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# The physico-chemistry of adhesions of protein resistant and weak polyelectrolyte brushes to cells and tissues†

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The non-specific adhesion of polymers and soft tissues is of great interest to the field of biomedical engineering, as it will shed light on some of the processes that regulate interactions between scaffolds, implants and nanoparticles with surrounding tissues after implantation or delivery. In order to promote adhesion to soft tissues, a greater understanding of the relationship between polymer chemistry and nanoscale adhesion mechanisms is required. In this work, we grew poly(dimethylaminoethyl methacrylate) (PDMAEMA), poly(acrylic acid) (PAA) and poly(oligoethylene glycol methacrylate) (POEGMA) brushes from the surface of silica beads, and investigated their adhesion to a variety of substrates via colloidal probe-based atomic force microscopy (AFM). We first characterised adhesion to a range of substrates with defined surface chemistry (self-assembled monolayers (SAMs) with a range of hydrophilicities, charge and hydrogen bonding), before studying the adhesion of brushes to epithelial cell monolayers (primary keratinocytes and HaCaT cells) and soft tissues (porcine epicardium and keratinized gingiva). Adhesion assays to SAMs reveal the complex balance of interactions (electrostatic, van der Waals interactions and hydrogen bonding) regulating the adhesion of weak polyelectrolyte brushes. This resulted in particularly strong adhesion of PAA brushes to a wide range of surface chemistries. In turn, colloidal probe microscopy on cell monolayers highlighted the importance of the glycocalyx in regulating non-specific adhesions. This was also reflected by the adhesive properties of soft tissues, in combination with their mechanical properties. Overall, this work clearly demonstrates the complex nature of interactions between polymeric biomaterials and biological samples and highlights the need for relatively elaborate models to predict these interactions.

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## 1. Introduction

The non-specific interactions between polymeric biomaterials and soft tissues play an important role in the performance of implants, tissue engineering scaffolds and drug delivery systems.<sup>1–5</sup> Physico-chemical interactions are particularly important in regulating the strength of bioadhesives, such as skin adhesives for surgical applications,<sup>6–8</sup> hydrogels for epicardial placement and stem cell delivery,<sup>9</sup> soft tissue adhesion for tissue regeneration<sup>3,4</sup> or mucoadhesives for dental adhesion.<sup>10,11</sup> Although a range of chemical functions and molecules have been introduced in biomaterials to promote covalent coupling and adhesion to soft

tissues, the control and regulation of non-specific physico-chemical interactions with surrounding tissues post-implantation and delivery can play an important role and even dominate adhesion performance.<sup>11,12</sup> In addition, polymeric coatings introduced around nanomaterials for drug and gene delivery, or for imaging, can significantly impact their biodistribution and systemic persistence/clearance.<sup>1,2,13</sup> Soft tissue adhesion is a complex process involving non-specific interactions, inter-penetration of polymer networks and soft tissue interfaces, and molecular coupling; these are regulated by the mechanical properties of the bulk biomaterial, the soft tissue and their interface.<sup>12,14</sup> Typically, after bringing a polymer into contact with a soft tissue, initial physical entanglement will be followed by secondary bonding to strengthen adhesive interactions.<sup>15</sup> As is often the case in Nature, these secondary bonds will be formed by multiple individually weak non-covalent bonds that combine to form a stable, highly specific, reversible and strong intermolecular connection. These intermolecular interactions typically involve a combination of hydrogen bonds, electrostatic interactions and van der Waals interactions.<sup>16</sup> The design of

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interfaces with controlled chemistry and intermolecular interactions is particularly important to the modelling and understanding of biointerface adhesion.

In addition, tissue adhesion is further complicated by the secretion of biomacromolecules from cells constituting the corresponding soft tissues, modulating their interactions with contacting polymers. For example, salivary glands in the oral mucosa secrete mucins, a variety of antimicrobial substances, salts, water, and intestinal proteins that modulate mucosal adhesion. Hence mucoadhesives must adhere to the moist surface of the oral mucosa whilst resisting the flushing action of the saliva and competition with associated biomacromolecules.<sup>17,18</sup>

Polymer brushes are interesting model biointerfaces for the study of physico-chemical parameters affecting bioadhesion. Indeed, polymer brushes offer unique control of the chemistry of interfaces, enabling the formation of both very hydrophobic and hydrophilic polymer coatings, as well as polyelectrolytes. Furthermore, the morphology and mechanics of these interfaces can be regulated *via* the control of brush grafting density, thickness, swelling and conformation.<sup>19,20</sup> The wealth of monomers that can be incorporated in polymer brushes has enabled a wide range of properties for these coatings, including protein resistance,<sup>21</sup> thermoresponsiveness,<sup>22,23</sup> electrolyte responsiveness<sup>24,25</sup> and bacterial resistance.<sup>26,27</sup> In addition, brushes can be readily grown from a wide range of substrates, making these coatings attractive for a range of biomedical applications from implant design<sup>28</sup> to gene delivery.<sup>29,30</sup>

The adhesive and anti-fouling properties of polymer brushes and their nanoscale mechanics have been characterised relatively extensively by atomic force microscopy (AFM). Colloidal probe microscopy (CPM) utilises an AFM cantilever with a spherical particle attached at the apex in order to study the nano-scale forces between a particle and substrate with defined physico-chemical properties. The most widely used microspheres for CPM is silica, as they have a low roughness, are commercially available in a variety of sizes, and their surfaces can be readily chemically modified.<sup>31,32</sup> The use of CPM has enabled the characterisation of protein adhesion to polymer brushes,<sup>33–35</sup> as well as bacterial adhesion.<sup>27</sup> In these studies, neutral and zwitterionic brushes that are particularly well solvated were shown to effectively limit adhesion of unwanted biomacromolecules and components of the bacterial membranes such as lipopolysaccharides. In addition, CPM was used to quantify adhesion and friction to hydrophobic and fluorophilic brushes; the brushes displaying increasing degrees of fluorination displayed reduced adhesion and friction (silica colloidal probes).<sup>36</sup> The conformation and swelling of polymer brushes also strongly affect adhesion strength and contact mechanics, therefore highlighting the impact of environmental factors on non-specific adhesion.<sup>37</sup> Such impact of the environment is also strikingly illustrated by the response of polyelectrolyte brushes to pH, electrolyte chemistry and ionic strength. Hence the strength of adhesion of silica microparticles to poly(2-vinylpyridine) brushes is enhanced by phosphate and sulfate electrolytes, compared to chloride, and is reduced at higher ionic strength.<sup>38</sup> Overall, although the non-specific adhesion of solid substrates, bacteria and proteins

to polymer brushes has been widely studied, limited studies have focused on non-specific adhesion to cells and tissues.

In this report, we study the adhesion of three types of polymer brushes to cells and tissues. We selected poly(acrylic acid) (PAA), poly(dimethylaminoethyl methacrylate) (PDMAEMA) and poly(oligoethylene glycol methacrylate) (POEGMA) as three important types of polymer brushes with distinct physico-chemical properties (anionic, cationic and neutral hydrophilic polymers, respectively). PAA displays a high density of carboxylic acid groups and negative charges (at high pH), is biocompatible and is widely used as biomaterials<sup>39</sup> to promote mucoadhesion,<sup>40–42</sup> as drug delivery systems,<sup>43</sup> for surface coating and modification<sup>44,45</sup> and for microdevice and sensor design.<sup>39</sup> PDMAEMA is a weak polybasic polymer with pH-responsive properties<sup>46</sup> that has applications in gene delivery.<sup>30,47</sup> Finally, POEGMA is known for its excellent protein resistance arising from the high density of ethylene glycol moieties on the polymer backbone, its ability to grow at high grafting densities and its hydrophilicity.<sup>48</sup> As such, its applications include surface modifications and coatings that resist protein absorption and thus prevent bacterial adhesion and biofilm formation.<sup>48–50</sup> We first examine the adhesion of these three brushes to a range of model monolayers presenting a range of chemistries (surface charge, hydrogen bonding and hydrophilicity). We then explore the impact of brush chemistry on adhesion to model cell monolayers (epithelial cell sheets) and investigate the impact of their glycocalyx on adhesion. Indeed, most epithelial cells are surrounded by a glycoprotein and glycolipid layer associated with their membrane and known as the glycocalyx; this ranges from 7 nm in thickness for red blood cells up to several 100 nm in some cases.<sup>51</sup> Consequently, when an implanted material comes into contact with epithelial cell sheets, often lining the surface of tissues, the glycocalyx is the first point of contact and its interaction with the material is of significant importance. Finally, we explore brush adhesion to tissues (gingival epithelium and epicardium) and discuss the impact of brush chemistry on adhesive behaviour.

## 2. Experimental section

### 2.1. Chemicals and materials

*tert*-Butyl acrylate (tBA,  $M_n = 128.17$ , 98%), 2-(dimethylamino)-ethyl methacrylate (DMAEMA,  $M_n = 157.21$ , 98%), oligo(ethylene glycol methyl ether methacrylate) (OEGMA,  $M_n = 300$ ), poly(L-lysine) (PLL), *N,N,N',N'',N'''*-pentamethyldiethylenetriamine (PMDETA, 99%), 2,2'-bipyridyl (bipy,  $\geq 99\%$ ), triethylamine ( $\geq 99.5\%$ ), copper chloride (Cu(I)Cl), copper bromide (Cu(I)Br and Cu(II)Br<sub>2</sub>), trifluoroacetic acid (TFA, 99%), toluene (anhydrous, 99.8%), iodomethane (99%), (3-aminopropyl)triethoxysilane (APTES, 99%), triethoxy(octyl)silane (97%), trichloro(1*H*,1*H*,2*H*,2*H*-perfluorooctyl)silane (97%), 11-mercaptoundecanoic acid (MUDA, 95%), 11-mercapto-1-undecanol (97%), Dulbecco's phosphate buffered saline (PBS), neuraminidase (from *Clostridium Perfringens*), heparinase III (from *Flavobacterium Heparinum*), wheat germ agglutinin (WGA, lectin from *Triticum vulgaris*, FITC conjugate), formaldehyde solution (4% in PBS) (PFA)



and 4',6-diamidino-2-phenylindole dihydrochloride (DAPI) were purchased from Sigma-Aldrich and used as received. Cu(I)Cl and Cu(I)Br were kept under vacuum until use. The silane initiator (3-trimethoxysilyl)-propyl 2-bromo-2-methylpropionate was purchased from Gelest. Acetone (technical), versene solutions (0.2 g L<sup>-1</sup> EDTA in PBS), keratinocyte serum-free medium (KSFM), Dulbecco's modified Eagle's medium (DMEM), trypsin (0.25%), L-glutamine, penicillin-streptomycin, bovine pituitary extract (BPE), human recombinant epidermal growth factor (EGF), and goat anti-mouse IgG (H + L) secondary antibodies (conjugated to AlexaFluor 488) were purchased from Thermo Fisher Scientific. Fibronectin solution (from human plasma) and the anti-heparin/heparan sulfate antibody were purchased from Merck Millipore. Foetal bovine serum (FBS) was purchased from Labtech. Collagen I (type I) was purchased from Corning. Dichloromethane (DCM, ≥ 99.8%) was purchased from Honeywell. Dimethylformamide (DMF, ≥ 99.9%) was purchased from VWR Chemicals. AFM probes (ORC8-10) were purchased from Bruker. Silica particles (unfunctionalised) were purchased from microParticles GmbH (mean diameter of 19.59 μm) and Bangs Laboratories (mean diameter of 300 nm). SEMGLU was purchased from Kleindiek Nanotechnik GmbH, Germany. Silicon wafers (100 mm diameter, <100> orientation, polished on one side/reverse etched) were purchased from Compant Technology Ltd. All Plasma treatment was carried out using a Henniker Plasma Vacuum System HPT-200. Human keratinocyte HaCaT cells and Human primary epidermal keratinocytes were kindly provided by Dr John Connelly, Blizzard Institute, Barts and The London School of Medicine and Dentistry, Queen Mary University of London.

## 2.2. Polymer brush synthesis on planar silicon substrates

The brushes were synthesized by the "grafting from" method using atom transfer radical polymerization (ATRP).

**Deposition of monolayers of silane initiator for ATRP on silicon substrates.** Samples of silicon wafer (1 × 1 cm) were plasma-oxidized (10 min, air), followed by immersion in a solution of silane initiator (3-trimethoxysilyl)-propyl 2-bromo-2-methylpropionate (30 μL) and triethylamine (50 μL) in anhydrous toluene (30 mL) and incubation in this solution at room temperature overnight. Samples were then washed with ethanol and dried under a stream of nitrogen. The dry thickness of the silane initiator layer was near 2 nm, as measured using spectroscopic ellipsometry.

**Synthesis of poly(acrylic acid) (PAA) brushes.** PAA brushes were fabricated through a two-step process: the protected monomer *tert*-butyl acrylate (tBA) was polymerized first, followed by deprotection to afford brushes of PAA. PtBA brushes were generated following a method adapted from Lego *et al.*<sup>52</sup> Prior to use, tBA was purified by passing through a column of basic alumina and freshly distilled under vacuum (78 mbar) at a temperature of 61–63 °C. Freshly purified and distilled tBA (30 mL, 205 mmol), PMDETA (122 μL, 584 μmol), CuBr<sub>2</sub> (3 mg, 13 μmol), and acetone (16 mL) were degassed *via* argon bubbling for 30 min. CuBr (65 mg, 453 μmol) was then very quickly added to this flask before degassing for a further 30 min.

Initiator-coated silicon substrates were placed in reaction vessels which were subsequently degassed *via* three cycles of high vacuum/argon gas refilling and heated to 60 °C. The monomer solution was then transferred to these reaction vessels under inert atmosphere and polymerization was allowed to proceed at 60 °C. To stop polymerization samples were immersed in acetone, rinsed with ethanol, deionised water, and dried under a stream of nitrogen. If necessary, silicon wafers were further cleaned through brief sonication in acetone. For the deprotection of PtBA brushes samples were immersed overnight in a DCM/TFA solution (10 : 1 (v/v)) at room temperature. The samples were then washed thoroughly in ethanol, followed by deionised water, and then dried under a stream of nitrogen.

**Synthesis of poly(dimethylaminoethyl methacrylate) (PDMAEMA) brushes.** PDMAEMA brushes were synthesised following a similar protocol as PtBA brushes except for a few differences: a monomer solution of DMAEMA (6.6 g, 42 mmol), bipy (320 mg, 2.05 mmol), CuBr<sub>2</sub> (18 mg, 81 μmol) and CuCl (84 mg, 849 μmol) in water/ethanol (4 : 1 (v/v), 15 mL) was used. The monomer was used as purchased without any need for initial purification, and polymerization was carried out at room temperature and was stopped by immersing samples in deionised water, rinsing with ethanol and drying under a stream of nitrogen.

**Synthesis of poly(oligoethylene glycol methacrylate) (POEGMA) brushes.** The procedure for synthesis of POEGMA brushes was identical to that of PDMAEMA brushes, except that a monomer solution of OEGMA (12.6 g, 42 mmol), bipy (320 mg, 2.05 mmol), CuBr<sub>2</sub> (18 mg, 81 μmol) and CuCl (84 mg, 845 μmol) in water/ethanol (4 : 1 (v/v), 15 mL) was used. For PAA, PDMAEMA and POEGMA brushes polymerization times were adjusted accordingly to give dry ellipsometric thicknesses of 30 nm; these thicknesses equated to polymerization times of 5 h, 20 min and 30 min, respectively.

**Characterisation of planar polymer brushes.** Dry brush thicknesses were measured using an  $\alpha$ -SE ellipsometer from J. A. Woolam Co., Inc., Ellipsometry solutions.  $\phi$  and  $\Delta$  were measured at wavelengths between 400 and 900 nm and a fixed incidence angle of 70°. For measurements of brush thicknesses when submerged in solution, substrates were placed in a sealed chamber fitted with a quartz window normal to the incident beam path. Different solutions were then passed through the chamber and the effect on the brush thickness was measured *via* the ellipsometer. Samples were incubated for 10 min prior to each *in situ* measurement. Measurements were taken in triplicate and were taken once the brush thicknesses had maintained equilibrium. For data analysis, all coatings were treated using a Cauchy model built on top of a silicon/native oxide or gold substrate, depending on the system studied.<sup>53</sup> For dry measurements of PtBA and PAA brushes, refractive indices measured were 1.53 and 1.52, respectively (measured at  $\lambda$  = 632.8 nm). For measurements in solution, refractive indices varied between 1.38 in PBS and 1.41 in deionised water (depending on swelling and associated hydration). The swelling behaviour of PDMAEMA and POEGMA brushes was previously studied by ellipsometry and reported by our group as well as others.<sup>22,54</sup>



Fourier Transform Infrared-Attenuated Total Reflectance Spectroscopy (ATR-FTIR) was carried out using a Bruker Tensor 27 with an MCT detector (liquid N<sub>2</sub> cooled). Spectra were acquired at a resolution of 4 cm<sup>-1</sup> with a total of 256 scans per run. Nitrogen was run through the system during measurements in order to limit the effects of environmental fluctuations. Contact angle measurements (using the sessile drop method) were taken using a Kruss DSA100 instrument using 3 µL droplets of deionised water. Measurements were carried out on three independent samples and for each sample an average reading was taken using measurements taken from three separate regions of the sample.

XPS was carried out on a Thermo Scientific K-Alpha X-ray XPS System using a monochromatic AlK $\alpha$  source. This was operated at 150 W under ultrahigh vacuum conditions (10<sup>-9</sup> mbar). As the surfaces were not electrically conductive, a built-in spectrometer charge neutralizing system was used which compensates for sample charging during measurements by flooding the surface with low energy electrons (<20 eV). To be able to compare data, the C<sub>1s</sub> hydrocarbon peak at 285.0 eV was used as an external standard for calibration of the binding energy (BE) scale. In order to identify the main peaks by their binding energies, survey spectra were collected from 1200 to 0 eV. In addition, high-resolution XPS spectra were acquired for O<sub>1s</sub>, C<sub>1s</sub>, and Si<sub>2p</sub> signals. An analyzer pass energy of 1 eV and 0.1 eV was used for the survey spectra and high resolution spectra, respectively.

### 2.3. Synthesis of polymer brush-coated silica nanoparticles

**Initiator deposition.** 20 µm silica beads are dispersed in deionised water for storage. To transfer them to a new solvent, beads are centrifuged at 4000 rpm for a few min until the beads fully precipitate. For the smaller beads used, with sizes in the nanometre range, higher centrifugation speeds (5000 rpm) and longer times were needed to fully precipitate the beads. The solvent was then carefully aspirated as much as possible using a syringe and the new solvent was added to the vial. The vial was then sonicated for 5 to 10 min until the suspension becomes cloudy. The sonication and centrifugation sequence is repeated three times to transfer the beads to the new solvent.

For grafting of the silane initiator the beads were transferred from water to ethanol and then to toluene using the aforementioned method. 50 mg of the beads were dispersed in a solution of anhydrous toluene (1 mL), triethylamine (50 µL) and silane initiator (10 µL) and left stirring overnight. The beads were then washed in toluene three times using the sonication-centrifugation method and then transferred to acetone and left in 1 mL of the solvent in the fridge ready for the polymerization step.

**Synthesis of polymer brushes from silica nanoparticles.** The protocol for PtBA polymer brush synthesis is a slight adaptation of the polymerization protocol used for planar silicon substrates. An extra 2 mL of acetone was initially added to the 1 mL silica dispersion and this was then degassed for 30 min with argon bubbling. Freshly purified and distilled tBA (3.0 mL, 21 mmol), acetone (1.6 mL), PMDETA (12 µL, 58 µmol) and CuBr<sub>2</sub> (0.3 mg, 1.34 µmol) were degassed for 30 min whilst mixing. The top was opened and CuBr (6.5 mg, 45.3 µmol) was

quickly added to the monomer solution followed by a further 30 min of degassing. Next, 2.5 mL of this monomer mixture was extracted and added to the reaction vessel containing the 3 mL particle suspension. The syringe used to extract it was flushed with argon three times prior to use.

Polymerization was then carried out at 60 °C whilst stirring. Degassing was continued for a further 30 min and the reaction vessel was then covered with parafilm. To terminate polymerization, the reaction mixture was bubbled with oxygen for a couple of min. The SiO<sub>2</sub>-polymer suspension was then centrifuged, washed in acetone three times and transferred into DCM/TFA (10 : 1 (v/v), 1 mL) and left stirring overnight at room temperature. The resulting PAA beads were washed in DCM three times and transferred into acetone for storage in the fridge.

For synthesis of PDMAEMA polymer brushes, 5 mL of the DMAEMA monomer solution (see Section 2.2 for preparation protocol) was added to a 5 mL silica dispersion (50 mg SiO<sub>2</sub>-silane beads in water/ethanol (4 : 1 (v/v))), degassed under argon for 30 min). Argon bubbling was continued and polymerization was allowed to proceed at room temperature. The reaction was stopped with oxygen bubbling and the beads were washed in water/ethanol (4 : 1 (v/v)). Synthesis of POEGMA polymer brushes was identical except that 5 mL of the OEGMA monomer solution (see Section 2.2) was instead added to the silica bead dispersion. The grafting density of polymer brushes that were previously grown from silica nanoparticles using this protocol was 0.5 chains per nm<sup>2</sup>, with dry thicknesses of 30 nm (and a *M<sub>n</sub>* of 50 kDa).<sup>30</sup>

#### Characterisation of polymer brush-coated silica particles.

The sizes and zeta potentials of functionalised silica particles were measured using a Malvern Zetasizer Nano ZS. For measurements, PtBA and PAA brushes were grown from 300 nm silica beads, then dispersed in ethanol or deionised water respectively, and then sonicated whilst shaking at regular intervals until a cloudy solution was obtained. Measurements of non-functionalised silica beads were carried out with beads dispersed in deionised water. Three independent repeats were obtained at room temperature for each sample and the average result was taken.

### 2.4. Formation of self-assembled monolayers (SAMs) on silicon substrates

The Table S1 (ESI<sup>†</sup>) gathers the contact angle data and ellipsometric thicknesses for all SAMs generated on silicon and gold-coated substrates.

**3-Aminopropyl triethoxysilane (APTES) SAMs.** A plasma-oxidized (10 min, air) silicon substrate was immersed in a solution of ethanol (1 mL) and 3-aminopropyl triethoxysilane (APTES) (10 µL) and left at room temperature for 30 min. The sample was then rinsed with ethanol and dried under a stream of nitrogen. Subsequently, the APTES-coated silicon substrate was immersed in a solution of 1 mM HCl for 5 min and then washed thoroughly with deionised water followed by ethanol, and then dried in a stream of nitrogen.

**Quaternized APTES (QAPTES) SAMs.** QAPTES-functionalised silicon substrates were prepared similarly to the APTES samples, except that, following APTES deposition, they were immersed in





a solution of 5 mM NaOH for 5 min, followed by incubation in a solution of iodomethane (13.3  $\mu\text{L}$ ) in DMF (1.5 mL) overnight at room temperature. The samples were then washed with ethanol and dried under nitrogen.

**Triethoxy(octyl)silane SAMs.** A plasma-oxidized silicon substrate was immersed in a solution of anhydrous toluene (1 mL), triethylamine (20  $\mu\text{L}$ ), and triethoxy(octyl)silane (20  $\mu\text{L}$ ) and left at room temperature overnight. The sample was then rinsed with ethanol and dried under a stream of nitrogen.

**Trichloro(1H,1H,2H,2H-perfluorooctyl)silane SAMs.** A solution of trichloro(1H,1H,2H,2H-perfluorooctyl)silane (30  $\mu\text{L}$ ) in anhydrous toluene (1 mL) was prepared. This solution was placed in a desiccator, in an open vial, adjacent to a plasma-oxidized (10 min, air) silicon substrate (but protected from splash by aluminium foil to avoid solution droplets contacting the substrate). The pressure was reduced using a vacuum pump until the toluene started evaporating (ebullition). The desiccator was closed and the vapour phase deposition was allowed to proceed overnight. The sample was then rinsed with ethanol and dried under a stream of nitrogen.

**Formation of hydroxyl and carboxylic acid (MUDA) SAMs on gold-coated substrates.** Gold-coated silicon substrates were prepared by initial cleaning of the silicon substrate in plasma (5 min, air), followed by the evaporation of a chromium layer (20 nm followed by the evaporation of a gold layer (200 nm) using an Edwards Auto 500 evaporator). The resulting gold-coated silicon substrates were then plasma-oxidized (10 min, air) and then directly immersed in 5 mM thiol ethanolic solutions (hydroxyl and carboxylic acid SAMs were generated from 11-mercapto-1-undecanol and 11-mercaptopundecanoic acid, respectively) at room temperature overnight. The substrates were then washed with ethanol and dried with nitrogen.

## 2.5. Colloidal probe force microscopy

**Preparation of colloidal probe AFM cantilevers.** Cantilevers were selected with a nominal spring constant of  $0.38 \text{ N m}^{-1}$  and this value was more accurately determined using the Sader calibration method.<sup>55</sup> Considering the proper alignment of beads with the longitudinal axis of a cantilever, recent work has shown that the spring constant of the cantilever is not significantly affected by colloidal attachment,<sup>56</sup> and it was therefore decided to measure the spring constant of cantilevers prior to bead attachment. A small volume of polymer brush functionalised-bead suspension was deposited onto a silicon wafer and the solvent was allowed to evaporate overnight; this resulted in a sparse arrangement of functionalised-beads at the silicon surface. In order to attach individual beads to the apex of an AFM cantilever, a custom built AFM (Attocube GmbH, Germany) integrated within an SEM (Quanta 3D FEG, FEI, EU/USA) was used, as described in previous work.<sup>57,58</sup> An integrated focused ion beam (FIB) was used to etch off a portion of the AFM tip to produce a blunted surface to which a bead can be attached. Fig. S1A and B (ESI<sup>†</sup>) show the AFM tip before and after exposure to the FIB. Next, the AFM tip was translated to a droplet of vacuum compatible glue (SEMGLU). Removal of the tip from the glue resulted in the deposition of a small amount

of glue at the apex of the tip, which was translated into contact with a single bead. This SEM system, combined with a high-resolution piezoelectric stage, allows the accurate positioning of the bead onto the apex of the tip, resulting in the accurate alignment of the bead with the longitudinal axis of the AFM cantilever. Focusing a high current electron beam of 1 nA or more for 5 min causes curing of the glue and subsequent firm attachment of the bead to the tip. Fig. S1C (ESI<sup>†</sup>) shows the AFM tip in contact with the bead on the silicon surface and Fig. S1D (ESI<sup>†</sup>) shows the resulting colloidal AFM tip after the curing of the adhesive.

**Operational set up of force probe microscope.** Force measurements were acquired using an NT-MDT Ntegra AFM rig operated in conventional force mode. The cantilever was approached towards the substrate surface and retracted away from the surface at a constant speed ( $1 \mu\text{m s}^{-1}$  for all experiments) by the piezoelectric stage. Silicon and gold substrates, glued to the bottom of a Petri dish, were submerged in the corresponding solution and allowed to equilibrate for a minimum of 15 min. AFM tips were washed in deionised water before and after any testing was carried out, or before the ionic concentration of the testing solution was changed. The pH of solutions was kept at  $7.0 \pm 0.4$  during all testing (Mettler Toledo, SG2 – SevenGo pH Meter), except for experiments carried out in PBS, which displayed a pH of 7.4. For the characterisation of cell monolayers, confluent monolayers grown on glass coverslips (fixed with PFA) were carefully glued to the Petri dish using Loctite Super Glue Precision. The glue was left to dry for a couple of minutes, during which time a small amount of PBS was pipetted onto the upper surface to ensure the cells remained hydrated. The cells were then submerged in PBS and left in the fridge until testing was carried out. Soft tissue samples were cut to a surface area of approximately  $20 \text{ mm}^2$ , with a thickness of approximately 5 mm. These were blotted dry with tissue paper and then glued to a Petri dish; the glue was left to set for a couple of minutes and then samples were submerged in PBS and left in the fridge until testing. All adhesion tests on cell monolayers and soft tissues were carried out in PBS. In between tests on different samples AFM tips were submerged in deionised water and then submerged in a versene solution (0.48 mM) for 5 min, and again washed in deionised water for 2 min. For every sample and condition tested, a minimum of three independent samples were characterised with a minimum of two different functionalised colloidal probes. Each repeat involved probing at least three different areas of a substrate, and for each of these scanned areas 100 indentation curves were performed in a grid pattern with areas ranging from 20 to  $60 \mu\text{m}^2$ .

**Quantitative analysis of AFM lift profiles.** Calibrations were carried out for each AFM tip on a hard non-functionalised silicon substrate, using a custom-built Matlab script for conversions. The parameters extracted are illustrated in Fig. 1. The detachment force is the maximum negative force reached on the cantilever. The detachment work is the total work required to fully detach the colloid from the substrate. The detachment length is defined as the distance between the colloidal probe and the substrate at the maximum negative force value



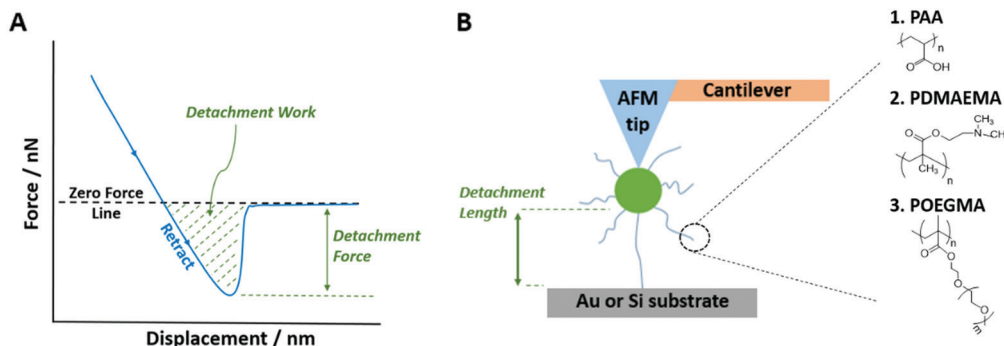


Fig. 1 Quantitative analysis of colloidal probe adhesion profiles. (A) Extraction of detachment work and force from AFM retraction curves. (B) Physical representation of the detachment length, taken from the retraction portion of the AFM curves, and the three different polymers which are used to functionalise the colloidal probes. The detachment length is defined as the distance between the colloidal probe and the substrate at the maximum negative force value.

(the detachment force), taking into account both the piezo displacement and cantilever deflection.

For measurements on soft tissues, the Young's modulus was additionally characterised from the AFM curves. This parameter was quantified using a custom-built Matlab script based on the Oliver–Pharr method for nanoindentation.<sup>59</sup>

## 2.6. Preparation of cell monolayers

**HaCaT cell culture and seeding.** HaCaT cells (human keratinocyte cell line) were cultured in DMEM containing 10% foetal bovine serum, 1% L-glutamine (200 mM) and 1% penicillin-streptomycin (5000 U mL<sup>-1</sup>). HaCaT cells were harvested with trypsin and versene solutions in a ratio of 1/9, centrifuged, counted and resuspended in DMEM at the desired density before seeding onto substrates, in a 24-well plate, at a density of 20 000 cells per well (13 000 cells per cm<sup>2</sup>), and left to form a confluent monolayer for 24 h in an incubator (37 °C and 5% CO<sub>2</sub>). After three aspirations with PBS, confluent monolayer samples were fixed with 4% formaldehyde solution in PBS (PFA) for 10 min, washed with PBS and left in the fridge ready for AFM measurements.

**Primary keratinocyte culture and seeding.** Human primary keratinocytes (HPKs) isolated from neonatal foreskin were cultured on collagen I (type I, 20 µg mL<sup>-1</sup> in PBS for 20 min) treated T75 flasks in keratinocyte serum-free medium (KSFM), supplemented with bovine pituitary extract (BPE) and EGF (Human Recombinant). Keratinocytes were harvested with trypsin and versene solutions in a ratio of 1/9, centrifuged, counted and re-suspended in KSFM at the desired density before seeding onto substrates. Glass coverslips (13 mm) were sterilized by autoclave and put in 24 well-plates. Glass coverslips were first treated with poly(L-lysine) (PLL, final concentration: 100 µg mL<sup>-1</sup>) followed by treatment with fibronectin solution (final concentration: 10 µg mL<sup>-1</sup>). Cells were seeded at a density of 20 000 cells per well (13 000 cells per cm<sup>2</sup>) and left to form a confluent monolayer for 24 h in an incubator (37 °C and 5% CO<sub>2</sub>). After three aspirations with PBS, confluent monolayer samples were fixed with 4% PFA for 10 min, washed with PBS and left in the fridge ready for AFM measurements.

**Neuraminidase and heparinase III treatments.** Neuraminidase was used to cleave the glycosidic linkages of neuraminic acids.

Keratinocyte monolayers (cultured on coated glass coverslips) were treated with neuraminidase (1 U mL<sup>-1</sup>) in KSFM medium for 1.5 h in a 24 well-plate. Heparinase III was used to cleave 1–4 linkages between hexosamine and glucuronic acid residues in heparan sulphate. Keratinocyte monolayers (cultured on coated glass coverslips) were left in heparinase III solutions (0.2 U mL<sup>-1</sup>, in KSFM) for 1.5 h in a 24 well-plate.

**Staining and microscopy.** Wheat germ agglutinin (WGA, 10 µg mL<sup>-1</sup>) and anti-heparin/heparan sulfate antibodies (10 µg mL<sup>-1</sup>) were used to stain the glycocalyx on the cell membrane. Dapi solutions (stock concentration: 5 mM, 1:1000) were prepared by dissolving 5 mg 4',6-diamidino-2-phenylindole dihydrochloride in 1750 µL deionised water. Goat anti-mouse IgG (H + L) secondary antibodies conjugated to AlexaFluor 488 (1 µg mL<sup>-1</sup>) were used for immunofluorescence staining. Fluorescence microscopy images were acquired with a Leica DMi8 fluorescence microscopy (CTR compact lamp; 63 × 1.30 NA, oil lens; DFC9000 GT camera). For each sample and condition three fluorescence microscopy images were taken. Quantification of the fluorescence intensities and profiles was then performed using ImageJ.

## 2.7. Preparation of tissue samples

Fresh soft tissue samples were obtained from C Humphreys & Sons Abattoir in Chelmsford, Essex. Both the epicardial and gingival samples were extracted from freshly slaughtered 6 to 7-month-old pigs. The animals were sacrificed and the relevant parts were extracted at the abattoir; these were then delivered by courier to the lab on the same day as the slaughter. The samples were then washed and stored in PBS overnight at 5 °C. All AFM adhesion testing was then carried out within 48 hours from the initial sacrifice of the animals.

Gingival samples were extracted using a scalpel from the lingual side of the lower mandible. These samples were taken from the region of the keratinized attached gingiva which is the gingival portion that lies between the free gingival groove and the mucogingival junction (Fig. S2B, ESI†). Epicardial samples were extracted from the wall of the left and right ventricles. These samples were taken from the healthy areas of the myocardium away from any major adipose tissue deposits or



prominent blood vessels (Fig. S2C, ESI†). For AFM testing, results were obtained across samples from at least two different animals for each condition tested. Fig. S2A (ESI†) shows an example of loaded epicardial samples in the AFM.

## 2.8. Statistics

A one-way ANOVA test with Tukey's *post hoc* analysis was used to determine statistical significance. In the case of the quantification of the glycocalyx, the analysis was carried out pairwise. For box and whisker diagrams the box represents the 1st, 2nd and 3rd quartiles, as standard, and the whiskers represent the standard deviations of data sets. Mean values for the data sets are also shown. In all other figure types (and for in-text referencing) standard errors are reported.

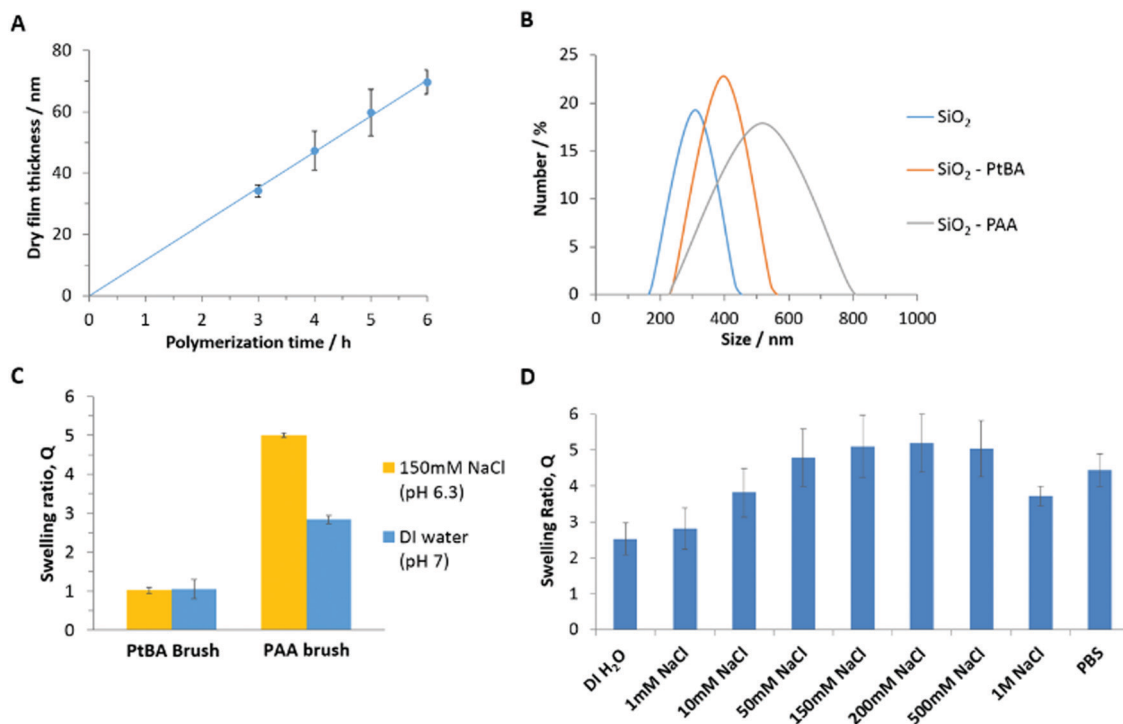
# 3. Results and discussion

## 3.1. Development of polymer-brush functionalised colloidal probes

To study polymer brush interactions with cell monolayers and tissues, we grew polymer brushes from silica microparticles (19.6  $\mu\text{m}$ ), *via* atom transfer radical polymerization (ATRP). The growth of PDMAEMA and POEGMA brushes from silica particles was directly adapted from protocols previously reported.<sup>29,60</sup> To generate PAA brushes, we opted for the controlled growth of PtBA brushes, prior to their conversion upon mild treatment

with dilute trifluoroacetic acid.<sup>52,61</sup> The growth kinetics of the PtBA brushes (monitored *via* ellipsometry) was 3–4 fold faster than that previously reported in similar conditions, but appeared relatively linear (Fig. 2A). However, Lego *et al.* had monitored brush growth *via* AFM, potentially underestimating thicknesses measured, due to the compression of the brush by the AFM tip.<sup>62</sup> In addition, the silane monolayers generated in the present study may display a higher density, resulting in higher brush densities and thickness growth profiles, although direct comparison is not possible.

The deprotection of PtBA brushes was subsequently monitored by ellipsometry (Fig. S3, ESI†), indicating a rapid decrease in the dry thickness of the brush upon incubation in DCM/TFA solutions, due to the loss of *tert*-butyl groups and associated chain relaxation. After overnight incubation in the DCM/TFA solution, the polymer thickness dropped to around 50% of its original value. This is in line with previous reports<sup>61</sup> and with the predicted reduction in molar mass of repeat units and associated dry film thickness (52%).<sup>63</sup> The FTIR spectra of PtBA brushes before and after deprotection confirmed the full conversion to PAA brushes (Fig. S4, ESI†), with the clear disappearance of the typical *tert*-butyl bending bands at 2976, 1392 and 1367  $\text{cm}^{-1}$ , and the shift (and broadening) of the carbonyl band from 1730 to 1712  $\text{cm}^{-1}$ .<sup>61,64</sup> This conversion is further confirmed by XPS spectra of the PtBA brushes before and after deprotection (Fig. S5, ESI†). The  $\text{C}_{1s}$  range of the spectra was deconvoluted into three peaks corresponding to



**Fig. 2** Functionalisation of silicon substrates and silica microparticles with PAA brushes. (A) Kinetics of the growth of PtBA polymer brushes from silicon substrates, monitored by ellipsometry. (B) Dynamic light scattering data of functionalised and non-functionalised 300 nm silica microparticles.  $\text{SiO}_2$  beads and PtBA functionalised beads were dispersed in ethanol and PAA in deionised water. (C) Ellipsometric swelling ratio measured for PtBA and PAA brushes in deionised water (DI) and 150 mM NaCl aqueous solutions. (D) Ellipsometric swelling ratio of PAA brushes (pH kept at  $7.0 \pm 0.4$ , except for PBS which had a pH of 7.4).



carbon atoms associated with C–C, C–O and C=O peaks. The peak associated with carbon atoms involved in C–O bonds was found to shift and broaden upon deprotection. In addition, the atomic percentages and corresponding binding energies of the  $C_{1s}$  and  $O_{1s}$  signals were extracted from the XPS survey spectra (Fig. S5C, ESI†), clearly indicating a reduction in the carbon content relative to oxygen following the deprotection of tBA groups. Both observations are in accordance with the expected change in the chemical structure of the brush, and in agreement with previously reported XPS characterisation data for these surfaces.<sup>65–67</sup>

The hydrodynamic diameter of nanoparticles coated with PtBA and PAA brushes was compared to that of uncoated nanoparticles (see Fig. 2B). The diameter of uncoated nanoparticles was found to be  $320 \pm 10$  nm and increased to  $390 \pm 10$  nm and  $530 \pm 10$  nm for  $SiO_2$ –PtBA and  $SiO_2$ –PAA particles, respectively. This is in good agreement with the expected dry thickness of PtBA brushes (not swollen in ethanol), which were found to grow to 60 nm in 5 h, from silicon substrates (Fig. 2A). The significant increase in hydrodynamic diameter observed for PAA-coated nanoparticles is in agreement with their pronounced swelling in deionised water (the diameters measured correspond to a swelling greater than 3) and the decrease in hydrophobicity associated with the deprotection of tBA groups. In addition, the  $\zeta$ -potentials measured for  $SiO_2$ –PtBA and  $SiO_2$ –PAA particles were  $23.1 \pm 0.1$  mV and  $-29.3 \pm 0.1$  mV, respectively, although these values are only qualitatively indicating the charge of the associated particles (PtBA particles did not resuspend in aqueous solutions, therefore preventing direct comparison with PAA-coated particles). Hence the charge reversal of the  $\zeta$ -potential is consistent with a significant change in surface chemistry of these particles and the introduction of negatively charged carboxyl groups following the deprotection step.

The swelling of the corresponding brushes was further characterised *via in situ* ellipsometry. The swelling ratio ( $Q$ ), defined as the ratio between the wet ( $L$ ) and dry ( $D$ ) ellipsometric thicknesses ( $Q = L/D$ ), was close to unity prior to deprotection of PtBA brushes (Fig. 2C), as expected from the hydrophobicity of *tert*-butyl acrylate repeat units. In contrast, PAA brushes were characterised by high swellings in deionised water and 150 mM NaCl solutions; this is in agreement with the expected behaviour of weak polyelectrolytes in the osmotic regime, characterised by high proton dissociation at neutral pH, substantial electrostatic repulsion between repeat units and the associated stretching of polymer chains.<sup>39,63,68</sup> The swelling ratio of PAA brushes was further characterised in PBS and in aqueous NaCl solutions with a range of ionic strengths (Fig. 2D). The swelling was observed to initially increase with the ionic strength, due to increasing proton dissociation and associated hydrophilicity. As the salt concentration increased further, the brush appeared to enter the neutral brush and salted brush regime and electrostatic interactions became largely screened, resulting in a decrease in brush thickness.<sup>63,68</sup> Finally, contact angle measurements were in agreement with the increase in hydrophilicity and brush swelling. Following the deprotection of PtBA brushes, average contact angles shifted from

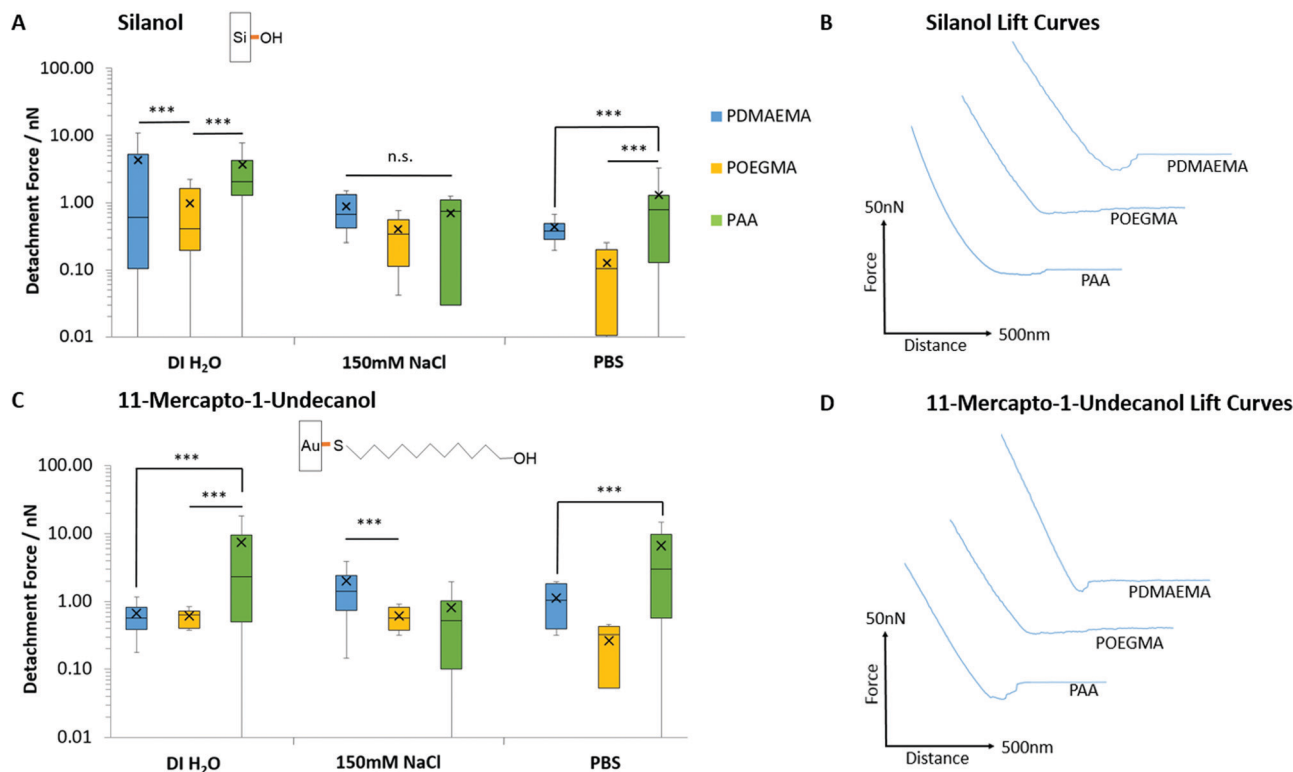
$87.0 \pm 0.6^\circ$  to  $37.4 \pm 0.7^\circ$  for PtBA and PAA brushes, respectively (Table S1, ESI†).

### 3.2. Polymer brush adhesion to SAMs

Prior to studying polymer brush adhesion to cells and tissues, we first investigated their adhesion to model substrates (self-assembled monolayers) presenting chemical moieties with a range of hydrophilicities, hydrogen bonding and charge. Fig. 3 presents the detachment forces observed for the adhesion of PAA, PDMAEMA and POEGMA brushes to hydroxyl-terminated monolayers alongside the corresponding representative lift curves. Overall, similar adhesions were measured to both 11-mercapto-1-undecanol monolayers assembled to gold and silicon oxide interfaces. As the surface packing density for monolayers assembled on silica<sup>24,69</sup> and gold<sup>70</sup> substrates are reported to be in the region of 5 molecules per  $nm^2$ , the similarity of these adhesion profiles is justified. POEGMA displayed little adhesion to these interfaces, in good agreement with its neutral structure, lacking strong proton acceptor or donor functions. POEGMA brushes indeed typically display moderate hydrophilicity and excellent anti-fouling properties.<sup>71,72</sup> In contrast, PAA and PDMAEMA displayed moderate adhesions to both surfaces. At the near-neutral pH at which these measurements were carried out (the pH of deionised water and NaCl solutions was in the range of  $7.0 \pm 0.4$ , whereas that of PBS solutions was 7.4), PAA brushes are globally negatively charged (we found a  $\zeta$ -potential of  $-29$  mV)<sup>73,74</sup> and PDMAEMA brushes are positively charged ( $\zeta$ -potential of 40 mV),<sup>29,75</sup> whereas silica substrates are negatively charged ( $\zeta$ -potential of  $-40$  mV).<sup>74,76</sup> Hence, in the case of PDMAEMA brushes, electrostatic interactions should lead to increased adhesion. However, PAA and PDMAEMA brush adhesion (detachment force) to silanol and 11-mercapto-1-undecanol monolayers are typically comparable and, in some cases, increased in the case of PAA (in deionised water and in PBS, in the case of 11-mercapto-1-undecanol monolayers; Fig. 3A and C). In addition, higher ionic strength (150 mM NaCl) did not lead to substantial reduction in adhesion force, as would be predicted in the case of oppositely charged surfaces. These changes were qualitatively reflected in the corresponding detachment work (see Fig. S6A and B, ESI†). Therefore, weak adhesion of PAA and PDMAEMA brushes to moderately hydrophilic and moderately charged silanol and 11-mercapto-1-undecanol monolayers appeared to be dominated by weak hydrogen bonding. In PBS solutions, adhesion forces measured for PDMAEMA and POEGMA brushes remained comparable to those measured in 150 mM NaCl solutions. Indeed, PBS has a NaCl concentration of 137 mM, similar to that of the NaCl solution used, and the pH of both solutions were relatively close (pH 7.4 and pH  $7.0 \pm 0.4$ , for PBS and NaCl, respectively). For PAA brushes, adhesion forces were increased in PBS compared with 150 mM NaCl (forces increased from 0.69 to 1.29 nN and 0.82 to 6.60 nN for silanol and 11-mercapto-1-undecanol SAMs, respectively). This phenomenon is associated with a reduction of the swelling of PAA brushes in PBS, compared to 150 mM NaCl, and may indicate the contribution of phosphate to hydrogen bonding at the surface.







**Fig. 3** Detachment forces and corresponding representative lift curves for the adhesion of PAA, PDMAEMA and POEGMA brushes to hydroxyl-terminated monolayers. Representative curves are taken from adhesion experiments performed in deionised water. (A) Detachment forces to silicon oxide interfaces. (B) Representative lift curves for adhesion to silanol monolayers. (C) Detachment forces to 11-mercapto-1-undecanol monolayers assembled on gold-coated silicon substrates. (D) Representative lift curves for adhesion to 11-mercapto-1-undecanol monolayers. Data is plotted as means, with box plots. \*\*\* $p \leq 0.001$ . n.s., non-significant.

Fig. 4 presents the adhesion forces measured for the three different polymer brushes to charged model substrates (associated detachment work values are given in Fig. S6, ESI†). As expected, negatively charged PAA brushes strongly adhered to positively charged quaternary ammonium monolayers (QAPTES); this is further evidenced in Fig. 5A where the jump-to-contact is indicative of a strong attractive interaction. In the corresponding representative lift curves (Fig. 5B), it is also clear that interactions between PAA brushes and oppositely charged monolayers (APTES and QAPTES) are strong (adhesion forces  $> 10.16$  nN in deionised water) and decrease at higher ionic strength (Fig. 4D), as would be expected from the corresponding screening of coulombic forces. Indeed, at neutral pH we found  $\zeta$ -potentials of 13 mV<sup>77,78</sup> and 37 mV for APTES and QAPTES, respectively, whereas  $\omega$ -mercaptoundecanoic acid (MUDA) displays a  $\zeta$ -potential of  $-40$  mV.<sup>79</sup> This behaviour contrasts with the adhesion of poly(2-vinylpyridine), which displayed increased adhesion forces at higher ionic strength.<sup>80,81</sup> This was attributed to changes in brush conformation from the salted regime to the osmotic regime. We also note that the ionic strength range at which brush adhesion was maximised was above the range of ionic strength presently tested (matching that of physiological buffers such as PBS). The neutral POEGMA, in contrast, displayed interactions below 3.45 nN, comparable to what was measured on neutral hydroxyl functionalised

interfaces (detachment forces were reduced for hydroxyl SAMs in comparison to charged monolayers, however, from the detachment work (Fig. S6, ESI†), interactions are of a similar scale), in agreement with its neutral charge. The adhesive response of PDMAEMA to these three charged monolayers was more surprising as, although it adhered weakly to APTES surfaces ( $< 4.05$  nN), its adhesion to MUDA was not as significantly increased ( $< 5.93$  nN), as was observed in the case of PAA adhering to APTES interfaces. This indicates that other interactions dominate the adhesive behaviour of PDMAEMA to the charged monolayers studied.

In addition to electrostatic interactions, hydrogen bonding may significantly impact the adhesion strength of PAA and PDMAEMA brushes to charged monolayers, owing to their acid/base character. Indeed, in contrast to PAA brushes, which showed a marked decrease in interactions at increasing ionic strength, the adhesion of PDMAEMA brushes to MUDA monolayers remained unaffected by increasing ionic strength (150 mM NaCl vs. deionised water; Fig. 4C). At neutral pH (and slightly below), PDMAEMA is only partially charged and a substantial proportion of amine moieties are not protonated.<sup>29,82</sup> Similarly, APTES and MUDA monolayers and PAA brushes will only be partially deprotonated at the neutral pH of the solutions used in this study. Therefore, hydrogen bonding between polymer brushes and APTES and MUDA monolayers may significantly



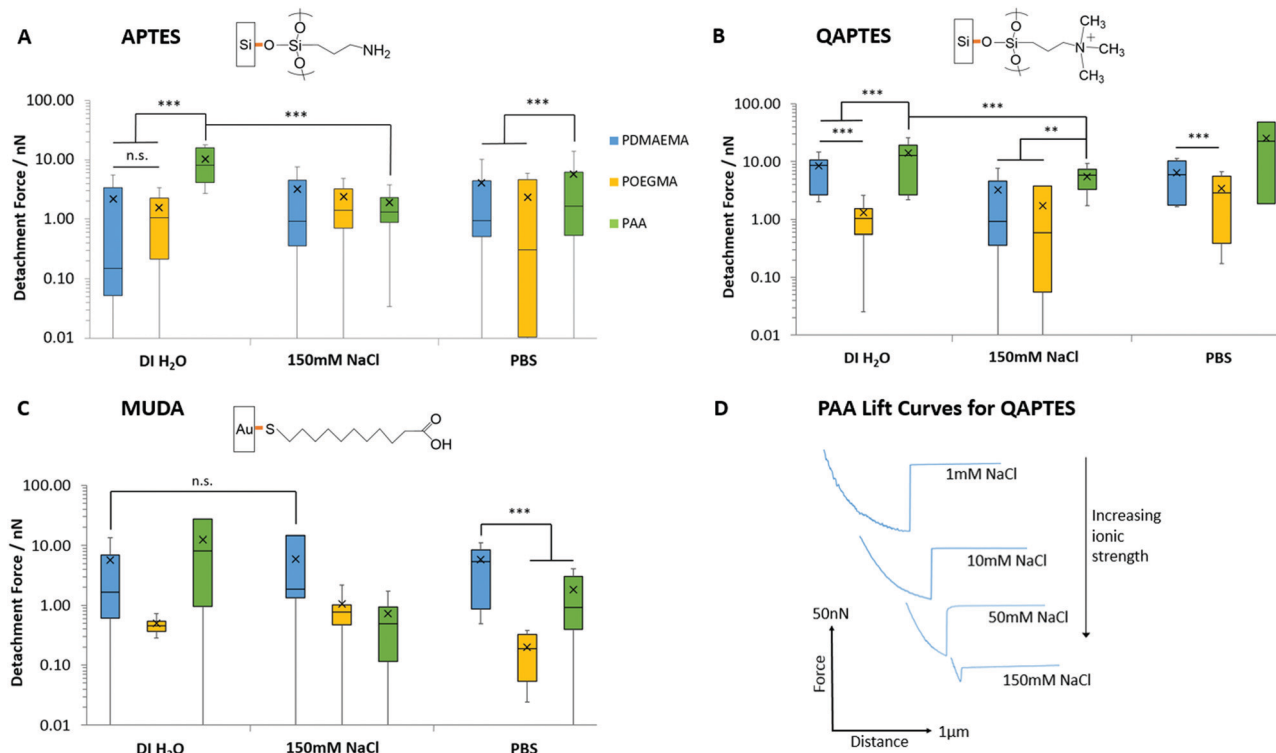


Fig. 4 Detachment forces measured for the adhesion of PAA, PDMAEMA and POEGMA brushes to charged monolayers. (A) Detachment forces to ammonium (APTES) monolayers. (B) Detachment forces to quaternary ammonium (QAPTES) monolayers. (C) Detachment forces to undecanoic acid (MUDA) monolayers assembled on gold-coated silicon substrates. (D) Representative lift curves for adhesion of PAA to QAPTES monolayers in solutions of increasing ionic strengths. Data is plotted as means, with box plots. \*\* $p \leq 0.01$ . \*\*\* $p \leq 0.001$ . n.s., non-significant.

contribute to bonding and adhesion profiles. Furthermore, although hydrogen bonding is reported to be influenced by the ionic strength of the medium, predicting the impact of electrolytes on hydrogen bonding remains difficult. Hence, cations are known to alter the melting temperature of double-stranded oligonucleotides, although this is *via* their combined impact on coulombic repulsion between phosphates and on hydrogen bonding between bases.<sup>83</sup> Electrolytes were shown to reinforce hydrogen bonding<sup>84</sup> and to perturb networks of intramolecular hydrogen bonds of water molecules.<sup>85</sup> In addition, electrolytes such as phosphates are particularly prone to hydrogen bond and alter interactions with biomacromolecules.<sup>86,87</sup> The relatively high interactions of PAA and PDMAEMA to MUDA and APTES, respectively (Fig. 4 and 5), in particular in PBS solutions, may therefore be explained by hydrogen bonding between the corresponding interfaces, perhaps stabilised by phosphate ions.

Further to the evidence for strong hydrogen bonding between polymer brushes and monolayers, the adhesion behaviour of PDMAEMA brushes to QAPTES also implies some impact of hydrophobic interactions on adhesion strength (Fig. 4B and 5B). Indeed, QAPTES is highly positively charged and cannot directly be involved in hydrogen bonding with other molecules. Therefore, the high charge density of cationic PDMAEMA (at neutral pH) should result in repulsive forces. In contrast to this expected behaviour, we observed no evidence of repulsion in the corresponding landing traces (Fig. 5A) and relatively strong adhesions can be observed in the retraction traces (Fig. 5B).

Quaternary ammoniums such as those of QAPTES are known to display some level of hydrophobicity, enabling the solubility of salts in some organic solvents. In addition, PDMAEMA also shows significant hydrophobicity and a clear pH-responsive behaviour, as detailed above, especially close to its pK<sub>a</sub> and above.<sup>29</sup> Therefore, hydrophobic interactions between these two interfaces are likely to play an important role in determining adhesion profiles.

In order to probe further into the impact of hydrophobic forces in regulating adhesion of PAA, PDMAEMA and POEGMA brushes, we generated octyl and perfluorooctyl SAMs on silicon substrates (with contact angles of  $96.3 \pm 1.3^\circ$  and  $103.7 \pm 1.4^\circ$ , respectively). Significant adhesion was measured for PDMAEMA and PAA brushes, whilst POEGMA brushes displayed lower adhesions, but higher than what was reported for hydrophilic SAMs (Fig. 6; detachment works follow similar trends, see Fig. S6, ESI†). In particular, PDMAEMA displayed strong adhesion to octyl SAMs ( $>5.13$  nN) and the perfluorooctyl SAMs (10.30 nN in 150 mM NaCl solutions). PAA displayed overall weaker adhesion to octyl SAMs, especially at high ionic strength, but relatively high adhesion to perfluorooctyl SAMs. The relatively strong adhesion of weak polyelectrolyte brushes to hydrophobic surfaces is likely due to their partial protonation/deprotonation at neutral pH and associated moderate hydrophobicity.<sup>29,82</sup> Hence, PDMAEMA brushes were found to adhere relatively strongly to PDMAEMA surfaces at neutral pH,<sup>88</sup> whereas little adhesion was observed between two symmetrical



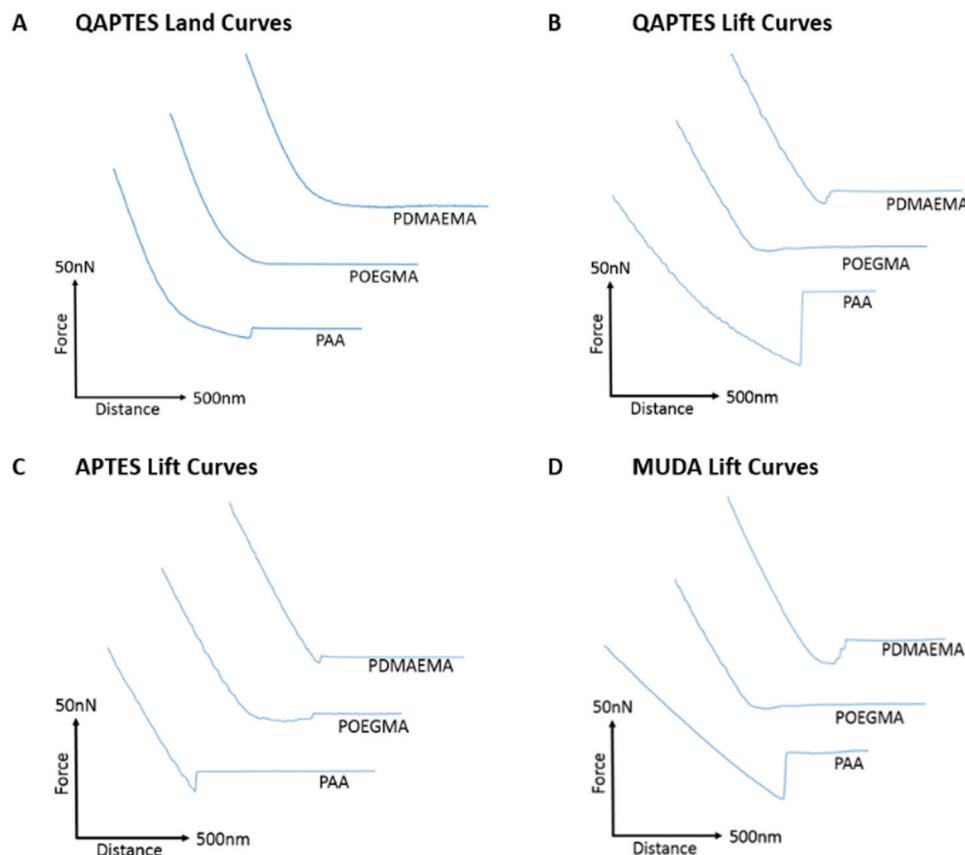


Fig. 5 Representative lift and land curves for the adhesion of PAA, PDMAEMA and POEGMA brushes to charged monolayers. All curves are taken from adhesion experiments performed in deionised water. (A) Land curves representative of adhesion of brushes to QAPTES monolayers. (B) Lift curves representative of adhesion to quaternary ammonium (QAPTES) monolayers. (C) Lift curves representative of adhesion to ammonium (APTES) monolayers. (D) Lift curves representative of adhesion to undecanoic acid (MUDA) monolayers assembled on gold.

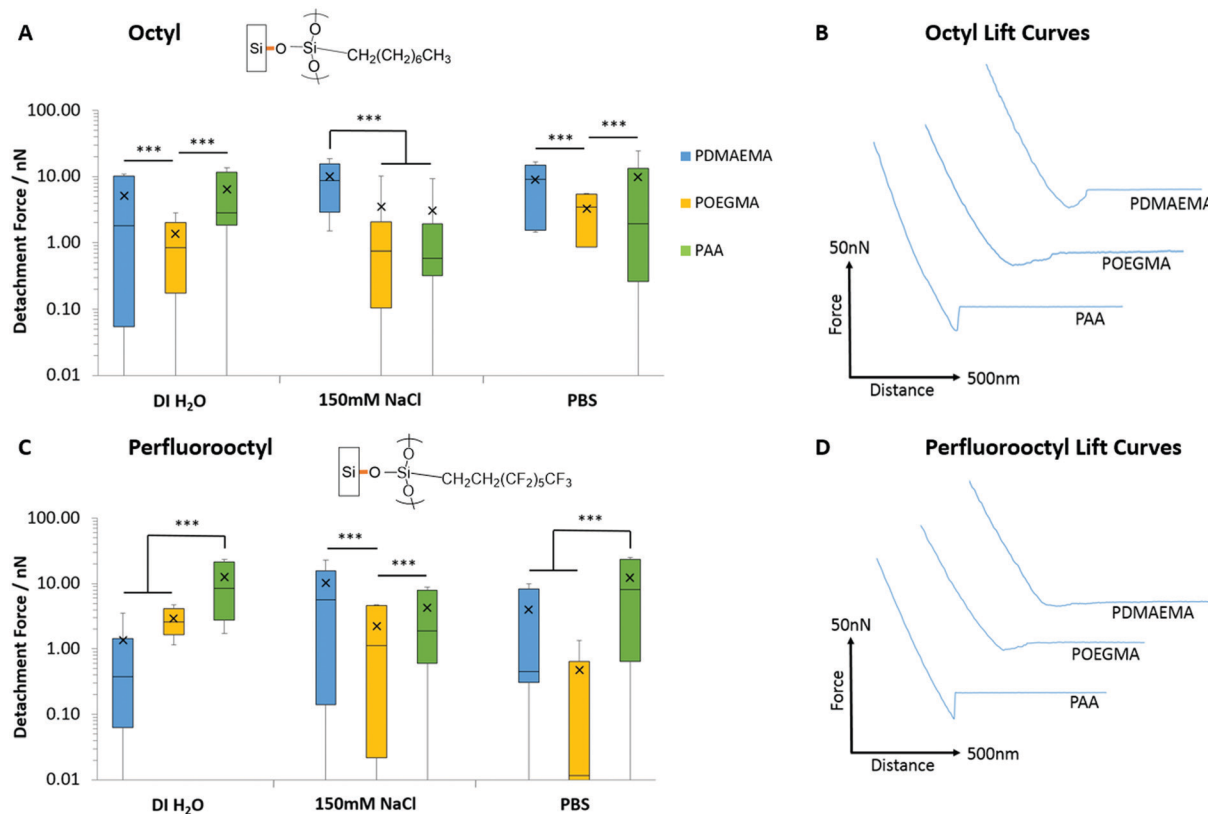
hydrophilic polymer brushes.<sup>89</sup> Similarly, polymer brushes displaying a Lower Critical Solution Temperature (LCST) were reported to strongly adhere (symmetrical interface bonding) above their LCST, but displayed weak interactions below their LCST.<sup>90</sup> Interestingly, the adhesion force of sparse (grafted to) PAA brushes was also reported to be significantly higher to alkyl monolayers than corresponding hydroxyl and carboxylic acid-terminated SAMs,<sup>91,92</sup> although this was for single desorption events rather than the cumulative desorption forces associated with full detachment of the tip. To account for such strong interactions between PAA brushes and hydrophobic SAMs, Friedsam *et al.* proposed that the structure of water close to these interfaces (and the lack of hydrogen bonding of interfacial water molecules, in contrast to the networks formed with hydroxyl and carboxylic acid terminated SAMs) led to easier displacement of water molecules closely associated with the hydrophobic SAMs, compared to hydrophilic SAMs. Indeed, the profiles of desorption of our PAA brushes displayed strong adhesion forces with a sharp detachment step in the case of alkyl SAMs ( $>6.43$  nN detachment force,  $>2.78$  fJ detachment work and  $<22.3$  nm detachment lengths; Fig. S7, ESI†), whereas detachment from hydrophilic silanol SAMs displayed an overall weak adhesion force, but a more gradual detachment profile and increased detachment length (3.69 nN detachment

force, 0.48 fJ detachment work and 68.4 nm detachment length; Fig. S7, ESI†). Such difference in the structure of water at hydrophobic and hydrophilic SAMs is also supported by molecular dynamics studies that give evidence for the occurrence of 0.3 nm gaps between vicinal water and hydrophobic SAMs.<sup>93</sup> Similarly, such water structuring was found to impact the adhesion of moderately hydrophobic peptides presenting catechol residues (DOPA), which showed increased bonding forces to hydrophobic SAMs, despite increased hydrogen bonding to hydrophilic SAMs.<sup>94</sup> Hence, our results also support the occurrence of an aqueous interfacial layer that differentially regulates adhesion of moderately hydrophobic polymer brushes to hydrophobic SAMs (Fig. 7).

### 3.3 Polymer brush adhesion to cell monolayers

Having studied the impact of substrate chemistry in a set of monolayers with a range of hydrophobicity/hydrophilicity, charge and hydrogen bonding potential, we next turned our attention to the adhesion profile of weak polyelectrolyte brushes and POEGMA to epithelial cell sheets. These biointerfaces can be regarded as simplified systems to the understanding of tissue bonding due to the greater cellular homogeneity within cultured cell sheets, their planarity and relative rigidity (at the macroscale, owing to the rigidity of the underlying substrate). We focused on two epithelial





**Fig. 6** Detachment forces and corresponding representative lift curves for the adhesion of PAA, PDMAEMA and POEGMA brushes to hydrophobic monolayers. Representative curves are taken from adhesion experiments performed in deionised water. (A) Detachment forces to octyl monolayers. (B) Representative lift curves for adhesion to octyl monolayers. (C) Detachment forces to perfluorooctyl monolayers. (D) Representative lift curves for adhesion to perfluorooctyl monolayers. Data is plotted as means, with box plots. \*\*\* $p \leq 0.001$ .

models: HaCaT cells (a human epidermal cell line) and primary keratinocytes (stem cells responsible for the homeostasis of the interfollicular epidermis).<sup>95</sup>

Fig. 8A and B present the adhesion profiles and quantification of adhesion forces of POEGMA, PDMAEMA and PAA brushes to cell sheets of keratinocytes and HaCaT cells. Due to the high density and the length of their oligo(ethylene glycol) side chains, POEGMA brushes are known for their excellent protein and bacterial resistance.<sup>71,72</sup> The exact detailed mechanism of their protein resistance remains only partially understood, but was proposed to result from their combined hydration shell and steric hindrance (and high chain densities), restricting the infiltration of biomacromolecules.<sup>49,50</sup> In agreement with these reports, POEGMA brushes were found to display very low adhesion to both HaCaT and primary keratinocyte monolayers, with adhesion forces below 0.29 nN (Fig. 8A and B). Such adhesions are lower than those measured for POEGMA, in the case of hydrophobic or charged SAMs, indicating relatively hydrophilic and weakly charged cell membranes. This is in contrast with the moderate adhesion forces measured between POEGMA brushes and bacteria,<sup>27</sup> which typically display relatively charged and hydrophobic membranes (with lipopolysaccharides).<sup>26</sup>

PDMAEMA displayed relatively weak interactions with both cell monolayers; this is associated with low adhesion forces and detachment work (<1.14 nN and <0.87 fJ, respectively;

Fig. 8A and B and Fig. S8A, ESI†). Although higher than that measured for POEGMA, such low adhesion is surprising considering the high positive  $\zeta$ -potential of PDMAEMA brush-based colloids and their rapid uptake by cells, for example for gene delivery applications.<sup>30,96,97</sup> Hence, it is possible that the rapid fouling of cationic polymer brushes by proteins and components found in the medium, results in the substantial modification of the PDMAEMA brush surface and associated decrease in  $\zeta$ -potential,<sup>98</sup> leading to a masking of short-range hydrophobic interactions and hydrogen bonding.

In contrast, PAA brushes displayed strong adhesions to primary keratinocytes (3.91 nN and 4.41 fJ, respectively; Fig. 8A and B and Fig. S8A, ESI†). This behaviour was associated with substantially longer detachment lengths than those reported for SAMs ( $752 \pm 17$  nm, compared to lengths typically <100 nm for SAMs), suggesting that the retraction of PAA-coated colloids is associated with substantial deformation of the cell membrane, contributing to the overall retraction profile. Strikingly, the adhesion of PAA brushes to HaCaT cells was very low (0.33 nN). To test whether fouling occurred at the brush surface, repeated measurements (600 adhesion and retraction events) were carried out and plotted as a function of cycle number (Fig. S8B, ESI†). The scatter of the data as function of cycle number, compared to the overall average, clearly indicates no significant positive or negative deviations as a function of time. To account for the





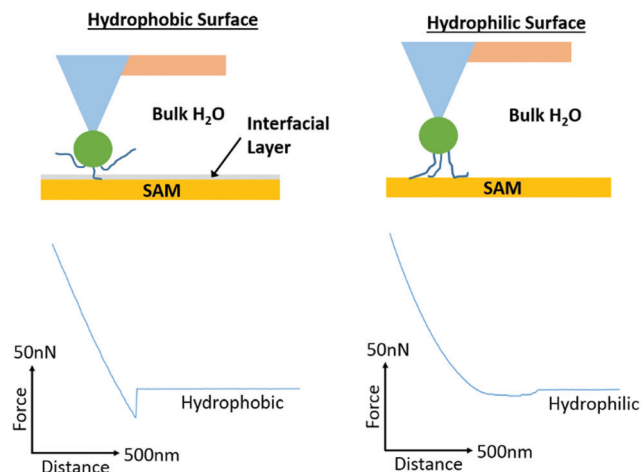


Fig. 7 Proposed mechanism of adhesion of PAA brushes to hydrophobic and hydrophilic monolayers. Representative lift curves are taken from adhesion measurements for alkyl (octyl and perfluorooctyl) and hydrophilic (silanol) SAMs submerged in deionised water. The desorption profiles indicate strong adhesion forces, with a sharp detachment step in the case of alkyl SAMs but overall weak adhesion forces with a more gradual detachment profile for hydrophilic SAMs. These profiles support the occurrence of an aqueous interfacial layer that differentially regulates adhesion of moderately hydrophobic polymer brushes to hydrophobic SAMs.<sup>91,92</sup>

high adhesion of PAA brushes to primary keratinocytes, we proposed that the glycocalyx (a proteoglycan brush-like layer that coats the cell membrane)<sup>99</sup> of the epithelial cell sheets studied differed. Staining of the glycocalyx with wheat germ agglutinin (WGA) and specific immunostaining of the heparin component of the glycocalyx indeed revealed striking differences in the abundance and localisation of the glycocalyx in primary keratinocytes and HaCaT cells (Fig. 9).

Heparin was found to be expressed at lower levels in HaCaT cells, compared to primary keratinocytes (Fig. 9A and B). In contrast, non-specific staining of the glycocalyx using WGA did not show the same trend; instead of a diffuse relatively homogenous staining, the glycocalyx was strongly localised at cell-cell junctions in the case of HaCaT monolayers (Fig. 9A). With primary keratinocytes, the glycocalyx was spread more uniformly on the apical membrane, with little sequestration at cell junctions. This was quantified by plotting the intensity profile of WGA sequestration in HaCaT cells and keratinocytes. This differential regulation of the localisation of the glycocalyx is proposed to result from the culture conditions: HaCaT cells were cultured in normal DMEM medium, which contains high levels of  $\text{Ca}^{2+}$ , enabling the formation of cell junctions, whereas keratinocytes were cultured in KSFM, a medium containing low levels of  $\text{Ca}^{2+}$ , in which cadherin-mediated junctions are typically not stabilised.<sup>100</sup> Overall, these results indicate that the localisation and abundance of the glycocalyx in HaCaT cells and keratinocytes differs significantly, and this correlates with the changes in adhesion measured for polyelectrolyte brushes to the corresponding cell monolayers.

To further test the impact of the glycocalyx on colloidal probe adhesion to primary keratinocytes, we treated cell sheets

with neuraminidase (broad-spectrum enzyme cleaving the glycocalyx) and heparinase (enzyme specifically cleaving heparin components).<sup>101</sup> The efficiency of such cleavage was confirmed by staining and fluorescence microscopy (Fig. 9B). Following such treatment, we found that PAA brush adhesion was significantly reduced (Fig. 8C and Fig. S8C, ESI†). In particular, heparinase treatment resulted in a reduction of adhesion forces to levels comparable to POEGMA adhesion on untreated cells. Neuraminidase treatment had a more modest impact, indicating that, although cleavage of heparin accounts for most of the adhesion strength of PAA brushes to cell membranes, full cleavage of the glycocalyx reveals buried domains and potentially directly exposes the phospholipid bilayer, accounting for the moderate adhesion measured. These observations are in good agreement with the work of Servais *et al.*, who reported that the adhesive force of pectin/CMC formulations to the mesothelium of several different tissues (investigated *via* uniaxial tensile strength tests using a customized apparatus for load/displacement measurements) was reduced after treatment of these samples with neuraminidase; in some cases this was by as much as 50%.<sup>102</sup> The dependence of weak polyelectrolyte adhesion to cells, mediated by the glycocalyx, is therefore demonstrated across multiple length scales. Similarly, enzymatic cleavage of the glycocalyx components had a significant impact on the adhesion of PDMAEMA brushes, indicating that steric repulsion may also be responsible for the weak adhesion profile of these brushes to cell monolayers; this further suggests that fouling of the PDMAEMA surface is responsible for such low adhesion, despite the absence of medium or serum in the testing conditions. Surprisingly, the adhesion of POEGMA brushes slightly increased after enzymatic treatment of primary keratinocytes (to 0.55 and 0.42 nN after heparinase and neuraminidase treatment, respectively). This suggests that such enzymatic treatment leads to the exposure of residues, perhaps with higher hydrophobicity, as this was a particular type of interaction that promoted stronger adhesion of POEGMA brushes with SAMs.

### 3.4. Polymer brush adhesion to soft tissue samples

We next studied the adhesion of polymer brushes to soft epithelial tissues: the gingiva (which structure and homeostasis is regulated by gingival keratinocytes) and the epicardium (a membrane to which adhesion is particularly relevant for epicardial placement strategies).<sup>103,104</sup> Fig. 10 presents the detachment force and work measured during the adhesion of PDMAEMA, POEGMA and PAA brushes to porcine gingiva and epicardium samples. Overall, interactions of polymer brushes were stronger with the epicardium compared to the gingiva. As expected, adhesion between POEGMA brushes and both tissue types was minimal, consistent with the protein resistance of this polymer brush. Adhesion forces and work of PAA and PDMAEMA brushes to gingival epithelium were increased compared to that of POEGMA but remained overall relatively weak (<2.72 nN). In contrast, the adhesion of PAA and PDMAEMA brushes to the epicardium increased significantly (3.58 and 5.67 nN, respectively). Hence PDMAEMA was found to



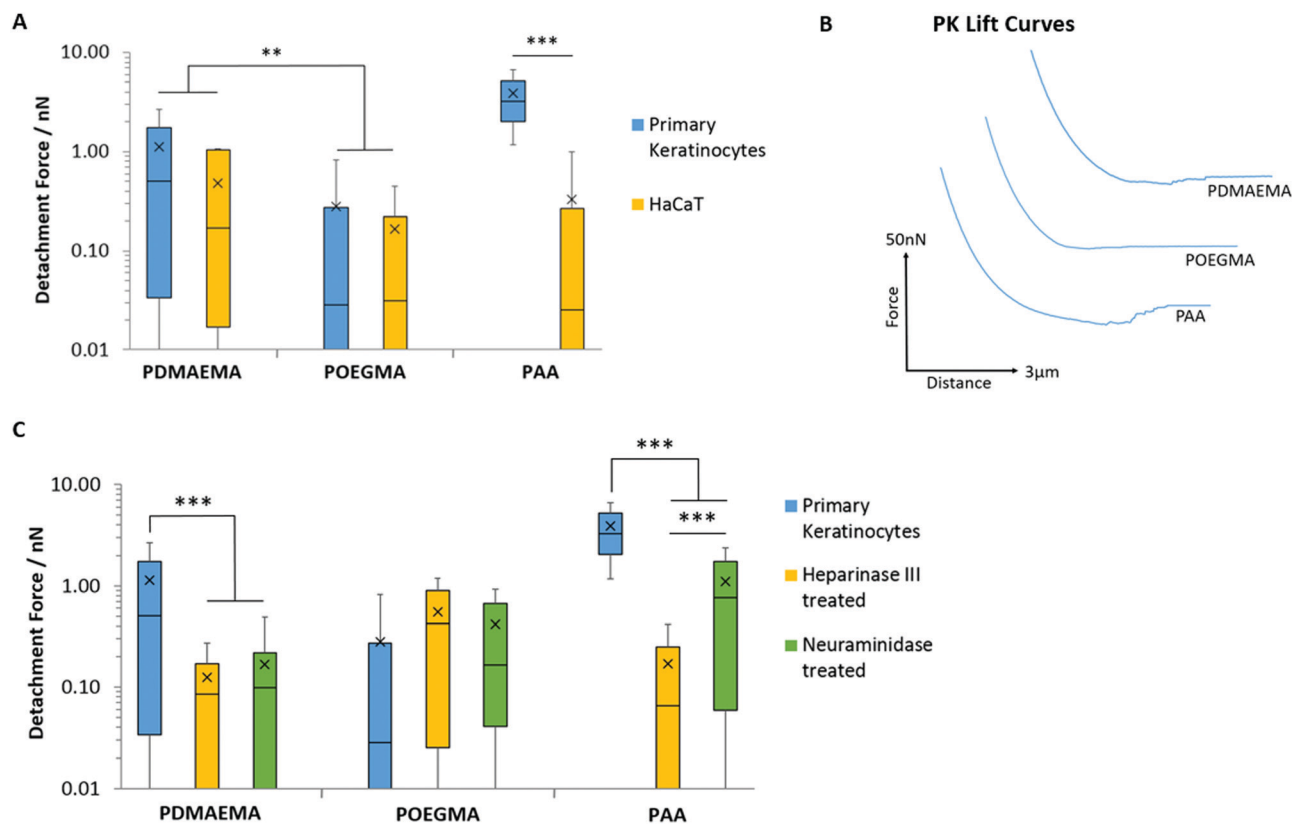


Fig. 8 Characterisation of adhesive interactions between polymer brushes and cell monolayers. Testing was carried out on samples submerged in PBS. (A) Detachment forces between polymers and primary keratinocyte (PK) and HaCaT cell monolayers. (B) Representative AFM lift curves displaying the interaction between the polymers and primary keratinocyte cells. (C) Detachment forces between polymers and primary keratinocyte monolayers with and without enzymatic treatment. Data is plotted as means, with box plots.  $**p \leq 0.01$ .  $***p \leq 0.001$ .

adhere relatively strongly to the epicardium, perhaps reflecting a higher coulombic attraction or hydrogen bonding with this tissue. The adhesion of PAA brushes to the epicardium was in line, although slightly lower than that measured to primary keratinocyte monolayers. These differences in adhesion to the epicardium and gingiva likely reflect differences in the composition of the cell surface, and in particular that of the glycocalyx in these two tissues.

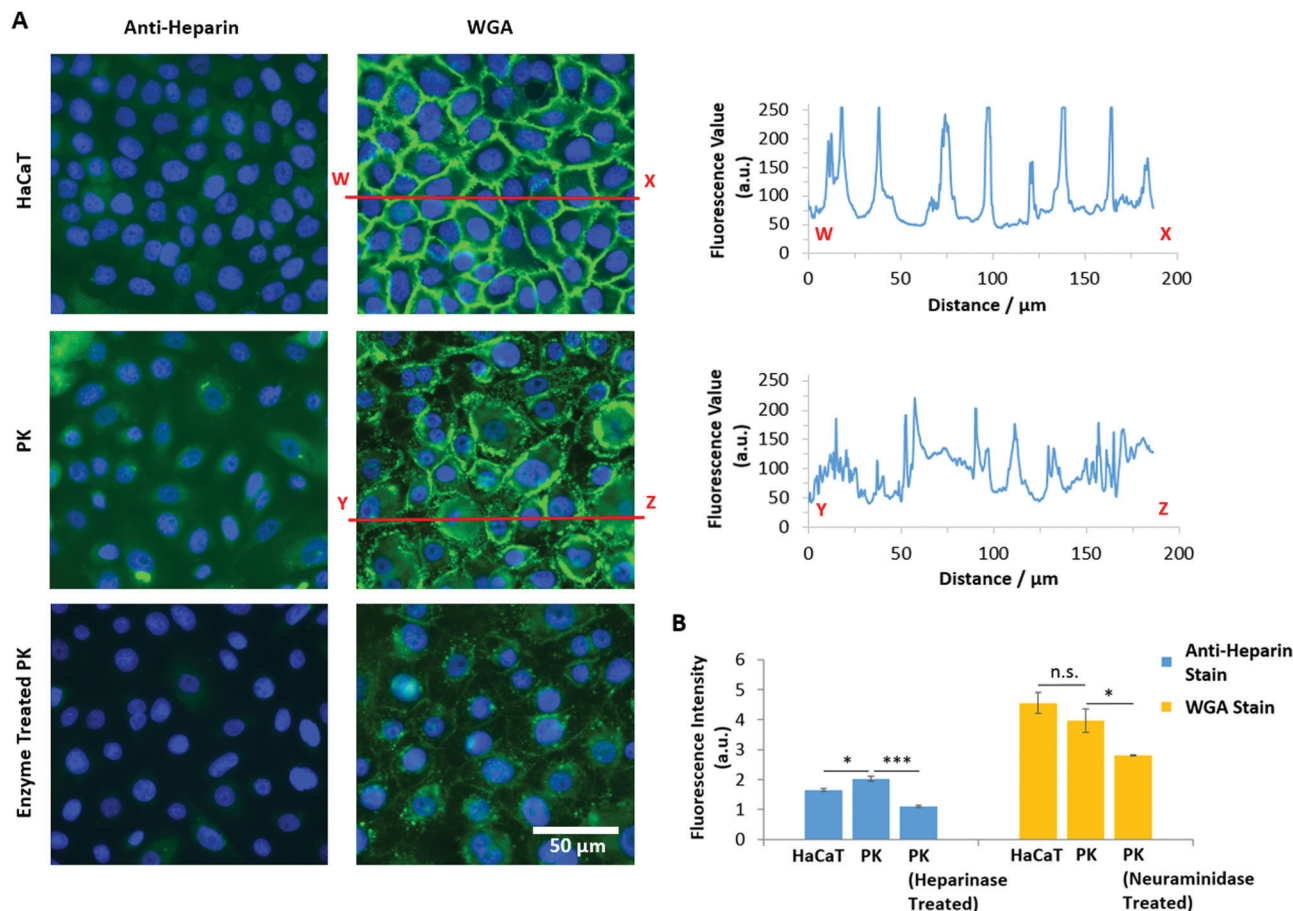
To the best of our knowledge, the epicardial glycocalyx has not yet been characterised. The parietal and visceral pericardium (epicardium) are known to be similar in structure, both comprising a serosal (mesothelial cell) component adjacent to a fibrous tissue layer.<sup>105</sup> The parietal pericardium has been shown to display a rich glycocalyx coating, in particular, rich in sialic acid residues.<sup>106</sup> Assuming there is similarity in the glycocalyx structure of both of these layers, the higher level of sialic acid content at the surface of the epicardium could qualitatively account for the strong adhesion observed for PDMAEMA brushes to this tissue.

In the oral mucosa, the epithelium forms a keratinized layer in which the glycocalyx interpenetrates with the salivary film.<sup>107</sup> The presence of a glycocalyx layer on the surface of corneocytes has been evidenced *via* electron microscope,<sup>108,109</sup> but to the best of our knowledge the exact composition of this layer is yet to be determined. The salivary film, which is estimated to be

70–100  $\mu\text{m}$  thick,<sup>110</sup> contains several molecules which include a large concentration of mucins<sup>111</sup> and a number of different types of bacterial species. These bacteria have been shown to adhere to each other, as well as to the surface of the oral epithelium *via* a network of their glycocalyx.<sup>112,113</sup> As such, there are a number of potential molecules that could contribute to polymeric adhesion to the keratinized gingiva; additional enzymatic studies will be required to identify the specific molecules present and the role that each plays in adhesion.

However, it should be noted that significant differences in tissue stiffness could account for at least some of the differences in adhesion measured between these two tissues. Indeed, the epicardium was found to be significantly softer than the gingival epithelium. This can be clearly seen in the large retraction lengths measured for the epicardium, especially for PDMAEMA brushes, compared to those measured for the gingiva and even cell monolayers (Fig. 10C and D). To confirm such differences in stiffness at the microscale, we measured the Young's modulus of these tissues, based on the curves obtained from the AFM adhesion data, using the Oliver–Pharr method to quantify corresponding moduli.<sup>59</sup> The gingiva was found to be significantly stiffer than the epicardium ( $1020 \pm 130$  and  $20.7 \pm 0.5$  kPa, respectively), suggesting that the colloidal probe indentation may result in substantial conformal deformation of the epicardium, and therefore an associated increase in





**Fig. 9** Quantification of the glycocalyx in primary keratinocytes (PK) and HaCaT cells through staining with wheat germ agglutinin (WGA) and specific immunostaining of the heparin component of the glycocalyx using anti-heparin. (A) Representative fluorescence images (blue, DAPI; green, anti-heparin or WGA) for HaCaT cells and primary keratinocytes. Primary keratinocytes were treated with heparinase III and neuraminidase and staining was carried out using anti-heparin and WGA, respectively. Cross-sections in the fluorescence values illustrate the differing localisation of the glycocalyx for HaCaT and primary keratinocyte cells. (B) Fluorescence intensity values for HaCaT cells, primary keratinocytes and enzyme treated primary keratinocytes, stained with either anti-heparin or WGA. Data is plotted as means.  $*p \leq 0.05$ .  $***p \leq 0.001$ . n.s., non-significant.

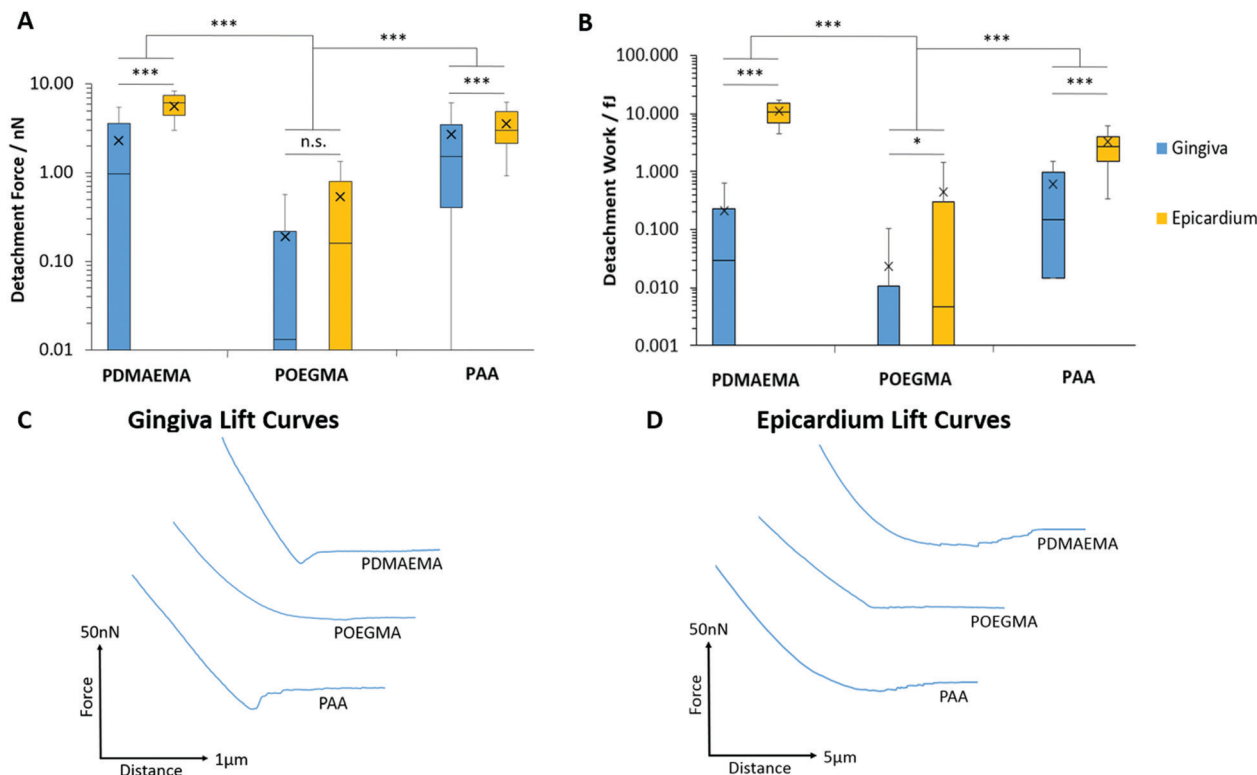
contact area. As such, our data suggests that tissue biochemistry and stiffness combine to regulate the strength of adhesion of soft polymeric interfaces.

## 4. Conclusions

Overall, our results indicate that the adhesion profile of weak polyelectrolyte brushes is relatively complex, arising from combinations of electrostatic and van der Waals interactions, as well as hydrogen bonding. The response of such adhesive behaviour to electrolytes and changes in the Debye length is in turn equally complex, and is further modulated by the formation of hydrogen bonds with electrolytes (such as phosphates). In this respect, the comparison of adhesion profiles to model substrates with those observed with cells and tissues is particularly insightful. Specifically, based on our data, we propose that: (1) adhesion to cells and tissues is primarily mediated by hydrogen bonding and electrostatic forces, based on the differential adhesion profiles observed for the three

brushes tested on biointerfaces and their comparison with the response to SAMs; (2) the glycocalyx has a profound impact on such adhesion and its composition (and abundance) at the surface of tissues will modulate soft matter adhesion; (3) the mechanics of the tissue/substrate (non-deformable in the context of SAMs) has a strong impact on the measured adhesion, in particular at the microscale, since it will dictate contact areas promoting adhesion. Therefore, our results demonstrate that simple considerations of electrostatic and hydrophobic interactions between polymer brushes, and even relatively simple interfaces (self-assembled monolayers), cannot fully account for adhesive profiles. In this context, in studies of brush adhesion to cells and tissues, the prediction of interactions remains particularly difficult to establish; relatively elaborate models, such as molecular dynamics simulations based on atomistic and coarse grain models, will be essential to capture the complex nature of polymer brush-interactions with biological samples, whether bacteria, eukaryotic cells or tissues. Our study also points to the need for an improved understanding and characterisation of the glycocalyx of the epithelium, or the





**Fig. 10** Characterisation of adhesive interactions between polymer brushes and porcine tissue samples. Testing was carried out on samples submerged in PBS. (A) Detachment force of epicardium and keratinized gingiva to polymers. (B) Detachment work between polymers and soft tissue samples. (C) Representative AFM lift curves between polymers and gingival samples. (D) Representative AFM lift curves between polymers and epicardium. Data is plotted as means, with box plots. \* $p \leq 0.05$ . \*\*\* $p \leq 0.001$ . n.s., non-significant.

surface chemistry of tissues, to which biomaterials bonding is required. Some studies have explored the structure and morphology of the glycocalyx in epithelial layers, but this is not systematic across all tissues, and often remains incomplete.

## Conflicts of interest

The authors declare no competing financial interests.

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## References

- J. L. Markman, A. Rekechenetskiy, E. Holler and J. Y. Ljubimova, *Adv. Drug Delivery Rev.*, 2013, **65**, 1866–1879.
- M. Kanamala, W. R. Wilson, M. Yang, B. D. Palmer and Z. Wu, *Biomaterials*, 2016, **85**, 152–167.
- V. Bhagat and M. L. Becker, *Biomacromolecules*, 2017, **18**, 3009–3039.
- S. Tiwari, R. Patil and P. Bahadur, *Polymers*, 2018, **11**, 1.
- T. N. Dinh, S. Hou, S. Park, B. A. Shalek and K. J. Jeong, *ACS Biomater. Sci. Eng.*, 2018, **4**, 3471–3477.
- C. Ghobril and M. W. Grinstaff, *Chem. Soc. Rev.*, 2015, **44**, 1820–1835.
- N. Annabi, A. Tamayol, S. R. Shin, A. M. Ghaemmaghami, N. A. Peppas and A. Khademhosseini, *Nano Today*, 2014, **9**, 574–589.
- E. Bressan, V. Favero, C. Gardin, L. Ferroni, L. Iacobellis, L. Favero, V. Vindigni, M. Berengo, S. Sivoletta and B. Zavan, *Polymers*, 2011, **3**, 509–526.
- J. R. Venugopal, M. P. Prabhakaran, S. Mukherjee, R. Ravichandran, K. Dan and S. Ramakrishna, *J. R. Soc., Interface*, 2012, **9**, 1–19.
- W. Samprasit, R. Kaomongkolgit, M. Sukma, T. Rojanarata, T. Ngawhirunpat and P. Opanasopit, *Carbohydr. Polym.*, 2015, **117**, 933–940.
- J. Tsibouklis, A. M. Middleton, N. Patel and J. Pratten, *J. Biomed. Mater. Res., Part A*, 2013, **101**, 3327–3338.
- M. L. B. Palacio and B. Bhushan, *Philos. Trans. R. Soc., A*, 2012, **370**, 2321–2347.
- J. S. Suk, Q. Xu, N. Kim, J. Hanes and L. M. Ensign, *Adv. Drug Delivery Rev.*, 2016, **99**, 28–51.
- M. Mehdizadeh and J. Yang, *Macromol. Biosci.*, 2013, **13**, 271–288.
- Y.-C. Nho, J.-S. Park and Y.-M. Lim, *Polymers*, 2014, **6**, 890–898.





- 16 J. Wong, A. Chilkoti and V. T. Moy, *Biomol. Eng.*, 1999, **16**, 45–55.
- 17 C. a Squier and M. J. Kremer, *J. Natl. Cancer Inst. Monogr.*, 2001, **52242**, 7–15.
- 18 M. Derrien, M. W. van Passel, J. H. van de Bovenkamp, R. Schipper, W. de Vos and J. Dekker, *Gut Microbes*, 2010, **1**, 254–268.
- 19 M. Krishnamoorthy, S. Hakobyan, M. Ramstedt and J. E. Gautrot, *Chem. Rev.*, 2014, **114**, 10976–11026.
- 20 J. O. Zoppe, N. C. Ataman, P. Mocny, J. Wang, J. Moraes and H. A. Klok, *Chem. Rev.*, 2017, **117**, 1105–1318.
- 21 C. Rodriguez-Emmenegger, E. Brynda, T. Riedel, M. Houska, V. Šubr, A. B. Alles, E. Hasan, J. E. Gautrot and W. T. S. Huck, *Macromol. Rapid Commun.*, 2011, **32**, 952–957.
- 22 A. J. Hackett, J. Malmström, P. J. Molino, J. E. Gautrot, H. Zhang, M. J. Higgins, G. G. Wallace, D. E. Williams and J. Travas-Sejdic, *J. Mater. Chem. B*, 2015, **3**, 9285–9294.
- 23 A. M. Jonas, K. Glinel, R. Oren, B. Nysten and W. T. S. Huck, *Macromolecules*, 2007, **40**, 4403–4405.
- 24 K. Y. Tan, J. E. Gautrot and W. T. S. Huck, *Langmuir*, 2011, **27**, 1251–1259.
- 25 D. E. S. Santos, D. Li, M. Ramstedt, J. E. Gautrot and T. A. Soares, *Langmuir*, 2019, **35**, 5037–5049.
- 26 O. Rzhapishevska, S. Hakobyan, R. Ruhel, J. Gautrot, D. Barbero and M. Ramstedt, *Biomater. Sci.*, 2013, **1**, 589–602.
- 27 C. Rodriguez-Emmenegger, S. Janel, A. de los Santos Pereira, M. Bruns and F. Lafont, *Polym. Chem.*, 2015, **6**, 5740–5751.
- 28 T. A. Petrie, J. E. Raynor, D. W. Dumbauld, T. T. Lee, S. Jagtap, K. L. Templeman, D. M. Collard and A. J. Garcia, *Sci. Transl. Med.*, 2010, **2**, 45ra60.
- 29 J. E. Gautrot, J. M. Rosenholm, D. Li, T. Gulin-Sarfraz, A. S. Sharili and M. Krishnamoorthy, *Biomacromolecules*, 2017, **18**, 4121–4132.
- 30 D. Li, A. S. Sharili, J. Connelly and J. E. Gautrot, *Biomacromolecules*, 2018, **19**, 606–615.
- 31 D. J. D'Sa, H.-K. Chan and W. Chrzanowski, *J. Colloid Interface Sci.*, 2014, **426**, 190–198.
- 32 H.-J. Butt, B. Cappella and M. Kappl, *Surf. Sci. Rep.*, 2005, **59**, 1–152.
- 33 P. Schön, E. Kutnyanszky, B. ten Donkelaar, M. G. Santonicola, T. Tecim, N. Aldred, A. S. Clare and J. J. G. Vancso, *Colloids Surf., B*, 2013, **102**, 923–930.
- 34 Y. Inoue, T. Nakanishi and K. Ishihara, *Langmuir*, 2013, **29**, 10752–10758.
- 35 Y. Inoue, T. Nakanishi and K. Ishihara, *React. Funct. Polym.*, 2011, **71**, 350–355.
- 36 N. S. Bhairamadgi, S. P. Pujari, C. J. M. Van Rijn and H. Zuillhof, *Langmuir*, 2014, **30**, 12532–12540.
- 37 Y. Yu, B. D. Kieviet, E. Kutnyanszky, G. J. Vancso and S. De Beer, *ACS Macro Lett.*, 2015, **4**, 75–79.
- 38 M. M. Elmahdy, A. Drechsler, E. Bittrich, P. Uhlmann and M. Stamm, *Colloid Polym. Sci.*, 2014, **292**, 1999–2012.
- 39 M.-J. Yin, M. Yao, S. Gao, A. P. Zhang, H.-Y. Tam and P.-K. A. Wai, *Adv. Mater.*, 2016, **28**, 1394–1399.
- 40 N. Salamat-Miller, M. Chittchang and T. P. Johnston, *Adv. Drug Delivery Rev.*, 2005, **57**, 1666–1691.
- 41 G. P. Andrews, T. P. Lavery and D. S. Jones, *Eur. J. Pharm. Biopharm.*, 2009, **71**, 505–518.
- 42 A. V. Dubolazov, Z. S. Nurkeeva, G. A. Mun and V. V. Khutoryanskiy, *Biomacromolecules*, 2006, **7**, 1637–1643.
- 43 Y. Huang, H. Yu and C. Xiao, *Carbohydr. Polym.*, 2007, **69**, 774–783.
- 44 E. De Giglio, S. Cometa, N. Cioffi, L. Torsi and L. Sabbatini, *Anal. Bioanal. Chem.*, 2007, **389**, 2055–2063.
- 45 W. Lee, T. G. Lee and W. G. Koh, *J. Ind. Eng. Chem.*, 2007, **13**, 1195–1200.
- 46 S. Sanjuan, P. Perrin, N. Pantoustier and Y. Tran, *Langmuir*, 2007, **23**, 5769–5778.
- 47 C. V. Synatschke, A. Schallon, V. Jérôme, R. Freitag and A. H. E. Müller, *Biomacromolecules*, 2011, **12**, 4247–4255.
- 48 A. Hucknall, S. Rangarajan and A. Chilkoti, *Adv. Mater.*, 2009, **21**, 2441–2446.
- 49 S. Swar, V. Zajíčová, J. Müllerová, P. Šubrtová, J. Horáková, B. Dolenský, M. Řezanka and I. Stibor, *J. Mater. Sci.*, 2018, **53**, 14104–14120.
- 50 M. Heuberger, T. Drobek and N. D. Spencer, *Biophys. J.*, 2005, **88**, 495–504.
- 51 D. Boettiger and B. Wehrle-Haller, *J. Phys.: Condens. Matter*, 2010, **22**, 194101.
- 52 B. Lego, M. François, W. G. Skene and S. Giasson, *Langmuir*, 2009, **25**, 5313–5321.
- 53 F. Jenkins and H. White, *Fundamentals of Optics*, McGraw-Hill Primis Custom Publishing, 4th edn, 2001, pp. 474–496.
- 54 F. Qu, D. Li, X. Ma, F. Chen and J. E. Gautrot, *Biomacromolecules*, 2019, **20**, 2218–2229.
- 55 J. E. Sader, I. Larson, P. Mulvaney and L. R. White, *Rev. Sci. Instrum.*, 1995, **66**, 3789–3798.
- 56 L. O. Heim, T. S. Rodrigues and E. Bonaccorso, *Colloids Surf., A*, 2014, **443**, 377–383.
- 57 Y. An, D. Li, N. Roohpour, J. E. Gautrot and A. H. Barber, *Dent. Mater.*, 2016, 1–9.
- 58 F. Hang, D. Lu, R. J. Bailey, I. Jimenez-Palomar, U. Stachewicz, B. Cortes-Ballesteros, M. Davies, M. Zech, C. Bödefeld and A. H. Barber, *Nanotechnology*, 2011, **22**, 365708.
- 59 W. C. Oliver and G. M. Pharr, *J. Mater. Res.*, 1992, **7**, 1564–1583.
- 60 K. Y. Tan, H. Lin, M. Ramstedt, F. M. Watt, W. T. S. Huck and J. E. Gautrot, *Integr. Biol.*, 2013, **5**, 899–910.
- 61 B. Lego, W. G. Skene and S. Giasson, *Macromolecules*, 2010, **43**, 4384–4393.
- 62 R. Barbey, L. Lavanant, D. Paripovic, N. Schüwer, C. Sugnaux, S. Tugulu and H.-A. Klok, *Chem. Rev.*, 2009, **109**, 5437–5527.
- 63 T. Wu, P. Gong, I. Szleifer, P. Vlček, V. Šubr and J. Genzer, *Macromolecules*, 2007, **40**, 8756–8764.
- 64 K. A. Davis and K. Matyjaszewski, *Macromolecules*, 2000, **33**, 4039–4047.
- 65 R. Dong, S. Krishnan, B. A. Baird, M. Lindau and C. K. Ober, *Biomacromolecules*, 2007, **8**, 3082–3092.
- 66 P. Louette, F. Bodino and J.-J. Pireaux, *Surf. Sci. Spectra*, 2005, **12**, 22–26.
- 67 N. D. Treat, N. Ayres, S. G. Boyes and W. J. Brittain, *Macromolecules*, 2006, **39**, 26–29.



- 68 R. Israels, F. A. M. Leermakers and G. J. Fleer, *Macromolecules*, 1994, **27**, 3087–3093.
- 69 T. von Werne and T. E. Patten, *J. Am. Chem. Soc.*, 1999, **121**, 7409–7410.
- 70 H. Hinterwirth, S. Kappel, T. Waitz, T. Prohaska, W. Lindner and M. Lämmerhofer, *ACS Nano*, 2013, **7**, 1129–1136.
- 71 H. Ma, J. Hyun, P. Stiller and A. Chilkoti, *Adv. Mater.*, 2004, **16**, 338–341.
- 72 J. E. Gautrot, B. Trappmann, F. Ocegüera-Yanez, J. Connelly, X. He, F. M. Watt and W. T. S. Huck, *Biomaterials*, 2010, **31**, 5030–5041.
- 73 A. Drechsler, A. Synytska, P. Uhlmann, M. M. Elmahdy, M. Stamm and F. Kremer, *Langmuir*, 2010, **26**, 6400–6410.
- 74 S. E. Burke and C. J. Barrett, *Langmuir*, 2003, **19**, 3297–3303.
- 75 G. Liu, D. Wu, C. Ma, G. Zhang, H. Wang and S. Yang, *ChemPhysChem*, 2007, **8**, 2254–2259.
- 76 L. N. Nagy, R. Mizsei, T. Kiss, Z. Varga, I. C. Szegvártó, M. Pálmai, A. Bóta, T. Kremmer, G. Tárkányi and J. Mihály, *J. Colloid Interface Sci.*, 2012, **390**, 34–40.
- 77 Z. Wu, H. Xiang, T. Kim, M. S. Chun and K. Lee, *J. Colloid Interface Sci.*, 2006, **304**, 119–124.
- 78 S. Vaidya, P. Thaplyal and A. Ganguli, *Nanoscale Res. Lett.*, 2011, **6**, 169.
- 79 S. Ansar, S. Chakraborty and C. Kitchens, *Nanomaterials*, 2018, **8**, 339.
- 80 M. M. Elmahdy, A. Drechsler, E. Bittrich, P. Uhlmann and M. Stamm, *Colloid Polym. Sci.*, 2014, **292**, 1999–2012.
- 81 M. Stamm, F. Kremer, A. Synytska, P. Uhlmann and A. Drechsler, *Langmuir*, 2012, **28**, 15555–15565.
- 82 W. M. De Vos, F. A. M. Leermakers, A. De Keizer, M. A. C. Stuart and J. M. Kleijn, *Langmuir*, 2010, **26**, 249–259.
- 83 M. Schlierf, S. Ramakrishnan, A. Keller, G. Krainer and G. Grundmeier, *Small*, 2017, **13**, 1702100.
- 84 T. Urbic, *Chem. Phys. Lett.*, 2014, **610–611**, 159–162.
- 85 J. T. O'Brien and E. R. Williams, *J. Am. Chem. Soc.*, 2012, **134**, 10228–10236.
- 86 Y. Yang and Q. Cui, *J. Phys. Chem. B*, 2007, **111**, 3999–4002.
- 87 G. R. J. Thatcher, D. R. Cameron, R. Nagelkerke and J. Schmitke, *J. Chem. Soc., Chem. Commun.*, 1992, 386.
- 88 M. Raftari, Z. J. Zhang, S. R. Carter, G. J. Leggett and M. Geoghegan, *Macromolecules*, 2015, **48**, 6272–6279.
- 89 S. Sakata, Y. Inoue and K. Ishihara, *Langmuir*, 2015, **31**, 3108–3114.
- 90 A. Synytska, E. Svetushkina, N. Puretskiy, G. Stoychev, S. Berger, L. Ionov, C. Bellmann, K. J. Eichhorn and M. Stamm, *Soft Matter*, 2010, **6**, 5907–5914.
- 91 C. Friedsam, A. D. C. Bécares, U. Jonas, M. Seitz and H. E. Gaub, *New J. Phys.*, 2004, **6**, 9.
- 92 C. Friedsam, M. Seitz and H. E. Gaub, *J. Phys.: Condens. Matter*, 2004, **16**, S2369–S2382.
- 93 R. Godawat, S. N. Jamadagni and S. Garde, *Proc. Natl. Acad. Sci. U. S. A.*, 2009, **106**, 15119–15124.
- 94 Z. A. Levine, M. V. Rapp, W. Wei, R. G. Mullen, C. Wu, G. H. Zerze, J. Mittal, J. H. Waite, J. N. Israelachvili and J.-E. Shea, *Proc. Natl. Acad. Sci. U. S. A.*, 2016, **113**, 4332–4337.
- 95 F. M. Watt and H. Green, *Nature*, 1982, **295**, 434–436.
- 96 A. P. Majewski, U. Stahlschmidt, V. Jérôme, R. Freitag, A. H. E. Müller and H. Schmalz, *Biomacromolecules*, 2013, **14**, 3081–3090.
- 97 S. Agarwal, Y. Zhang, S. Maji and A. Greiner, *Mater. Today*, 2012, **15**, 388–393.
- 98 K. Y. Tan, H. Lin, M. Ramstedt, F. M. Watt, W. T. S. Huck and J. E. Gautrot, *Integr. Biol.*, 2013, **5**, 899–910.
- 99 S. Weinbaum, J. M. Tarbell and E. R. Damiano, *Annu. Rev. Biomed. Eng.*, 2007, **9**, 121–167.
- 100 D. W. Owens, V. G. Brunton, E. K. Parkinson and M. C. Frame, *Biochem. Biophys. Res. Commun.*, 2000, **269**, 369–376.
- 101 A. Singh, S. C. Satchell, C. R. Neal, E. A. McKenzie, J. E. Tooke and P. W. Mathieson, *J. Am. Soc. Nephrol.*, 2007, **18**, 2885–2893.
- 102 A. B. Servais, A. Kienle, C. D. Valenzuela, A. B. Ysasi, W. L. Wagner, A. Tsuda, M. Ackermann and S. J. Mentzer, *Tissue Eng., Part A*, 2018, **24**, 199–206.
- 103 M. Araña, J. J. Gavira, E. Peña, A. González, G. Abizanda, M. Cilla, M. M. Pérez, E. Albiasu, N. Aguado, M. Casado, B. López, S. González, M. Soriano, C. Moreno, J. Merino, J. M. García-Verdugo, J. Díez, M. Doblaré, B. Pelacho and F. Prosper, *Biomaterials*, 2014, **35**, 143–151.
- 104 N. Tano, T. Narita, M. Kaneko, C. Ikebe, S. R. Coppen, N. G. Campbell, M. Shiraishi, Y. Shintani and K. Suzuki, *Mol. Ther.*, 2014, **22**, 1864–1871.
- 105 E. R. Rodriguez and C. D. Tan, *Prog. Cardiovasc. Dis.*, 2017, **59**, 327–340.
- 106 A. Ohtsuka, S. Yamana and T. Murakami, *Histochem. Cell Biol.*, 1997, **107**, 441–447.
- 107 Y. Collan and A.-M. Raeste, *Eur. J. Oral Sci.*, 1978, **86**, 374–385.
- 108 V. Nguyen, A. Ndoye and L. Hall, *J. Cell Sci.*, 2001, **114**, 1189–1204.
- 109 K. Wolff and E. Schreiner, *J. Invest. Dermatol.*, 1968, **51**, 418–430.
- 110 L. M. C. Collins and C. Dawes, *J. Dent. Res.*, 1987, **66**, 1300–1302.
- 111 A. Sengupta, D. Valdramidou, S. Huntley, S. J. Hicks, S. D. Carrington and A. P. Corfield, *Arch. Oral Biol.*, 2001, **46**, 529–538.
- 112 L. Vitkov, W. D. Krautgartner, M. Hannig and K. Fuchs, *FEMS Microbiol. Lett.*, 2001, **202**, 25–30.
- 113 M. Chevalier, C. Sakarovitch, I. Precheur, J. Lamure and V. Pouyssegur-Rougier, *Acta Odontol. Scand.*, 2015, **73**, 267–273.

