

Isomerization and fluorescence depolarization of merocyanine 540 in polyacrylic acid. Effect of pH

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MS received 26 April 2002; revised 8 July 2002

Abstract. Dynamics of isomerization and fluorescence depolarization of merocyanine 540 (MC540) in an aqueous solution of polyacrylic acid (PAA) have been studied using picosecond time resolved fluorescence spectroscopy. It is observed that the dynamics of isomerization and depolarization are sensitive enough to monitor the uncoiling of PAA at high pH (> 6). At low pH (< 3), when the polymer remains in a hypercoiled form, polymer bound MC540 experiences very high microscopic friction and, hence, the isomerization and depolarization processes are very slow. At high pH (> 6) a polyanion is formed and the polymer assumes an extended configuration due to electrostatic repulsion. At high pH (> 6), the anionic probe MC540 is expelled from the polyanion to bulk water and the dynamics of isomerization and fluorescence depolarization become faster by 12 and 5 times respectively, compared to those at low pH.

Keywords. Photoisomerization; fluorescence depolarization; uncoiling.

1. Introduction

Many chemical processes involve motion of a portion of the molecule against another (e.g. in isomerization) or the molecule as a whole (e.g. in reorientation). Such processes are affected by the friction of the medium¹. While effect of friction in bulk liquid is known, there is much less information on friction in organized environments. The microscopic friction of organized assemblies towards photoisomerization and reorientational dynamics has been the subject of several recent studies. Photoisomerization i.e. *cis-trans* isomerization in excited electronic state, has been studied in a cyclodextrin cavity², at the water surface^{3,4}, in the water pool of microemulsion⁵, at micellar interface⁶, lipids⁷, sol-gel glass⁸ and polymer environments⁹. Reorientational dynamics has been studied using fluorescence anisotropy decay in DNA¹⁰, micelles¹¹⁻¹³, polymer-surfactant aggregates¹⁴. In the case of reorientational dynamics, the motion of the macromolecules becomes superimposed on the bound fluorescence probes. Several authors have discussed in detail how one can delineate the motion of macromolecules or surfactant assemblies and the fluorescent probe¹⁰⁻¹⁴.

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In the present work, we focus on the effect of *pH*-induced structural change of polyacrylic acid on isomerization and reorientation. The major aim is to demonstrate that the dynamics of isomerization and reorientation is a sensitive probe to monitor this structural change. For acidic polyelectrolytes such as polyacrylic acid (PAA), polymethacrylic acid (PMAA), DNA or poly-glutamic acid, at low *pH* the acidic (–COOH) groups do not dissociate. At low *pH*, the polyacids remain in very compact conformations with hydrophobic microenvironments. In such conformations, these polymers resemble globular proteins and can encapsulate hydrophobic solutes. At high *pH*, the –COOH group becomes ionized. Because of the repulsion between the anionic carboxylate side chains, the polymer assumes an extended rod-like shape. At low *pH*, a fluorescent probe experiences an extremely hydrophobic environment with very large friction inside the hypercoiled PAA polymer. However, at high *pH*, when the polyanion PAA opens up due to electrostatic repulsion, the fluorescent probe becomes exposed to water and experiences bulk water-like environment with low friction. The *pH*-dependent conformational transition of such polymers have been studied by fluorescence^{9,15–19}, atomic force microscopy¹⁹, and has been theoretically analysed using the Ising model¹⁷. Previous fluorescence studies of this interesting phenomenon relied on the steady state intensities of the vibronic bands of pyrene^{17,18}, electron quenching of aromatic singlets^{15b} and the efficiency of energy transfer^{9b,20}.

The *pH*-dependent uncoiling of the polymers is likely to affect the dynamics of photophysical processes, e.g. *cis*–*trans* isomerization and fluorescence depolarization due to friction. Jones and Oh studied photoisomerization of a cyanine dye, pseudo-isocyanine iodide in PMMA using the phase modulation technique^{9a}. The main nonradiative pathway of cyanine dyes is photoisomerization about a central double bond^{1,21–24}. Jones and Oh reported that at low *pH* when the microscopic friction is high, the photoisomerization is suppressed^{9a}. This is manifested in the increase in the emission intensity and lifetime at low *pH*^{9a}. However, they did not provide the rate constants of isomerization of the dye bound to PMMA at different *pH*. The rate of fluorescence anisotropy decay also depends on microscopic friction. The uncoiling of PAA with increase in *pH* should reduce microscopic friction towards reorientational motion. Jones and Oh did not study the orientational dynamics of PMMA bound dye.

In the present work, we have chosen another cyanine dye, merocyanine 540 (MC540, chart 1) as a probe^{25–29}. The main non-radiative pathway for MC540 is photoisomerization, as the other common non-radiative pathways, e.g. intersystem crossing to the triplet manifold and the internal conversion, are extremely inefficient^{26a,b}. The ground state conformation of MC540 is exclusively all-*trans*^{26b}. Following excitation to the first singlet state, the system moves freely along the torsional coordinate as there is practically no barrier for *cis*–*trans* isomerization in the excited state. The potential surface of this system is such that the perpendicular geometry in the $\pi\pi^*$ excited state is almost isoenergetic with the ground state^{1a}. Thus at the perpendicular geometry, it undergoes rapid transition from the excited electronic state to the nearly isoenergetic ground state. In the ground state, the perpendicular geometry corresponds to the peak of the barrier between the *cis* and the *trans* isomer in the ground state. Once the system reaches the peak it can undergo transition to either of them in the ground state with equal probability. The dynamics of isomerization and fluorescence depolarization of MC540 has earlier been used to probe many organized media such as micelles, reverse micelles and proteins^{11,27–30}.

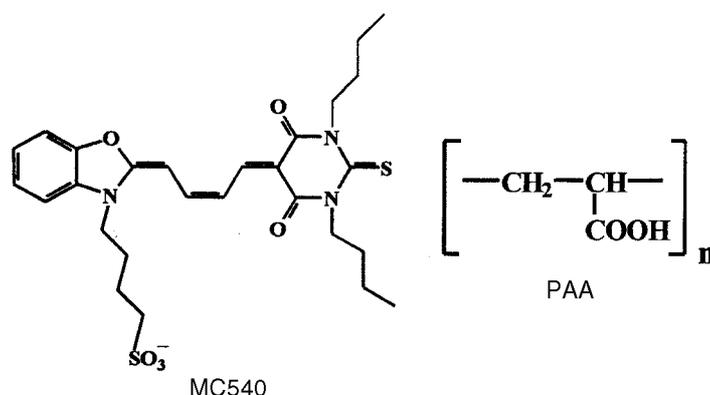


Chart 1. Structures of MC540 and PAA (acid form).

In this work we plan to investigate how the variation of friction in PAA with pH affects the dynamics of isomerization and fluorescence depolarization of MC540.

2. Experimental

The dye, merocyanine 540 (Sigma) and the polymer, PAA, sodium salt (Aldrich, $M_w = 5100$) were used as received. The sample was excited at 570 nm using a synchronously pumped rhodamine 6G dye laser (Coherent 702-1) pumped by a CW mode locked Nd:YAG laser (Antares 76S). The exciting laser light is cut off by a Corning filter (2-60). The fluorescence at 630 nm was detected at magic angle polarization, using a MCP photomultiplier (Hamamatsu 2809U). For rotational relaxation studies, the polarization of the exciting light was rotated by 90° at regular intervals using a half-wave plate to get the perpendicular (I_\perp) and parallel (I_\parallel) components. Then the anisotropy function, $r(t)$ was calculated using the formula,

$$r(t) = \frac{I_\parallel(t) - GI_\perp(t)}{I_\parallel(t) + 2GI_\perp(t)}. \quad (1)$$

The G value of the setup was determined using a probe with very fast rotational relaxation time, e.g. Nile red in methanol.

3. Results

3.1 Steady state results

In aqueous solution, MC540 exhibits two broad absorption peaks at 500 nm and 535 nm^{28,29}. The broad absorption peak at 500 nm has been assigned to the non-fluorescent dimer of MC540 and the peak at 535 nm to the monomer^{28,29}. The absorption spectra of an aqueous solution containing 4×10^{-6} M MC540 and 10 mM PAA at different pH values are shown in figure 1a. The individual contributions of the dimer and the monomer to the total absorption spectrum are obtained by decomposing the absorption spectrum. By decomposing the absorption spectrum of MC540 in 10 mM

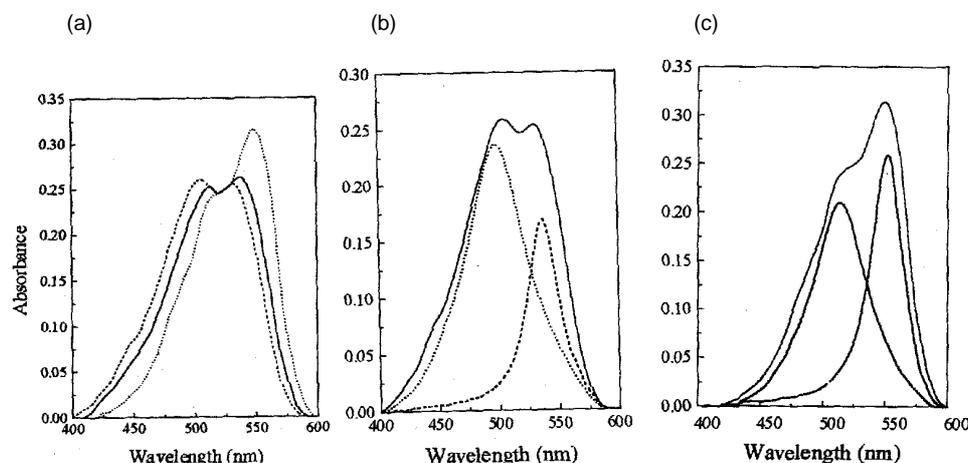


Figure 1. (a) Absorption spectra of an aqueous solution containing 4×10^{-6} M MC540 and 10 mM PAA at (i) pH = 6 (----), (ii) pH = 4 (—), (iii) pH = 2 (.....). (b) Decomposition of the absorption spectra of merocyanine 540 in 10 mM PAA at pH = 6, dimer (.....), monomer (----), total (—). (c) Decomposition of the absorption spectra of merocyanine 540 in 10 mM PAA at pH = 2, dimer (.....), monomer (----), total (—).

PAA at pH = 6, two bands are located at 500 nm and 535 nm (figure 1b). Evidently the 500 nm absorption band may be attributed to the dimer of MC540 and the 535 nm band to the monomer. At pH = 2, the decomposed absorption spectrum of MC540 in 10 mM PAA exhibits two absorption bands at 510 nm and 550 nm (figure 1c). We have earlier shown that the monomer absorption band exhibits a red shift with decrease in solvent polarity^{28,29}. While absorption maximum of the monomer is at 535 nm in water, it shifts to 565 nm in dioxane²⁹. Thus the 550 nm peak at pH = 2 is assigned to MC540 monomer bound to relatively less polar and hydrophobic environment of neutral PAA.

It is readily seen that at pH = 6, the height of dimer absorption peak of MC540 is nearly 1.4 times that of the monomer peak (figure 1b). However, at pH = 2, the monomer peak becomes 1.25 times that of the dimer peak (figure 1c). This shows that with decrease in pH, inside the PAA the dimers of MC540 breakup and polymer-bound MC540 exists predominantly as a monomer. In summary, absorption spectra suggest that at low pH, when MC540 molecules are entrapped inside the hypercoiled conformation of PAA, the dye molecules remain largely as monomers.

With decrease in solvent polarity, MC540 exhibits a red shift in emission maximum and a marked increase in emission quantum yield (ϕ_f) and lifetime²⁹. Emission maximum of MC540 shifts from 575 nm in water to 590 nm in dioxane. ϕ_f of MC540 increases from 0.04 in water to 0.52 in dioxane. Figure 2 shows the variation in the emission spectra of MC540 with pH in 10 mM PAA. It is readily seen, with increase in pH, that the emission maximum exhibits a blue shift from 585 nm at pH = 3 to 575 nm at pH = 6. It may be noted that in aqueous solution, the emission spectrum of 4×10^{-6} M MC540 does not change with the variation in pH.

With increase in pH, the emission quantum yield (ϕ_f) of MC540 in 10 mM PAA decreases nearly 7 times from 0.28 at pH = 2 to 0.04 at pH 7 (figure 2a and table 1). The

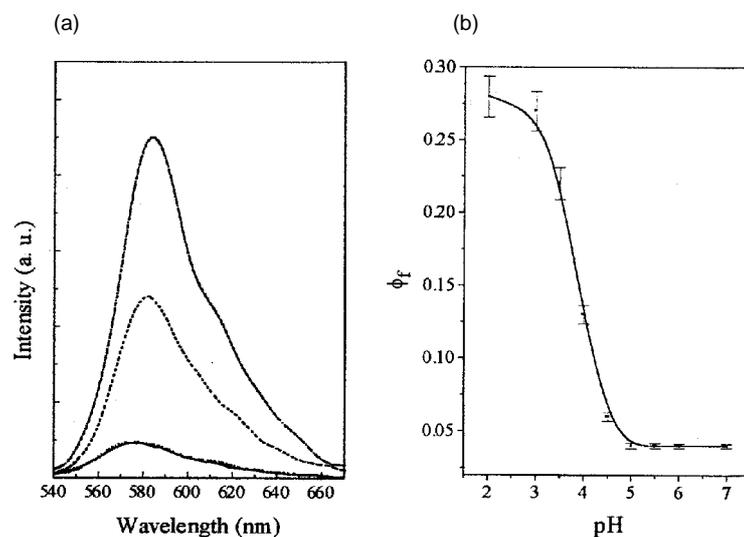


Figure 2. (a) Emission spectra of an aqueous solution containing 4×10^{-6} M MC540 and 10 mM PAA at (i) $pH = 6$ (—), (ii) $pH = 5$ (.....), (iii) $pH = 4$ (----) and (iv) $pH = 3$ (-·-·-). (b) pH dependence quantum yield (ϕ_f) of MC540 in an aqueous solution containing 4×10^{-6} M MC540 and 10 mM PAA.

Table 1. Variation of quantum yield (ϕ_f), average life time (τ_f) and isomerization rate constant (k_{iso}) of merocyanine 540 (MC 540) with change in pH in an aqueous solution of 10 mM polyacrylic acid (PAA).

pH	ϕ_f^*	$\langle \tau_f \rangle^*$ (ps)	$k_{iso} (\times 10^{-9}) (s^{-1})$
2.0	0.28	1000	0.72
3.0	0.27	975	0.75
3.5	0.22	900	0.87
4.0	0.13	700	1.25
4.5	0.06	350	2.70
5.0	0.04	170	5.65
5.5	0.04	130	7.40
6.0	0.04	120	8.00
7.0	0.04	110	8.70

* $\pm 5\%$

variation of ϕ_f of MC540 with pH in 10 mM PAA is summarized in table 1 and displayed in figure 2b. From figure 2b it is apparent that the emission quantum yield of MC540 displays a sigmoidal dependence on pH , shows sharp decrease at $pH > 4$ and levels off at $pH \approx 5$. The marked increase in ϕ_f of MC540 in 10 mM PAA at low pH suggests significant retardation of the non-radiative pathways of MC540. As noted earlier, the main nonradiative pathway in the excited state of MC540 is isomerization. The increase in ϕ_f at low pH indicates that the isomerization process of MC540 is markedly suppressed when it is trapped inside the 'hypercoiled' undissociated acid, PAA.

3.2 Time resolved fluorescence

In aqueous solution, fluorescence decay of MC540 is a single exponential with a lifetime (τ_f) of 110 ps and independent of pH of the solution²⁹. The fluorescence decay of MC540 in 10 mM PAA differs markedly from that in pure water. We fitted the decays to a biexponential, e.g. $a_1\exp(-t/\tau_1) + a_2\exp(-t/\tau_2)$ and used the average fluorescence lifetime, $\langle\tau_f\rangle = a_1\tau_1 + a_2\tau_2$.

Fluorescence decays of MC540 in 10 mM PAA at different pH values are shown in figure 3a and the average fluorescence lifetimes ($\langle\tau_f\rangle$) are listed in table 1. Figure 3b shows variation of $\langle\tau_f\rangle$ with pH. It is readily seen that with increase in pH, τ_f of MC540 decreases about 9 times from 1000 ps at pH = 2 to a bulk water like value of 110 ps at pH > 6. The increase in $\langle\tau_f\rangle$ at low pH provides additional support to the contention that the isomerization process of MC540 is inhibited inside the hypercoiled PAA. With increase in pH, as the polymer uncoils and becomes negatively charged, it repels the anionic probe MC540 to bulk water. As a result, the dynamics of isomerization becomes faster and becomes very similar to that in bulk water.

From the observed values of ϕ_f and $\langle\tau_f\rangle$, the rate constant (k_{iso}) of the nonradiative isomerization process of MC540 has been determined using the relation,

$$k_{\text{iso}} = (1 - \phi_f)/\langle\tau_f\rangle. \quad (2)$$

The values of k_{iso} of MC540 in a 10 mM aqueous solution of PAA at various pH levels are given in table 1. Variation of k_{iso} of MC540 as a function of pH is shown in figure 3c. It is obvious that the rate constant of the isomerization process increases about 12 times as pH is increased from 2 to 7. The plot of k_{iso} against pH exhibits a sharp transition at a pH \approx 4.5. The remarkably small magnitude of k_{iso} of MC540 in PAA at low pH, may be attributed to the restrictions imposed on the isomerization process of MC540 by the polymer chains in the hypercoiled form.

3.3 Fluorescence depolarization

The rotational or reorientational motion of a fluorescent probe gives rise to fluorescence depolarization. As a result, decay of the fluorescence anisotropy function, $r(t)$ is a measure of the rotational mobility of the probe. In water, the decay of fluorescence anisotropy of MC540 is found to be a single exponential with the time constant 600 ± 30 ps. This is similar to the reported anisotropy decay of MC540 in ethanol¹¹ and dioxane³⁰. The anisotropy decay of MC540 in water was found to be independent of pH.

In the presence of 10 mM PAA, fluorescence anisotropy decay of MC540 is found to be very different from that in pure water. At low pH, fluorescence anisotropy decay is found to be biexponential with much longer time constants (figure 4 and table 2). For instance at pH = 2, anisotropy decay of MC540 exhibits two components of $\tau_{1R} = 1500$ ps ($a_{1R} = 0.33$, i.e. 33%) and $\tau_{2R} = 4000$ ps ($a_{2R} = 0.67$, i.e. 67%). Thus in 10 mM PAA at pH = 2, the average rotational relaxation time of MC540, $\langle\tau_{\text{rot}}\rangle = a_{1R}\tau_{1R} + a_{2R}\tau_{2R}$, is 3200 ± 150 ps. This is about 5 times slower than the rotational relaxation time of MC540 in water (600 ps). At pH \geq 6, the anisotropy decay of MC540 in 10 mM PAA is found to be identical to that in pure water. The plot of $\langle\tau_{\text{rot}}\rangle$ of MC540 in 10 mM PAA against pH exhibits a sigmoidal dependence with sharp transition at pH \approx 4 (figure 5). The very long rotational relaxation time of MC540 in PAA at low pH, indicates that the rotational mobility of the probe decreases markedly when the polymer wraps around the probe.

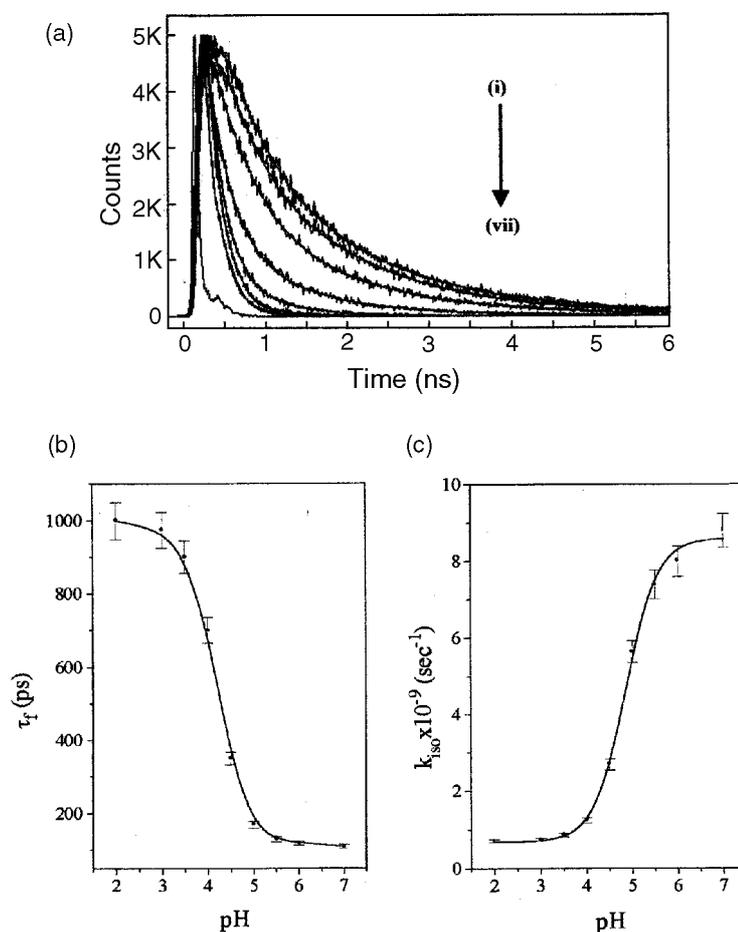


Figure 3. (a) Fluorescence decays of MC540 in an aqueous solution containing 4×10^{-6} M MC540 and 10 mM PAA at (i–vii) pH = 2.0, 3.0, 3.5, 4.0, 4.5, 5.5 and 7.0. (b) Variation of average fluorescence lifetime ($\langle \tau_f \rangle$) of MC540 with pH in 10 mM PAA. (c) pH dependence k_{iso} of MC540 in 10 mM PAA.

Table 2. Variation of fluorescence anisotropy decay time (τ_{rot}) of merocyanine 540 (MC540) with change of pH in a 10 mM polyacrylic acid solution.

pH	a_{1R}	τ_{1R} (ps)	a_{2R}	τ_{2R} (ps)	$\langle \tau_{\text{rot}} \rangle^*$ (ps)
2.0	0.33	1500	0.67	4000	3200
3.0	0.15	500	0.85	3600	3100
3.5	0.12	450	0.88	3100	2800
4.0	0.13	140	0.87	2400	2100
4.5	0.18	170	0.82	1800	1500
5.0	0.30	250	0.70	1500	1100
5.5	0.50	150	0.50	1400	780
6.0	1.0	600			600
7.0	1.0	600			600

$$*\langle \tau_{\text{rot}} \rangle = a_{1R}\tau_{1R} + a_{2R}\tau_{2R}; \pm 5\%$$

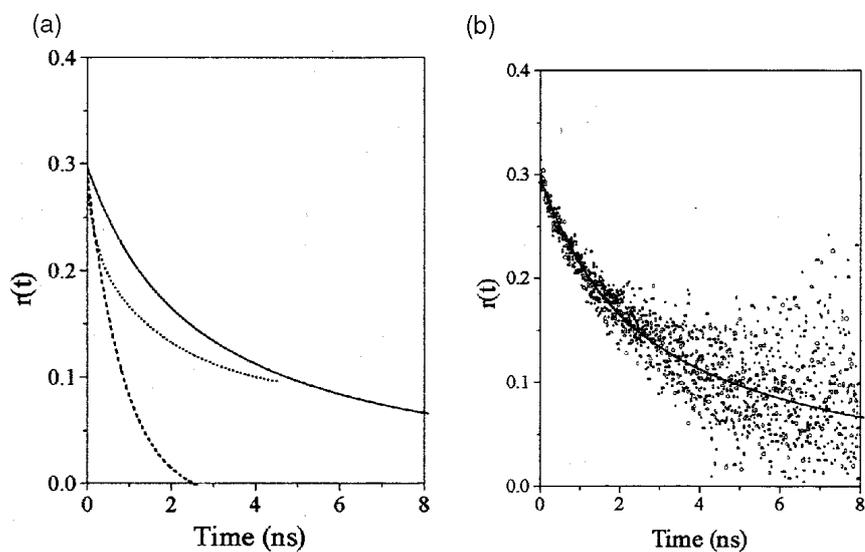


Figure 4. (a) Decays of fluorescence anisotropy of MC540 in an aqueous solution containing 4×10^{-6} M MC540 and 10 mM PAA at (i) pH = 7 (---), (ii) pH = 5 (.....) and (iii) pH = 2 (—). (b) Fitted curve along with raw data of fluorescence anisotropy decay of MC540 in 10 mM PAA at a pH = 2.

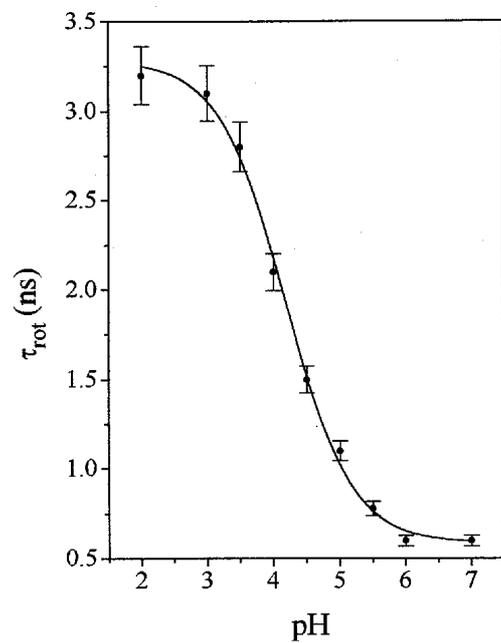


Figure 5. pH dependence of average rotational relaxation time ($\langle\tau_{rot}\rangle$) of MC540 in an aqueous solution containing 4×10^{-6} M MC540 and 10 mM PAA.

4. Discussions

The most important finding of the present work is that the pH -dependent structural change (uncoiling) of the polymer PAA, significantly influences the dynamics of isomerization and reorientation of MC540. At low pH , when the polymer wraps around the probe MC540 molecule in a compact hypercoiled form, the dynamics of the two processes are markedly slowed down compared to bulk water. However, at high pH the polyanion expands and because of its electrostatic repulsion the anionic probe, MC540, is expelled from the hydrophobic pocket of the polymer to bulk water. As a result, at high pH the probe MC540, exhibits bulk water-like fast isomerization and reorientational dynamics. For both the processes, the pH dependence of all the dynamic parameters displays a sigmoidal curve with a sharp transition at $pH \approx 4$. It may be recalled that similar sharp transition at $pH \approx 4$ was also reported in other studies on PAA^{9,15-20}, and has been attributed to ionization of PAA and consequent uncoiling. Thus, isomerization and reorientational dynamics are sensitive probes to monitor the pH dependent uncoiling of the polymer.

In 10 mM PAA, the dynamics of isomerization of MC540 at low pH is found to be 12 times slower than that at high pH (> 6) or that in water. It has been demonstrated earlier that for MC540 the photoisomerization dynamics depends on polarity as well as friction²⁹. For PAA, however, it has been shown that even at low pH the polarity of the microenvironment remains quite high¹⁷. The observed emission maximum (585 nm) of MC540 in 10 mM PAA corresponds to a water-dioxane mixture containing 60% water²⁹. For this mixture, the average lifetime ($\langle \tau_f \rangle$) and quantum yield ϕ_f of MC540 are 350 ps and 0.2 respectively²⁹. However, in the present work for MC540 in 10 mM PAA at $pH = 2$ we observed a higher ϕ_f (≈ 0.28) and longer lifetime (≈ 1000 ps). This indicates that the observed retardation of the isomerization process of MC540 in PAA at low pH arises only partly due to reduced polarity and mainly from the high microscopic friction in the hypercoiled polymer.

The very high microscopic friction in PAA at low pH is also responsible for the slow rotational relaxation time of MC540 at low pH . The observed multiexponential decay of $r(t)$ for MC540 in PAA at low pH may arise from various factors³¹. First, in such a micro-heterogeneous system, the probe MC540 might experience different magnitudes of friction in different regions. Second, the motion of the polymer chains may become superimposed on the rotational motion of the probe MC540 and as a result, the decay of $r(t)$ may be due to different kinds of independent motions¹⁰⁻¹⁴. In the case of spherical micelles, the multiexponential decay of $r(t)$ is analysed in terms of three independent motions – (i) wobbling of the probe in a cone; (ii) translational diffusion of the probe along the spherical micellar particles, and (iii) overall rotation of the micelle¹⁰⁻¹⁴. In the present case of the hypercoiled PAA at low pH shape, hydrodynamic radius, location of the probe and many other parameters are not known. Thus quantitative analysis of the decay of $r(t)$ is difficult at this stage. Nevertheless, it is evident that the average rotational relaxation time of MC540 in PAA, very clearly indicates the uncoiling of the polymer with increase in pH . One should note that the measurement of microscopic friction (microviscosity) is not straightforward even in the case of simple liquids^{1,13,14}. Even the dependence of translational diffusion coefficient on viscosity is found to be nonlinear in many cases³². Because of these complications, many authors have used reorientational relaxation time (τ_{rot}) as a measure of microscopic friction. One can also use isomerization

rate as a measure of microviscosity. Thus both the plots of τ_{rot} vs $p\text{H}$ or k_{iso} vs $p\text{H}$ should be considered as a plot of microscopic friction vs $p\text{H}$.

5. Conclusion

Dynamics of isomerization and fluorescence depolarization (i.e. rotational motion) of MC540 are found to be sensitive indicators to monitor the change of microenvironment of polyacrylic acid as a function of $p\text{H}$. With increase in $p\text{H}$ as the polymer (PAA) undergoes a structural change from hypercoiled to extended rod-like configuration, the rate constant of photoisomerization of MC540 increases about 12 times as $p\text{H}$ of the solution increases from 2 to 7. This leads to decrease of emission intensity and lifetime of MC540 by similar magnitude. The decrease in friction with $p\text{H}$ is also manifested in the nearly 5-fold decrease in the average rotational relaxation time of MC540 in PAA.

Acknowledgements

Thanks are due to the Council of Scientific and Industrial Research (CSIR), New Delhi and Department of Science and Technology, Government of India for generous research grants. DS, SS and PD thank the CSIR for fellowships.

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