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# Influence of the Interaction Strength between Supramolecular Complexes on the Topography of Neutral Polymer Multilayer Films

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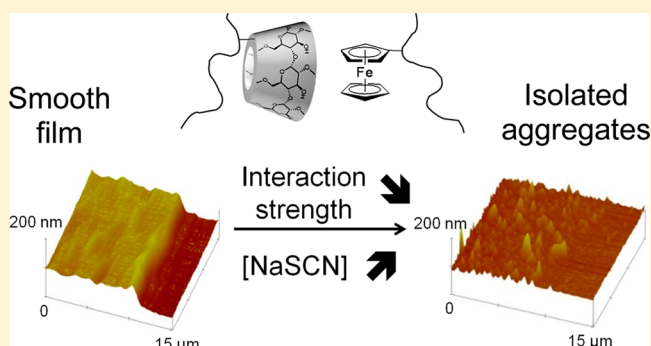
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## Supporting Information

**ABSTRACT:** Step-by-step polymer film buildup processes lead to polymer coatings, e.g., polyelectrolyte multilayers, of various structures ranging from continuous smooth films to droplet like discontinuous coatings. Yet, the origin of these different behaviors depending upon the system is not yet known. This study is a first attempt to rationalize the evolution of the coating structure as a function of the strength of the interactions between the polymers constituting the film. We investigated the influence of the strength of noncovalent host–guest interactions between cyclodextrin (CD) and pyrene (Py), ferrocene (Fc) or adamantane (Ad) on the structure of neutral poly(*N*-hydroxypropylmethacrylamide) (PHPMA) multilayers films formed in a step-by-step manner. In solution, the strength of the inclusion complex (measured by  $\log K$  where  $K$  is the complex association constant) is increasing in the order  $\text{Py}/\beta\text{-CD} < \text{Fc}/\beta\text{-CD} < \text{Ad}/\beta\text{-CD}$  and can be further varied in the presence of different sodium salts at different ionic strengths. Depending upon this strength, the buildup process is limited to the formation of isolated aggregates for PHPMA-CD/PHPMA-Py, leading to smooth continuous films for PHPMA-CD/PHPMA-Fc and to droplet-like films, not entirely covering the substrate, for PHPMA-CD/PHPMA-Ad. To study the influence of the strength of the host–guest interactions on the film topography, PHPMA-CD/PHPMA-Fc films were built in the presence of different sodium salts at different ionic strengths. For low host–guest interactions, only isolated aggregates are formed on the substrate. As the strength of the host–guest interactions increases (increase of  $\log K$ ), the formed films go through a droplet-like structure, before becoming continuous but rough for stronger interactions. When the interaction strength is further increased, the roughness of the films decreases, leading to a smooth continuous film before becoming rough again at still higher interaction strength. Smooth continuous multilayers seem thus to be obtained for an optimal range of the interaction strength.



## INTRODUCTION

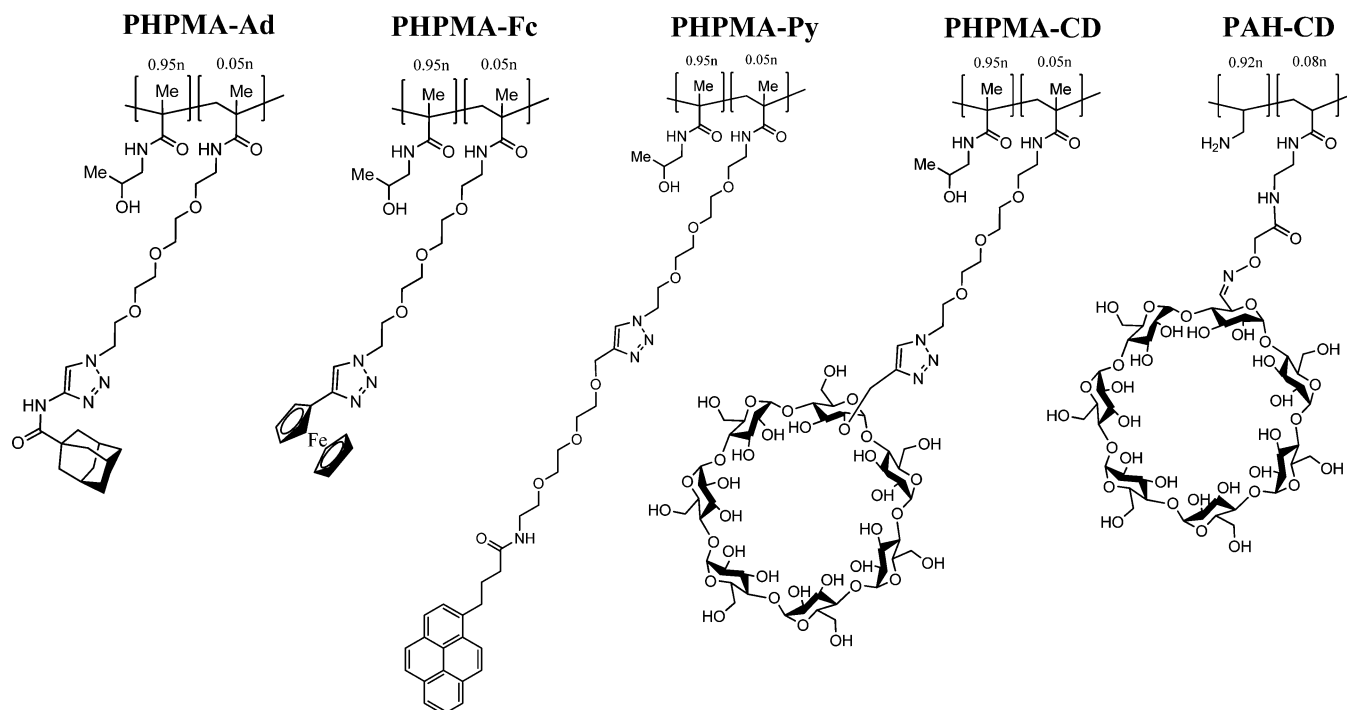
Step-by-step film buildup processes constitute a very general way to functionalize surfaces.<sup>1,2</sup> Multilayer films are obtained by alternating adsorption of two types of interacting species on inorganic<sup>3–5</sup> or organic<sup>6,7</sup> substrates. Deposition of polymer

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Scheme 1. Chemical Structures of PHPMA-Ad, PHPMA-Fc, PHPMA-Py, PHPMA-CD, and PAH-CD



layers through electrostatic interactions<sup>8–10</sup> is the most prominent example of these types of coatings. Such multilayer films have become extremely popular because it is often believed that step-by-step deposition of polymers leads to a thin coating covering the whole substrate with a nicely stratified structure. Such stratification has been proven for several polyelectrolyte systems, rendering them very appealing for multifunctionalization.<sup>1,11–13</sup> Yet, full coverage of the substrate is not systematically obtained. Grain- or island-like topographies are often encountered during the multilayer buildup. In 2001, Picart et al. showed that the step-by-step deposition of hyaluronic acid (HA) and poly(L-lysine) (PLL) leads to the formation of islands on silica surfaces during the initial deposition steps.<sup>14,15</sup> As the number of deposition steps increases, these islands merge and form a gel-like film whose thickness increases exponentially with the number of deposition steps. Similarly poly(vinyl sulfate)/poly(allylamine hydrochloride) (PAH) gives rise to a grainy structure during the initial deposition steps.<sup>4</sup> Other systems of polymers lead to the formation of islands or grains beyond the early deposition steps. Cini et al. found that poly(sodium phosphate)/PAH leads to an island-like deposit on silicon wafers up to at least 150 deposition steps.<sup>16</sup> Despite its crucial importance for the entire subsequent multilayer film, only few studies have tried to explain the topography observed (grain- or island-like, droplets, gel-like, etc.) during the initial deposition steps. A recent study by Guillaume-Gentil et al. has focused on this aspect comparing a weakly interacting polyelectrolyte couple (HA/PLL) and a strongly interacting polyelectrolyte couple (polystyrene sulfonate (PSS)/PAH). After a given number of deposition steps, a HA/PLL multilayer can be either a continuous film or an assembly of droplets, depending on the nature of the substrate.<sup>17</sup> In the case of the PSS/PAH system, a continuous film is obtained independently of the type of the substrate. Comparing their results to recent simulations, Guillaume-Gentil et al. proposed an explanation based on the dielectric

constant of the substrate, which directly affects the strength of the interactions between the polyelectrolytes and the substrate. According to these simulations, the strength of the monomer–monomer interactions strongly influences the topography of the polyelectrolyte multilayer films.<sup>18–20</sup> Complexation processes between polycations and polyanions can induce a dewetting of the polyelectrolytes from the substrate. For weakly interacting monomers, this dewetting induced the formation of small complexes on the surface leading to droplet or island morphologies.

Multilayer films based on polymers interacting through supramolecular interactions such as hydrogen bonding,<sup>21–23</sup> ligand–receptor<sup>24</sup> or host–guest interactions<sup>25–28</sup> are also widely reported. In particular, the use of  $\beta$ -cyclodextrins ( $\beta$ -CD) is an attractive approach because this natural 7-membered sugar ring is cheap, readily available, and is an effective host for several hydrophobic guests. Suzuki et al. showed that electrostatic repulsions between PAH chains can be overcome through host–guest interactions between ferrocene (Fc) moieties grafted on the PAH chains and  $\beta$ -CD homobifunctionalized molecules.<sup>25</sup> Huskens and co-workers developed layer-by-layer (LbL) assemblies of adamantane (Ad)-functionalized dendrimers and  $\beta$ -CD-functionalized gold nanoparticles, obtaining well-defined thin multilayer films.<sup>29</sup> Using nanoimprint lithography<sup>30,31</sup> or transfer printing,<sup>32,33</sup> they created 3D supramolecular assemblies with controlled size and geometry. Van der Heyden et al. showed that chitosan chains onto which were grafted  $\beta$ -CD and Ad moieties, respectively, can be alternately deposited on a gold surface covered with Ad-functionalized self-assembly monolayers. Yet, the film buildup seems to stop after the third deposition step.<sup>26</sup> Fc- and  $\beta$ -CD-modified PAH were reported to form multilayer films and capsules at least until 8 bilayers.<sup>27</sup> The alternated incubation of the substrate in PAH-CD aqueous and PAH-Fc methanol solution allowed decreasing both the charge repulsion and the polymer dissolution during the film formation. More recently,

Dubacheva et al. succeeded in an unlimited host–guest multilayer film buildup in aqueous solution by using alternately neutral poly(*N*-hydroxypropylmethacrylamide) (PHPMA) chains respectively bearing  $\beta$ -CD (PHPMA-CD) and Fc (PHPMA-Fc) moieties.<sup>28</sup> Yet, none of these studies focused on the topography of the formed film. It is well-known that the association constant of  $\beta$ -CD/Ad,  $\beta$ -CD/Fc and  $\beta$ -CD/pyrene (Py) complexes is decreasing in this order.<sup>34</sup> Several studies pointed out that the stability and the association constant of  $\beta$ -CD/Ad<sup>35</sup> and  $\beta$ -CD/hydrophobic molecules<sup>36–39</sup> complexes can be modulated by cosolutes such as salts. Thus, host–guest interactions offer a good opportunity to investigate the influence of the interaction strength between the two polymers on the structure of the resulting step-by-step film.

Here, we investigate the buildup and the topography of PHPMA films formed step-by-step by the deposition of PHPMA bearing  $\beta$ -CD hosts and hydrophobic guests (G), i.e., Ad, Fc and Py. Depending upon the strength of the inclusion complex, the buildup process is limited for PHPMA-CD/PHPMA-Py, leading to the formation of a continuous film for PHPMA-CD/PHPMA-Fc and of isolated islands, which do not cover the whole substrate, for PHPMA-CD/PHPMA-Ad. The  $\beta$ -CD/Fc complex association constant can be tuned by the presence of different types and concentrations of salts, as can be shown by isothermal titration microcalorimetry. Thus, the influence of the inclusion complex strength was studied by building up PHPMA-CD/PHPMA-Fc films in the presence of different sodium salts at different ionic strengths.

## MATERIALS AND METHODS

**Chemicals.** Poly(allylamine hydrochloride) bearing 8%  $\beta$ -CD moieties (named PAH-CD) and poly(*N*-hydroxypropylmethacrylamide) derivatives bearing 5% guest moieties ( $G = \text{Ad, Fc or Py}$ ) and host moieties  $\beta$ -CD were synthesized as described in the Supporting Information (SI). The PHPMA derivatives were named PHPMA-G (i.e., PHPMA-Fc, PHPMA-Py or PHPMA-Ad) and PHPMA-CD, respectively. These functionalized neutral polymers are represented in Scheme 1. All these PHPMA polymers were prepared by chemical modification of a PHPMA- $\text{N}_3$  precursor having the following molecular features:  $M_n = 34040$  g/mol and  $M_w = 92640$  g/mol. The length of the ethylene glycol spacer between the pyrene group and the PHPMA backbone is not similar to that of the other PHPMA derivatives. It was necessary to use a functionalized Py with six ethylene glycol groups to get a PHPMA-Py sufficiently soluble in water for our study. The preparation and characterization of PHPMA-G and PHPMA-CD are given in the SI.

Sodium chloride ( $\text{NaCl}$ ), sodium sulfate ( $\text{Na}_2\text{SO}_4$ ), sodium perchlorate ( $\text{NaClO}_4$ ), sodium thiocyanate ( $\text{NaSCN}$ ),  $\beta$ -CD, and 1,1'-ferrocenedimethanol ( $\text{Fc}(\text{MeOH})_2$ ) were purchased from Sigma-Aldrich. Sodium fluoride ( $\text{NaF}$ ) was purchased from Merck, and adamantane-1-carboxylic acid ( $\text{Ad-COOH}$ ) was obtained from Alfa Aesar. All chemicals were used as received. Buffer solutions were prepared at 10 mM HEPES with ultra pure water (Milli-Q Plus system, Millipore, Billerica, MA) with added sodium salts at different ionic strengths. All polymer solutions were prepared at a concentration of 0.1 mg/mL with HEPES buffer solution containing a sodium salt at different ionic strengths. HEPES buffer and polymer solutions were adjusted at pH 7.4 using a 0.1 M NaOH solution.

**Film Buildup.** The PHPMA-G/PHPMA-CD film buildup was performed using polymer solutions and HEPES buffer solution containing the same added sodium salt at the same fixed ionic strength. In order to obtain an anchoring layer, the substrate was first put in contact with a PAH-CD solution for 5 min followed by a rinsing step of 5 min with HEPES buffer solution. PAH-CD is electrostatically adsorbed on the substrate and so allows immobilizing  $\beta$ -CD moieties. The PHPMA-G/PHPMA-CD LbL assembly was performed by

alternately exposing the substrate for 5 min to PHPMA-G and PHPMA-CD solutions. An intermediate rinsing step after each polymer deposition was done with HEPES buffer solution for 5 min. The temperature was fixed at 22 °C during all the experiments.

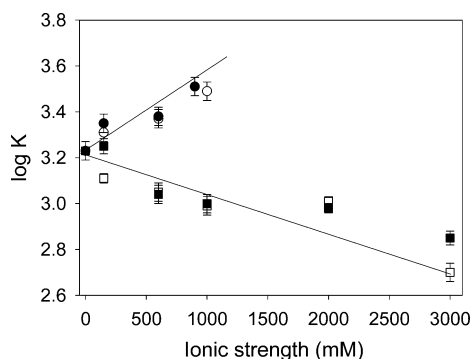
**Quartz Crystal Microbalance with Monitored Dissipation (QCM-D) Measurements.** All experiments were performed using quartz crystal microbalance with dissipation monitoring (Q-Sense E4 system equipped with four flow chambers from Q-Sense, Sweden). Monitoring the resonance behavior of piezoelectric oscillators allows measuring the mass adsorbed at the surface of the oscillator in real time. Gold-coated crystals with 5 MHz ( $\nu = 1$ ) resonance frequency were used. The third, fifth, and seventh harmonics ( $\nu = 3, 5$  and 7) at 15, 25, and 35 MHz were also recorded to have a better stability and a higher sensitivity than obtained with the fundamental resonance frequency.

**Atomic Force Microscopy (AFM) Measurements.** AFM images were obtained in contact mode in liquid conditions with Nanoscope IV from Veeco (Santa Barbara, CA). The samples were kept under liquid (Milli-Q water) for all experiments. The images were carried out with silicon nitride cantilevers, spring constant 0.03 N/m (model MSCT-AUHW, Veeco, CA). Deflection and height mode images were scanned simultaneously at a fixed scan rate (2 Hz) with a resolution of  $512 \times 512$  pixels. The film buildup was made on a gold-coated crystal, and the film thickness was measured by using the “scratch” method. The scratches were achieved with a plastic cone tip and were always imaged perpendicular to the fast scan axis. Profilometric section analysis of a scratched film allowed us to precisely determine the thickness of the film over the scanned area. The film thickness, i.e., the  $z$  distance between the bare substrate and the surface of the film, is considered as the minimal film thickness covering all the substrate. The mean thickness of the scratched film was determined by measuring the thickness at least on three areas. The film roughness is the root mean square (RMS) given by the AFM software on  $15 \times 15 \mu\text{m}^2$  images. Data evaluations were performed with the NanoScope software version 5.31r1 (Digital Instruments, Veeco).

**Isothermal Titration Microcalorimetry (ITC) Measurements.** The “heat of reaction” between  $\beta$ -CD and  $\text{Fc}(\text{MeOH})_2$  (respectively  $\text{Ad-COOH}$ ) was measured at 25 °C by means of isothermal titration microcalorimetry (Nano Isothermal Titration Calorimeter, TA Instruments, New Castle, USA). This microcalorimeter is fitted with two thermostated cells with a volume of 1 mL: a reference cell containing Milli-Q water, and a sample cell containing the  $\text{Fc}(\text{MeOH})_2$  (or  $\text{Ad-COOH}$ ) solution to be titrated. All solutions were degassed 30 min under vacuum before use. The titration cell was regulated at 25 °C. A typical experiment involved 20 sequential 12.29  $\mu\text{L}$  injections of the  $\beta$ -CD solution (8.8 mM) into the ITC cell containing the  $\text{Fc}(\text{MeOH})_2$  (or  $\text{Ad-COOH}$ ) solution (1.2 mM). The concentrations were chosen so that the “mixing molar ratio” between  $\beta$ -CD and the hydrophobic molecules is equal to 2 at the end of the titration. The injections were made by an automated microsyringe (250  $\mu\text{L}$ ) ended with a propeller, which stirred the solution at 250 rpm. Two consecutive injections were separated by a resting period of 400 s to allow the microcalorimeter trace to come back to a baseline corresponding to the absence of any heat flow between the sample cell, in which host–guest interactions took place, and the reference cell, filled with Milli-Q water. For each sodium salt and each salt concentration, the dilution heats were determined by injecting  $\beta$ -CD,  $\text{Fc}(\text{MeOH})_2$ , and  $\text{Ad-COOH}$  solutions into HEPES buffer solution. The binding enthalpy was obtained by subtracting the integrated heat measured during the injection of  $\beta$ -CD in  $\text{Fc}(\text{MeOH})_2$  (respectively  $\text{Ad-COOH}$ ) from the dilution heat of  $\beta$ -CD in HEPES, measured at the same salt concentration and pH value. Finally, the resulting “heat of reaction” was divided by the provided number of  $\beta$ -CD moles. The first injection was not taken into account in the data analysis. The peak integration was performed with Origin 7.5 after establishment of a linear baseline over 50 points, taken in the time intervals between consecutive  $\beta$ -CD injections, over the points where the heat flow reached a steady value. In order to obtain the association constant of the complex, the measured heat data were processed assuming the formation of a complex 1:1.<sup>40</sup> Two independent



experiments were done, and the reproducibility was on the order of 10%.



**Figure 1.** Association constant ( $\log K$ ) of  $\beta$ -CD/Fc(MeOH)<sub>2</sub>, measured by ITC, as a function of the ionic strength for NaF (black circles), Na<sub>2</sub>SO<sub>4</sub> (white circles), NaSCN (black squares), and NaClO<sub>4</sub> (white squares). ITC experiments were performed in 10 mM HEPES buffer at pH 7.4 and 25 °C. The data represents the mean and the standard deviation of at least two independent experiments. The lines are there to help guide the eye.

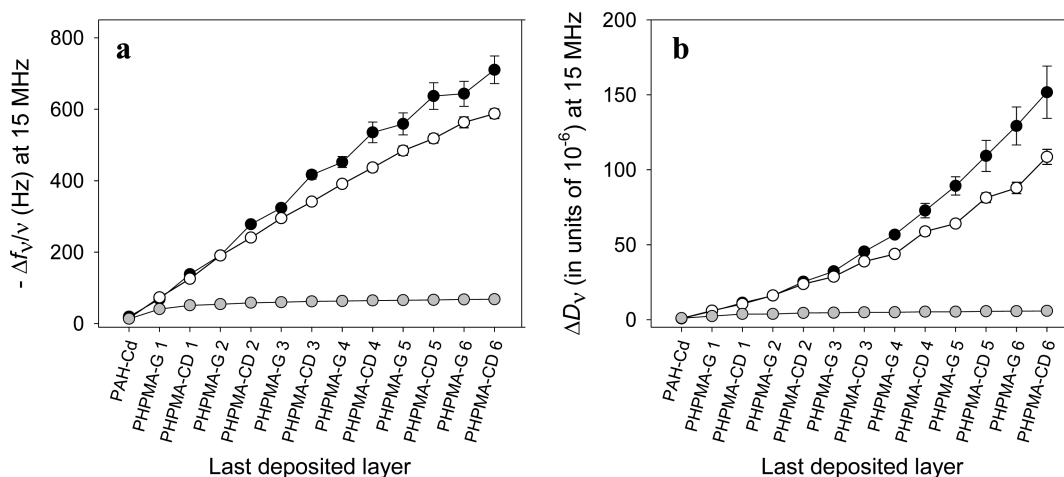
## RESULTS AND DISCUSSION

### Association Constant of the $\beta$ -CD/Guest Complexes.

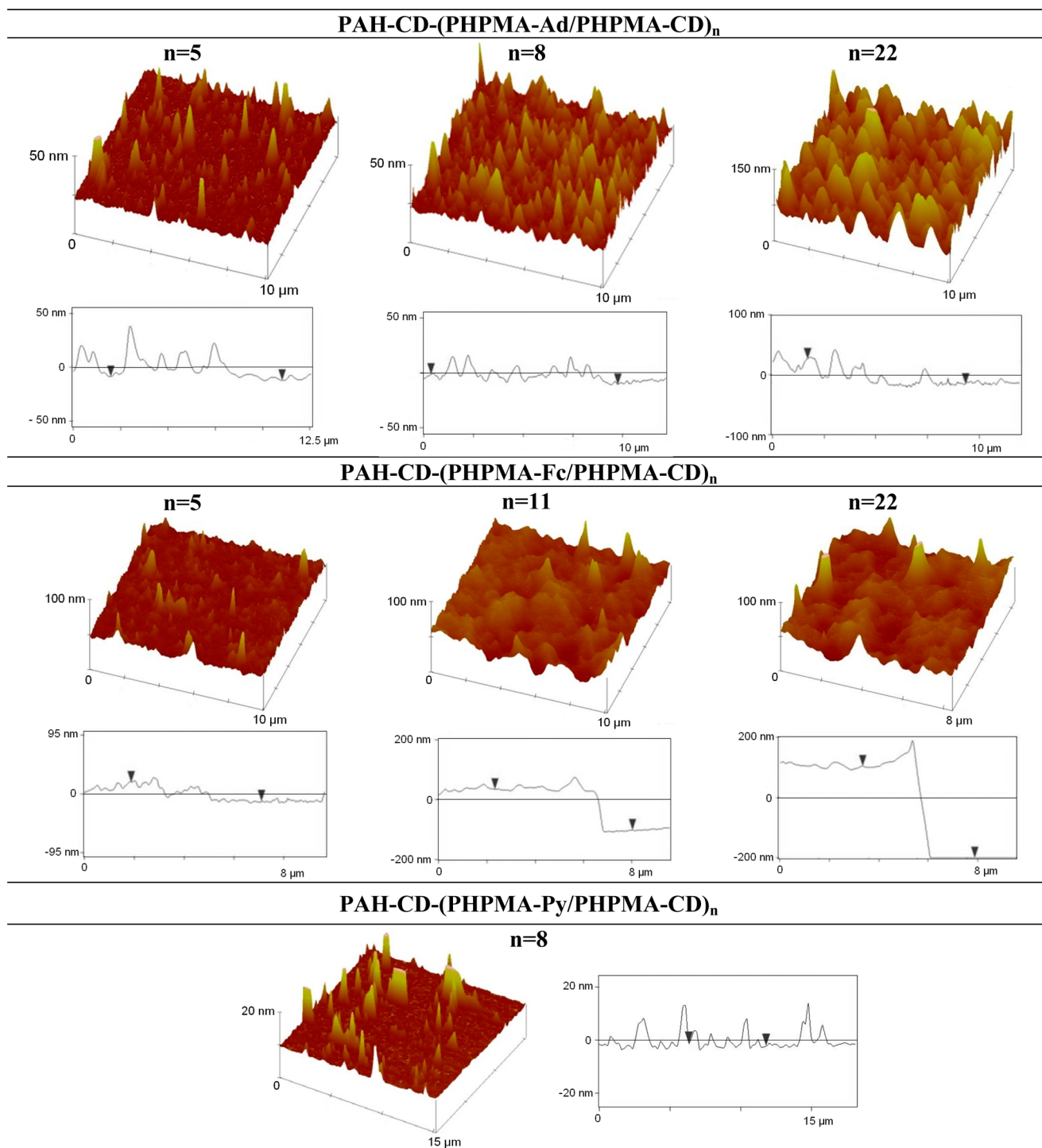
Before the study of the buildup of PHPMA-CD and PHPMA-G multilayer films with different guests, we first determined the association constant of the  $\beta$ -CD/G complexes in solution. It is well-known that the stability constant of  $\beta$ -CD/Ad,  $\beta$ -CD/Fc, and  $\beta$ -CD/Py complexes decreases in this order. Ad-COOH and Fc(MeOH)<sub>2</sub> were used for ITC experiments. ITC allows gaining access to the association constants ( $K$ ) of  $\beta$ -CD/Fc(MeOH)<sub>2</sub> and  $\beta$ -CD/Ad-COOH, which can be compared to that of the literature.<sup>34</sup> In 10 mM HEPES buffer solutions at pH 7.4 and 25 °C (without additional salt), the constants were estimated at  $\log K = 4.27 \pm 0.02$  for  $\beta$ -CD/Ad-COOH and  $\log K = 3.23 \pm 0.04$  for  $\beta$ -CD/Fc(MeOH)<sub>2</sub>. These values are in good agreement with the values found in the literature for similar systems:  $\log K = 4.26$  for  $\beta$ -CD/Ad-COOH in pure

water at pH 8.5 and 25 °C,<sup>41</sup> and  $\log K = 3.33$  for  $\beta$ -CD/ferrocene carboxylate in 50 mM phosphate buffer at pH = 8.6 with 0.1 M NaCl at 25 °C.<sup>34</sup> Due to the poor water solubility of Py, the stability constant of  $\beta$ -CD/Py complex was not determined in our conditions. This constant is estimated in the literature at 2.69 in water at 25 °C.<sup>42</sup> To confirm the ability of salt to tune the strength of the host/guest interaction, we determined the association constant of the  $\beta$ -CD/Fc(MeOH)<sub>2</sub> complex in different types of salts varying the ionic strength. We used four different sodium salts Na<sub>2</sub>SO<sub>4</sub>, NaF, NaClO<sub>4</sub>, and NaSCN. The results are gathered in Figure 1, where  $\log K$  is presented as a function of the ionic strength. The ionic strength was increased up to 3 M for NaSCN and NaClO<sub>4</sub> but limited to 900 mM for NaF and 1 M for Na<sub>2</sub>SO<sub>4</sub> due to the limited solubility of the respective salts (Table S-1 in the SI) or of  $\beta$ -CD in the salt. We observed two different trends according to the salt present in solution. On one hand, by increasing the ionic strength of NaClO<sub>4</sub> and NaSCN concentration from 0 up to 3 M, the association constant of the  $\beta$ -CD/Fc(MeOH)<sub>2</sub> complex decreases from 3.23 down to 2.85 and 2.70, respectively. The influence of the perchlorate (ClO<sub>4</sub><sup>−</sup>) and the thiocyanate (SCN<sup>−</sup>) anions on the stability constant of  $\beta$ -CD/Fc(MeOH)<sub>2</sub> measured in solution can be rationalized as follows: both ClO<sub>4</sub><sup>−43–45</sup> and SCN<sup>−44,46</sup> anions form 1:1 complexes with  $\beta$ -CD (with an association constant of  $\log K = 1.42^{44}$  and  $0.99^{44}$ , respectively) and compete with the formation of the  $\beta$ -CD/Fc(MeOH)<sub>2</sub> complex. Indeed, it has been proposed that the inside of the CD cavity is composed of positively polarized carbon atoms leading to the inclusion of hydrophobic anionic guests<sup>47</sup> such as the ClO<sub>4</sub><sup>−</sup> and SCN<sup>−</sup> anions. A similar decrease of the association constant of  $\beta$ -CD/nabumetone,<sup>36</sup>  $\beta$ -CD/phenylbenzothiazole derivatives,<sup>38</sup> and  $\beta$ -CD/3-hydroxy-2-naphthoic acid<sup>43</sup> complexes has been reported in the presence of ClO<sub>4</sub><sup>−</sup> anions.

On the other hand, the increase of the Na<sub>2</sub>SO<sub>4</sub> and NaF ionic strength up to, respectively, 1 and 0.9 M induces an increase of  $\log K$  of the  $\beta$ -CD/Fc(MeOH)<sub>2</sub> complex value up to about 3.5 (Figure 1). Similarly, Harries and co-workers obtained a linear increase of the  $\beta$ -CD/Ad association constant when increasing the K<sub>2</sub>SO<sub>4</sub> and KF concentration.<sup>35</sup> Sulfate



**Figure 2.** Typical evolution of the normalized frequency shift  $-\Delta f_v/\nu$  (a) and of the energy dissipation  $\Delta D_v$  (b), measured at 15 MHz ( $\nu = 3$ ) by QCM-D, as a function of the last deposited layer for PHPMA-CD/PHPMA-Ad (black circles), PHPMA-CD/PHPMA-Fc (white circles), and PHPMA-CD/PHPMA-Py (gray circles). Polymer solutions were prepared at 0.1 mg/mL in 10 mM HEPES - 150 mM NaCl buffer at pH 7.4. The data represents the mean and the standard deviation of three independent experiments. The values of the standard deviation are smaller or similar size compared to the reported symbol.



**Figure 3.** Morphologies and profilometric sections, obtained by AFM in contact mode in the liquid state, of PAH-CD-(PHPMA-Ad/PHPMA-CD)<sub>n</sub>, of PAH-CD-(PHPMA-Fc/PHPMA-CD)<sub>n</sub> at different numbers of deposited bilayers (*n*), and of PAH-CD-(PHPMA-Py/PHPMA-CD)<sub>8</sub>. Profilometric sections are obtained on scratched films. Polymer solutions were prepared at 0.1 mg/mL in 10 mM HEPES/150 mM NaCl buffer at pH 7.4.

(SO<sub>4</sub><sup>2-</sup>) and fluoride (F<sup>-</sup>), strongly hydrated anions, do not bind to β-CD.<sup>43,45</sup> The increase of the association constant is correlated with the water molecules released from both the β-CD cavity and Fc(MeOH)<sub>2</sub> upon the formation of the complex. The presence of SO<sub>4</sub><sup>2-</sup> and F<sup>-</sup> disturbs the internal structure of water around the β-CD and its guest, rendering the releasing step of water easier and thus displacing the equilibrium toward the complex formation.<sup>48</sup> By changing the nature and the

concentration of the anion in solution, the association constant between β-CD and Fc(MeOH)<sub>2</sub> can be modulated. After checking this effect in the bulk, we investigated the influence of this interaction strength on the buildup of neutral polymer multilayers based on cyclodextrin/guest interactions.

**Neutral Polymer Multilayer Buildup Based on β-CD/Guest Interactions.** Multilayer assemblies were prepared by alternating depositions of PHPMA-G (i.e., PHPMA-Ad,

**Table 1.** Evolution of the Thicknesses and the Roughnesses of PHPMA-CD/PHPMA-Ad and PHPMA-CD/PHPMA-Fc Based Films (AFM Data Extracted from Figure 3) Allowing the Denomination of the Topography of the Films

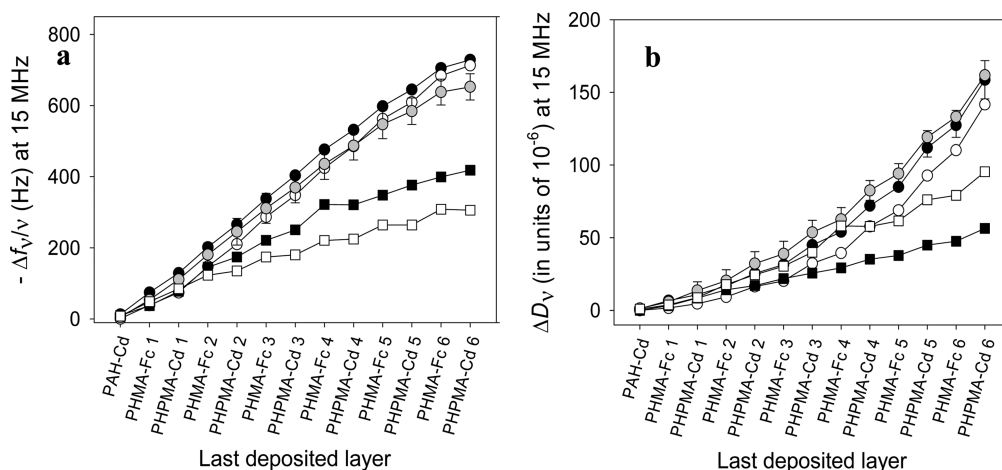
	<i>n</i>	<i>d</i> <sub>AFM</sub> (nm)	RMS roughness (nm)	film topography
PAH-CD-(PHPMA-Ad/ PHPMA-CD) <sub><i>n</i></sub>	5	11	6	rough
	8	13	11	continuous
	22	20	26	droplet-like
PAH-CD-(PHPMA-Fc/ PHPMA-CD) <sub><i>n</i></sub>	5	31	6	smooth
	11	136	13	continuous
	22	337	21	

PHPMA-Fc, or PHPMA-Py) and PHPMA-CD. The PHPMA backbone being a neutral polymer, the cohesion of the resulting assemblies is only due to the supramolecular host–guest interactions that occur between cyclodextrins and adamantane, ferrocene, or pyrene moieties. Functionalized PHPMA polymers were prepared in two steps: the synthesis of PHPMA-N<sub>3</sub> (*M*<sub>w</sub> = 34 040 g/mol, *M*<sub>n</sub> = 92 640 g/mol, polydispersity index (PDI) = 2.72) with a grafting ratio of 5% and its functionalization by the guest through click-chemistry. Using the same batch of PHPMA-N<sub>3</sub> allows us to obtain functionalized PHPMA with different guests and β-CD having roughly the same length of polymer chains and the same grafting ratio (5%). The chemical structure of all polymers can be found in Scheme 1, and their preparation is described in the SI.

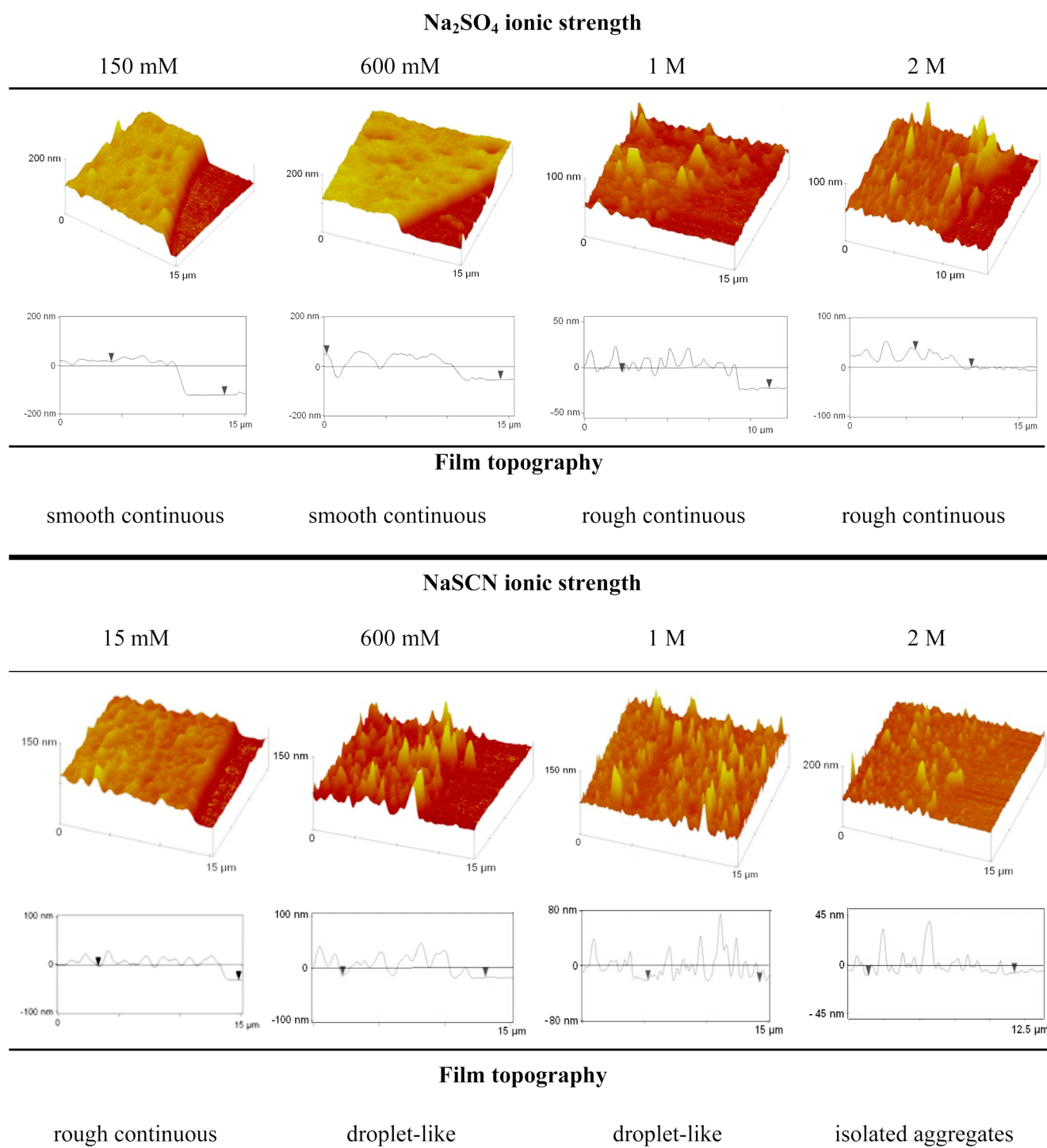
The buildup of the multilayer films was followed by QCM-D (Figure 2). An anchoring layer of polycationic PAH-CD was first adsorbed. A strong and almost linear evolution of the normalized frequency shift ( $-\Delta f_v/\nu$ ), measured at 15 MHz, is observed for both PHPMA-CD/PHPMA-Ad and PHPMA-CD/PHPMA-Fc systems, reaching roughly 750 and 600 Hz at the end of the sixth bilayer, respectively (Figure 2a). In contrast, the PHPMA-CD/PHPMA-Py buildup is limited to one bilayer (40 Hz). The association constant between β-CD and pyrene ring ( $\log K = 2.69$  in water at 25 °C<sup>42</sup>) is probably too weak.

After 6 bilayers, the dissipation values reach  $150 \times 10^{-6}$  and  $110 \times 10^{-6}$  for PHPMA-CD/PHPMA-Ad and PHPMA-CD/PHPMA-Fc, respectively. These values are relatively large, suggesting that the assemblies are highly hydrated, as was shown in our former study. Indeed, PHPMA-CD/PHPMA-Fc films were found to have 95% content in water.<sup>28</sup> To show that the buildup is obtained only due to host–guest interactions, we performed two sets of control experiments: one where only PHPMA-CD was used and the second one where at least one of the host or guest functionalized PHPMA was replaced by PHPMA-N<sub>3</sub>. We thus followed by QCM the buildup of the following systems: PHPMA-CD/PHPMA-CD, PHPMA-CD/PHPMA-N<sub>3</sub>, PHPMA-N<sub>3</sub>/PHPMA-Fc and PHPMA-N<sub>3</sub>/PHPMA-N<sub>3</sub> (Figure S-1 in SI). We notice that no buildup of films is obtained in all this cases. The host/guest couple is thus necessary showing that PHPMA-CD/PHPMA-Fc and PHPMA-CD/PHPMA-Ad buildups are only obtained due to the host/guest interactions.

In order to determine the topography of PAH-CD-(PHPMA-G/PHPMA-CD)<sub>*n*</sub> based films (with G = Ad, Fc and Py), AFM images were performed in contact mode and in liquid state at different numbers of bilayers ranging from 5 up to 22 bilayers. The films were scratched to determine their thicknesses by a profilometric section (Figure 3). The film thickness, i.e. the *z* distance between the bare substrate and the surface of the film, is considered as the minimal film thickness completely covering the substrate. Although similar QCM-D signal evolutions were observed along the buildup of PHPMA-CD/PHPMA-Ad and PHPMA-CD/PHPMA-Fc films, the topography evolutions are not similar. On one hand, the PHPMA-CD/PHPMA-Ad system leads to extremely rough films mainly constituted of droplets that grow in number and in size with the number of deposition steps (Figure 3, first row). On the other hand, the PHPMA-CD/PHPMA-Fc system forms fairly flat and smooth films whose thickness increases with the number of deposition steps (Figure 3, second row). Concerning the PHPMA-CD/PHPMA-Py system, only few and isolated aggregates are visible on the substrate showing almost no adsorbed polymer (Figure 3, third row). After 22 deposited bilayers, PHPMA-CD/PHPMA-Fc films reach a thickness of about 300 nm with a roughness of 21 nm when a



**Figure 4.** Evolution of (a) the normalized frequency shift  $-\Delta f_v/\nu$  and (b) the dissipation factor  $\Delta D_v$ , measured at 15 MHz ( $\nu = 3$ ) by QCM, as a function of the sequence of adsorbed polymer layers during the buildup of PAH-CD-(PHPMA-Fc/PHPMA-CD)<sub>7</sub> in 10 mM HEPES at pH 7.4 without added salt (gray circles) and with NaF (black circles), Na<sub>2</sub>SO<sub>4</sub> (white circles), NaSCN (black squares), and NaClO<sub>4</sub> (white squares) at the ionic strength of 150 mM.



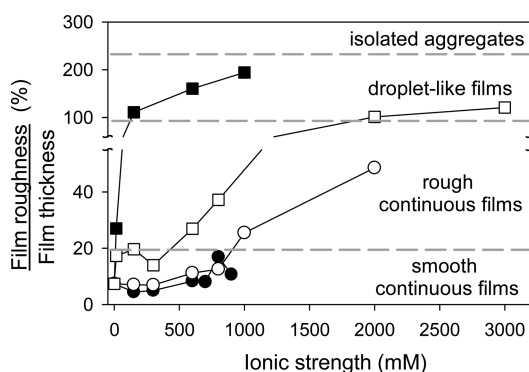
**Figure 5.** 3D AFM images and profilometric sections, obtained by AFM in contact mode in liquid environment, of a scratched PAH-CD-(PHPMA-Fc/PHPMA-CD)<sub>7</sub> film built in a 10 mM HEPES buffer solution at pH 7.4 in the presence of Na<sub>2</sub>SO<sub>4</sub> and NaSCN at different ionic strengths. Polymer solutions were prepared at 0.1 mg/mL.

PHPMA-CD/PHPMA-Ad film has a thickness of 20 nm, with the largest extension of its islands of about 50 nm (Figure 3 and Table 1). The roughness of PHPMA-CD/PHPMA-Ad films is similar to or higher than the thickness (Table 1), as it is shown in the profilometric section (Figure 3, first row).

All the results reported above were obtained with a polymer concentration of 0.10 mg/mL. To favor the buildup of the

films, we investigated the effect of the polymer concentration; two polymer concentrations were further used: 0.05 and 0.50 mg/mL. Figure S-2 in the SI shows the topography and the profilometric section of the PAH-CD-(PHPMA-Fc/PHPMA-CD)<sub>n</sub> and PAH-CD-(PHPMA-Ad/PHPMA-CD)<sub>n</sub> films built at polymer concentrations of 0.05 and 0.10 mg/mL up to  $n = 22$  bilayers and built at a polymer concentration of 0.50 mg/mL up





**Figure 6.** Evolution of the roughness to thickness ratio of a PAH-CD-(PHPMA-Fc/PHPMA- $\beta$ -CD)<sub>7</sub> film as a function of NaF (black circles), Na<sub>2</sub>SO<sub>4</sub> (white circles), NaSCN (black squares), and NaClO<sub>4</sub> (white squares) ionic strength. The film was built in 10 mM HEPES at pH 7.4 with a polymer concentration of 0.1 mg/mL.

to  $n = 11$  bilayers. The topography of these films does not depend on the concentration of the polymer solutions used during the buildup process (Figure S-2 in the SI). However, higher polymer concentrations result in larger islands in the case of PHPMA-CD/PHPMA-Ad and thicker films in the case of PHPMA-CD/PHPMA-Fc (Table S-1 in the SI). Based on these results, three types of film topography can be defined: smooth continuous films when the film roughness is below 20% of the film thickness, rough continuous films when the film roughness is above 20% of the film thickness, and island-like films when the film thickness is below the film roughness. PHPMA-Ad, PHPMA-Fc, and PHPMA-Py have the same length of polymer chains and the same grafting ratio in hydrophobic groups along the polymer backbone (5%). The differences in film topography and buildup can only be attributed to the host–guest interaction strength between Ad, Fc, or Py groups and  $\beta$ -CD. The association constant of Ad and  $\beta$ -CD being one of the highest reported in the literature, we will qualify the strength of the interaction between PHPMA-Ad and PHPMA-CD as “strong”. According to the AFM images, whatever the polymer concentration, strong interactions between neutral polymers through host–guest association lead to a rough continuous film or an island-like film, whereas, regardless of the number of bilayers or of the concentration of the polymers used, intermediate strength interactions (involving Fc groups) lead to a smooth continuous film. Too weak interactions (involving Py derivatives) no longer allow film construction but rather the formation of isolated aggregates.

**Influence of the Host–Guest Affinity on Neutral Polymer Multilayer Buildup.** First, using ITC, we showed that the interaction of  $\beta$ -CD with Fc(MeOH)<sub>2</sub> can be modulated by changing the nature and the concentration of anions. Then, we observed that PHPMA-CD/PHPMA-Ad multilayers are droplet-like, whereas PHPMA-CD/PHPMA-Fc ones are rather smooth. Based on this observation, our idea was

to modulate the affinity of the  $\beta$ -CD interaction with Fc by varying the nature of the sodium salt anions and their concentration in the PHPMA solutions used to build PAH-CD-(PHPMA-Fc/PHPMA-CD)<sub>n</sub> films. The polymer concentration was fixed at 0.1 mg/mL. First, the film buildup was performed in 10 mM HEPES buffer at pH 7.4 and at an ionic strength of 150 mM using different sodium salts, namely NaF, Na<sub>2</sub>SO<sub>4</sub>, NaSCN, and NaClO<sub>4</sub> (Figure 4). It appears that the normalized frequency shifts measured by QCM-D corresponding to F<sup>−</sup> and SO<sub>4</sub><sup>2−</sup> are significantly higher than those corresponding to films constructed in the presence of SCN<sup>−</sup> and ClO<sub>4</sub><sup>−</sup> as the anions of the supporting electrolyte. Without added salt, the buildup of the film is close to the one obtained in the presence of F<sup>−</sup> and SO<sub>4</sub><sup>2−</sup>. The QCM measurements confirm that the buildup of PHPMA-CD/PHPMA-Fc films can be modulated by the choice of the supporting electrolyte anion.

Furthermore, the topography and the thickness of PAH-CD-(PHPMA-Fc/PHPMA-CD)<sub>7</sub> films were determined by AFM when built in the different types of salts at different ionic strengths. In the absence of salt, the buildup process leads to a smooth continuous film with a thickness of 116 nm and a roughness of 9 nm (Figure S-3 in the SI).

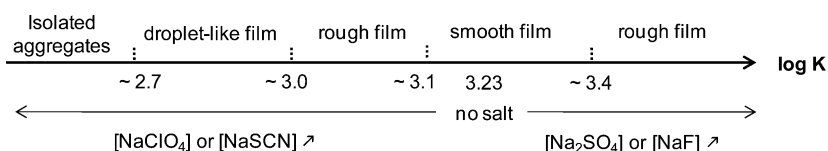
By increasing the concentration of Na<sub>2</sub>SO<sub>4</sub> (or NaF), corresponding to an increase of the  $\beta$ -CD/Fc interaction strength (Figure 1), smooth and continuous films are formed until 800 mM (Figure 5 and Figure S-4 in the SI). A decrease of the film thickness is observed keeping almost the same roughness (Table S-3 in the SI). By increasing the concentration of NaSCN (or NaClO<sub>4</sub>), corresponding to a decrease in the  $\beta$ -CD/Fc interaction strength (Figure 1), the film buildup becomes less and less effective, leading to a smaller film thickness (Figure 5 and Figure S-4 in the SI) and a higher roughness (Table S-3 in the SI). The film topography changes from a smooth continuous film toward a droplet-like film.

The roughness to thickness ratio of PAH-CD-(PHPMA-Fc/PHPMA-CD)<sub>7</sub> films as a function of the ionic strength for the four studied salts is represented in Figure 6. By strengthening the host–guest interaction, the topography of the PHPMA-CD/PHPMA-Fc films changes from smooth and continuous to droplet-like. Whereas for the weakest host–guest interactions ( $\log K = 2.7$  at 3 M NaSCN), no PHPMA-CD/PHPMA-Fc film buildup is obtained. This was also found for PHPMA-CD/PHPMA-Py buildup where  $\log K = 2.69$  for  $\beta$ -CD/Py in water. This is represented schematically in Scheme 2. These results seem to confirm that the strength of the interaction between the two polymers constituting the film strongly influences the film buildup. Intermediate interaction strength seems to be required to form continuous films whereas if too low, the interaction strength limits the film buildup.

## CONCLUSION

We investigated the influence of the strength of noncovalent host–guest interactions between CD and Py, Fc, and Ad on the

**Scheme 2.** Schematic Representation of the Influence of the Ionic Strength on the Topography of PHPMA-CD/PHPMA-Fc Multilayers, via the Tuning of the  $\beta$ -CD/Fc Association Constant



structure of PHPMA-CD/PHPMA-Py, PHPMA-CD/PHPMA-Fc, and PHPMA-CD/PHPMA-Ad films formed in a step-by-step manner. In solution, the strength of the inclusion complex increases in the order  $\text{Py}/\beta\text{-CD} < \text{Fc}/\beta\text{-CD} < \text{Ad}/\beta\text{-CD}$ . Depending upon this strength, the buildup process is limited to the formation of isolated aggregates for PHPMA-CD/PHPMA-Py, leads to smooth continuous films for PHPMA-CD/PHPMA-Fc, and to droplet-like films, which do not entirely cover the substrate, for PHPMA-CD/PHPMA-Ad. To study in a systematic way the influence of the strength of the host–guest interactions on the topography of the films, PHPMA-CD/PHPMA-Fc films were built in the presence of different sodium salts at different ionic strengths. For weak host–guest interactions, only isolated aggregates are formed on the substrate. As the interaction strength of the host–guest interactions increases (increase of  $\log K$ ), the formed films go through a droplet-like structure, before becoming continuous but rough for stronger interactions. When the interaction is further increased, the films become smooth before becoming rough again at still higher interaction strength. Further studies are needed to verify whether these conclusions apply to other step-by-step constructed polymer films. Moreover, the interactions between the polymers and the substrate should be taken into account to get a general picture of the initial steps of LbL polymer film constructions. Nevertheless, this study constitutes, to our knowledge, the first attempt to rationalize the comprehension of the initial steps of a polymer multilayer buildup as a function of the polymer interaction strength. This study should also trigger new studies on polyelectrolyte multilayers.

## ■ ASSOCIATED CONTENT

### ■ Supporting Information

Synthesis of the different PHPMA polymers; table of the solubility of the salts used in water; table of thicknesses and roughnesses of PHPMA-Ad/PHPMA- $\beta$ -CD and PHPMA-Fc/PHPMA- $\beta$ -CD films built at different polymer concentrations; Buildup of PHPMA-CD/PHPMA-N3, PHPMA-N3/PHPMA-Fc, PHPMA-N3/PHPMA-N3 and PHPMA-CD/PHPMA-CD films; AFM images and profilometric sections of scratched PAH- $\beta$ -CD-(PHPMA-Fc/PHPMA- $\beta$ -CD)<sub>7</sub> films built at different polymer concentrations in 150 mM NaCl; AFM images and profilometric sections of scratched PAH- $\beta$ -CD-(PHPMA-Fc/PHPMA- $\beta$ -CD)<sub>7</sub> films without added salt and in the presence of NaF or NaClO<sub>4</sub>; table of PAH- $\beta$ -CD-(PHPMA-Fc/PHPMA- $\beta$ -CD)<sub>7</sub> film thickness and roughness measured by AFM without added salt and in the presence of added salt at different ionic strengths. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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### Notes

The authors declare no competing financial interest.

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