fluctuations in potential and electric field occur at the 1 ps time scale, for example, as discussed, a change in pH was shown to correspond to approximately 0.007 V nm^{-1} which can be compared with a typical standard deviation for the electric field of 0.025 V nm^{-1} †. Calculation of potentiometric (bio-)sensor response therefore requires the ability to distinguish a small change in a noisy signal; this problem is somewhat alleviated in experimental systems by measurement equipment averaging over macroscopic timescales (microseconds to seconds). Potential and electric field calculations are shown as a function of simulation duration in Supplementary Information 10. The results show the duration used in this work can provide precise calculation of mean electrostatic properties and evidences thermodynamic convergence thereby demonstrating the merits of the longer duration of simulations in the present work compared to other reported calculations of the silica-water interfacial potential^{38–41,43,44}.

† Calculated over 180 ns, with snapshots taken at 100 ps intervals for the -0.041 Cm^{-2} crystalline silica model



5 Conclusions

Understanding the electrostatics of the silica-water interface is vital to many modern technologies such as electrochemical fuel cells¹, water filtration³ and biosensing⁷⁷. In particular, the electrostatic potential and electric field are particularly relevant to understanding the response of potentiometric biosensors such as field-effect transistor-based sensors. With a particular focus on understanding the field-effect biosensors, in the present work, molecular dynamics simulations of the electrostatic properties at the silica-electrolyte-biomolecule interface have been presented. The effect of varying surface charge density and the addition of a highly charged model biomolecule (DNA) was investigated and the results compared against three commonly used analytical mean-field models of the electrical double layer. Both a crystalline and an amorphous silica surface model were investigated, and their differences compared and contrasted. Molecular dynamics simulations were performed for significantly longer duration than related literature, facilitating increased precision and accuracy of the electrostatic properties.

The first few molecular layers at the surface dominate the electrostatic properties of highly charged interfaces and therefore a layer close to the surface was analysed in more detail, referred to as the ‘Stern-like’ layer within the present work. Water formed three highly ordered surface layers in the crystalline silica model, in contrast to the amorphous model which showed much less water structuring. Sodium ion density in the Stern-like layer was found to increase approximately linearly with surface charge density. Water polarisation increased approximately linearly up to a surface charge density of -0.17 Cm^{-2} , but at the very high density of -0.38 Cm^{-2} , water polarisation was found to decrease due to displacement by the high density of sodium ions present at the interface. An empirical relationship was used to relate the surface charge density to measured pH for these systems. Using this relation, these simulations predict that, for both amorphous and crystalline silica systems, from pH 2–12, both sodium ion accumulation and water polarisation does not reach a maximum. At higher pH, and therefore at higher surface charge, maximal water polarisation occurs, however, at such high pH the effects of silica surface dissolution become significant. These effects cannot be simulated using the current methodology but attempts to describe them have been made in other ‘reactive’ forcefields³⁸.

The surface potential properties were calculated and compared to transferable models of the electrical double layer. The amorphous surface showed a larger change in surface potential for a given change in surface charge density or effective pH than the crystalline model, despite both systems being approximately equal hydroxyl density, charge density and electrolyte ionic strength. The greater surface potential shift of the amorphous system was explained as a result of increased sodium ion accumulation due to a higher surface area and increased availability of favourable sodium ion binding sites. This novel result suggests that for pH sensor design, amorphous surfaces will have enhanced pH response compared to more structured surfaces with lower surface area. By contrast, commonly used models for describing pH response of potentiometric sensors, such as site-binding models,

cannot describe such differences due to surface morphology.

The electric field below the silica substrate was calculated as a measure of the ‘field effect’, the phenomenon which drives the response of field-effect transistor-based sensors. A shift of 0.007 Vnm^{-1} per effective pH was calculated. When DNA was introduced, a similar magnitude shift of 0.007 Vnm^{-1} was calculated, and given typical IS-FET sensors can resolve changes of at least one pH unit, this result predicts that this DNA-system should be experimentally detectable. The response rapidly diminished with distance of the DNA from the surface, in agreement with expectations based on the Debye-Hückel model. This result provides a first proof-of-concept for this type of simulation applied to potentiometric biosensing applications. The effects of biomolecule dynamics, biomolecule-ion interactions (e.g. ion displacement by biomolecules), the finite size of the biomolecule and the surface morphology are all explicitly treated, providing a wealth of information unavailable in current potentiometric biosensor models. Thus we posit that molecular dynamics provides a novel complementary tool to existing potentiometric biosensor models.

Conventional mean-field models provide predictions that are often insufficient for rational design of potentiometric sensors with regards to binding of molecular analytes. For example, recent studies have shown potentiometric detection of neutral alkanes when applied in nitrogen gas³³ or humid vapour^{34,115}, both of which are predicted to produce no signal using Poisson-Boltzmann-based models due to the lack of charge on each alkane molecule. The measured response is likely due to changes in electric field induced by water polarisation and analyte dipole orientation, both of which would be described by the present molecular dynamics model and will be investigated in future work.

A unique capability of potentiometric biosensors is detection of electrostatic properties, in contrast to conventional biosensors which often operate via mass and optical detection^{116,117}. As a result of this capability, properties unmeasurable using conventional biosensors can be determined such as conformational changes of the analyte¹¹⁸. It is therefore anticipated that simulation of the response of potentiometric biosensors via methods which are capable of modelling the dynamics of molecules will become increasingly important as potentiometric sensing becomes more wide-spread in the form of point-of-care diagnostic devices and environmental sensing applications.

Conflicts of Interest

There are no conflicts to declare.

Acknowledgements

The authors acknowledge the use of both the Reedbush HPC facilities at the University of Tokyo and the IRIDIS HPC Facility, and associated HPC and linux support services at the University of Southampton, in the completion of the present work. The NAMD software was used in this work, which was developed by the Theoretical and Computational Biophysics Group in the Beckman Institute for Advanced Science and Technology at the University of Illinois at Urbana-Champaign. This work was completed by an overseas researcher under the Postdoctoral Fellowship of the



Japan Society for the Promotion of Science 2017/18. The authors acknowledge Dr. In-Chul for providing code to implement the EW3DC correction used in this work and Dr. Christoph Riedel for proofreading and typography corrections.

References

- J. M. Andújar and F. Segura, *Renewable and Sustainable Energy Reviews*, 2009, **13**, 2309–2322.
- A. K. Gupta and M. Gupta, *Biomaterials*, 2005, **26**, 3995–4021.
- M. Noked, A. Soffer and D. Aurbach, *J. Solid State Electrochem.*, 2011, **15**, 1563–1578.
- T. Metke, A. S. Westover, R. Carter, L. Oakes, A. Douglas and C. L. Pint, *Sci. Rep.*, 2016, **6**, 24680.
- P. Bergveld, *IEEE Trans. Biomed. Eng.*, 1972, **BME-19**, 342–351.
- P. Bergveld, *IEEE Trans. Biomed. Eng.*, 1970, **BME-17**, 70–71.
- Y. Cui, Q. Wei, H. Park and C. M. Lieber, *Science*, 2001, **293**, 1289–1292.
- B. M. Lowe, K. Sun, I. Zeimpekis, C.-K. Skylaris and N. G. Green, *Analyst*, 2017, **142**, 4173–4200.
- K. Y. Chumbimuni-Torres, Z. Dai, N. Rubinova, Y. Xiang, E. Pretsch, J. Wang and E. Bakker, *J. Am. Chem. Soc.*, 2006, **128**, 13676–13677.
- R. Thüerer, T. Vigassy, M. Hirayama, J. Wang, E. Bakker and E. Pretsch, *Anal. Chem.*, 2007, **79**, 5107–5110.
- M.-Y. Shen, B.-R. Li and Y.-K. Li, *Biosens. Bioelectron.*, 2014, **60**, 101–111.
- G. Shalev, Y. Rosenwaks and I. Levy, *Biosens. Bioelectron.*, 2012, **31**, 510–515.
- L. De Vico, L. Iversen, M. H. Sørensen, M. Brandbyge, J. Nygård, K. L. Martinez and J. H. Jensen, *Nanoscale*, 2011, **3**, 3635.
- P. M. Dove and S. F. Elston, *Geochim. Cosmochim. Ac.*, 1992, **56**, 4147–4156.
- P. M. Dove and C. J. Nix, *Geochim. Cosmochim. Ac.*, 1997, **61**, 3329–3340.
- P. M. Dove, *Am J Sci*, 1994, **294**, 665–712.
- J. V. Smith, *PNAS*, 1998, **95**, 3370–3375.
- P. Nair and M. Alam, *IEEE Trans. Electron Devices*, 2007, **54**, 3400–3408.
- C. Heitzinger, Y. Liu, N. J. Mauser, C. Ringhofer and R. W. Dutton, *J. Comput. Theor. Nanosci.*, 2010, **7**, 2574–2580.
- C. Heitzinger and G. Klimeck, *J. Comput. Electron.*, 2007, **6**, 387–390.
- C. Heitzinger, R. Kennell, G. Klimeck, N. Mauser, M. McLennan and C. Ringhofer, *J. Phys.: Conf. Ser.*, 2008, **107**, 012004.
- C. Heitzinger, N. J. Mauser and C. Ringhofer, *SIAM J. Appl. Math.*, 2010, **70**, 1634–1654.
- S. Baumgartner, C. Heitzinger, A. Vacic and M. A. Reed, *Nanotechnology*, 2013, **24**, 225503.
- S. Baumgartner, M. Vasicek, A. Bulyha and C. Heitzinger, *Nanotechnology*, 2011, **22**, 425503.
- T. Windbacher, V. Sverdlov and S. Selberherr, *Computational Electronics*, 2009. IWCE'09. 13th International Workshop On, 2009, pp. 1–4.
- T. Windbacher, V. Sverdlov, S. Selberherr, C. Heitzinger, N. Mauser and C. Ringhofer, *AIP Conf. Proc.*, 2010, **1199**, 507.
- P. R. Nair and M. A. Alam, *Nano Lett.*, 2008, **8**, 1281–1285.
- T. Windbacher, V. Sverdlov and S. Selberherr, *Biomedical Engineering Systems and Technologies*, Springer, New York City, 2010, pp. 85–95.
- M. H. Sørensen, N. A. Mortensen and M. Brandbyge, *Appl. Phys. Lett.*, 2007, **91**, 102105–102105–3.
- L. De Vico, M. H. Sørensen, L. Iversen, D. M. Rogers, B. S. Sørensen, M. Brandbyge, J. Nygård, K. L. Martinez and J. H. Jensen, *Nanoscale*, 2011, **3**, 706.
- N. Lloret, R. S. Frederiksen, T. C. Møller, N. I. Rieben, S. Upadhyay, L. De Vico, J. H. Jensen, J. Nygård and K. L. Martinez, *Nanotechnology*, 2013, **24**, 035501.
- R. R. Pethig and S. Smith, *Introductory Bioelectronics: For Engineers and Physical Scientists*, John Wiley & Sons, Hoboken, New Jersey, 2012.
- N. Swaminathan, A. Henning, T. Jurca, J. Hayon, G. Shalev and Y. Rosenwaks, *Sens. Actuators B Chem.*, 2017, **248**, 240–246.
- B. Wang and H. Haick, *ACS Appl. Mater. Interfaces*, 2013, **5**, 5748–5756.
- N. Gao, W. Zhou, X. Jiang, G. Hong, T.-M. Fu and C. M. Lieber, *Nano Lett.*, 2015, **15**, 2143–2148.
- G. Zheng, X. P. A. Gao and C. M. Lieber, *Nano Lett.*, 2010, **10**, 3179–3183.
- I.-Y. Chung, J. Lee, M. Seo and C. H. Park, *Jpn. J. Appl. Phys.*, 2016, **55**, 127001.
- J. C. Fogarty, H. M. Aktulga, A. Y. Grama, A. C. T. van Duin and S. A. Pandit, *J. Chem. Phys.*, 2010, **132**, 174704.
- Y. Maekawa, Y. Shibuta and T. Sakata, *Jpn. J. Appl. Phys.*, 2013, **52**, 127001.
- Y. Maekawa, Y. Shibuta and T. Sakata, *Chem. Phys. Lett.*, 2015, **619**, 152–157.
- Y. Maekawa, Y. Shibuta and T. Sakata, *ChemElectroChem*, 2014, **1**, 1516–1524.
- B. M. Lowe, Y. Maekawa, Y. Shibuta, T. Sakata, C.-K. Skylaris and N. G. Green, *Phys. Chem. Chem. Phys.*, 2017, **19**, 2687.
- I.-C. Yeh and A. Wallqvist, *J. Chem. Phys.*, 2011, **134**, 055109.
- C. D. Lorenz, P. S. Crozier, J. A. Anderson and A. Travesset, *J. Phys. Chem. C*, 2008, **112**, 10222–10232.
- R. Lavery, J. H. Maddocks, M. Pasi and K. Zakrzewska, *Nucleic Acids Res.*, 2014, **42**, 8138–8149.
- B. M. Lowe, C.-K. Skylaris, N. G. Green, Y. Shibuta and T. Sakata, *Jpn. J. Appl. Phys.*, 2018, **57**, 04FM02.
- I.-C. Yeh and M. L. Berkowitz, *J. Chem. Phys.*, 1999, **111**, 3155–3162.
- J. O. Bockris, A. K. Reddy and M. E. Gamboa-Aldeco, *Modern Electrochemistry 2A*, Kluwer Academic Publishers, Boston,

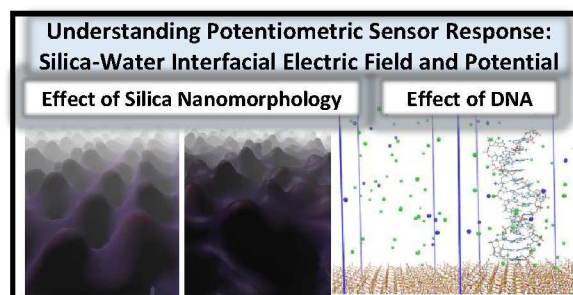


- 2002, vol. 2A.
- 49 X. Duan, Y. Li, N. K. Rajan, D. A. Routenberg, Y. Modis and M. A. Reed, *Nat. Nano.*, 2012, **7**, 401–407.
- 50 X. Wen, S. Gupta, Y. Wang, T. R. N. Iii, S. C. Lee and W. Lu, *Appl. Phys. Lett.*, 2011, **99**, 043701.
- 51 B. K. Wunderlich, P. A. Neff and A. R. Bausch, *Appl. Phys. Lett.*, 2007, **91**, 083904.
- 52 D. Landheer, G. Aers, W. R. McKinnon, M. J. Deen and J. C. Ranuarez, *J. Appl. Phys.*, 2005, **98**, 044701.
- 53 D. Landheer, W. McKinnon, G. Aers, W. Jiang, M. Deen and M. Shinwari, *IEEE Sens. J.*, 2007, **7**, 1233–1242.
- 54 S. Uno, M. Iio, H. Ozawa and K. Nakazato, *Jpn. J. Appl. Phys.*, 2010, **49**, 01AG07.
- 55 Y. Nishio, S. Uno and K. Nakazato, *Jpn. J. Appl. Phys.*, 2013, **52**, 04CL01.
- 56 H. Ohshima, *Biophysical Chemistry of Biointerfaces*, John Wiley & Sons, 2011.
- 57 R. J. Hunter, *Foundations of Colloid Science*, Oxford University Press, Oxford, UK, 2001.
- 58 J. Lyklema, *Fundamentals of Interface and Colloid Science: Solid-Liquid Interfaces*, Academic Press, Cambridge, Massachusetts, 1995.
- 59 D. Grahame, *Chem. Rev.*, 1947, **41**, 441–501.
- 60 J. O. Bockris and A. K. Reddy, *Modern Electrochemistry 1*, Kluwer Academic Publishers, Boston, 1998, vol. 1.
- 61 M. A. Brown, G. V. Bossa and S. May, *Langmuir*, 2015, **31**, 11477–11483.
- 62 M. S. Kilic, M. Z. Bazant and A. Ajdari, *Phys. Rev. E*, 2007, **75**, 021502.
- 63 M. S. Kilic, M. Z. Bazant and A. Ajdari, *Phys. Rev. E*, 2007, **75**, 021503.
- 64 P. W. Atkins, *Physical Chemistry*, W H Freeman & Co, New York, 1994.
- 65 T. Sakata and Y. Miyahara, *Angew. Chem. Int. Ed.*, 2006, **45**, 2225–2228.
- 66 F. N. Ishikawa, M. Curreli, H.-K. Chang, P.-C. Chen, R. Zhang, R. J. Cote, M. E. Thompson and C. Zhou, *ACS Nano*, 2009, **3**, 3969–3976.
- 67 F. Uslu, S. Ingebrandt, D. Mayer, S. Böcker-Meffert, M. Odenthal and A. Offenhäusser, *Biosens. Bioelectron.*, 2004, **19**, 1723–1731.
- 68 K. Jayant, K. Auluck, S. Rodriguez, Y. Cao and E. C. Kan, *Phys. Rev. E*, 2014, **89**, 052817.
- 69 J. Fritz, E. B. Cooper, S. Gaudet, P. K. Sorger and S. R. Manalis, *PNAS*, 2002, **99**, 14142–14146.
- 70 A. Poghosian, A. Cherstvy, S. Ingebrandt, A. Offenhäusser and M. J. Schöning, *Sens. Actuat. B-Chem.*, 2005, **111–112**, 470–480.
- 71 J. Bobacka, A. Ivaska and A. Lewenstam, *Chem. Rev.*, 2008, **108**, 329–351.
- 72 A. Tarasov, M. Wipf, R. L. Stoop, K. Bedner, W. Fu, V. A. Guzenko, O. Knopfmacher, M. Calame and C. Schönenberger, *ACS Nano*, 2012, **6**, 9291–9298.
- 73 R. E. G. van Hal, J. C. T. Eijkel and P. Bergveld, *Adv. Colloid Interface.*, 1996, **69**, 31–62.
- 74 D. E. Yates, S. Levine and T. W. Healy, *J. Chem. Soc., Faraday Trans. 1*, 1974, **70**, 1807–1818.
- 75 R. E. G. van Hal, J. C. T. Eijkel and P. Bergveld, *Sens. Actuat. B-Chem.*, 1995, **24**, 201–205.
- 76 S. Bermejo, *Chemical Sensors: Comprehensive Sensor Technologies Volume 5: Electrochemical and Optical Sensors*, Momentum Press, New York, 1st edn, 2011, pp. 201–247.
- 77 P. Bergveld, *Sens. Actuators B*, 2003, **88**, 1–20.
- 78 A. Rimola, D. Costa, M. Sodupe, J.-F. Lambert and P. Ugliengo, *Chem. Rev.*, 2013, **113**, 4216–4313.
- 79 L. Zhuravlev, *Colloids Surf. A Physicochem. Eng. Asp.*, 2000, **173**, 1–38.
- 80 B. M. Lowe, C.-K. Skylaris and N. G. Green, *J. Colloid Interf. Sci.*, 2015, **451**, 231–244.
- 81 E. Tombácz, *Chem. Eng.*, 2009, **53**, 77–86.
- 82 G. H. Bolt, *J. Phys. Chem.*, 1957, **61**, 1166–1169.
- 83 V. Puchol, J. El Haskouri, J. Latorre, C. Guillem, A. Beltrán, D. Beltrán and P. Amorós, *Chem. Commun.*, 2009, 2694.
- 84 B. B. Owen and S. R. Brinkley, *Chem. Rev.*, 1941, **29**, 461–474.
- 85 F. S. Emami, V. Puddu, R. J. Berry, V. Varshney, S. V. Patwardhan, C. C. Perry and H. Heinz, *Chem. Mater.*, 2014, **26**, 2647–2658.
- 86 Y. Duval, J. A. Mielczarski, O. S. Pokrovsky, E. Mielczarski and J. J. Ehrhardt, *J. Phys. Chem. B*, 2002, **106**, 2937–2945.
- 87 K. Leung, I. M. B. Nielsen and L. J. Criscenti, *J. Am. Chem. Soc.*, 2009, **131**, 18358–18365.
- 88 W. Siu and R. Cobbold, *IEEE Trans. Electron Devices*, 1979, **26**, 1805–1815.
- 89 T. Sakata, K. Nishimura, Y. Miyazawa, A. Saito, H. Abe and T. Kajisa, *Anal. Chem.*, 2017, **89**, 3901–3908.
- 90 J. Y. Oh, H.-J. Jang, W.-J. Cho and M. S. Islam, *Sensors and Actuators B: Chemical*, 2012, **171–172**, 238–243.
- 91 S. Dastidar, A. Agarwal, N. Kumar, V. Bal and S. Panda, *IEEE Sens. J.*, 2015, **15**, 2039–2045.
- 92 N. Kumar, J. Kumar and S. Panda, *ECS J. Solid State Sci. Technol.*, 2015, **4**, N18–N23.
- 93 M. J. Schöning, F. Ronkel, M. Crott, M. Thust, J. W. Schultze, P. Kordos and H. Lüth, *Electrochimica Acta*, 1997, **42**, 3185–3193.
- 94 M. J. Schöning, Ü. Malkoc, M. Thust, A. Steffen, P. Kordoš and H. Lüth, *Sensors and Actuators B: Chemical*, 2000, **65**, 288–290.
- 95 F. K. Perkins, L. M. Tender, S. J. Fertig and M. C. Peckerar, *Nanostructure Science, Metrology and Technology*, Gaithersburg, MD, United States, 2002, pp. 251–265.
- 96 F. Wei, B. Sun, Y. Guo and X. S. Zhao, *Biosensors and Bioelectronics*, 2003, **18**, 1157–1163.
- 97 A. W. Peterson, R. J. Heaton and R. M. Georgiadis, *Nucleic Acids Res*, 2001, **29**, 5163–5168.
- 98 A. Bandiziol, P. Palestri, F. Pittino, D. Esseni and L. Selmi, *IEEE Trans. Electron Devices*, 2015, **62**, 3379–3386.



- 99 F. S. Emami, V. Puddu, R. J. Berry, V. Varshney, S. V. Patwardhan, C. C. Perry and H. Heinz, *Chem. Mater.*, 2016, **28**, 406–407.
- 100 K. Hart, N. Foloppe, C. M. Baker, E. J. Denning, L. Nilsson and A. D. MacKerell, *J. Chem. Theory Comput.*, 2012, **8**, 348–362.
- 101 H. Heinz, T.-J. Lin, R. Kishore Mishra and F. S. Emami, *Langmuir*, 2013, **29**, 1754–1765.
- 102 F. S. Emami, V. Puddu, R. J. Berry, V. Varshney, S. V. Patwardhan, C. C. Perry and H. Heinz, *Chem. Mater.*, 2014, **26**, 5725–5734.
- 103 S. Miyamoto and P. A. Kollman, *J. Comput. Chem.*, 1992, **13**, 952–962.
- 104 A. A. Gurtovenko and I. Vattulainen, *J. Chem. Phys.*, 2009, **130**, 215107.
- 105 F. Musso, P. Mignon, P. Ugliengo and M. Sodupe, *Phys. Chem. Chem. Phys.*, 2012, **14**, 10507.
- 106 J. W. Lee, A. Mani and J. A. Templeton, *Langmuir*, 2015, **31**, 7496–7502.
- 107 J. W. Murray, *Journal of Colloid and Interface Science*, 1974, **46**, 357–371.
- 108 C. C. Bueno, A. M. Amarante, G. S. Oliveira, D. K. Deda, O. Teschke, E. d. F. Franca and F. L. Leite, *IEEE Sens. J.*, 2014, **14**, 1467–1475.
- 109 M. A. Brown, Z. Abbas, A. Kleibert, R. G. Green, A. Goel, S. May and T. M. Squires, *Phys. Rev. X*, 2016, **6**, 011007.
- 110 J. Diot, J. Joseph, J. Martin and P. Clechet, *J. Electroanal. Chem. Interfacial Electrochem.*, 1985, **193**, 75–88.
- 111 C. Fung, P. Cheung and W. Ko, *Electron Devices Meet.*, 1980, 689–692.
- 112 K. Sun, I. Zeimpekis, C. Hu, N. M. J. Ditshego, O. Thomas, M. R. R. de Planque, H. M. H. Chong, H. Morgan and P. Ashburn, *Nanotechnology*, 2016, **27**, 285501.
- 113 L. Bousse, N. F. D. Rooij and P. Bergveld, *IEEE Trans. Electron Devices*, 1983, **30**, 1263–1270.
- 114 S. Chen, J. G. Bomer, E. T. Carlen and A. van den Berg, *Nano Lett.*, 2011, **11**, 2334–2341.
- 115 N. Shehada, J. C. Cancilla, J. S. Torrecilla, E. S. Pariente, G. Brønstrup, S. Christiansen, D. W. Johnson, M. Leja, M. P. A. Davies, O. Liran, N. Peled and H. Haick, *ACS Nano*, 2016, **10**, 7047–7057.
- 116 H. H. Nguyen, J. Park, S. Kang and M. Kim, *Sensors*, 2015, **15**, 10481–10510.
- 117 S. Sang, Y. Wang, Q. Feng, Y. Wei, J. Ji and W. Zhang, *Crit. Rev. Biotechnol.*, 2015, 1–17.
- 118 D. Cahen, R. Naaman and Z. Vager, *Adv. Funct. Mater.*, 2005, **15**, 1571–1578.



Graphical Abstract: (8cm x 4cm)

A molecular dynamics methodology for detailed potentiometric response prediction is presented and silica amorphicity was found to affect pH sensitivity.

