Effect of the supramolecular interactions on the nanostructure of halloysite/biopolymer hybrids: a comprehensive study by SANS, fluorescence correlation spectroscopy and electric birefringence†

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The structural properties of halloysite/biopolymer aqueous mixtures were firstly investigated by means of combining different techniques, including small-angle neutron scattering (SANS), electric birefringence (EBR) and fluorescence correlation spectroscopy (FCS). Among the biopolymers, non-ionic hydroxypropylcellulose and polyelectrolytes (anionic alginate and cationic chitosan) were selected. On this basis, the specific supramolecular interactions were correlated to the structural behavior of the halloysite/biopolymer mixtures. SANS data were analyzed in order to investigate the influence of the biopolymer adsorption on the halloysite gyration radius. In addition, a morphological description of the biopolymer-coated halloysite nanotubes (HNTs) was obtained by the simulation of SANS curves. EBR experiments evidenced that the orientation dynamics of the nanotubes in the electric field is influenced by the specific interactions with the polymers. Namely, both variations of the polymer charge and/or wrapping mechanisms strongly affected the HNT alignment process and, consequently, the rotational mobility of the nanotubes. FCS measurements with fluorescently labeled biopolymers allowed us to study the aqueous dynamic behavior of ionic biopolymers after their adsorption onto the HNT surfaces. The combination of EBR and FCS results revealed that the adsorption process reduces the mobility in water of both components. These effects are strongly enhanced by HNT/polyelectrolyte electrostatic interactions and wrapping processes occurring in the halloysite/chitosan mixture. The attained findings can be useful for designing halloysite/polymer hybrids with controlled structural properties.

Introduction

In the last decades, supramolecular hybrids based on inorganic nanoparticles and organic macromolecules have attracted great interest as a consequence of their potential applications in several technological fields, including packaging,1,2 catalysis,3–8 pharmaceutics,8–14 and remediation.15–19 As evidenced in a recent review,20 the adsorption of sustainable polymers can confer functional properties to the nanoparticles depending on their peculiar nanoarchitecture. It was proved that the modification of carbon21 and boron22 nanotubes with poly(N-isopropylacrylamide) (PNIPAAm) generates hybrid molecular nanorods with assembly/disassembly behaviors in response to temperature variations. Recently, thermo-sensitive nanocarriers were obtained by...
supramolecular functionalization of halloysite with oppositely charged PNIPAAm polymers, such as amine-terminatated PNIPAAm and PNIPAAm/methacrylic acid copolymer. Electrostatic interactions were successfully exploited to fabricate pH-sensitive drug delivery hybrids based on Laponite nanodisks and poly(ethylene glycol)-poly(lactic acid) diblock copolymer. Efficient nanocarriers for drugs were prepared by the adsorption of alginate onto calcium carbonate and silica nanoparticles.

Among inorganic nanoparticles, halloysite clay nanotubes (HNTs) are suitable for the preparation of smart nanohybrids because of their peculiar surface properties in terms of chemical composition and electrical charge. In particular, the chemical composition of the HNT internal surface is based on alumina, while the external shell is formed by silica. As a consequence of the different acid–base equilibria of alumina and silica groups, the HNT inner and external surfaces are positively and negatively charged, respectively, within an extended pH interval and, in addition, the charge conditions can be tuned by pH. Rheological measurements evidenced that HNT aqueous suspensions can form a lyotropic liquid crystalline phase depending on the pH conditions. The HNT dispersions exhibited stronger shear-thinning behavior by the addition of microcrystalline cellulose. The HNT surfaces can be selectively modified by ionic molecules through electrostatic interactions. The adsorption of cationic alkyltrimethylbromides onto the halloysite outer lumen was controlled by pH conditions, which influence the characterization of polymer/HNT hybrids, with the aim of exploring the influence of the specific supramolecular interactions on the nanoarchitecture of the HNT/biopolymer hybrids. These investigations were conducted by using a comprehensive approach based on combining different methods (small-angle neutron scattering, fluorescence correlation spectroscopy and electric birefringence) not employed before together for the characterization of polymer/HNT hybrids, with the aim of gaining an improved understanding of their interactions and structures and the resulting stability in aqueous solutions.

Accordingly, studies on the structure and dynamics of polymer/HNT hybrids in water could be crucial to understand the stabilization mechanism controlling the aqueous colloidal stability of the nanotubes. In this work, we investigated the structural behavior of aqueous mixtures based on halloysite and biopolymers with different charges, including non-ionic hydroxypropylcellulose and biopolyelectrolytes (anionic alginate and cationic chitosan), in order to explore the influence of the specific supramolecular interactions on the nanoarchitecture of the HNT/biopolymer hybrids. These investigations were conducted by using a comprehensive approach based on combining different methods (small-angle neutron scattering, fluorescence correlation spectroscopy and electric birefringence) not employed before together for the characterization of polymer/HNT hybrids, with the aim of gaining an improved understanding of their interactions and structures and the resulting stability in aqueous solutions.

Materials and methods

Chemicals

Hydroxypropyl cellulose (HPC; average molecular weight = 80 kg mol⁻¹), sodium alginate (average molecular weight = 90 kg mol⁻¹), chitosan (deacetylation degree = 75–85%, average molecular weight = 120 kg mol⁻¹), fluorescein isothiocyanate (FITC), 5-[4,6-dichlorotiazin-2-yl]amino]fluorescein (DTAF), phosphate-buffered saline (PBS), sodium bicarbonate (NaHCO₃), sodium hydroxide (NaOH) and glacial acetic are Sigma products. Halloysite (HNT, purity ≥ 99.5%) from Matauri Bay was provided by Imerys. SEM images of HNTs are presented in the ESL. All the chemicals were used without further purification. Water was of Millipore grade. D₂O was purchased from Eurisotop in 99.9% isotopic purity.
Preparation of HNT/biopolymer dispersions

HNT/biopolymer dispersions in aqueous solvents were prepared as reported elsewhere.\(^{36}\) Firstly, stable aqueous solutions of each polymer were obtained by magnetically stirring at 25 °C for 3 h. Chitosan was solubilized in acidic solvent (pH = 4.5) because of its low solubility under neutral conditions. The pH of the aqueous solvent was adjusted by adding glacial acetic acid. The final concentration of acetic acid was set at 5 g dm\(^{-3}\) (0.083 mol dm\(^{-3}\)). Then, HNT/biopolymer dispersions with various compositions were obtained by direct addition of appropriate amounts of halloysite powder into the polymer solutions. The HNT/biopolymer mixtures were homogenized by ultrasonication for 10 min and subsequent magnetic stirring at 25 °C for 24 h.

Small-angle neutron scattering (SANS)

SANS measurements were carried out at Institut Laue-Langevin (ILL), Grenoble (France), on the instrument D11.\(^{42}\) The experiments were conducted at four different configurations with sample-to-detector (and collimation in parenthesis) distances of 1.5 m (8 m), 8 m (8 m), and 34 m (34 m), using a wavelength \(\lambda\) of 6.0 Å (fwhm of 10%) and 39 m (40.5 m) with \(\lambda = 13.0\ \)Å and fwhm of 10%. Based on these experimental conditions, the investigated scattering vector \((q = 4\pi\sin(\theta/2))/\lambda, \theta\) being the scattering angle) ranged between 0.007 and 4.20 nm\(^{-1}\). The two-dimensional patterns were corrected for the detector efficiency using the scattering of a 1 mm H\(_2\)O sample and for the dark current signal; the contribution from the empty cell was subtracted; and finally, the patterns were radially averaged as all dimensional patterns were corrected for the detector efficiency. The experiments were performed in full contrast conditions (D\(_2\)O as solvent) on pristine halloysites,\(^{45}\) the HNT concentration was set at 80 g dm\(^{-3}\). Then, HNT/biopolymer dispersions with various compositions were obtained by direct addition of appropriate amounts of halloysite powder into the polymer solutions. The HNT/biopolymer mixtures were homogenized by ultrasonication for 10 min and subsequent magnetic stirring at 25 °C for 24 h.

Scanning electron microscopy (SEM)

The surface morphology of halloysite nanotubes was investigated using a microscope, ESEM FEI QUANTA 200F. Before each experiment, the surface of the sample was coated with gold in argon by means of an Edwards Sputter Coater S150A to avoid charging under the electron beam. The measurements were carried out in high vacuum mode (<6 x 10\(^{-14}\) Pa) for simultaneous secondary electrons; the energy of the beam was 25 kV and the working distance was 10 mm.

Results and discussion

SANS data analysis: structural characterization of HNT/biopolymer hybrids

Fig. 1 shows the scattering curves in full contrast for HNT and HNT/biopolymer dispersions with a mass ratio of biopolymer/HNT of 0.1. As observed for pure halloysite from different sources,\(^{45}\) SANS curves of the HNT/biopolymer mixtures did not evidence any oscillations, in agreement with the large polydispersity of the HNT radii. Similar observations were detected for HNT/surfactant systems.\(^{33}\)

According to the literature,\(^{50,51}\) two different Guinier regions can be considered for elongated particles, such as rigid rods. Within the low \(q\) Guinier interval, the scattered intensity of elongated objects varies as

\[
I(q) = I(0) \times \exp\left(-\frac{R_g^2 q^2}{3}\right)
\]  
(1)

where \(I(0)\) is the scattering intensity at the limit \(q \to 0\), while \(R_g\) is the gyration radius of the whole particle, which is related to the length \(L\) and the radius \(R\) of the scattered rod by the following relation

\[
R_g^2 = \frac{(L^2)}{12} + \frac{(R^2)}{2}
\]  
(2)
On this basis, we calculated that the theoretical $R_g$ of halloysite nanotubes is 294 nm by taking into account their average sizes in terms of length (1000 nm) and external radius (80 nm).45

According to eqn (1), $\ln(I(q))$ vs. $q^2$ plots describe linear trends allowing us to determine $R_g$ and $I(0)$ from their slopes and their intercepts, respectively. SANS data at small angles ($0.007 \text{ nm}^{-1} \leq q \leq 0.009 \text{ nm}^{-1}$) were analyzed by eqn (1). As displayed in Fig. 2, $\ln(I(q))$ was linearly dependent on $q^2$ for all of the investigated dispersions. Table 1 collects the $R_g$ and $I(0)$ data obtained from the Guinier analysis of SANS curves.

In addition, the Guinier analysis for elongated particles can be conducted in the intermediate $q$ range, where the dependence of the scattered intensity on the radial gyration radius is expressed as

$$q \cdot I(q) = I^*(0) \times \exp \left(-\frac{R_g^2 q^2}{2}\right)$$

(3)

where $I^*(0)$ is the prefactor and $R_g^2 = 2\gamma$ and, thus, we can state that $R_g$ determined from the intermediate $q$ region is related only to the cross section of the rod.52 Accordingly, we calculated $R_g = 53$ nm for cylinders with $R = 80$ nm, which is the average outer radius of halloysite nanotubes.45 By taking into account both the inner and the outer radii of halloysite (80 and 15 nm, respectively),15 we estimated $R_g = 57$ nm for the cross section of the hollow nanotube.

It should be noted that eqn (3) is valid for $qR_g \approx 1$, which indicates that $q$ values between 0.015 and 0.02 nm$^{-1}$ represent the proper intermediate $q$ range for scattered halloysite. As expected by eqn (3), $\ln(q \cdot I(q))$ vs. $q^2$ plots within the intermediate $q$ range (Fig. 2) were successfully fitted by linear equations that provided the corresponding $R_g$ and $I^*(0)$ values (Table 1).

From the Guinier linear analysis for elongated objects, we find that the halloysite gyration radii are hardly affected by the adsorption of both anionic alginate and cationic chitosan (Table 1).

Regarding $I(0)$, the HNT/biopolymer composites showed slightly larger values with respect to that of pure HNT (Table 1) due to the polymer adsorption onto the halloysite surfaces.

As reported for silica/polyelectrolytes systems,53 the actual amount of biopolymer adsorbed onto the nanotubes can be calculated from $I(0)$ values, which are related to the density ($\lambda N$) and the volume ($V$) of the nanoparticles by the following equation

$$I(0) = \lambda N V^2 \Delta SLD^2 S(0)$$

(4)

where $S(0)$ is the structure factor at $q \rightarrow 0$, whereas $\Delta SLD$ is the difference between the scattering length density of the nanoparticle ($SLD_{\text{nanoparticle}}$) and that of the solvent ($SLD_{\text{solvent}}$). $\lambda N$ is the number density of HNT and was determined by the concentration and HNT geometry considering the corresponding polydispersity. Based on eqn (4), we calculated $I(0) = 27.6 \times 10^5$ cm$^{-1}$ for the pure HNT solution, which is close to the experimental result (Table 1). Details on the calculation of the number density ($\lambda N$) and the volume ($V$) of HNT are presented in the ESL†.

As concerns the biopolymer coated nanotubes, the volume ($V_{\text{HNT/Biop}}$) and the scattering length density ($SLD_{\text{HNT/Biop}}$) of the hybrid nanoparticles can be expressed as

$$V_{\text{HNT/Biop}} = V_{\text{HNT}} + \chi(\phi_{\text{Biop}}) N_{\text{HNT}}$$

(5)

$$SLD_{\text{HNT/Biop}} = (SLD_{\text{HNT}} V_{\text{HNT}} + SLD_{\text{Biop}} \chi(\phi_{\text{Biop}}) N_{\text{HNT}})/(V_{\text{HNT}} + \chi(\phi_{\text{Biop}}) N_{\text{HNT}})$$

(6)
where φ_Biop and V_HNT are the total volume fraction of biopolymer and the volume of uncoated HNT, while χ corresponds to the fraction of biopolymer adsorbed onto halloysite with respect to all the biopolymer in the dispersion.

Assuming S(0) = 1, the numerical solution of eqn (2)–(4) allowed us to estimate χ from the I(0) values. Based on the calculated χ values (Table 2), we calculated the biopolymer/HNT volume ratio (R_V(Biop/HNT)) of the coated nanotubes, as detailed in the ESI†.

The biopolymer surface coverage (Z) onto the HNTs was calculated by using the R_V(HNT/Biop) values and taking into account the specific surface area (SSA) of halloysite (28.3 m² g⁻¹) through the following equation

\[ Z = \rho_{Biop}(\rho_{HNT-SSA} - R_V(Biop/HNT)) = C_{HNT}/C_{Biop} \cdot \chi \cdot SSA_{HNT} \]  

(7)

As evidenced in Table 2, we observed larger Z values for the ionic biopolymers compared to that of HPC indicating that the electrostatic interactions play a crucial role in the formation of complex systems based on halloysite. For chitosan and alginate, very similar values for the surface coverage are observed, being slightly higher for cationic chitosan, which preferentially interacts with the HNT outer surface, wrapping the nanotubes. On the other hand, anionic alginate should mostly be confined within the positively charged HNT cavity. It might also be noted that the obtained Z values for the ionic biopolymers correspond to a “dry” (neglecting hydration water, which, of course, will be there at the real surface layer) polymer layer thickness of ~1.6–1.8 nm, which is a realistic value for dense polymer coverage.

Similar to our previous work on HNT/surfactant hybrids, SANS curves of HNT/biopolymer mixtures were simulated by using a hollow cylinder with a uniform SLD profile (Fig. 3).

According to this model, the scattering intensity can be expressed as

\[ I(q) = \frac{1}{N} \times P(q, R_0, \Delta R, \sigma, L, SLD_{HNT/Biop}, SLD_{s}) + I_{back} \]  

(8)

where the form factor is \( P(q, R_0, \Delta R, \sigma, L, SLD_{HNT/Biop}, SLD_{s}) \) and \( R_0, \Delta R \) and L are the geometrical parameters (internal radius, shell thickness and length, respectively), I_{back} is the incoherent scattered background (evaluated by Porod analysis applied to the larger q range), and SLD_{HNT/Biop} and SLD_{s} are the scattering length densities of the nanotube shell and solvent, respectively. Namely, we assumed that the biopolymer is included in the shell of the hollow cylinder, while the core is based on pure D₂O.

The same assumption was considered for the SANS data analysis of halloysite nanotubes modified with anionic surfactants, which are selectively adsorbed within their cavity. SLD_{HNT/Biop} was estimated on the basis of the composition of the biopolymer coated HNT determined from the Guinier analysis of SANS data in the low q range (Table 2). The SLD_{HNT/Biop} values used for the simulation of the SANS curves of the HNT/biopolymer mixtures are presented in the ESI†. Based on the Schulz–Zimm distribution, a polydispersity (σ) defined as \( (\Delta R^2)/|\Delta R|^2 \) − 1 was considered for the shell thickness. As reported for the simulation of pure HNT from the Matauri Bay deposit,\textsuperscript{43} R_i and L were fixed at 15 and 1000 nm, respectively.

**Electric birefringence (EBR): the effect of biopolymer adsorption on the HNT rotational mobility**

The analysis of EBR results allowed us to investigate the influence of the biopolymer adsorption on the rotational mobility of halloysite nanotubes. As an example, Fig. 4a displays the relaxation of the birefringence signal for the HNT/HPC aqueous mixture (mass ratio = 0.1). It can be noted that the electric field pulse was chosen to be so short that no saturation of the signal took place, but only an initial orientation was imposed.

As observed for the pure HNT\textsuperscript{25} and for HNT/surfactant hybrids,\textsuperscript{13} a transient birefringence was induced by the electric field applied in the form of a rectangular pulse. This phenomenon is related to the HNT polarizability that causes a partial alignment of the nanotubes. Once the voltage pulse is terminated, the nanotubes are free to reorient and, consequently, the magnitude of the birefringence (Δn) exponentially decreases with time following the equation

\[ \Delta n = \Delta n_0 \exp(-t/\tau) \]  

(9)

where \( \tau \) is the characteristic relaxation time of the nanotubes, while \( \Delta n_0 \) represents the maximum of the birefringence signal. As an example, the fitting analysis of Δn vs. t decay for the HNT/HPC aqueous mixture (mass ratio = 0.1) is reported in the ESI†. Fig. 5 shows the dependence of \( \tau \) on the biopolymer/HNT mass ratio (R_V(Biop/HNT)) of the mixtures. The corresponding rotational diffusion coefficient (D_rot) reported in the ESI† was calculated as D_rot = (6τ)^{-1}. **Table 1** Fitting parameters obtained by the Guinier analysis of SANS data

<table>
<thead>
<tr>
<th></th>
<th>I(0)×10⁴ cm⁻¹ (from low q region)</th>
<th>R_0/nm (from low q region)</th>
<th>I(0)×10⁴ cm⁻¹ (from intermediate q region)</th>
<th>R_0/nm (from intermediate q region)</th>
</tr>
</thead>
<tbody>
<tr>
<td>HNT</td>
<td>27.4 ± 1.0</td>
<td>223 ± 18</td>
<td>166 ± 2</td>
<td>41 ± 5</td>
</tr>
<tr>
<td>HNT/alginate</td>
<td>31.7 ± 1.1</td>
<td>222 ± 19</td>
<td>240 ± 2</td>
<td>39 ± 5</td>
</tr>
<tr>
<td>HNT/chitosan</td>
<td>30.8 ± 1.1</td>
<td>226 ± 21</td>
<td>187 ± 2</td>
<td>40 ± 4</td>
</tr>
<tr>
<td>HNT/HPC</td>
<td>30.5 ± 1.2</td>
<td>209 ± 20</td>
<td>281 ± 3</td>
<td>40 ± 5</td>
</tr>
</tbody>
</table>

**Table 2** Quantitative adsorption parameters for HNT/biopolymer hybrids at a biopolymer/HNT mass ratio of 0.1, as determined by the Guinier analysis of SANS data in the low q range

<table>
<thead>
<tr>
<th></th>
<th>X</th>
<th>V(Biop/HNT)</th>
<th>Z/mg m⁻²</th>
</tr>
</thead>
<tbody>
<tr>
<td>HNT/alginate</td>
<td>2.97×10⁻²</td>
<td>7.56×10⁻⁴</td>
<td>3.44±0.06</td>
</tr>
<tr>
<td>HNT/chitosan</td>
<td>1.78×10⁻²</td>
<td>4.21×10⁻⁴</td>
<td>3.19±0.07</td>
</tr>
<tr>
<td>HNT/HPC</td>
<td>9.61×10⁻³</td>
<td>2.38×10⁻⁴</td>
<td>1.50±0.03</td>
</tr>
</tbody>
</table>
As a general result, the presence of the biopolymer induced an enhancement indicating that the rotational mobility of the nanotubes was reduced. This effect can be attributed to the biopolymer adsorption onto the halloysite surfaces. In particular, $\tau$ vs. $R_w(Biop/HNT)$ plots exhibited increasing trends for $R_w(Biop/HNT) \leq 0.3$, while further addition of the biopolymer did not affect the relaxation time of the nanotubes. This indicates that here the attachment of the biopolymer is effectively saturated. In this regard, it should be noted that the reduction of the HNT rotational mobility cannot be simply attributed to the increase of mass upon adsorption because the amount of the biopolymer bound to the halloysite surfaces is less than 2% for all the investigated systems (Table 2). It should be noted that the presence of biopolymer causes a slow enhancement of the viscosity, which could contribute to the reduction of the HNT rotational mobility. However, this effect is negligible because the biopolymer concentration range is within a dilute regime (up to ca. 0.12 wt%). As an example, a relative $D_{rot}$ variation of only 7% is expected for the aqueous HPC/HNT mixture (with concentrations of 1 and 0.1 wt% for halloysite and polymer, respectively), because its intrinsic viscosity is 1.078, as reported in our previous paper.36 In addition, the influence of the polymer concentration on $D_{rot}$ would be more important if the solvent viscosity significantly affected the HNT rotational mobility. Accordingly, overlapping and/or bridging of the polymer-coated nanotubes could be expected for the investigated mixtures. Compared to non-ionic HPC, ionic biopolymers induced stronger effects on the HNT rotational mobility due to the electrostatic interactions. The selective
interactions of the polyelectrolytes with the charged HNT surfaces affected their influence on the rotational diffusion of the nanotubes. The effect of chitosan on the HNT rotational mobility is stronger with respect to that of alginate, which is mostly confined to the positively charged clay lumen. These results could indicate that chitosan is bound at the outside surface interconnecting different HNTs, thus substantially slowing down their rotation.

In addition, $D_{\text{rot}}$ values allowed us to determine the length of the nanotubes in the biopolymer/halloysite mixtures by using Broersma theory, which is valid for rigid rods with length/diameter ratios between 2 and 30. This approach was successfully employed for halloysite nanotubes modified with anionic surfactants. Details on the calculation of the HNT length using Broersma theory are presented in the ESI†. We estimated that the HNT length in water is 809 ± 7 nm. As a general result, the presence of biopolymers induced an increase of the halloysite length (see ESI†). In particular, the chitosan adsorption generated the strongest effect on the HNT length, which reached the largest value of 1228 ± 11 nm for the biopolymer/HNT mass ratio of 0.67. According to the EBR results, we estimated the overlap concentration for HNTs by assuming a simple cubic model where the contact distance corresponds to the length of the nanotubes. Specifically, the critical volume fraction ($\phi_c$) for the HNT overlapping was calculated as $\pi R^2/L^3$. As concerns pure HNTs, we determined that the overlap concentration is ca. 500 times higher with respect to the concentration of the investigated dispersion, confirming that the EBR results reflect the free rotation of the nanotubes. We observed that the overlap concentrations for biopolymer-coated nanotubes are ca. 2 orders higher with respect to the concentration of the investigated mixtures. Therefore, the influence of the biopolymer adsorption on the HNT rotational mobility can be explored by the EBR data.

Fluorescence correlation spectroscopy (FCS): the influence of the adsorption onto the HNT surfaces on biopolymer dynamic behavior in aqueous medium

The influence of the HNT/biopolymer electrostatic interactions on the dynamic behaviour in water of the biopolyelectrolytes was explored by FCS studies. Biopolymers were fluorescently labelled with proper probes, such as FITC and DTAF for chitosan and alginate, respectively. Fig. 6 shows the effect of the HNT addition on the correlation function of the FITC/chitosan aqueous solution.

According to the adsorption process, a significant reduction of the biopolymer mobility was detected in the presence of HNT. As evidenced by Fig. 6, the correlation functions of both FITC/chitosan and HNT/FITC/chitosan systems were successfully described by the stretched model expressed by the following equation55

$$G(t) = G(0) \cdot \left[ 1 + \left( \frac{t}{\tau} \right)^{\alpha} \right] \cdot \left( \frac{1 + \left( \frac{\tau_c}{\tau} \right)}{1 + \left( \frac{\tau_c}{\tau} \right)^{2.5}} \right)^{0.5}$$

where $T$ is the fraction of the molecules in the triplet state, $\alpha$ is the stretched parameter and $\tau_{\text{T}}$ is their relaxation time. $S$ is given by the anisotropy (ratio of the vertical and lateral extensions), while $\tau_c$ and $G(0)$ are the intercept and the decay time, respectively. As reported in the literature,56 the contribution of the triplet state cannot be neglected for DTAF molecules, while we did so for FITC.

It should be noted that the pure diffusion model was successfully employed for the FCS data analysis of HNT/surfactant hybrids containing Nile red.33 Table 3 reports the fitting parameters ($G(0)$ and $\tau_c$) for FITC/chitosan and HNT/FITC/chitosan ($\alpha$ values were 0.770 ± 0.011 and 0.904 ± 0.015, respectively).

In addition, we calculated the diffusion coefficient ($D$) as

$$D = \omega_0^2/4\tau_c$$

where $\omega_0$ is the lateral extension (590 nm) of the confocal volume.

The decay of FITC/chitosan reflects the dynamic behavior of the polymer, confirming that the fluorescent probe was successfully attached to the chitosan molecule. It is noteworthy that the diffusion coefficient of the HNT/FITC/chitosan mixture is much

![Fig. 5](image-url) Relaxation time determined by the fitting of the EBR data as a function of the biopolymer/HNT mass ratio at a HNT concentration of 0.15 g dm$^{-3}$ (SANS and FCS experiments were done at 0.1).

![Fig. 6](image-url) Normalized FCS decay curves for FITC/chitosan and HNT/FITC/chitosan mixtures in water. The concentration of the labeled chitosan (FITC/chitosan) was set at 0.05 mg dm$^{-3}$, while the overall concentration of chitosan was fixed at 5 g dm$^{-3}$. Accordingly, the labelled chitosan/HNT mass ratio was 10$^{-4}$ and the chitosan/HNT mass ratio was 0.1. Red lines represent the fitting according to the pure diffusion model (eqn (10)). The FCS curve for HNT/FITC/chitosan was arbitrarily shifted along the y-axis by adding a constant 0.2 to the experimental data.
larger (ca. 50 times) with respect to that of the pure labeled polymer, which proves the strong binding of chitosan. Table 3 compares the corresponding hydrodynamic radius ($R_h$) calculated by using the Stoke–Einstein equation. In contrast, the presence of HNT led only to a small reduction of mobility of DTAF/alginate (Fig. 7), which indicates here only a rather weak extent of binding.

The ESR† reports the fitting parameters obtained by the fitting through the triplet state model56 of FCS curves for DTAF/alginate and HNT/DTAF/alginate suspensions. The presence of HNT induced a reduction by a factor of 2 for the diffusion coefficient of the DTAF/alginate. Based on the FCS results, we can assert that dynamic behavior in water of both biopolyelectrolytes decreases as a consequence of their adsorption onto HNT surfaces. This effect is much stronger for cationic chitosan, which is wrapped onto the nanotubes. We can assume that the $D$ values for HNT/labeled polymers are given by two contributions: (1) the fast diffusion process ($D_{fast}$), which is related to the unbound polymer; (2) the slow diffusion ($D_{slow}$) due to the diffusion of the polymer adsorbed onto HNT surfaces. Based on the SANS data analysis (Table 2), only ca. 2 wt% of both chitosan and alginate are bound onto HNT. Therefore, the fast process should be predominant in the experimental diffusion coefficient. This consideration is valid for HNT/DTAF/alginate, while the much slower diffusion of HNT/FITC/chitosan could indicate that the biopolymer becomes immobilized by bridging different HNTs. In conclusion, FCS findings agree with the EBR data, which evidenced that chitosan adsorption induced the most significant reduction of the HNT rotational diffusion coefficient. Based on these results, we can argue that bridging between chitosan-coated nanotubes can be hypothesized.

**Conclusions**

We investigated the structural behavior of aqueous mixtures composed of halloysite nanotubes (HNTs) and differently charged biopolymers, such as cationic chitosan, anionic alginate and non-ionic hydroxypropylcellulose. The simulation of SANS curves by a hollow cylinder model evidenced that the biopolymer coated nanotubes possesses similar geometrical features (in terms of sizes and polydispersity) as those previously observed for pure HNT. SANS data at low and intermediate $q$ were successfully analyzed by the Guinier approach for rod-like objects. In addition, the SANS analysis showed that charged biopolymers exhibit larger adsorption efficiencies that can be attributed to the stronger electrostatic interactions. In agreement, EBR results showed that the decrease of the HNT rotational mobility is more affected and reduced for halloysite/ionic biopolymer mixtures. In this respect, chitosan caused a somewhat stronger alteration of the halloysite rotational mobility because of the HNT wrapping driven by the attractive forces between the positively charged biopolymer and the halloysite external surface, which is positively charged. The analysis of FCS curves evidenced that the adsorption process decreases the aqueous diffusion coefficients of both polyelectrolytes. The stronger effect is observed for the HNT/chitosan mixture, which showed a reduction by ca. 50% compared to that of the pure biopolymer as a consequence of the wrapping process. In conclusion, a systematic correlation between the structure of the HNT/biopolymer hybrid and the structure of the biopolymer was demonstrated by the investigation of aqueous mixtures by applying a comprehensive set of characterisation techniques (SANS, EBR, and FCS) to these composites.

**Conflicts of interest**

The authors have no conflicts of interest to declare.

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Table 3  Intercept and the decay time (with the corresponding diffusion coefficient and hydrodynamic radius) obtained from the fitting analysis of FCS curves. Pure diffusion and triplet state models were employed.

<table>
<thead>
<tr>
<th>Fitting model</th>
<th>$G(0)$</th>
<th>$\tau_0$/ms</th>
<th>$D$/µm² s⁻¹</th>
<th>$R_h$/nm</th>
</tr>
</thead>
<tbody>
<tr>
<td>FITC/chitosan</td>
<td>0.0943 ± 0.0003</td>
<td>1.62 ± 0.03</td>
<td>53.6 ± 1.0</td>
<td>4.56 ± 0.09</td>
</tr>
<tr>
<td>HNT/FITC/chitosan</td>
<td>0.0782 ± 0.0004</td>
<td>54.8 ± 1.6</td>
<td>1.59 ± 0.05</td>
<td>154 ± 4</td>
</tr>
<tr>
<td>DTAF/alginate</td>
<td>0.119 ± 0.003</td>
<td>1.80 ± 0.29</td>
<td>48 ± 8</td>
<td>5.1 ± 0.8</td>
</tr>
<tr>
<td>HNT/DTAF/alginate</td>
<td>0.078 ± 0.007</td>
<td>3.5 ± 1.0</td>
<td>24 ± 7</td>
<td>10 ± 2</td>
</tr>
</tbody>
</table>

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Fig. 7 Normalized FCS decay curves for DTAF/alginate and HNT/DTAF/alginate mixtures in water. The concentration of DTAF/alginate was set at 0.1 mg dm⁻³, while the overall concentration of alginate was fixed at 10 g dm⁻³. Accordingly, the labelled alginate/HNT mass ratio was 10⁻³ and the mass ratio between the overall alginate and HNT was 0.1. Red lines represent the fitting according to the triplet state model.56

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