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Light, temperature, and pH control of aqueous azopyridine-terminated poly(*N*-isopropylacrylamide) solutions†

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Azopyridines (AzPy) act as light-sensitive groups that undergo reversible *cis*–*trans* isomerization upon UV irradiation, as hydrogen-bond acceptors, and as ionizable moieties. The kinetics of the thermal *cis*- to *trans*-AzPy deactivation are slow except when the Py nitrogen is H-bonded or cationic. The properties of AzPy were used here to control the phase transition of aqueous solutions of α -azopyridine- ω -*n*-dodecyl-poly(*N*-isopropylacrylamides) (C12-PN-AzPy) with the molar mass (M_n) ranging from 5800 to 19 700 g mol^{−1}. The C12-PN-AzPy polymers form cationic star-micelles in solutions of pH 3 and flower-micelles in neutral and basic solutions. This diversity of micelle morphology underlies the temperature-, pH- and UV-irradiation-driven phase transition of aqueous C12-PN-AzPy solutions as demonstrated by turbidimetry, ¹H NMR spectroscopy, and microcalorimetry. Unlike azobenzene, the commonly used photoresponsive moiety to actuate amphiphilic polymers, AzPy can affect the thermoresponsive behavior of polymers in response to three orthogonal triggers: pH, through changes in ionization; light, via *trans*–*cis* photoisomerization; and time, from hours to a few ms, via the kinetics of the dark *cis*–*trans* relaxation. The study leads the way to responsive sensors or actuators in the form of aqueous fluids, hydrogels, or films by the application of light and changes of temperature and pH in permutable sequences.

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Introduction

Aromatic azo compounds form an important class of dyes and pigments used for centuries as brilliant dyestuffs for textiles, paints, and many other applications.¹ In the 1970s, polymers containing azobenzenes, commonly referred to as “azo-polymers”, caught the attention of photochemists and polymer scientists alike, driven by the pioneering work of Neckers among others.² Major advances within the field of azo-polymers occurred with the emergence of the concept of azobenzenes as photosensitive “triggers” of macroscopic changes of the properties of materials and solutions, such as phase

changes,³ and photo-induced motion and deformation of polymer films.⁴ In most cases, the photo-response is linked to the *trans*-to-*cis* photoisomerization of azobenzene chromophores attached to the polymeric structures. UV-irradiation of azobenzene causes the conversion of the *trans* isomer to the less stable *cis* isomer, which in the dark slowly relaxes to the *trans* form. The *trans*-to-*cis* photoisomerization of azobenzene changes the photophysical properties of the chromophore, as detected by steady-state and transient UV-Vis absorbance spectroscopy, both of which are well documented.^{5–7} Other physical properties of the azobenzene moiety are affected as well, most notably its dipole moment: it increases or decreases upon photoisomerization, depending on the type and position of phenyl ring substituents.^{8,9} Thus, photoisomerization of azobenzenes linked to a thermosensitive amphiphilic polymer affects the phase transition temperature of the polymer in water as a consequence of the different hydrophilicity of the *cis* and *trans* isomers.^{10,11}

For practical applications that require fast reversibility, it is advantageous to use azobenzenes that exhibit an extremely fast thermal *cis* to *trans* relaxation, so that a single UV light beam can induce continuous *trans* ↔ *cis* isomerizations.¹² The rate of the dark *cis* to *trans* isomerization of azobenzene can be enhanced significantly through manipulations of the π electron distribution, such as *para*-substitution of the phenyl rings¹³ or the replacement of one phenyl ring of azobenzene

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by a pyridine ring.^{14,15} The dark *cis-trans* isomerization of azopyridine (AzPy) in the neutral form is nearly as slow as that of azobenzene. It is extremely fast (relaxation time ~ 1 ms or less) for the quaternized azopyridinium (AzPy⁺) group. The formation of hydrogen bonds between the AzPy nitrogen and H-bond donors also greatly accelerates the thermal *cis-trans* isomerization.

Azopyridine-substituted water-soluble polymers are amphiphilic. They tend to self-assemble in water, adopting various morphologies as a function of their chemical composition and their environment, in particular the solution pH and the presence of H-bond acceptors.^{16–18} This phenomenon is exemplified in a recent study of aqueous solutions of poly(*N*-isopropylacrylamides) bearing an AzPy on one chain end and an *n*-dodecyl group on the other end (C12-PN-AzPy) (Scheme 1a).¹⁹ In neutral or basic water, they form “flower-like” core-shell micelles with a core of *n*-dodecyl chains and AzPy groups surrounded by a PNIPAM corona. In solutions of pH 3, the micelle morphology changes as a result of the protonation of the AzPy group. C12-PN-AzPyH⁺ polymers form star micelles consisting of a hydrophobic core made solely of assembled *n*-dodecyl groups, surrounded by extended PNIPAM chains terminated with the cationic azopyridinium group.

Aqueous solutions of PNIPAM and its derivatives undergo a reversible phase transition upon heating past a temperature of ~ 32 °C that corresponds to the coil-to-globule collapse of the polymer chains and subsequent aggregation of the dehydrated PNIPAM globules into larger mesoglobules.^{20–22} This property is exploited commonly to actuate temperature sensitive materials. The phase transition temperature of PNIPAM derivatives in water can be adjusted over a broad range of temperatures by linking to PNIPAM hydrophilic/hydrophobic comonomers^{22–24} or end groups.^{25,26} The introduction of stimuli sensitive components responsive to stimuli, such as light,²⁷ pH,^{28,29} salinity, redox conditions,^{30,31} or CO₂,³² expands even wider the range of possible applications of PNIPAM containing polymers. End-modified PNIPAM deriva-

tives are particularly useful as they can drive large changes of the physical or chemical properties of the materials through minor changes of the polymer chemical composition.³³

Generally, non-reactive chain ends do not affect the phase transition temperature of PNIPAM of $M_n \geq 50$ kDa. In the case of shorter PNIPAM chains, as a rule, hydrophilic end groups increase the solution phase transition temperature and hydrophobic end groups decrease it.³⁴ These effects result from changes in the hydrophilic/lipophilic balance of the modified polymer chains. Charged end groups, such as carboxylates^{35,36} and quaternary ammonium salts,^{37,38} tend to prevent the aggregation of the dehydrated PNIPAM globules through electrostatic stabilization. If the size of carboxylated PNIPAM globules is sufficiently small, compared to the wavelength of light, the solution remains clear at temperatures higher than the phase transition temperature.

We report here a study by microcalorimetry, turbidimetry, and NMR spectroscopy of the phase transition of aqueous C12-PN-AzPy solutions as a function of changes in temperature and solution pH, and upon UV-light irradiation. The study demonstrates also that through their ionizability and pH-/H-bond-dependent dark relaxation kinetics, azopyridine groups are simple, versatile, and powerful tools to modulate the temperature-induced phase transition of amphiphilic polymers, such as PNIPAM upon the application of two or three orthogonal parameters, a key outcome of this investigation.

Experimental section

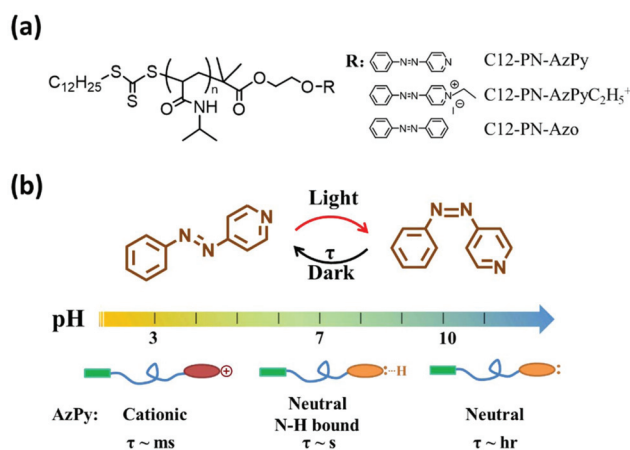
Materials

Water was deionized using a Millipore Milli-Q system. The C12-PN-AzPy samples were prepared by reversible addition-fragmentation transfer radical polymerization using a chain transfer agent bearing an *n*-dodecyl group and an azopyridine moiety, as described in detail previously.¹⁹ Their molar mass ranges from M_n 5800 g mol⁻¹ to 19 700 g mol⁻¹ (Table 1). Their chemical structure is presented in Scheme 1.

Methods

Turbidity measurements. The cloud point temperatures of polymer aqueous solutions (1.0 mg mL⁻¹) were determined using a UV-Vis spectrometer (Agilent 8452A) equipped with a temperature controller (Agilent 89090A). The heating rate was 0.5 °C min⁻¹. Samples were equilibrated for 2 min at each temperature before measurement. The solution transmittance at 550 nm was measured as a function of the temperature from 15 °C to 60 °C. To determine the cloud point of polymer aqueous solutions subjected to UV light irradiation, the same protocol was performed with solutions subjected to constant irradiation at 365 nm with a LED-UV curing lamp (10% intensity, 30 mW cm⁻²). The cloud point (T_{cp}) was taken as the temperature of the onset of the decrease of the solution transmittance.

Differential Scanning Calorimetry (DSC) measurements were performed on a VP-DSC microcalorimeter (MicroCal Inc.) with



Scheme 1 (a) Chemical structure of the polymers investigated in this study; (b) schematic representation of the azopyridine end group of C12-PN-AzPy in solutions of pH 3, 7 and 10.¹⁹



Table 1 Molar mass data of the polymers examined here and properties of their aqueous solutions (pH 7 and pH 3, polymer concentration: 1.0 mg mL⁻¹)

Sample name	Mn (g mol ⁻¹) ^a	M _w /M _n ^a	T _{cp} /°C		T _m /°C		ΔH/kJ	
			pH = 7	pH = 3	pH = 7	pH = 3	pH = 7	pH = 3
C12-PN-AzPy 5 K	5800	1.25	14.0	23.9	18.9	— ^b	2.7	— ^b
C12-PN-AzPy 7 K	7800	1.03	20.0	27.5	24.2	26.2	3.2	3.9
C12-PN-AzPy 12 K	12 900	1.09	22.7	28.2	26.8	28.1	4.3	4.9
C12-PN-AzPy 20 K	19 700	1.03	27.0	32.4	29.2	30.1	6.0	7.3
C12-PN-AzPyC ₂ H ₅ ⁺ 12 K	12 900	1.09	30.0	28.4	31.1	31.0	4.7	4.2
C12-PN-Azo 12 K	13 600	1.23	22.5	19.5	26.6	— ^b	4.2	— ^b

^a M_n and M_w/M_n were obtained by GPC with MALS detection (see ref. 14). ^b Data not available.

a cell volume of 0.520 mL. Three consecutive heating and cooling scans from 10–70 °C were performed with a heating and cooling rate of 1.0 °C min⁻¹ using polymer solutions in water at a concentration of 1.0 mg mL⁻¹.

¹H NMR spectra were recorded on a Bruker AMX-400 (400 MHz) spectrometer. Polymer solutions in D₂O (2.0 mg mL⁻¹) were brought to pH 10 by the dropwise addition of a 0.1 M NaOD solution. The samples were kept in a refrigerator for 24 h prior to measurement. For irradiation-dependent ¹H NMR analysis, the polymer solutions were irradiated with a LED-UV curing lamp (365 nm, 10% intensity, 30 mW cm⁻²) for 30 s immediately prior to measuring their ¹H NMR spectrum.

Evaluation of the photo/pH/temperature responsiveness of the polymer solutions

The pH- and photo-controlled thermoresponsive properties of polymer solutions were evaluated on a Mettler Toledo UV-Vis spectrophotometer (UV5 bio) equipped with a CuveT temperature controller. For experiments at constant temperature in the absence of UV light illumination, the absorbance of a solution of C12-PN-AzPy (1.0 mg mL⁻¹) at 550 nm was recorded during a series of consecutive changes of the solution pH from pH 7 to pH 3 and from pH 7 to 10 using, respectively, 0.2 mM HCl or 0.2 mM NaOH aqueous solutions to adjust the solution pH. The same measurements were performed also for polymer solutions under constant UV irradiation (365 nm, 30 mW cm⁻²) at a constant temperature.

Results and discussion

3.1 pH Dependence of the phase transition of aqueous *trans*-C12-PN-AzPy solutions

Table 1 presents key parameters of the heat-induced phase transition of C12-PN-AzPy aqueous solutions as a function of pH and the molar mass of the polymers in the absence of UV light irradiation. The T_{cp} values of aqueous C12-PN-AzPy solutions of pH 7 increase with increasing polymer molar mass (Fig. 1a), as a consequence of the increase of the hydrophilic polymer chain length while the length of the hydrophobic end group remains constant.^{26,39,40} The T_{cp} of C12-PN-Azo 12 K, a control sample that has an azobenzene end group instead of the AzPy group, is nearly identical to that of C12-PN-AzPy 12 K

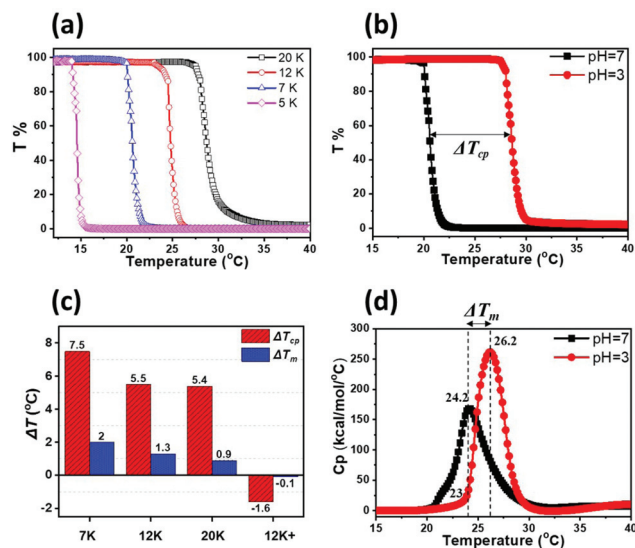


Fig. 1 (a). Plots of the % transmittance at 550 nm as a function of temperature for solutions of C12-PN-AzPy of various molar mass in water (pH 7); (b) plots of the % transmittance as a function of temperature for solutions of C12-PN-AzPy 7 K of pH 7 (black squares) and pH 3 (red circles). ΔT_{cp} is defined as T_{cp} (pH 3) – T_{cp} (pH 7); (c) ΔT_{cp} (red bar) and ΔT_m (blue bar) values for C12-PN-AzPy of various molar mass; (d) thermograms of solutions of C12-PN-AzPy 7 K of pH 7 and pH 3, ΔT_m is defined as T_m (pH 3) – T_m (pH 7).

(22.5 °C vs. 22.7 °C), see Fig. S1.† The cloud points of C12-PN-AzPy solutions of pH 3, for which the azopyridine end group is protonated (pK_a of azopyridine ~4.52),⁴¹ are higher than those of the corresponding neutral solutions, as seen in Fig. 1b for solutions of C12-PN-AzPy 7 K. The amplitude of the increase in T_{cp} upon acidification, ΔT_{cp} = T_{cp} (pH 3) – T_{cp} (pH 7) decreases with increasing polymer molar mass (see Fig. 1c, red bars). The T_{cp} of C12-PN-AzPyC₂H₅⁺ 12 K, which bears the cationic *N*-ethyl azopyridinium end group, is (28.4 °C, Fig. S1.†) a value similar to that of C12-PN-AzPy 12 K in a pH 3 solution (28.2 °C). As a comparison, the T_{cp} values of aqueous solutions of unmodified PNIPAM of pH 3, 7, and 11 were determined as well (Fig. S2.†). The T_{cp} values of the pH 3 and 11 solutions were ~0.5 °C lower than those of the pH 7 solution. It is worth mentioning that the amide group of the NIPAM repeat units and the ester group that links the AzPy end group to the main



chain withhold the experimental conditions, as reported previously according to previous reports.^{19,42}

High sensitivity microcalorimetry (HS-DSC) thermograms recorded for aqueous solutions of C12-PN-AzPy 7 K of pH 7.0 (black squares) and pH 3.0 (red circles) are presented in Fig. 1d, where T_m denotes the temperature corresponding to the maximum of the endotherm. The shift to a higher temperature of the transition maximum upon acidification ($0.9\text{ °C} < \Delta T_m < 2.0\text{ °C}$, with $\Delta T_m = T_m(\text{pH} 3) - T_m(\text{pH} 7)$) (Fig. 1c, blue bars) is smaller than ΔT_{cp} . The enthalpy of the phase transition (ΔH , expressed in kJ mol^{-1} of NIPAM units) decreases with decreasing C12-PN-AzPy molar mass, especially in the case of neutral solutions (Table 1). This trend was observed also in earlier studies of aqueous solutions of neutral α , ω di-*n*-octadecyl-PNIPAM that forms core-shell flower micelles.^{43,44} Since the transition enthalpy corresponds to the release of polymer-bound water molecules into bulk water,⁴⁵ a decrease in enthalpy indicates a lower degree of hydration of the PNIPAM corona, especially in the corona section close to the alkyl core as a result of the chain confinement. The ΔH values recorded for aqueous solutions of the longest polymer, C12-PN-AzPy 20 K in neutral and acidic solutions, 6.0 kJ mol^{-1} and 7.3 kJ mol^{-1} , respectively, are similar to those of unmodified linear PNIPAM.^{44,46} Hence, C12-PN-AzPy 20 K exists in water in the form of free chains or loose aggregates, rather than core-shell micelles.

Comparing the turbidity curve recorded for the aqueous acidic C12-PN-AzPy 7 K (Fig. 1b, red curve) to the thermogram of this solution (Fig. 1d, red curve), we observe that the T_{cp} value of this solution, 27.5 °C , is higher than T_m (26.2 °C). The azopyridinium ions stabilize the dehydrated/collapsed chains against macroscopic aggregation,³⁶ but their stabilizing effect is confined within a narrow temperature window ($\sim 1\text{--}2\text{ °C}$ above T). Upon further heating, solutions become turbid indicating the formation of large mesoglobules. We surmise, that as the temperature increases, the hydrophobic AzPy⁺ groups relocate within the collapsed chains decreasing the electrostatic repulsion between collapsed globules. In the case of the neutral C12-PN-AzPy (pH 7) solutions, T_{cp} nearly coincides with the endotherm onset temperature, which confirms that under neutral conditions the clouding of the solution and the dehydration/collapse of the PNIPAM chains occur at about the same temperature (Fig. 1d). The T_{cp} values of polymer solutions of pH 10 are slightly lower than the corresponding values for solutions of pH 7, which can be attributed to the presence of OH[−] ions.^{47,48} (see turbidity plots in Fig. S3†).

3.2. Combined effects of pH and UV light irradiation on the phase transition of aqueous C12-PN-AzPy solutions

UV light photoirradiation of *trans* AzPy yields the less stable *cis* AzPy that thermally relaxes back to the *trans* form. As described above, the *cis*–*trans* thermal relaxation of AzPyH⁺ in C12-PN-AzPyH⁺ micelles (pH 3) is on the order of a few ms as determined by transient absorption spectroscopy (Scheme 1b, box pH = 3 inset). In neutral solutions, the *cis*–*trans* dark relaxation is on the order of a few seconds, since the flower-like morphology of C12-PN-AzPy enables the formation of H-bonds

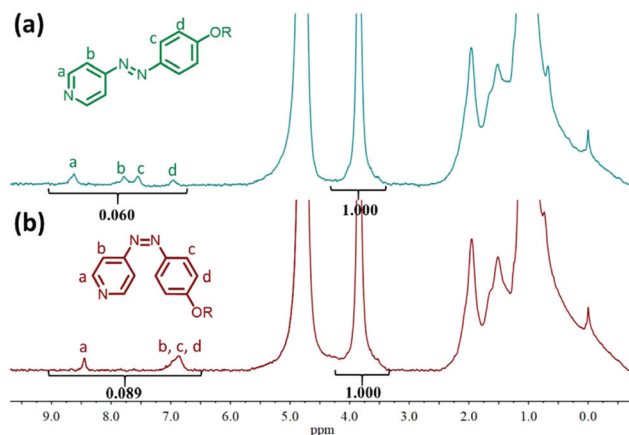


Fig. 2 ¹H-NMR spectrum of C12-PN-AzPy 7 K (a) before and (b) after UV irradiation at pH = 10 (1.0 mg mL^{−1} in D₂O, 18.5 °C).

between the AzPy nitrogen and the amide hydrogen of the surrounding PNIPAM repeat units. In a basic solution, the dark *cis* to *trans* relaxation of AzPy takes several hours since the hydrogen bonds between the AzPy nitrogen and the PNIPAM amide groups are broken by the excess hydroxyl ions competing with the AzPy nitrogen as hydrogen bond acceptors.¹⁹ *Cis*-C12-PN-AzPy is sufficiently long-lived in alkaline aqueous solutions for the characterization by NMR spectroscopy, turbidimetry, and steady state absorption spectroscopy (Fig. S4†).

The ¹H NMR spectra of C12-PN-AzPy in D₂O (pH 10) before irradiation and after 30 s irradiation at 365 nm are presented in Fig. 2a and b, respectively. Then *trans*–to–*cis* isomerization induces significant changes of the aromatic proton resonances. The protons *meta* to the pyridine nitrogen and the protons on the phenyl ring experience large upfield shifts from 7.79, 7.54, and 6.95 ppm (labeled b, c, and d) to ~ 6.86 ppm. The protons *ortho* to the pyridine nitrogen (labeled a in Fig. 2) undergo a small upfield shift from 8.63 to 8.45 ppm. In addition, the total area of the signals due to the aromatic protons, normalized to the area of the signal at 3.85 ppm due to the NIPAM methine protons increases by 48.3% (from 0.060 for *trans*-C12-PN-AzPy to 0.089 for the *cis* isomer). The upfield shift of the aromatic proton resonances reflects the decrease of the π -electron delocalization in the non planar *cis* AzPy, compared to *trans* AzPy.³⁷ The apparent increase of the aromatic resonance intensity is an indication of the enhanced mobility of the aromatic rings of *cis*-AzPy compared to *trans*-AzPy, possibly as a result of a higher level of hydration of the *cis* AzPy vs. the *trans* AzPy. Similar ¹H NMR results were observed in aqueous solutions of C12-PN-AzPy 12 K (Fig. S5†).

The cloud points of pH 10 aqueous solutions of *cis*-C12-PN-AzPy were obtained by measuring the changes in the solution transmittance at 550 nm as a function of temperature under constant irradiation at 365 nm. The resulting turbidity plots of solutions of *cis*-C12-PN-AzPy 7 K (full symbols) and *trans*-C12-PN-AzPy 7 K (open symbols) are presented in Fig. 3b. The cloud point of the *cis* form is significantly higher than



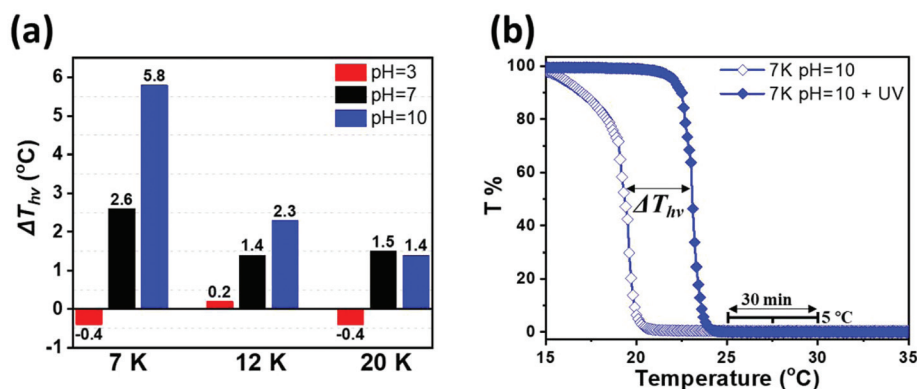


Fig. 3 (a) ΔT_{hv} values for C12-PN-AzPy of various molar mass at pH 3 (red bar), pH 7 (black bar) and pH 10 (blue bar), where $\Delta T_{hv} = T_{cp}$ (under irradiation) – T_{cp} (before irradiation). (b) Plots of the % transmittance as a function of temperature for solutions of C12-PN-AzPy 7 K of pH 10 before (open squares) and under continuous UV irradiation (full squares) at 365 nm.

Table 2 UV induced cloud point shift for the polymers under different pH values, where $\Delta T_{hv} = T_{cp}$ (under irradiation) – T_{cp} (before irradiation) (polymer concentration: 1.0 mg mL⁻¹)

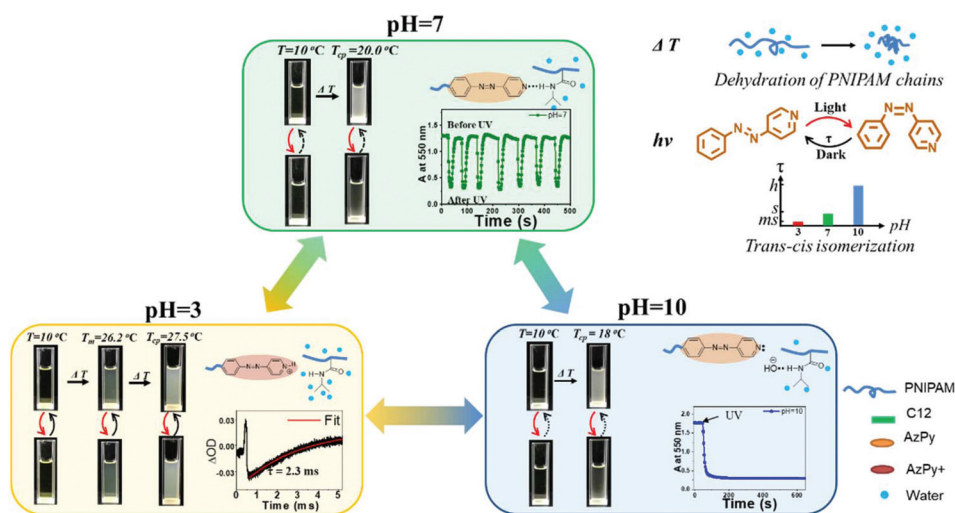
Sample name	pH	T_{cp} before UV/°C	T_{cp} after UV/°C	$\Delta T_{hv}/^{\circ}\text{C}$
C12-PN-AzPy 7 K	3	27.5	27.1	-0.4
	7	20.0	22.6	2.6
	10	16.0	21.8	5.8
C12-PN-AzPy 12 K	3	28.2	28.4	0.2
	7	22.7	24.1	1.4
	10	22.0	24.3	2.3
C12-PN-AzPy 20 K	3	32.4	32.0	-0.4
	7	27.0	28.5	1.5
	10	26.8	28.2	1.4
C12-PN-AzPyC ₂ H ₅ ⁺ 12 K	7	30.0	30.4	0.4
C12-PN-Azo 12 K	7	22.5	24.5	2.0

that of the *trans* form, with $\Delta T_{hv} = T_{cp}(\text{cis}) - T_{cp}(\text{trans}) \sim 5.8$ °C. The ΔT_{hv} value decreases with increasing PNIPAM molar mass, due to the increase of the NIPAM units/AzPy ratio

(see Table 2) presented in Fig. S6.† The increase in the hydration, observed by ¹H NMR spectroscopy, and larger dipole moment of *cis*-AzPy, compared to *trans*-AzPy, are contributing factors causing the increase of $T_{cp}(\text{cis})$.⁴⁹

We measured also turbidity plots of C12-PN-AzPy aqueous solutions of pH 3 and 7 before irradiation (*trans* form) and under continuous UV irradiation. For solutions of pH 3, the T_{cp} values recorded during irradiation were identical to the corresponding $T_{cp}(\text{trans})$, as anticipated since the short-lived *cis* isomer cannot be detected by turbidity measurements. The T_{cp} of C12-PN-AzPyC₂H₅⁺ during irradiation, also, is identical to that recorded for *trans*-C12-PN-AzPyC₂H₅⁺ (see Table 2). A small increase of T_{cp} upon irradiation was observed for C12-PN-AzPy in neutral aqueous solutions, which indicates that the irradiated solutions contain a small, but detectable, fraction of *cis*-C12-PN-AzPy throughout the measurement.

Scheme 2 presents an overview of the properties and appearance of aqueous solutions (pH 3, pH 7, and pH 10) of



Scheme 2 pH- and UV-induced thermoresponsive properties of C12-PN-AzPy (the temperatures correspond to a polymer of $M_n \sim 7000$ g mol⁻¹, ΔOD : difference in the optical density before and after irradiation).

AzPy capped PNIPAMs upon UV irradiation and/or upon heating and the different time scales of the phenomena involved.

In neutral or basic aqueous solutions of C12-PN-AzPy (blue and green frames) the polymers adopt a core-shell micelle morphology. Upon heating in the dark, the PNIPAM chains of the shell dehydrate and collapse resulting in the formation of smaller hydrophobic micelles that aggregate into larger particles, observed visually by the appearance of turbidity (top row vials in each frame). Upon short UV irradiation, the turbid solutions become transparent, since light converts *trans*-C12-PN-AzPy into *cis*-C12-PN-AzPy, which has a higher T_{cp} . In the dark, solutions of pH 7 become turbid again within a few seconds. The recovery is very slow in the case of solutions of pH 10.

In solutions of pH 3 (yellow frame), C12-PN-AzPyH⁺ chains assemble in star cationic micelles. The heat induced dehydration of PNIPAM results in the formation of small aggregates of collapsed micelles stabilized electrostatically, barely detectable visually, that increase in size upon further heating.³⁶ Constant UV irradiation has no effect on the visual appearance of pH 3 C12-PN-AzPyH⁺ solutions at a given temperature, since the *trans* and *cis* isomers undergo fast exchange.

Based on the above, we can design complex “isothermal” phase transition systems by simply changing the solution pH or applying UV irradiation. For example, we can control the solution turbidity at a constant temperature by consecutive changes of the solution pH from pH 7 to pH 3 or to pH 10 using 0.2 mM HCl or 0.2 mM NaOH aqueous solutions, see Fig. S7.† An increase of the pH from 7 to 10 will instantly stop the UV on/off reversible isothermal transition, and the reversibility is recovered upon decreasing the solution pH from 10 to 7. Various permutations of the trigger sequences can be envisaged in specific situations.

Conclusion

The ionization of AzPy end groups, the *trans*–*cis* photoisomerization of AzPy, and the kinetics of the *cis*–*trans* relaxation over time ranging from hours to a few ms influence the thermo-responsive behavior of the polymers, each in a specific way. Ionization of the azopyridine end group stabilizes the dehydrated chains against macroscopic aggregation within a specific temperature range (~1–2 °C above T_m). Of particular importance from the practical view point, the UV induced turbidity of the polymeric solutions changes as a function of pH, as a consequence of the pH-dependent kinetics of the *cis* to *trans* dark relaxation of AzPy. This study contributes to the fundamental understanding of the effects of end group ionization and photosensitivity on the macroscopic phase transition of modified PNIPAMs. It leads the way to simple strategies for the use of AzPy functional amphiphilic polymers in aqueous fluids, hydrogels, or films as sensors or actuators controllable by the application of light and changes of temperature and pH in permutable sequences.

Conflicts of interest

There are no conflicts of interest to declare.

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