



Cite this: *J. Mater. Chem. B*, 2018,
6, 3770

Nanoparticle-supported polymer brushes for temperature-regulated glycoprotein separation: investigation of structure–function relationship†

Huiting Ma,^{ab} Lingdong Jiang,^a Solmaz Hajizadeh,^a Haiyue Gong,^a
Bin Lu^{*b} and Lei Ye^{*a}

In this work, we synthesized a series of nanoparticle-supported boronic acid polymer brushes for affinity separation of glycoproteins. Polymer brushes were prepared by surface-initiated atom transfer radical polymerization of glycidyl methacrylate and *N*-isopropylacrylamide, followed by stepwise modification of the pendant as well as the end functional groups to introduce boronic acid moieties through a Cu(I)-catalyzed alkyne–azide cycloaddition reaction. We investigated the impact of the polymer structure on glycoprotein binding under different pH and temperature conditions, and established new methods that allow glycoproteins to be more easily isolated and recovered with minimal alteration in solvent composition. Our experimental results suggest that for the separation of glycoproteins, terminal boronic acids located at the end of polymer chains play the most important role. The thermo-responsibility of the new affinity adsorbents, in addition to the high capacity for glycoprotein binding (120 mg ovalbumin per g adsorbent), provides a convenient means to realize simplified bioseparation not only for glycoproteins, but also for other carbohydrate-containing biological molecules.

Received 6th March 2018,
Accepted 3rd May 2018

DOI: 10.1039/c8tb00627j

rsc.li/materials-b

Introduction

In recent years, polymer brushes have attracted much attention due to their unique structure and novel functional properties.^{1–4} Of particular interest is the grafting of macromolecules on the surfaces of different materials to improve their wettability, lubricity, hydrophilicity and colloidal stability.^{5,6} Among the different types of surface-initiated polymerization techniques, atom transfer radical polymerization (ATRP) is widely used because of its mild reaction conditions, versatile control of polymer structure and compatibility with a broad range of monomers.^{7–10} By using simple post-polymerization modification, polymer brushes made by ATRP can be easily converted into functional materials for use in bioseparation, enzyme catalysis, bioanalysis, *etc.*^{11,12}

Glycoproteins play an important role in many biological processes and are frequently used as biomarkers for different diseases. Effective separation of glycoproteins is one important step in many research areas and diagnostics.^{13–16} Lectin affinity

chromatography (LAC) is widely used in this regard, as lectins can bind carbohydrates specifically. LAC has been realized using monolithic capillary columns, silica microparticles packed in microcolumns, sometimes hyphenated with mass spectrometry.^{17–20} However, LAC has some limitations such as being time-consuming process, and having complicated operations and high cost.²¹ Other methods *e.g.* based on capillary electrophoresis (CE) also use complicated protocols and require specialized instruments.²² Among all the methods used for glycoprotein enrichment, boronic acid-based affinity separation has become increasingly popular because of its low bias, long shelf time and convenient handling.^{23–28} In particular, polymer materials containing boronate affinity ligands can be designed to offer selective glycoprotein enrichment and separation, as boronic acids can form stable ester bonds with *cis*-diol groups at the carbohydrate site of glycoproteins. The sensitivity of boronate ester bonds towards pH variation can be used to modulate glycoprotein binding and release, thereby decreasing the influence of non-specific protein binding.^{25,29} Different types of boronate affinity materials have been developed, for example, magnetic nanoparticles, monolithic capillary columns modified with boronic acids,^{30–32} and boronic acids immobilized on mesoporous silica³³ as well as on agarose gel.^{34,35} Considering the steric hindrance and diffusion barrier encountered in protein separation, nanoparticles are ideal supports for immobilization of boronic acids to achieve efficient bioseparation.

^a Division of Pure and Applied Biochemistry, Department of Chemistry, Lund University, Box 124, 221 00 Lund, Sweden. E-mail: lei.ye@tbiokem.lth.se

^b Key Laboratory of Environment and Health, Ministry of Education & Ministry of Environmental Protection, State Key Laboratory of Environmental Health (Incubating), School of Public Health, Tongji Medical College, Huazhong University of Science and Technology, Wuhan, Hubei 430030, China. E-mail: lubin@mails.tjmu.edu.cn

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c8tb00627j

In previous studies, we developed clickable boronic acids that could be easily conjugated to silica-supported polymer brushes for glycoprotein separation.^{13,36,37} The nanoparticle-supported polymer brushes displayed high capacity for glycoprotein binding, and the protein binding could be controlled by adjusting the solution temperature. In order to gain further insights into the structure–function relationship of nanoparticle-supported boronic acid polymer brushes, in this work we carried out a systematic investigation to study the impact of the polymer structure on glycoprotein binding. Glycidyl methacrylate (GMA), with and without *N*-isopropylacrylamide (NIPAm), was used as a monomer to prepare silica-supported polymer brushes by surface-initiated atom transfer radical polymerization (SI-ATRP). The copolymerization with NIPAm resulted in temperature-responsive polymer chains grafted on the nanoparticle surface. When the solution temperature was raised to above the lower critical solution temperature (LCST) of polyNIPAm (32 °C), the polymer brushes became dehydrated and collapsed on the surface,^{38,39} thereby affecting glycoprotein binding. The GMA monomer was used to introduce epoxide groups into the polymer brush that could be converted into pendant azide groups,^{40,41} which could be easily modified through a Cu(I)-catalyzed alkyne–azide cycloaddition reaction (click chemistry) to give the desired boronic acid polymer brushes.⁴² This work focused on studying the effect of the density and location of boronic acid ligands, and the influence of temperature on glycoprotein binding with the nanoparticle-supported polymer brushes. Besides studying the structure–function relationship, we were also interested in developing new methods for protein separation that require minimal alteration of the chemical composition of the protein solution, a condition more favourable for the final step in protein purification. For this purpose we intended to use the thermo-responsive properties of the polymer brushes to regulate the binding and release of glycoproteins.

Experimental

Materials

Tetraethylorthosilicate (TEOS), (3-aminopropyl)triethoxysilane (APTES), triethylamine (TEA), 2-bromoisobutyl bromide (BIBB), *N*-isopropylacrylamide (NIPAm), glycidyl methacrylate (GMA), propargylamine, CuSO₄, sodium ascorbate, sodium azide, *N,N,N',N',N''*-pentamethyldiethylenetriamine (PMDETA), tris[2-(dimethylamino)ethyl]amine (Me₆TREN), Alizarin Red S (ARS), 3-aminophenylboronic acid hemisulfate (3-APBA hemisulfate), sodium hydrogen carbonate, propargyl chloroformate, ovalbumin (OVA), bovine serum albumin (BSA), horseradish peroxidase (HRP), sodium dodecyl sulfonate (SDS), sulphuric acid, acetonitrile (ACN), methanol, ammonia, toluene, *N,N*-dimethylformamide (DMF), tetrahydrofuran (THF), *D*-fructose, isopropanol and acetone were purchased from Sigma-Aldrich and were used without further purification. 2,2'-Azinobis[3-ethylbenzthiazoline-6-sulfonic acid]-diammonium salt (1-Step ABTS) was purchased from Thermo Scientific. CuBr was stirred for 24 h in acetic acid, collected by centrifugation, washed with water and methanol, and dried in a

vacuum before use. Ultrapure water (18.2 MΩ cm) was obtained from an ELGA LabWater System (Vivendi Water Systems Ltd). The clickable boronic acid, 3-(prop-2-ynyloxy-carbonylamino)-phenylboronic acid (PCAPBA), was synthesized according to our previously published procedure.³⁶

Preparation of nanoparticles and nanoparticle-supported polymer brushes

Silica nanoparticles (Si). Silica nanoparticles were synthesized by the Stöber method,⁴³ as described in our previous work.^{13,37} Ammonia (25% aqueous solution, 22.4 mL) was added into a mixture of 100 mL of methanol and 33 mL of water in a 1 L glass beaker and agitated using a magnetic stirrer. To this solution, a mixture of 13.8 mL of TEOS and 130 mL of methanol was added quickly. The suspension obtained was stirred at room temperature for 8 h. Finally, the product was collected by centrifugation, washed 3 times with water, 3 times with methanol and dried in a vacuum. The obtained product is denoted as Si.

Amino-functionalized silica nanoparticles (Si@NH₂). To a round bottom flask (100 mL), Si (3.0 g) and 1% APTES solution (0.5 mL APTES dissolved in 49.5 mL anhydrous toluene) were added. The mixture was stirred at 110 °C for 24 h. After the reaction, the particles were collected by centrifugation, washed 3 times with acetone, 3 times with methanol, and dried in a vacuum. The obtained product is denoted as Si@NH₂.

Initiator-functionalized silica nanoparticles (Si@initiator). To a round bottom flask (25 mL), Si@NH₂ (0.25 g), triethylamine (0.40 mL, 2.9 mmol) and THF (12 mL) were added. The mixture was cooled on an ice-water bath. To the suspension, BIBB (0.31 mL, 5.0 mmol) was slowly added. The reaction suspension was then warmed to room temperature and stirred for 24 h. The nanoparticles were collected by centrifugation, washed 3 times with water, 3 times with methanol, and dried in a vacuum. The obtained product is denoted as Si@initiator.

Silica-supported polyGMA brushes (Si@pGMA). To a round bottom flask (25 mL), Si@initiator (0.20 g), GMA (0.4 mL, 3 mmol), CuBr₂ (1.1 mg, 0.005 mmol), CuBr (7.2 mg, 0.05 mmol) and 2-propanol (6 mL) were added. The mixture was purged with nitrogen gas for 15 min before PMDETA (10 μL, 0.05 mmol) was added. After another 15 min purging with nitrogen gas, the flask was sealed and polymerization was carried out under magnetic stirring at 60 °C for 24 h. After the reaction, the nanoparticles were collected by centrifugation, washed 3 times with water, 3 times with methanol, and dried in a vacuum. The obtained product is denoted as Si@pGMA.

Silica-supported GMA-NIPAm copolymer brushes (Si@pco). To a round bottom flask (25 mL), Si@initiator (0.20 g), GMA (0.4 mL, 3 mmol), NIPAm (0.68 g, 6 mmol), CuBr₂ (1.1 mg, 0.005 mmol), CuBr (7.2 mg, 0.05 mmol) and 2-propanol (6 mL) were added. The mixture was purged with nitrogen gas for 15 min before Me₆TREN (14 μL, 0.05 mmol) was added. After another 15 min purging with nitrogen gas, the flask was sealed and the reaction mixture was stirred at 60 °C for 24 h. After the reaction, the nanoparticles were collected, washed 3 times with water, 3 times with methanol, and dried in a vacuum. The obtained product is denoted as Si@pco.



Silica-supported polyNIPAm brushes (Si@pNIPAm). To a round bottom flask (25 mL), Si@initiator (0.20 g), NIPAm (0.68 g, 6 mmol), CuBr₂ (1.1 mg, 0.005 mmol), CuBr (7.2 mg, 0.05 mmol) and 2-propanol (6 mL) were added. The mixture was purged with nitrogen gas for 15 min before Me₆TREN (14 μ L, 0.05 mmol) was added. After another 15 min purging with nitrogen gas, the flask was sealed and the reaction mixture was stirred at 60 °C for 24 h. After the reaction, the nanoparticles were collected, washed and dried using the procedures described above. The obtained product is denoted as Si@pNIPAm.

Azide-modified polymer brushes on silica nanoparticles (Si@pGMA-N₃@N₃). Si@pGMA (150 mg) was mixed with sodium azide (78 mg), ammonium chloride (64.5 mg) and DMF (7.5 mL), and stirred at 60 °C for 24 h. The obtained product was then isolated and purified following the procedures as described above. The obtained particle is denoted as Si@pGMA-N₃@N₃.

Azide-terminated polymer brushes on silica nanoparticles (Si@pGMA-OH@N₃). Si@pGMA particles (160 mg) were immersed in 6 mL of 0.5 M sulphuric acid and stirred at 70 °C for 12 h. After washing 3 times with water and 3 times with methanol, the particles were dried under vacuum. The acid-treated particles (150 mg) were mixed with sodium azide (78 mg) and ammonium chloride (64.5 mg) in DMF (7.5 mL), and stirred at 60 °C for 24 h. The obtained product was then isolated and purified following the procedures as described above. The obtained product is denoted as Si@pGMA-OH@N₃.

Azide-modified copolymer brushes on silica nanoparticles (Si@pco-N₃@N₃). Si@pco (150 mg) was mixed with sodium azide (78 mg) and ammonium chloride (64.5 mg) in DMF (7.5 mL), and stirred at 60 °C for 24 h. The obtained product was then isolated and purified following the procedures as described above. The obtained product is denoted as Si@pco-N₃@N₃.

Azide-terminated copolymer brushes on silica nanoparticles (Si@pco-OH@N₃). Si@pco (160 mg) was immersed in 6 mL of 0.5 M sulphuric acid, stirred at 70 °C for 12 h. After washing 3 times with water and 3 times with methanol, the particles were dried under vacuum. The acid-treated particles (150 mg) were mixed with sodium azide (78 mg) and ammonium chloride (64.5 mg) in DMF (7.5 mL), and stirred at 60 °C for 24 h. The particles were isolated and purified following the procedures as described above. The obtained product is denoted as Si@pco-OH@N₃.

Azide-terminated polyNIPAm brushes on silica nanoparticles (Si@pNIPAm@N₃). To a round bottom flask (25 mL), Si@pNIPAm (150 mg), sodium azide (78 mg), ammonium chloride (64.5 mg) and DMF (7.5 mL) were added. After 5 min of sonication, the suspension was stirred at 60 °C for 24 h. The nanoparticles were then isolated and purified following the procedures as described above. The obtained product is denoted as Si@pNIPAm@N₃.

Boronic acid-modified polymer brushes (Si@pGMA-BA@BA). Si@pGMA-N₃@N₃ particles (50 mg) and PCAPBA (6 mg) were suspended in 6 mL of methanol/water (1/1). After 5 min of sonication, CuSO₄ (100 mM, 20 μ L) and sodium ascorbate (100 mM, 100 μ L) were added. The reaction mixture was shaken on a rocking table for 24 h at room temperature. After the reaction, the obtained particles were washed 3 times with water,

3 times with methanol, and dried in a vacuum. The obtained product is denoted as Si@pGMA-BA@BA.

Using the same procedure, other boronic acid-modified polymer brushes, Si@pGMA-OH@BA, Si@pco-BA@BA, Si@pco-OH@BA and Si@pNIPAm@BA, were synthesized from Si@pGMA-OH@N₃, Si@pco-N₃@N₃, Si@pco-OH@N₃ and Si@pNIPAm@N₃, respectively.

Characterization of nanoparticle-supported polymer brushes

Infrared spectroscopy, thermogravimetry and elemental analysis. Attenuated total reflection (ATR) infrared spectra of the composite particles were collected using a Nicolet IS5 FT-IR instrument (Thermo Fisher-Scientific Inc., Waltham, USA). All the spectra were collected at 25 °C within a range of 4000–375 cm⁻¹ with a resolution of 4 cm⁻¹ and 32 scans. The particle morphology was inspected using a JEOL scanning electron microscope (SEM, JSM-6700F, JEOL, Japan). Thermal gravimetric analysis (TGA) was carried out in synthetic air, with a sample heating rate of 10 °C min⁻¹. Elemental analysis (C, H, N, Br, and B) was performed by Mikroanalytisches Laboratorium Kolbe (Germany).

Binding of low molecular weight *cis*-diols. Boronic acid-functionalized nanoparticles (10 mg) were added into 2 mL of ARS solution (0.1 mM dissolved in 20 mM pH 7.4 PBS buffer containing 0.5 M NaCl) and sonicated for 5 min. The fluorescence emission (500–650 nm) of the sample was measured using an excitation wavelength at 469 nm on a QuantaMaster C-60/2000 spectrofluorometer (Photon Technology International, Lawrenceville, NJ). The boronic acid-functionalized particles were Si@pGMA-BA@BA, Si@pGMA-OH@BA, Si@pco-BA@BA, Si@pco-OH@BA and Si@pNIPAm@BA. For comparison, the azide-functionalized particles Si@pGMA-N₃@N₃, Si@pGMA-OH@N₃, Si@pco-N₃@N₃, Si@pco-OH@N₃, and Si@pNIPAm@N₃ were tested under the same conditions.

Protein binding analysis

The amount of protein bound to the boronic acid-functionalized particles was investigated under three different conditions. The amount of protein binding was calculated using the equation:

$$Q = \frac{(C_0 - C_t)V}{m} \times 10^3$$

where C_0 (mg mL⁻¹) is the initial protein concentration, C_t (mg mL⁻¹) is the concentration of the unbound protein, V (mL) is the volume of the protein solution, and m (in mg) is the mass of the composite particles. All the binding tests were carried out in triplicate, and the data are reported as mean value \pm standard deviation.

Equilibrium protein binding analysis. In a series of 2 mL microcentrifuge tubes, 3 mg of boronic acid-functionalized particles was mixed with 1 mL of protein solution (0.2–1.4 mg mL⁻¹) prepared in different buffers. The samples were shaken on a rocking table at 20 °C or shaken at 40 °C with a speed of 300 rpm using a thermal mixer (Eppendorf, Hamburg, Germany) for 2 h. After incubation the samples were immediately centrifuged. The free protein in the supernatant was measured using a fluorescence spectrometer. OVA was quantified by measuring the fluorescence



emission at 330 nm (E_{x} : 284 nm) and HRP by measuring the emission at 340 nm (E_{x} : 284 nm).

Verification of protein separation by sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE). In a series of 2 mL microcentrifuge tubes, 10 mg of composite particles (Si@pNIPAm@BA) was suspended in 1 mL of protein solution containing 1 mg mL⁻¹ OVA and 1 mg mL⁻¹ BSA dissolved in 20 mM PBS buffer (pH 9.0, containing 0.5 M NaCl). After incubation at 20 °C for 2 h, the samples were centrifuged and 20 µL of the supernatant was taken for SDS-PAGE analysis. The remaining samples were then shaken at 40 °C for 2 h. After immediate centrifugation, all the supernatant was removed, from which 20 µL was used for SDS-PAGE analysis. The settled particles were then washed with the same PBS buffer 3 times, before the proteins were eluted by treating the particles with 1 mL of 0.2 M acetate buffer (pH 4.0) for 2 h. The proteins collected in the washing and elution steps were also analyzed by SDS-PAGE.

Regulation of glycoprotein binding in the presence of excess BSA. A solution of HRP (0.1 mg mL⁻¹) and BSA (1 mg mL⁻¹) dissolved in 20 mM PBS buffer (pH 9.0 containing 0.5 M NaCl) was prepared. This protein solution was used to test the capability of the boronic acid-functionalized composite particles to bind the glycoprotein HRP under different conditions. The binding experiments were carried out in three steps:

(1) In a series of 2 mL microcentrifuge tubes, 3 mg of boronic acid-functionalized particles was suspended in 1 mL of the protein solution. The mixture was shaken at 20 °C for 2 h, at a speed of 300 rpm using a thermal mixer. After centrifugation, 20 µL of the supernatant was taken to determine the free HRP by measuring the enzyme activity using 1-Step ABTS. Briefly, the 20 µL supernatant containing HRP was mixed with 150 µL of 1-Step ABTS in a microplate well. The microplate was incubated at 20 °C for 5 min, and then 100 µL of 1% SDS (in 20 mM PBS buffer, pH 9.0) was added to stop the enzymatic reaction. The absorbance of the reaction mixture at 410 nm was measured to calculate the amount of the unbound HRP.

(2) The remaining 980 µL mixture from step 1 (containing the nanoparticles and proteins) was heated to 40 °C and shaken at 300 rpm for 2 h. After centrifugation, 20 µL of the supernatant was taken to measure the amount of free HRP, using the procedure as described above.

(3) The remaining 960 µL mixture from step 2 was cooled to 20 °C before 1 mg of D-fructose was added. The mixture was shaken at 20 °C and at 300 rpm for 2 h. After centrifugation, 20 µL of the supernatant was taken to measure the amount of free HRP, using the procedure as described above.

The boronic acid-functionalized particles used were Si@pco-BA@BA, Si@pco-OH@BA, and Si@pNIPAm@BA.

Re-cycling of temperature-controlled protein binding and dissociation. In 2 mL microcentrifuge tubes, 3 mg of composite particles was suspended in 1 mL of protein solution containing 0.1 mg mL⁻¹ HRP and 1 mg mL⁻¹ BSA dissolved in 20 mM PBS buffer (pH 9.0, containing 0.5 M NaCl). The mixture was shaken at 20 °C for 2 h at 300 rpm using a thermal mixer. After centrifugation, 20 µL of the supernatant was taken to measure

the amount of free HRP using the 1-Step ABTS substrate. The remaining 980 µL of mixture was re-suspended, heated to 40 °C and shaken at 300 rpm for 2 h. After centrifugation, 20 µL of the supernatant was taken to measure the amount of free HRP. In the third step, the remaining 960 µL of mixture was re-suspended and shaken at 20 °C for 2 h. After centrifugation, 20 µL of the supernatant was taken to measure the amount of free HRP. In the last step, the remaining 940 µL of mixture was re-suspended, heated to 40 °C and shaken at 300 rpm for 2 h. After centrifugation, 20 µL of the supernatant was taken to measure the amount of free HRP.

Elution of glycoproteins from boronic acid-functionalized particles. In a series of 2 mL microcentrifuge tubes, 3 mg of composite particles was suspended in 1 mL of protein solution containing 0.1 mg mL⁻¹ HRP and 1 mg mL⁻¹ BSA dissolved in 20 mM PBS buffer (pH 9.0, containing 0.5 M NaCl). After shaking at 20 °C and at 300 rpm for 2 h, the samples were centrifuged, and the supernatant was removed. The composite particles were then washed 3 times with the same PBS buffer before the particles were separated by centrifugation. The glycoprotein bound to the particles was eluted using four different conditions:

(1) The composite particles were mixed with 1 mL of 0.2 M acetate buffer (pH 4.0) and shaken at 20 °C and at 300 rpm for 2 h. After centrifugation, 20 µL of the supernatant was taken to measure the amount of free HRP.

(2) The remaining 980 µL mixture from step 1 (containing the nanoparticles and proteins) was heated to 40 °C and shaken at 300 rpm for 2 h. After centrifugation, 20 µL of the supernatant was taken to measure the amount of free HRP.

(3) Similar to step 1, the initial composite particles with the bound glycoprotein were mixed with 1 mL of 0.2 M acetate buffer (pH 4.0) containing 1 mg mL⁻¹ D-fructose, and shaken at 20 °C and at 300 rpm for 2 h. After centrifugation, 20 µL of the supernatant was taken to measure the amount of free HRP.

(4) The remaining 980 µL mixture from step 3 (containing the nanoparticles and proteins) was heated to 40 °C and shaken at 300 rpm for 2 h. After centrifugation, 20 µL of the supernatant was taken to measure the amount of free HRP.

Results and discussion

Synthesis and characterization of nanoparticle-supported polymer brushes

In this work we designed a series of nanoparticle-supported polymer brushes functionalized with boronic acid at different locations in the polymer chains. The main idea was to find out the most suitable structure of nanoparticle-supported polymer brushes for glycoprotein separation. In addition to understanding this structure–function relationship, we were also interested in using thermodesorption to elute glycoproteins from the affinity materials, as this method might not involve the dramatic change in solvent conditions of protein samples, which can be beneficial for protein purification. The intermediate polymer brushes were synthesized from GMA and/or NIPAm according to our previous publications,^{13,36,37,42} and were further functionalized with the

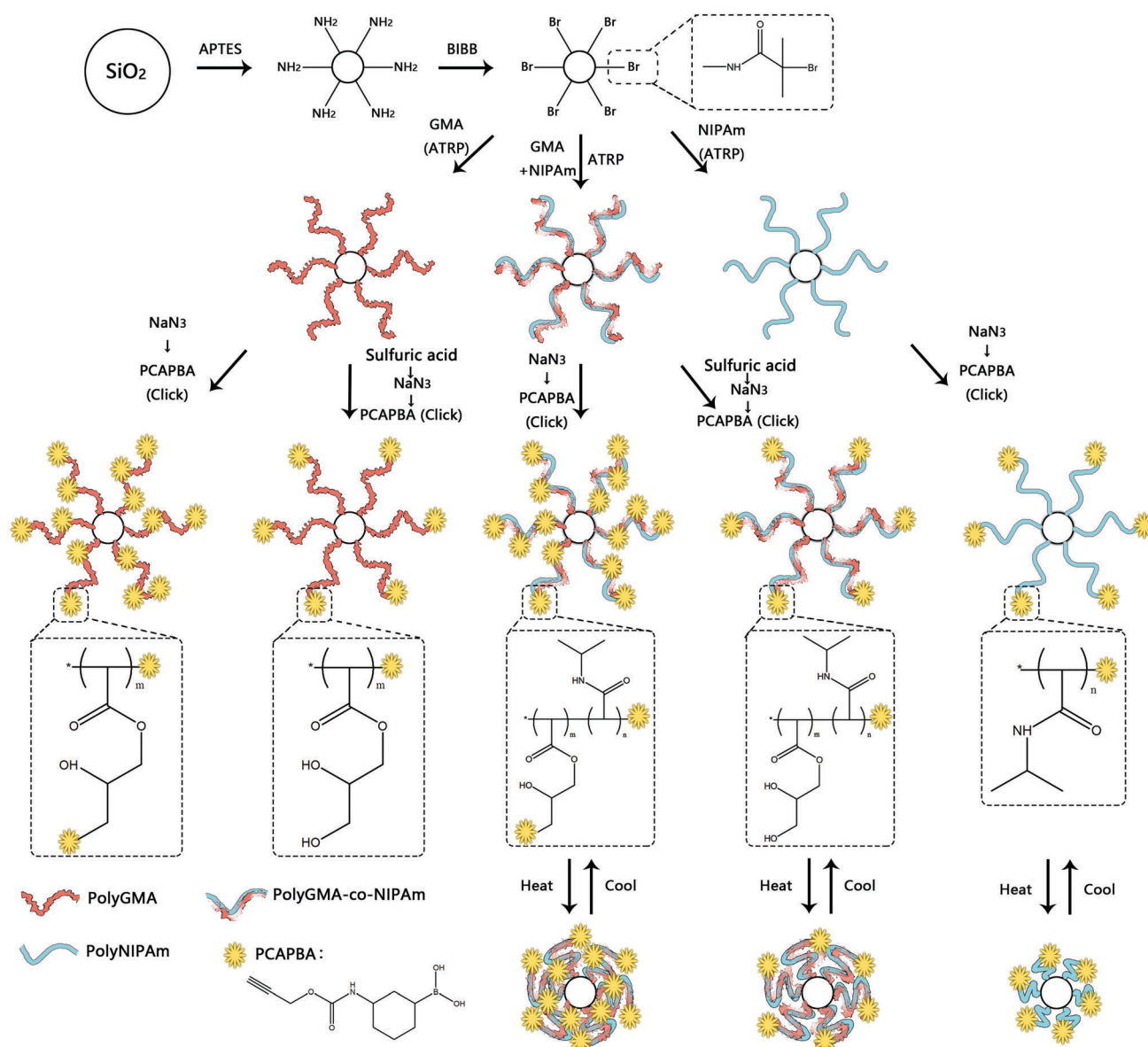


clickable boronic acid through a Cu(I)-catalyzed click reaction. The synthetic steps are illustrated in Scheme 1. Firstly, silica nanoparticles were prepared using the one-step Stöber process.⁴³ Amino groups were then introduced onto the silica nanoparticles to allow the ATRP initiator to be immobilized. The initiator-functionalized nanoparticles, Si@initiator, were then used to prepare three types of nanoparticle-supported polymer brushes: Si@pGMA, Si@pco and Si@pNIPAm. In the next step, the epoxide rings in the GMA moieties were converted into either pendant azide or diol groups by treatment with NaN₃ or sulfuric acid, and the terminal Br was converted into azide by reacting with NaN₃. Finally, the azide groups in the polymer brushes were used to introduce the clickable boronic acid with the high efficiency click reaction.

The nanoparticles synthesized in this work (including Si, Si@NH₂, Si@initiator, and the azide-functionalized nanoparticles

(Si@pGMA-N₃@N₃, Si@pGMA-OH@N₃, Si@pco-N₃@N₃, Si@pco-OH@N₃, Si@pNIPAm@N₃) were analyzed using SEM. As seen from the SEM images (Fig. 1 and Fig. S1, ESI†), the original Si nanoparticles were spherical with a smooth surface (Fig. 1a) and had a diameter of 100–150 nm. After grafting the polymer brushes on the surface, the obtained composite particles maintained a spherical shape, but the roughness of the particles increased obviously (Fig. 1b–f).

The chemical composition of the nanoparticles obtained after different synthetic steps was analysed by FT-IR. In Fig. 2, the most intense absorption peak appears at 1100 cm⁻¹, characteristic of the Si–O–Si vibration band. The peak at 1720 cm⁻¹ is attributed to the carbonyl group (C=O) in Si@polyGMA (Fig. 2b). After treatment with sodium azide and conjugation with the clickable boronic acid, a residual azide signal at



Scheme 1 Synthesis of boronic acid-functionalized polymer brushes on silica nanoparticles.



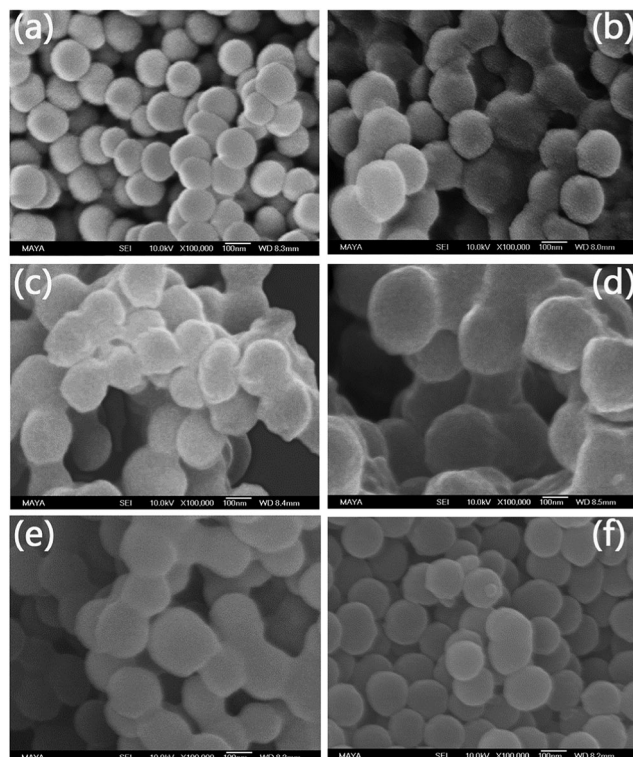


Fig. 1 SEM images of nanoparticles (a) Si, (b) Si@pGMA- N_3 @ N_3 , (c) Si@pGMA-OH@ N_3 , (d) Si@pco- N_3 @ N_3 , (e) Si@pco-OH@ N_3 , (f) Si@pNIPAm@ N_3 .

2100 cm^{-1} was still observed from Si@pGMA-BA@BA (Fig. 2c), which could be explained as the unreacted azide groups remaining in the polymer brushes, possibly due to the steric hindrance encountered during the click reaction. In the case of Si@pGMA-OH@BA, no azide signal was found (Fig. 2d), which can be explained by the low abundance of azide in its precursor Si@pGMA-OH@ N_3 , as well as complete conversion of the terminal azide into boronic acid. For Si@pco, in addition to the IR signals from the polymerized GMA monomer, new absorption bands appeared at 1650 cm^{-1} and 1540 cm^{-1} , which are the characteristic amide signals due to the polymerized NIPAm (Fig. 2e). After Si@pco was reacted with sodium azide and conjugated to the clickable boronic acid, the product Si@pco-BA@BA also displayed a residual azide signal at 2100 cm^{-1} (Fig. 2f), similar to that observed from Si@pGMA-BA@BA (Fig. 2c), possibly due to the incomplete conversion of azide into boronic acid caused by the steric hindrance during the click reaction. Like for Si@pGMA-OH@BA, the low abundance azide in Si@pco-OH@BA could not be detected after the terminal azide was converted into boronic acid *via* the click reaction (Fig. 2g). From the two nanoparticle-supported polyNIPAm brushes, Si@pNIPAm and Si@pNIPAm@BA, the characteristic amide I and amide II bands at 1650 cm^{-1} and 1540 cm^{-1} are obvious but are relatively weak, suggesting a lower content of the organic polymer in these composite particles.

The organic content of the different composite particles was assessed by TGA, and the chemical composition of the composite particles was measured by elemental analysis. These experimental results are presented in Table 1 and Fig. S3 (ESI[†]). In the TGA data,

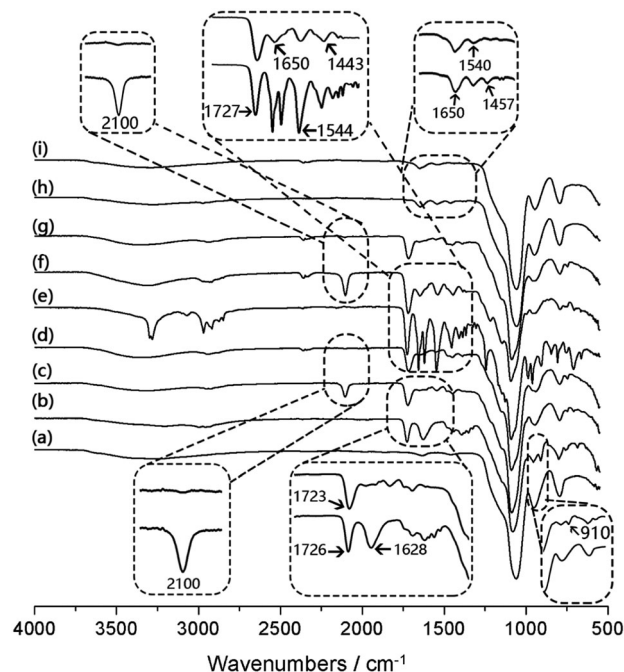


Fig. 2 IR spectra of Si (a), Si@pGMA (b), Si@pGMA-BA@BA (c), Si@pGMA-OH@BA (d), Si@pco (e), Si@pco-BA@BA (f), Si@pco-OH@BA (g), Si@pNIPAm (h) and Si@pNIPAm@BA (i).

the weight loss under $200\text{ }^{\circ}\text{C}$ is attributed to the evaporation of water. The residual weight remaining after the TGA was used to calculate the content of the inorganic silica material.

Among all the boronic acid-functionalized composite particles, Si@pNIPAm@BA had the lowest organic content, which is in agreement with the weak amide signals in the IR spectra of the nanoparticle-supported polyNIPAm brushes, Si@pNIPAm and Si@pNIPAm@BA (Fig. 2h and i). In our previous work, we also observed that the efficiency of growing polyNIPAm by SI-ATRP is lower than that of polyGMA.

Based on the contents of boron and inorganic silica, the number of boronic acids conjugated to the silica-supported nanoparticle brushes was calculated. Thus the boronic acid-functionalized composite nanoparticles Si@pGMA-BA@BA, Si@pGMA-OH@BA, Si@pco-BA@BA, Si@pco-OH@BA and Si@pNIPAm@BA were found to contain 0.75, 0.27, 0.45, 0.25 and 0.10 mmol boronic acid per gram of silica, respectively. The variation of boronic acid contents in these materials was expected based on the abundance of azide groups in their precursor particles Si@pGMA- N_3 @ N_3 , Si@pGMA-OH@ N_3 , Si@pco- N_3 @ N_3 , Si@pco-OH@ N_3 and Si@pNIPAm@ N_3 .

The content of boron in the final polymer can be calculated from the elemental analysis and the TGA results (Table 1, the last column). The content of boron was controlled by varying the density of azide groups in the polymer brush. Briefly, treating the epoxy-containing polymer (*e.g.* Si@pGMA) with sulfuric acid before introducing the azide leads to a lower boron content. For other composite particles, the boron content can be further adjusted by varying the amount of NIPAm relative to GMA, as well as by controlling the length of the polymer chain (by virtue of ATRP).



Table 1 Results of elemental analysis and TGA measurements

Particles	C (%)	H (%)	N (%)	Br (%)	B (%)	Si (%)	B/Si (mmol g ⁻¹)	B/polymer (mmol g ⁻¹)
Si@NH ₂	0.21	1.49	1.22	n.d.	n.d.	94.0	—	—
Si@initiator	3.88	1.94	0.51	1.33	n.d.	87.5	—	—
Si@pGMA	35.16	5.26	1.33	0.12	n.d.	38.6	—	—
Si@pGMA-N ₃ @N ₃	34.7	5.15	12.58	0.02	n.d.	17.3	—	—
Si@pGMA-BA@BA	29.53	4.42	7.72	n.d.	0.34	42.0	0.75	0.54
Si@pGMA-OH	28.69	4.94	0.33	0.12	n.d.	37.8	—	—
Si@pGMA-OH@N ₃	28.44	4.8	0.71	0.02	n.d.	40.7	—	—
Si@pGMA-OH@BA	32.00	5.27	0.91	n.d.	0.13	43.9	0.27	0.21
Si@pco	45.92	6.67	3.95	0.12	n.d.	28.3	—	—
Si@pco-N ₃ @N ₃	36.48	5.63	10.25	0.003	n.d.	24.1	—	—
Si@pco-BA@BA	20.27	5.36	4.99	n.d.	0.13	26.5	0.45	0.16
Si@pco-OH	34.02	5.93	3.04	0.11	n.d.	29.0	—	—
Si@pco-OH@N ₃	31.7	5.08	2.08	0.002	n.d.	32.5	—	—
Si@pco-OH@BA	36.58	5.67	1.34	n.d.	0.08	29.5	0.25	0.1
Si@pNIPAm	7.59	2.39	1.49	0.13	n.d.	83.7	—	—
Si@pNIPAm@N ₃	7.32	2.16	2.36	0.004	n.d.	73.2	—	—
Si@pNIPAm@BA	8.33	2.29	1.86	n.d.	0.09	82.1	0.10	0.5

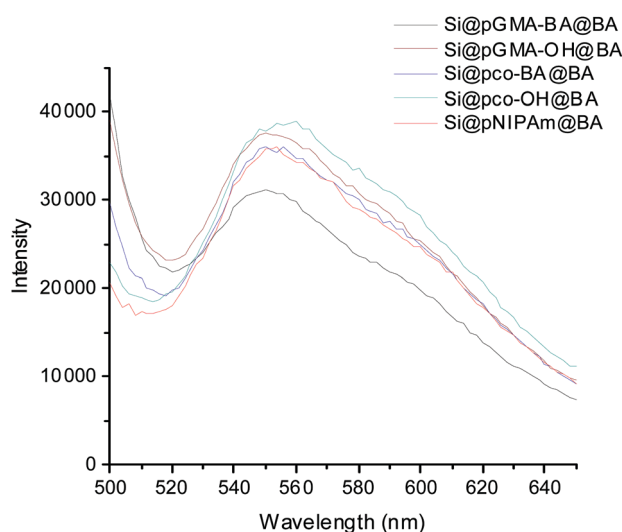


Fig. 3 Fluorescence emission of boronic acid-functionalized composite particles after exposure to ARS.

To confirm the successful conjugation of boronic acid into the polymer brushes, Si@pGMA-BA@BA, Si@pGMA-OH@BA, Si@pco-BA@BA, Si@pco-OH@BA and Si@pNIPAm@BA were treated with a fluorogenic *cis*-diol ARS to measure the binding-induced fluorescence emission.¹³ As shown in Fig. 3, all the boronic acid-functionalized particles emitted strong fluorescence after the particles were mixed with ARS in a basic buffer.

Binding of glycoproteins with different boronic acid-functionalized particles

To compare the protein-binding behaviours of the boronic acid-functionalized nanoparticles, we first carried out binding experiments in single protein solutions adjusted to different pH values and at different temperatures. OVA and HRP were used as model glycoproteins^{44,45} and BSA as a representative non-glycoprotein.⁴⁶

In order to compare the composite particles with different thermo-responsive properties more clearly, the equilibrium

binding data in Fig. 4 are presented in two sets: one for the non-thermo-responsive particles Si@pGMA-BA@BA and Si@pGMA-OH@BA, and another for the thermo-responsive particles Si@pco-BA@BA, Si@pco-OH@BA and Si@pNIPAm@BA.

From Fig. 4a and b, it is clear that the boronic acid-functionalized polymer brushes without the NIPAm component had a higher capacity for OVA binding (~ 125 mg g⁻¹) than the thermo-responsive polymer brushes (~ 100 mg g⁻¹), both higher than the capacity reported in our previous work.³⁷ The slightly higher capacity of Si@pGMA-BA@BA and Si@pGMA-OH@BA for glycoprotein binding can be explained as their higher boronic acid content than the thermo-sensitive copolymer brushes and polyNIPAm polymer brushes grafted on the silica nanoparticles (Si@pco-BA@BA, Si@pco-OH@BA and Si@pNIPAm@BA, Table 1). Interestingly, the polymer brushes functionalized with boronic acid at the chain termini displayed OVA capacity similar to that functionalized both at chain termini and at pendant side groups (*i.e.* Si@pGMA-OH@BA vs. Si@pGMA-BA@BA and Si@pco-OH@BA vs. Si@pco-BA@BA). It is likely that due to the high grafting density (as discussed in our previous work) and a steric effect, the terminal boronic acid played the most important role in binding the glycoprotein.

Glycoprotein binding with all the boronic acid-functionalized polymer brushes displayed clear pH-dependence. As seen in Fig. 4c and d, when the solution pH changed from 9 to 4, OVA binding to the composite particles decreased significantly. The pH-dependent OVA binding can be explained as the structural change of the appended phenylboronic acid upon pH variation. Under alkaline conditions the boron atom has a sp³ configuration and is able to form strong boronate ester bonds with *cis*-diols, whereas under acidic conditions the boron changes to a sp² configuration, leading to hydrolysis of the boronate ester bonds to release the bound glycoprotein. The sensitivity of the boronate ester bond against pH can therefore be used to control the binding and dissociation of glycoproteins from boronate affinity materials.^{37,47}

For the two types of glycoproteins tested, OVA and HRP, in general the uptake of OVA by the boronic acid-functionalized



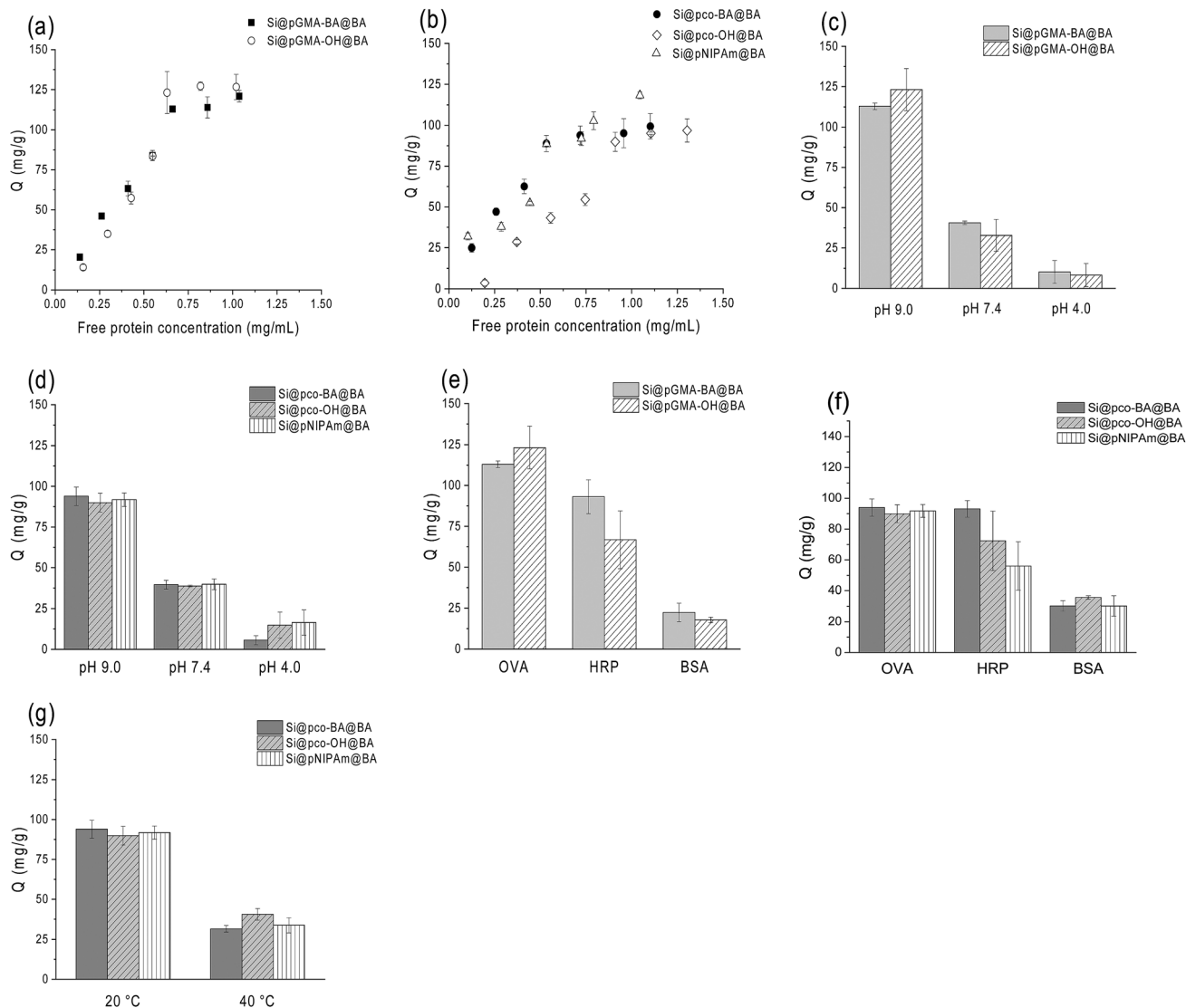


Fig. 4 Protein binding characteristics. (a and b) Binding isotherms of OVA measured at 20 °C and pH 9.0. (c and d) OVA binding to the composite particles measured at 20 °C and different pH values. The initial concentration of OVA was 1 mg mL⁻¹. (e and f) Binding of different proteins to the composite particles measured in pH 9 buffer at 20 °C. The initial concentration of the individual protein was 1 mg mL⁻¹. (g) Binding of OVA to the thermo-responsive polymer brushes measured in pH 9 buffer at different temperatures. The initial OVA concentration was 1 mg mL⁻¹.

polymer brushes was higher than that of HRP (Fig. 4e and f). Under the same conditions the non-specific protein binding, as estimated from the BSA binding, was much lower. The difference of binding between OVA and HRP by boronate affinity materials has been observed in our previous investigations and can be explained as the content of the accessible *cis*-diol structures in these two glycoproteins. Although HRP has more glycosylation sites, OVA has a higher saccharide content and therefore can present more accessible *cis*-diols to bind the boronic acid polymer brushes.

To evaluate the water-compatibility of the nanoparticle-supported polymer brushes, we measured the rate of particle sedimentation in aqueous buffer by monitoring the change of optical transmittance of the particle suspension. As shown in Fig. S4a and b (ESI[†]), at 20 °C the particle Si@pGMA-BA@BA precipitated significantly faster than Si@pGMA-OH@BA. The faster

sedimentation of Si@pGMA-BA@BA was due to the multiple phenylboronic acids in the polymer chains. The phenylboronic acids at low pH are relatively hydrophobic, making the composite particles form aggregates to sediment faster than at high pH. Because Si@pGMA-OH@BA contains abundant hydroxyl groups, the composite particle is hydrophilic in a wide pH range, and therefore it does not sediment even at low pH.

The three nanoparticle-supported polymer brushes containing NIPAm were designed to introduce thermo-responsive properties into the composite particles. In the literature it is well known that polyNIPAm has a lower critical solution temperature (LCST) of around 32 °C. Below the LCST polyNIPAm is hydrophilic and exists as an extended and hydrated chain, whereas above the LCST the polymer chain becomes hydrophobic and collapses. To verify that the two boronic acid-functionalized copolymer brushes have similar temperature-controlled structural changes, we studied the



rate of particle sedimentation at 20 °C and 40 °C. As shown in Fig. S4c and d (ESI†), the two composite particles Si@pco-BA@BA and Si@pco-OH@BA dispersed in pH 9 buffer at 40 °C settled slightly faster than in the same buffer at 20 °C. The faster sedimentation can be explained as a result of the thermo-induced dehydration and collapse of the polymer brushes. We should note, however, that at 40 °C although the NIPAm components tend to make the polymer brushes collapse, the higher pH (pH 9) makes the boronic acids become negatively charged, leading to stabilizing the colloidal suspension. The overall result is that the sedimentation of Si@pco-BA@BA and Si@pco-OH@BA became not very obvious when the temperature increased from 20 °C to 40 °C.

One of our intriguing questions is if the temperature-controlled structural transition can be utilized to control the binding of glycoproteins to thermo-responsive polymer brushes. In the next experiment, we therefore studied the effect of solution temperature on the uptake of OVA by the three composite particles Si@pco-BA@BA, Si@pco-OH@BA and Si@pNIPAm@BA. As shown in Fig. 4g, when the solution temperature changed from 20 °C to 40 °C, for all the three polymer brushes the amount of OVA binding in pH 9 buffer decreased more than 60% (from $\sim 90 \text{ mg g}^{-1}$ to $\sim 35 \text{ mg g}^{-1}$). This profound temperature effect provides an interesting approach to modulate glycoprotein binding with the new boronate affinity particles, and should provide a convenient means to enable more simplified protein separation.

Verification of protein separation by SDS-PAGE

In order to observe more directly the protein separation with the temperature-responsive nanoparticles, we used SDS-PAGE analysis to monitor a mixture of OVA and BSA after it was treated with Si@pNIPAm@BA. The protein mixture was first incubated with Si@pNIPAm@BA at 20 °C, and then at 40 °C. The unbound proteins at these two different temperatures were analysed by SDS-PAGE.

As shown in Fig. 5, at 20 °C both BSA ($\sim 66 \text{ kDa}$) and OVA ($\sim 45 \text{ kDa}$) were depleted from solution by the nanoparticles (Lane 2, 3). The small shift of the protein bands, compared to the marker (Lane 1), was caused by the difference in the buffer pH. The BSA binding can be attributed to nonspecific adsorption. When the temperature was raised to 40 °C, the majority of BSA was released from the nanoparticles, while the amount of OVA released was relatively less than BSA (Lane 4). The release of protein caused by the increased temperature can be explained as a result of the collapse of the polymer brushes on the particle surface, making the originally bound BSA be excluded from the composite particles. For OVA, because of its covalent binding to the terminal boronic acid, the effect of increased temperature is less than for BSA. After the washing steps, it is clear that the final protein solution eluted from the nanoparticles contained mainly OVA (Lane 6).

Controlling glycoprotein binding *via* temperature and addition of monosaccharides

In conventional boronate affinity separation, variation of pH is the most used method to control the binding and release of *cis*-diols from boronic acid-functionalized adsorbents. Alternative methods to release glycoproteins at a constant pH, *e.g.* by

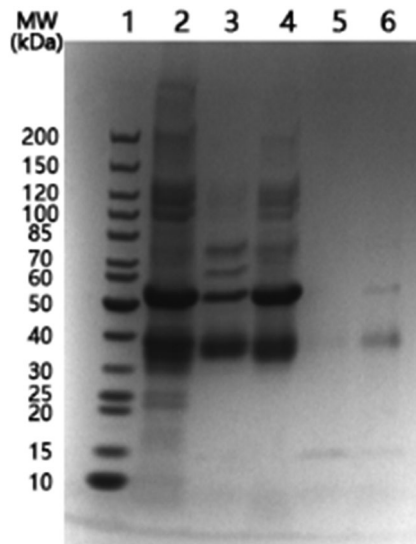


Fig. 5 SDS-PAGE analysis of the protein sample treated with Si@pNIPAm@BA. Lane 1, protein marker; Lane 2, protein mixture containing 1 mg mL^{-1} OVA and 1 mg mL^{-1} BSA; Lane 3, unbound protein mixture at 20 °C; Lane 4, unbound protein mixture at 40 °C; Lane 5, collected washing buffer; and Lane 6, proteins eluted in acetate buffer (pH 4).

adjusting temperature, can be useful because it does not require a significant adjustment of pH. As glycoprotein binding is mediated by the saccharide moieties, another way of releasing glycoprotein without changing the pH is by adding competing monosaccharides. As an example, under alkaline conditions D-fructose is able to form stable ester bonds with boronic acids, and it has been used as a competing *cis*-diol to displace glycoproteins from boronate affinity particles.^{42,48}

To investigate the effects of temperature and fructose on glycoprotein binding in a more realistic environment, we studied the binding of HRP on the three thermo-responsive particles in the presence of excess non-glycoprotein, BSA. In this experiment HRP was used as a model glycoprotein, as its concentration could be determined conveniently by measuring the enzyme activity. As shown in Fig. 6, all the three boronic acid-functionalized silica nanoparticles (containing NIPAm) showed a very clear response to temperature and D-fructose. It is noteworthy that temperature and fructose displayed a concerted effect on HRP binding: addition of fructose and increasing temperature resulted in the maximum release of HRP from the boronate affinity particles. The temperature and fructose response were not affected by the high concentration of BSA.

Using Si@pNIPAm@BA as an example, we investigated the reversibility of the temperature-controlled HRP binding to boronic acid-functionalized polymer brushes. Fig. 6d shows that after the solution temperature was increased from 20 °C to 40 °C, HRP binding decreased from 23 mg g^{-1} to 4 mg g^{-1} . After the solution temperature returned to 20 °C, in the same sample the HRP binding went back to the initial value. This result indicates clearly that the temperature-controlled glycoprotein binding and release are reversible. We should note that for the nanoparticle-supported polymer brushes that did not contain



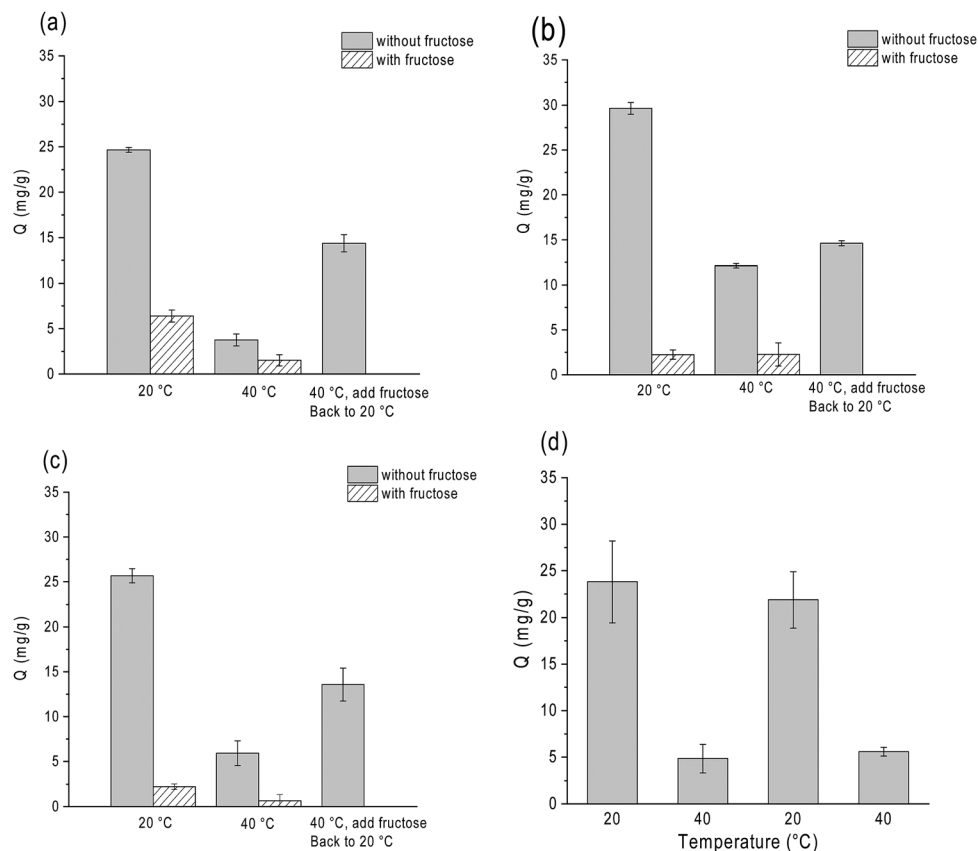


Fig. 6 Controlling HRP binding by temperature and fructose. The boronic acid-functionalized particles Si@pco-BA@BA (a), Si@pco-OH@BA (b) and Si@pNIPAm@BA (c) were mixed with the protein solution at 20 °C in pH 9 buffer to measure the HRP binding. After this step the samples were heated to 40 °C before the HRP binding was measured. For the last bar, 1 mg of fructose was added at 40 °C to the initially fructose-free mixture before the sample was cooled to 20 °C. In (d) the HRP binding to Si@pNIPAm@BA was measured from a sample undergoing repeated alteration of temperature between 20 and 40 °C.

the thermo-responsive NIPAm component (Si@pGMA-BA@BA and Si@pGMA-OH@BA), variation of temperature did not affect the HRP binding (Fig. S5, ESI†). The possibility of controlling protein binding *via* temperature is interesting for future exploration, for example to study selective loading and release of protein pharmaceuticals in thermo-responsive polymer brushes.

Boronate affinity chromatography is a proven technique for purification of glycoproteins.^{23–28} In general terms, separation of glycoproteins through reversible boronate ester bonds does not cause protein denaturation. In the protein binding experiment, we quantified the glycoprotein HRP by measuring its enzyme activity. The temperature-recycling experiment (Fig. 6d) demonstrated that the HRP binding and releasing could be controlled by temperature variation, and the binding-releasing event was truly reversible. As the enzyme activity of the released protein resumed after returning to the original temperature, it is very clear that the protein function before and after the binding was not affected.

Controlling the release of glycoprotein from boronate affinity particles

As mentioned already in previous sections, variation of pH is often used to control glycoprotein binding to boronate affinity materials. To investigate how several effectors can be combined to influence glycoprotein release, we studied the binding and

release of HRP on Si@pco-BA@BA in the presence of excess of BSA under different solvent conditions. The composite particles were first loaded with HRP in the binary protein solution in pH 9 buffer and washed to remove the non-specifically adsorbed protein. The glycoprotein was then eluted from the particles using pH 4 buffer at different temperatures, with and without the addition of fructose in the elution buffer. As shown in Fig. 7, although elution with pH 4 buffer could release HRP, adding fructose in the elution buffer increased the protein recovery by almost 50%. When the protein elution with pH 4 buffer was carried out at 40 °C, we obtained the highest HRP recovery irrespective of fructose. This result points to a more efficient means of glycoprotein separation by combining acidic buffer with thermo-desorption, which can be easily applied to thermo-responsive boronate affinity materials.

In this work the boronic acids were immobilized on the polymer brushes through stable covalent bonds. Consequently, no boronic acid will detach from the composite particles to associate with the protein in the elution step. As the binding of glycoprotein to the boronic acid-modified polymer brushes was through a carbohydrate moiety, the glycoproteins are expected to maintain their structure, as we have demonstrated in a previous work where a bound glycoenzyme (HRP) maintained its catalytic activity.⁴²



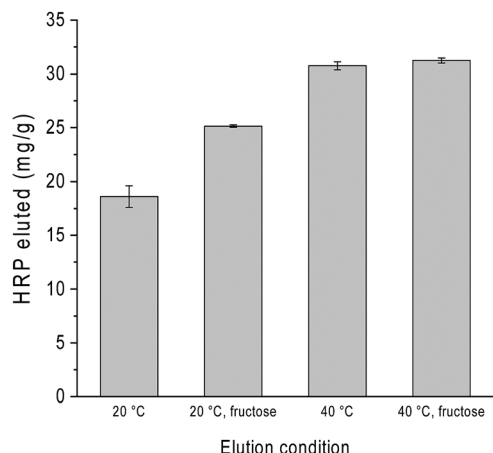


Fig. 7 Elution of HRP from Si@pco-BA@BA using acidic buffer under different conditions.

Conclusions

In this work, we synthesized five different types of boronic acid-functionalized polymer brushes on silica nanoparticles to study their structure–function relationship. We also investigated the use of temperature variation to control the binding and release of glycoproteins from boronate affinity particles. For polymer brushes designed to bind glycoproteins, our results suggest that the most important boronic acids contributing to protein binding are located at the chain termini of the polymer brushes. For the thermo-responsive polymer brushes, we demonstrated that it is possible to use temperature to control the glycoprotein binding and release. The reversible characteristic of the boronate ester bonds involved in the protein separation process has been revealed through temperature recycling experiments. For the first time, we have shown that glycoproteins bound on boronate affinity particles can be eluted by simply varying the solution temperature. When combined with acidic buffer, the new thermo-desorption provided the highest protein recovery. The new boronic acid-functionalized, nanoparticle-supported polymer brushes hold interesting potential to be exploited not only for protein separation, but also for other applications including controlled drug delivery and release.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work was supported by the Swedish Foundation for International Cooperation in Research and High Education (STINT, grant no. CH2015-6254), the Swedish Research Council FORMAS (grant no. 2016-00591) and the National Natural Science Foundation of China (NSFC, grant no. 21611130030). Huiting Ma thanks the China Scholarship Council (201606160072) for a visiting scholarship. Huiting Ma is a visiting student in Lund University

and a PhD candidate in the Huazhong University of Science and Technology.

References

- 1 S. G. Boyes, B. Akgun, W. J. Brittain and M. D. Foster, *Macromolecules*, 2003, **36**, 9539–9548.
- 2 X. Kong, T. Kawai, J. Abe and T. Iyoda, *Macromolecules*, 2001, **34**, 1837–1844.
- 3 K. Matyjaszewski, *Macromolecules*, 2012, **45**, 4015–4039.
- 4 Y.-P. Wang, X.-W. Pei and K. Yuan, *Mater. Lett.*, 2005, **59**, 520–523.
- 5 T. von Werne and T. E. Patten, *J. Am. Chem. Soc.*, 1999, **121**, 7409–7410.
- 6 L. Zhang and M. Fang, *Nano Today*, 2010, **5**, 128–142.
- 7 F. Zhou and W. T. Huck, *Phys. Chem. Chem. Phys.*, 2006, **8**, 3815–3823.
- 8 C. Perruchot, M. Khan, A. Kamitsi, S. V. Armes, T. von Werne and T. Patten, *Langmuir*, 2001, **17**, 4479–4481.
- 9 Z. Zhou, S. Zhu and D. Zhang, *J. Mater. Chem.*, 2007, **17**, 2428–2433.
- 10 S. Zhang, W. Guo, J. Wei, C. Li, X.-J. Liang and M. Yin, *ACS Nano*, 2017, **11**, 3797–3805.
- 11 I. Y. Galaev and B. Mattiasson, *Trends Biotechnol.*, 1999, **17**, 335–340.
- 12 G. Pasparakis and M. Vamvakaki, *Polym. Chem.*, 2011, **2**, 1234–1248.
- 13 L. Jiang, M. E. Messing and L. Ye, *ACS Appl. Mater. Interfaces*, 2017, **9**, 8985–8995.
- 14 J. Zhang, T. He, L. Tang and Z. Q. Zhang, *J. Sep. Sci.*, 2016, **39**, 1691–1699.
- 15 L. Zhang, Y. Xu, H. Yao, L. Xie, J. Yao, H. Lu and P. Yang, *Chem. – Eur. J.*, 2009, **15**, 10158–10166.
- 16 E. Robertson and J. Kennedy, *Bioseparation*, 1996, **6**, 1–15.
- 17 F. M. Okanda and Z. El Rassi, *Electrophoresis*, 2006, **27**, 1020–1030.
- 18 M. Madera, Y. Mechref and M. V. Novotny, *Anal. Chem.*, 2005, **77**, 4081–4090.
- 19 M. Bedair and Z. El Rassi, *J. Chromatogr. A*, 2004, **1044**, 177–186.
- 20 P. M. Drake, B. Schilling, R. K. Niles, M. Braten, E. Johansen, H. Liu, M. Lerch, D. J. Sorensen, B. Li and S. Allen, *Anal. Biochem.*, 2011, **408**, 71–85.
- 21 X. Mao, Y. Luo, Z. Dai, K. Wang, Y. Du and B. Lin, *Anal. Chem.*, 2004, **76**, 6941–6947.
- 22 C. Zhang, C. Bi, W. Clarke and D. S. Hage, *J. Chromatogr. A*, 2017, **1523**, 114–122.
- 23 Y. Qu, J. Liu, K. Yang, Z. Liang, L. Zhang and Y. Zhang, *Chem. – Eur. J.*, 2012, **18**, 9056–9062.
- 24 K. Sparbier, S. Koch, I. Kessler, T. Wenzel and M. Kostrzewa, *J. Biomol. Tech.*, 2005, **16**, 407–413.
- 25 Y. Liu, L. Ren and Z. Liu, *Chem. Commun.*, 2011, **47**, 5067–5069.
- 26 R. Nishiyabu, Y. Kubo, T. D. James and J. S. Fossey, *Chem. Commun.*, 2011, **42**, 1106–1123.
- 27 D. Li, Y. Chen and Z. Liu, *Chem. Soc. Rev.*, 2015, **44**, 8097–8123.
- 28 Z. Liu and H. He, *Acc. Chem. Res.*, 2017, **50**, 2185–2193.



- 29 L. Ren, Z. Liu, M. Dong, M. Ye and H. Zou, *J. Chromatogr. A*, 2009, **1216**, 4768–4774.
- 30 H. Wang, Z. Bie, C. Lü and Z. Liu, *Chem. Sci.*, 2013, **4**, 4298–4303.
- 31 X. Zhang, X. He, L. Chen and Y. Zhang, *J. Mater. Chem. B*, 2014, **2**, 3254–3262.
- 32 W. Xi, T. F. Scott, C. J. Kloxin and C. N. Bowman, *Adv. Funct. Mater.*, 2014, **24**, 2572–2590.
- 33 Y. Xu, Z. Wu, L. Zhang, H. Lu, P. Yang, P. A. Webley and D. Zhao, *Anal. Chem.*, 2008, **81**, 503–508.
- 34 B. J. Gould and P. M. Hall, *Clin. Chim. Acta*, 1987, **163**, 225–230.
- 35 Q. Zhang, A. A. Schepmoes, J. W. Brock, S. Wu, R. J. Moore, S. O. Purvine, J. W. Baynes, R. D. Smith and T. O. Metz, *Anal. Chem.*, 2008, **80**, 9822–9829.
- 36 T. Suksrichavalit, K. Yoshimatsu, V. Prachayasittikul, L. Bülow and L. Ye, *J. Chromatogr. A*, 2010, **1217**, 3635–3641.
- 37 L. Jiang, H. Bagán, T. Kamra, T. Zhou and L. Ye, *J. Mater. Chem. B*, 2016, **4**, 3247–3256.
- 38 J.-F. Lutz, Ö. Akdemir and A. Hoth, *J. Am. Chem. Soc.*, 2006, **128**, 13046–13047.
- 39 H. Tu, C. E. Heitzman and P. V. Braun, *Langmuir*, 2004, **20**, 8313–8320.
- 40 T. Kawai, K. Saito and W. Lee, *J. Chromatogr. B: Biomed. Sci. Appl.*, 2003, **790**, 131–142.
- 41 K. Ohno, T. Morinaga, S. Takeno, Y. Tsujii and T. Fukuda, *Macromolecules*, 2007, **40**, 9143–9150.
- 42 H. Bagán, T. Kamra, L. Jiang and L. Ye, *Macromol. Chem. Phys.*, 2017, **218**, 1600432.
- 43 W. Stöber, A. Fink and E. Bohn, *J. Colloid Interface Sci.*, 1968, **26**, 62–69.
- 44 J. A. Huntington and P. E. Stein, *J. Chromatogr. B: Biomed. Sci. Appl.*, 2001, **756**, 189–198.
- 45 N. C. Veitch, *Phytochemistry*, 2004, **65**, 249–259.
- 46 K. A. Majorek, P. J. Porebski, A. Dayal, M. D. Zimmerman, K. Jablonska, A. J. Stewart, M. Chruszcz and W. Minor, *Mol. Immunol.*, 2012, **52**, 174–182.
- 47 L. Li, Y. Lu, Z. Bie, H. Y. Chen and Z. Liu, *Angew. Chem., Int. Ed.*, 2013, **52**, 7451–7454.
- 48 W. Wang, X. Gao and B. Wang, *Curr. Org. Chem.*, 2002, **6**, 1285–1317.

