

# Reversible, high-affinity surface capturing of proteins directed by supramolecular assembly

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**ABSTRACT:** The ability to design surfaces with reversible, high affinity protein binding sites represents a significant step forward in the advancement of analytical methods for diverse biochemical and biomedical applications. Herein, we report a dynamic supramolecular strategy to directly assemble proteins on surfaces based on multivalent host-guest interactions. The host-guest interactions are achieved by one-step nanofabrication of a well-oriented  $\beta$ -cyclodextrin host-derived self-assembled monolayer on gold ( $\beta$ -CD-SAM) that forms specific inclusion complexes with hydrophobic amino acids located on the protein's surface. Cytochrome C, insulin,  $\alpha$ -chymotrypsin and RNase A are used as model guest proteins. Surface plasmon resonance (SPR) and static time of flight secondary ion mass spectrometry (ToF-SIMS) studies demonstrate that all four proteins interact with the  $\beta$ -CD-SAM in a specific manner via the hydrophobic amino acids on the surface of the protein. The  $\beta$ -CD-SAMs bind the proteins with high nanomolar to single-digit micromolar dissociation constants ( $K_D$ ). Importantly, while the proteins can be captured with high affinity, their release from the surface can be achieved under very mild conditions. Our results expose the great advantages of using a supramolecular approach for controlling protein immobilization, in which the strategy described herein provides unprecedented opportunities to create advanced bioanalytic and biosensor technologies.

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## Introduction

Protein immobilization on material surfaces is at the heart of many biochemical and biomedical applications, ranging from development of biosensors,<sup>1</sup> bioanalytical tools,<sup>2</sup> to biocatalysis<sup>3</sup> and drug delivery.<sup>4</sup> Along with the widespread application of protein immobilization technology, there are also significant challenges to overcome and practical difficulties to resolve. Notably, proteins are required to be strongly linked onto the surface with high density without adversely affecting their activity.<sup>5</sup> Concomitantly, suitable immobilization methods are required that facilitate the facile release of the bound proteins to regenerate and reuse the assay, material or device.<sup>6</sup>

Over the years, many different approaches have been developed for protein immobilization on surfaces utilizing physisorption,<sup>7</sup> bioaffinity interaction<sup>8</sup> and covalent coupling.<sup>9</sup> There are advantages and disadvantages to each of the aforementioned approaches, pertaining to simplicity of fabrication, reusability, stability of immobilized protein and retention of protein conformation and activity.<sup>9,10</sup> A relatively unexplored strategy, although with vast potential, involves the use of non-covalent supramolecular interactions.<sup>11-13</sup> Taking advantage of the highly stable chemistry of cucurbituril-ferrocene inclusion complexes, proteins have been site-specifically modified with ferrocene to be reversibly and stably immobilized onto a cucurbit[7]uril (CB[7]) monolayer.<sup>14</sup> In a recent example,<sup>11</sup> proteins have been tagged with a different number of hexahistidines (His6) at specific positions on the protein to understand

its effect on protein orientation and binding strength upon interaction with nickel nitrilotriacetic acid (Ni-NTA) self-assembled monolayers. This strategy gave access to a high degree of control not only over the orientation of the proteins on the surfaces but also to the binding strength of the proteins with the surface. Moreover the multivalency effect yielded dissociation constant values in the low nM regime. These and other examples in the literature<sup>15-17</sup> illustrate how supramolecular interactions and their predictability can be used to design protein immobilization systems with enhanced features and improved ability to tune their properties as required. Much though remains to be uncovered regarding the potential of supramolecular assembly for devising protein immobilization systems with emergent properties amenable to current needs. With this proviso in mind, in this work we investigated the ability of surface-tethered  $\beta$ -cyclodextrins ( $\beta$ -CDs) to promote facile, versatile and stable protein assembly but yet disassembly obviating the use of harsh conditions and protein denaturation.

CDs are macrocyclic oligosaccharide-derived non-symmetrical host molecules that selectively recognize, as a result of their apolar cone-shaped cavity, among other hydrophobic moieties, hydrophobic amino acids.<sup>18</sup> Notably,  $\beta$ -CDs have been demonstrated to form inclusion complexes with hydrophobic amino acids on the surfaces of proteins, namely phenylalanine (Phe), tyrosine (Tyr), leucine (Leu) and tryptophan (Trp).<sup>18-22</sup> These amino acids in the proteins can be

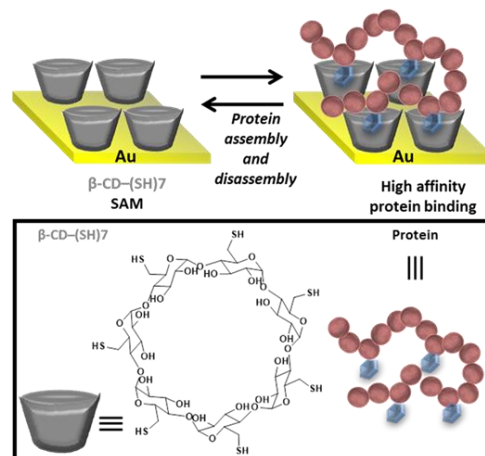
strongly and selectively included into the  $\beta$ -CD cavity, forming complexes with mM binding affinity.<sup>23</sup> This capability has been used to good advantage in drug delivery<sup>24</sup> and formulation of pharmaceutical proteins.<sup>25</sup> However, there is clear scope to broaden the ways in which the unique  $\beta$ -CD-protein non-covalent interactions can be utilized to the modulation and assembly of proteins. In this framework, we present a study, enabled by the non-covalent interactions between  $\beta$ -CD and proteins, which demonstrates previously unmatched simplicity, versatility and capability to create highly stable, yet reversible, protein assemblies on material surfaces. Considering that proteins display at their surfaces multiple hydrophobic amino acids, we have harnessed the concept of multivalency<sup>26</sup> to promote the simultaneous binding of multiple amino acids on the protein to multiple well-oriented  $\beta$ -CD moieties tethered onto a gold surface. The heptathiolated derivative of  $\beta$ -CD ( $\beta$ -CD-(SH)7), in which the seven thiol moieties are installed on the smaller primary rim of the  $\beta$ -CD, was used to directly create a  $\beta$ -CD-terminated self-assembled monolayer (SAM) via multiple thiolate-gold covalent bonds (Scheme 1). This approach allows the larger secondary rim to be exposed at the interface, facilitating the binding of the amino acid residues from the protein surface inside the cavity. Four model proteins, namely cytochrome C, insulin,  $\alpha$ -chymotrypsin and RNase A, which display randomly distributed hydrophobic amino acids throughout their surfaces (Figure 1) for  $\beta$ -CD binding, were investigated to assess the generality of our observations.

## Results and discussion

Since concentration and time play an important role on SAM formation and organization,<sup>27,28</sup> these parameters were investigated to evaluate the optimum conditions to create well-oriented surface tethered  $\beta$ -CDs.  $\beta$ -CD-(SH)7 SAMs were formed by immersing freshly cleaned gold substrates in either 0.1 mM or 1 mM DMF solutions of  $\beta$ -CD-(SH)7 for either 12, 24 or 48 h. In order to ascertain the properties of the formed  $\beta$ -CD-(SH)7 SAMs, contact angle, ellipsometry and X-ray photoelectron spectroscopy (XPS) analysis were carried out (Figure 2).

As illustrated by the water contact angle, all  $\beta$ -CD-(SH)7 SAMs exhibit hydrophilic properties (Figure 2a) due to the presence of the hydroxyl groups on the outer edge of the amphiphilic  $\beta$ -CD molecules. However, their hydrophilicity differs, with advancing contact angle ranging from 25° (0.1 mM-12 h) to 55° (1 mM-48 h). These differences in hydrophilicity can provide some indication of the orientation of the CD macrocycle on the surface, where a decrease in hydrophobicity of the SAM might suggest the presence of a more exposed hydrophobic cavity at the interface. While significant contact angle differences were observed between 0.1 mM-12 h and 1 mM-48 h, the intermediate conditions show comparable ad-

vancing and receding contact angles. The hysteresis ( $\theta_{Adv} - \theta_{Rec}$ ) values for all the  $\beta$ -CD-(SH)7 SAMs vary from 8° to 15°, suggesting the formation of relatively ordered and well defined monolayers.



Scheme 1. Schematic representation of an effective supramolecular strategy to promote high and strong protein binding, which can be reversed under mild conditions. The strategy involves formation of a well-oriented  $\beta$ -CD-(SH)7 SAM, which forms multivalent interactions with hydrophobic moieties (shown as blue hexagons) present on the protein surface.

Ellipsometry data (Figure 2b) provided further information regarding the organization of the SAMs, in which thickness dimensions close to the theoretical height of the  $\beta$ -CD molecule (0.78 nm)<sup>29</sup> were observed in all but one instance. These findings are consistent with the SAMs being packed with the CD orientated with the macrocycle parallel to the plane of the gold surface. The SAM fabricated with a 0.1 mM solution of  $\beta$ -CD-(SH)7 incubated for 12 h exhibited a layer thickness of 1.60 nm, which is comparable to the theoretical outer diameter of the  $\beta$ -CD (1.54 nm).<sup>29</sup> This suggesting that the macrocycle might be orientated perpendicular to the gold surface in this instance.

Further studies were carried out by XPS to probe the chemical composition and further understand the packing density of the SAMs. XPS data confirms the formation of the  $\beta$ -CD-(SH)7 SAMs, showing signals from the S (2p), O (1s) and C (1s) binding energies on all surfaces (data not shown). Interestingly, the ratios of S to Au (Figure 2c) and bound (S-Au) to unbound (SH) sulfur (Figure 2d) vary, depending mainly on the concentration used. The SAMs created by incubation of a 0.1 mM  $\beta$ -CD-(SH)7 solution exhibited a higher ratio of bound sulfur, even after only 12 h of incubation. The SAMs formulated with 0.1 mM solutions of  $\beta$ -CD-(SH)7 are attached to the surface via an average of 5 S-Au bonds, whereas 3-4 S-Au bonds are observed for the SAMs formed using 1 mM solutions of  $\beta$ -CD-(SH)7. As previously reported,<sup>30</sup> this

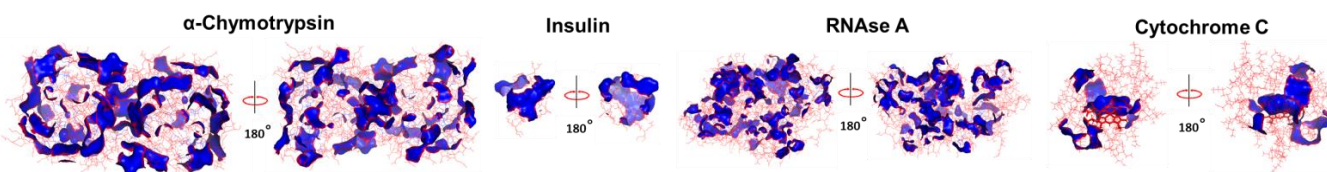


Figure 1. Overall structures of  $\alpha$ -chymotrypsin, insulin, RNase A and cytochrome C, with the hydrophobic amino acids on the surfaces of proteins (i.e. Phe, Tyr, Leu and Trp) highlighted in blue. The structural images were generated using PyMol.

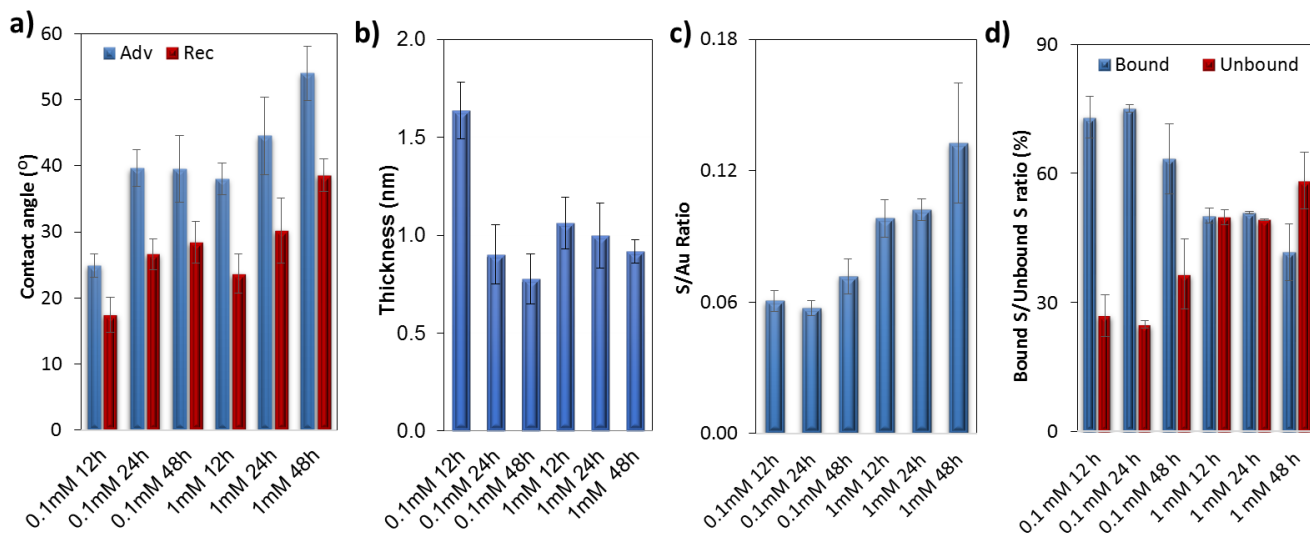


Figure 2. a) Advancing and receding water contact angles, b) ellipsometric thickness, c) S/Au ratio as determined by XPS analysis and d) Bound S/Unbound S ratio as determined by XPS analysis for the  $\beta$ -CD-(SH)7 SAMs formed using different concentrations and times.

difference in the number of covalent bonds formed can affect lateral diffusion during SAM formation and consequently surface molecular coverage. Indeed, the increased mobility restrictions imposed by the multiple thiol anchors associated with SAMs fabricated from 0.1 mM solutions could explain the lower sulfur content at these surfaces as established by calculating the XPS S/Au ratios. On the other hand, the higher sulfur content in SAMs formulated in a 1 mM solution provides an indication of a more tightly packed monolayer when compared with the SAMs formed from the 0.1 mM concentration. This characteristic is important for protein surface immobilization since sparsely packed CDs on the surface could lead to non-specific binding.

In order to further ascertain the density or permeability of the  $\beta$ -CD-(SH)7 SAMs, cyclic voltammetry (CV) studies were undertaken. Since  $\beta$ -CD moieties are known to form stable complexes with ferrocene and its derivatives,<sup>31,32</sup> water soluble ferrocene carboxylic acid (FCA) was used as a redox-active probe to quantify the  $\beta$ -CD surface density.  $\beta$ -CD-(SH)7 SAMs were immersed in a 0.1 M FCA solution for 4 hours and then rinsed with ultra-high quality (UHQ) water. Subsequently, the oxidation and the reduction potentials of FCA at the modified gold surfaces were investigated by cyclic voltammetry (Figure 3).

The cyclic voltammograms associated with surfaces modified with the 1 mM SAMs displayed the lowest anodic and cathodic current peak intensities. Interestingly the peak separation observed at CVs obtained for the 0.1 mM and 1 mM SAM-modified surfaces yielded peak separations of  $41 \pm 5$  mV and  $32 \pm 3$  mV, respectively. When peak separations fall below 59 mV, this is indicative of an electrochemical surface bound process occurring.<sup>33</sup> This is further supported by the data in Figure S1, in which anodic peak currents obtained at varying scan rates are reported. Proportionality between peak current and scan rate indicated that ferrocene was bound to the surface. If it were under diffusion control however, proportionality to the square root of scan rate would be predicted, which was not observed. Interestingly, the higher concentration of CD found on SAMs fabricated from 1 mM solutions resulted in a lower peak separation, which is indicative of faster electron transfer kinetics vs the surface modified at lower

concentrations. Based on the CV data obtained for all different SAM conditions (*i.e.* varying concentration and incubation time), the surface coverages ( $\Gamma$ ), as displayed in Figure 4, were determined according to the Laviron-derived equation for surface-confined electroactive species (Equation 1).<sup>34</sup>

$$\Gamma = \frac{I_p 4RT}{n^2 F^2 A \nu} \quad (1)$$

Where  $I_p$  represents the peak current,  $R$  the gas constant,  $T$  the temperature,  $n$  the number of electrons in the Fe redox reaction ( $n=1$ ),  $F$  the Faraday Constant,  $A$  the electroactive area ( $A = 0.785 \text{ cm}^2$ ) and  $\nu$  the scan rate ( $\nu = 0.3 \text{ V/s}$ ).

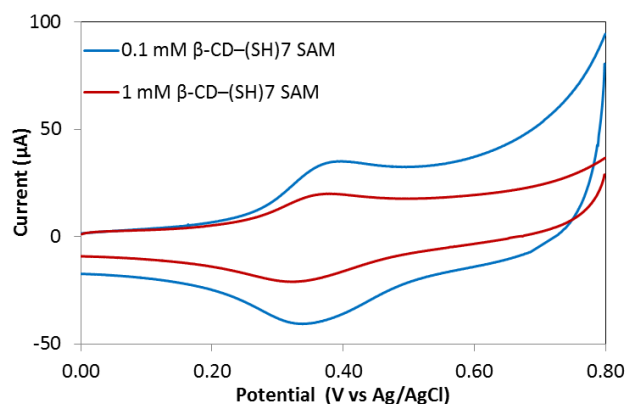


Figure 3. Typical cyclic voltammograms obtained in 0.2 M sodium sulfate solutions at a scan rate of 0.3 V/s of 0.1 mM SAMs and 1 mM SAMs, both formed for 24 hours. All the surfaces were exposed to a FCA solution for 4 hours.

For reference purposes, the maximum theoretical ferrocene coverage (MTFC) on a CD SAM was also determined. By considering the 1:1 complexation between the CD and ferrocene in addition to the theoretical outer diameter of the  $\beta$ -CD ( $1.54 \text{ nm}$ )<sup>29</sup> the MTFC was calculated to be  $8.9 \times 10^{-11} \text{ mol cm}^{-2}$ . When comparing this theoretical value with the experimental values obtained for the 0.1 mM SAMs, it is evident that ferrocene interacts directly with the gold surface in addi-

tion to the CD modified SAM, thus increasing the overall current associated with the ferrocene electrochemistry. This behavior is likely to be a consequence of entrapment of ferrocene between the CD moieties indicated by the relatively larger peak separation.

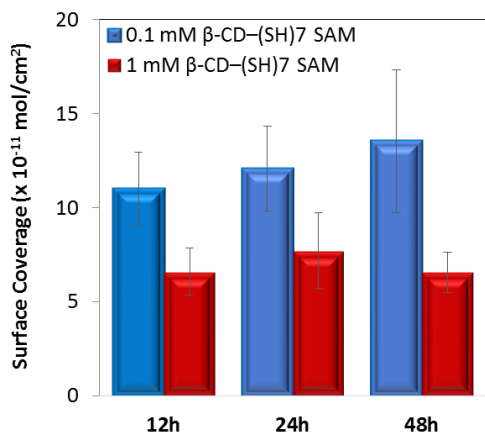


Figure 4. Ferrocene surface coverage obtained, as determined by CV analysis, for the  $\beta$ -CD-(SH)7 SAMs formed using different concentrations (0.1 mM and 1 mM) and times (12 h, 24 h and 48 h).

In cyclic voltammetry, the resultant current has two contributions. One is faradaic and another is capacitance (charging currents). The charging current was larger ( $\sim 16.2 \mu\text{A}$  at 0 V) for surface modified at the lower concentration of CD versus higher concentration ( $\sim 8.0 \mu\text{A}$  at 0 V) as a result of lower packing density of the SAM. Therefore, ions can migrate to and from the electrode surface more readily in a given time, giving rise to the observed increase in charging current.<sup>35-37</sup> The data suggest that the 0.1 mM SAMs form loosely-packed structures, wherein the spaces between CDs are large enough for the ferrocene molecules to partake in electrochemical reaction at the bare gold surface. This results in a larger electrochemically active area which is revealed by the greater currents. In contrast, the SAMs formulated from 1 mM solutions display ferrocene surface coverages below but closer to the MTFC on a CD SAM, thus suggesting that ferrocene primarily complexes with the closely packed CD monolayer on the surface.

Each of the analytical techniques provided information regarding the packing and orientation of the CDs on the different

$\beta$ -CD-(SH)7 SAMs. Based on the hydrophilicity, the thickness of the monolayers and the number of bound thiols as determined by XPS, the CDs in both 0.1 mM and 1 mM SAMs are predominantly oriented with the macrocycle's cavity exposed at the interface. This orientation allows for maximum contact between the hydrophobic amino acid in the protein and the apolar CD cavity. However, the concentration used for SAM formation has an effect on the packing of the CDs on the gold surface. As highlighted by the reduced amount of S/Au ratio and increased permeability as demonstrated by cyclic voltammetry, SAMs fabricated with 0.1 mM solutions exhibited a less well packed structure of CDs. Thus, based on these findings, 1 mM SAMs incubated for 24 hours were chosen as the optimum surface to promote high protein binding while simultaneously limiting non-specific binding through the presence of a packed CD surface.

Protein immobilization studies using surface plasmon resonance (SPR) spectroscopy demonstrated the assembly of the  $\alpha$ -chymotrypsin, insulin, RNase A and cytochrome C proteins on the  $\beta$ -CD-(SH)7 SAMs (Figure 5). Whilst the binding properties are somewhat dependent on the protein characteristics, all proteins bound to the CD functionalized surfaces with high nanomolar/low micromolar dissociation constants ( $K_D$ ) as shown in Table 1.

Table 1. SPR-derived  $K_D$  values for the interaction between the 1 mM  $\beta$ -CD-(SH)7 SAM and the different proteins and protein binding capacity of the surface.

Protein	$K_D$ ( $\mu\text{M}$ )	Protein binding capacity ( $\text{ng}/\text{nm}^2$ )
$\alpha$ -chymotrypsin	$0.63 \pm 0.20$	1.93
Insulin	$0.82 \pm 0.35$	0.55
RNase A	$1.12 \pm 1.04$	1.48
Cytochrome C	$3.21 \pm 0.90$	1.78

$$R_{\text{eq}} = \left( \frac{C_p}{C_p + K_D} \right) R_{\text{max}} \quad (2)$$

The  $K_D$ s were calculated based on the SPR responses at equilibrium ( $R_{\text{eq}}$ ), which were plotted against the concentra-

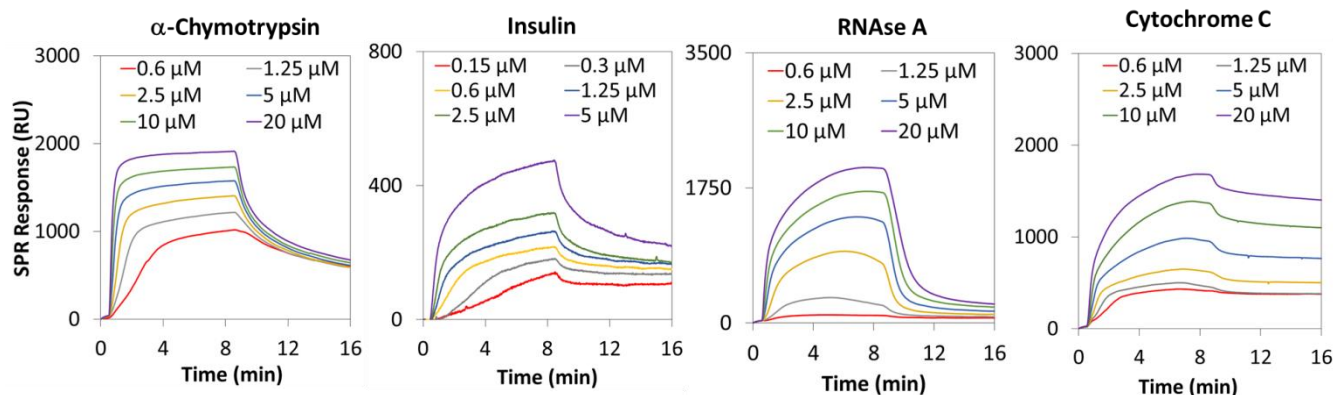


Figure 5. SPR sensorgram traces performed with 1 mM 24 h  $\beta$ -CD-(SH)7 SAMs and different concentrations of  $\alpha$ -chymotrypsin, insulin, RNase A and cytochrome C.

tion of injected protein ( $C_p$ ) and fitted to a 1:1 steady-state affinity model. The model utilizes a nonlinear least-squares regression method to fit data to the Langmuir adsorption isotherm (Equation 2). The equation not only allows calculating  $K_D$  but also the surface saturation response,  $R_{max}$ .

The interaction between a single CD moiety and a hydrophobic binding site on the surface of a protein is weak, with  $K_{DS}$  typically in the mM range.<sup>23</sup> Thus, the recognition events occurring between the studied proteins and the CD-(SH)7 SAMs are characterized by multiple interactions acting simultaneously, affording more than 1000-fold increase in binding affinity. Chymotrypsin, insulin and RNase A display, within the error, comparable binding affinities, while cytochrome C binds to the surface with the lowest affinity ( $K_b=3.21 \mu\text{M}$ ). These findings are consistent with the surface percentage of hydrophobic amino acids on the four proteins as determined by PyMol (chymotrypsin, 19.0%; insulin 25.5%; RNase A 19.9%; cytochrome C 15.6%). The lowest affinity obtained for cytochrome C might be correlated with the lowest surface percentage of hydrophobic amino acids in this protein. However, other parameters, which can possibly include type, orientation and accessibility of hydrophobic amino acids at the protein surface, might play a role in the overall affinity obtained.

It is, though, important to note that while the formation of the CD inclusion complexes with hydrophobic amino acids on the protein's surface has a large contribution to the SPR response and, thus binding affinity, other non-covalent interactions, such as hydrogen bonding, between the protein and the CD glucose units also play a role. This effect has been demonstrated by creating a gold-tethered SAM of glucose moieties as a control and conducting SPR spectroscopy analysis upon exposure to the different proteins. CDs are comprised of glucose monomers and thus the glucose-terminated SAM embodies similar functionalities at the interface but where the apolar cavity is not present. As shown in Figure S1, all the proteins bind to the glucose-terminated SAMs, however the SPR response is significantly lower than that seen for the  $\beta$ -CD-(SH)7 SAMs.

As anticipated, the SPR response for the  $\beta$ -CD-(SH)7 SAMs and the protein analytes are dependent on the protein molecular weight ( $M_w$ ), with the lower  $M_w$  insulin displaying the lowest response. By considering  $R_{max}$  and that 100 response units (RUs) is equivalent to  $0.1 \text{ ng/mm}^2$ ,<sup>2,38,39</sup> the maximum protein binding capacity of the  $\beta$ -CD-(SH)7 SAM can be determined. As illustrated in Table 1, the  $\beta$ -CD-(SH)7 SAM possess a high capacity for protein binding with protein densities ranging from *ca.*  $0.5 \text{ ng/nm}^2$  to *ca.*  $2 \text{ ng/nm}^2$ , which are largely dependent on the protein  $M_w$ .

Moreover, as a result of the noncovalent nature of the CD-amino acid interactions, the captured proteins could be easily released from the surface by exposure to sodium dodecyl sulfate (SDS) (Figure 6). This "use-regenerate" cycle could be repeated multiple times with minimal loss of binding capability.

Time of flight secondary ion mass spectrometry (ToF-SIMS) studies were also conducted to understand whether the  $\beta$ -CD-(SH)7 SAM can induce a particular protein orientation on the surface. Static ToF-SIMS, which allows analysing the outermost 2 nm of the surface with high chemical specificity, has been previously used<sup>40-42</sup> to investigate the identity and orientation of surface-tethered proteins and a similar strategy was employed herein.

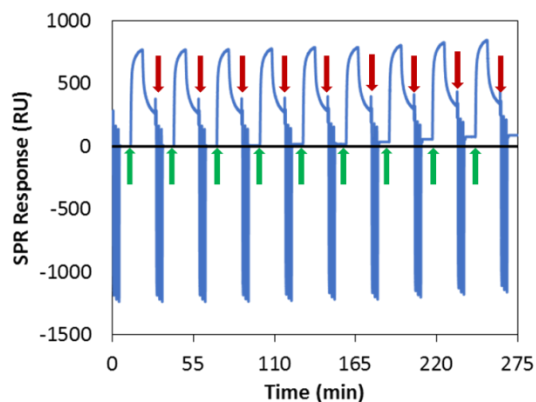


Figure 6. SPR responses from 9 SPR cycles that were performed using  $2.5 \mu\text{M}$   $\alpha$ -chymotrypsin PBS solution on  $1 \text{ mM}$   $24 \text{ h}$   $\beta$ -CD-(SH)7 SAMs. The green arrows indicate the protein injection time and the red arrows show the beginning of the SDS regeneration step.

Cytochrome C, which contains a rigid porphyrin ring coordinated with a single iron atom, was used as the model protein to be studied by ToF-SIMS. The SAM fabricated by incubation of a  $1 \text{ mM}$  solution of  $\beta$ -CD-(SH)7 for 24 hours was exposed to a  $1 \text{ mM}$  cytochrome C solution in PBS for 2 hours, and subsequently rinsed in UHQ water. As a control, glucose-terminated SAMs were also exposed to similar cytochrome C incubation conditions. For further control bare gold and both types of SAMs without protein exposure have been analyzed (Figure S3).

Since two types of chips: cytochrome C on glucose-terminated SAMs and cytochrome C on  $\beta$ -CD-(SH)7 SAMs have the same overall chemical composition, statistical analysis is required to identify more subtle differences between samples. Variance patterns within the ToF-SIMS secondary ion peak intensities between cytochrome C on the  $\beta$ -CD-(SH)7 SAM and the glucose-terminated SAM were analyzed by multivariate analysis in order to understand if differences in protein orientation could be inferred.<sup>43,44</sup> The ToF-SIMS relative ion intensities for cytochrome C on a  $\beta$ -CD-(SH)7 SAM and cytochrome C on a glucose-terminated SAM are significantly different from each other (Figure S4). To obtain more detailed information about protein orientation, the distinctive iron-porphyrin fragment ( $\text{C}_{34}\text{H}_{33}\text{N}_4\text{O}_4\text{Fe}^+$ ,  $617.27 \text{ m/z}$ ) of cytochrome C was examined on both protein surfaces (Figure 7a).

The ion intensity of the iron-porphyrin ( $M_w = 618 \text{ Da}$ ) in both protein surfaces are remarkably different, with a lower ion intensity for the protein immobilized on the  $\beta$ -CD-(SH)7 functionalized SAM (Figure 7). As static ToF-SIMS collects chemical information from outermost 2 nm of the surface, these results indicate that on the  $\beta$ -CD-(SH)7 functionalized SAM, the cytochrome C prevalently adopts an orientation with the porphyrin ring pointing down towards the CD molecules and less available to the primary ion beam. On the glucose-terminated SAMs, the cytochrome C porphyrin ring is more exposed to the analytical probe, resulting in a greater peak intensity. To support this hypothesis, other distinct fragments of cytochrome C were considered. The ion chosen was  $145.10 \text{ m/z}$  ( $\text{C}_5\text{H}_9\text{N}_2\text{O}_3^+$ ), indicating the Gly1-Asp2 residue, which is located at the opposite side of the cytochrome C compared to the porphyrin ring. In this case, the trend in both protein surfaces is inverted, with a higher ion intensity for the protein

immobilized on the  $\beta$ -CD-(SH)7 SAM (Figure 7b). These effects are not believed to have resulted from different protein amounts immobilized on the surfaces as although then overall peak intensities would change, normalized intensity patterns would be unaffected, as shown by Park *et al.*<sup>43</sup>. This suggests that the interaction between the hydrophobic amino acids and the CD cavities in the  $\beta$ -CD-(SH)7 functionalized SAM are capable of inducing a defined protein orientation, which is different from that induced by other non-covalent interactions. There is the likelihood that this orientation is dictated by the position of the hydrophobic amino acids on the protein surface that can form stronger interactions with the surface-tethered CDs.

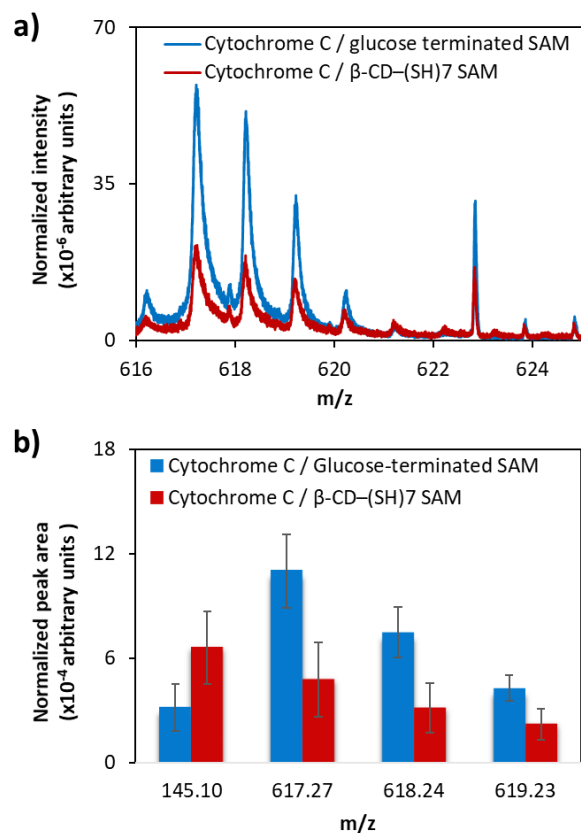


Figure 7. a) An overlay of the ToF-SIMS peak intensities and b) pattern peak intensities for cytochrome C iron-porphyrin fragment, wherein the protein has been immobilized on either a glucose-terminated SAM or a  $\beta$ -CD-(SH)7 SAM. Peak areas are normalized to total ion count. Each bar shows peak area average of 12 measurements and standard deviation within sample set. Significance of differences between samples was confirmed by T-test.

## Conclusions

An efficient strategy for the construction of protein assemblies through a bottom-up approach which harnesses non-covalent interfaces has been described. Furthermore, the facile approach does not require any protein modification process. Even though the exact binding properties and orientation of the protein on the CD-modified surface will always be dictated by the protein structure (i.e. unique distribution of hydrophobic groups on the surface of proteins), the strategy can be applied to a wide range of proteins and provide high nanomolar to low micromolar dissociation constants. Additionally, while

the proteins are tightly bound to the surface due to multivalency, the CD-terminated surfaces can be readily addressed to regenerate the free CDs. Moreover, the surface can be reused for protein immobilization post-regeneration. Considering all these attributes of broad applicability and versatility of CD-terminated surfaces to immobilize proteins, this work opens up new, efficient routes towards the realization of advanced bio-analytic and biosensor technologies.

## ASSOCIATED CONTENT

**Supporting Information.** Materials and Methods. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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### Author Contributions

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