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rsc.li/materials-advancesExploiting α -benzylated 1,4-butanediols to expedite the discovery of small-molecule, LCST-type sulfobetaine zwitterionic materials†

Yen-Ho Chu, * Pin-Hsuan Chen and Hsin-Heng Huang

Herein, we report both the concise synthesis and discovery of a small library of 16 sulfobetaine-like zwitterionic materials (ZMs) that exhibit LCST phase transitions in water and their structural optimization to further fine-tune the polarity of the benzyl substituent on the α -position of the sulfonate moiety. Ultimately, we achieved the preparation of novel ZMs carrying attractive T_c values, which demonstrated the real value of the structural tunability of these ZMs. Furthermore, we present a proof-of-concept application for the analysis of biomolecules. To the best of our knowledge, this is the first report on sulfobetaine-substituted ZMs as small-molecule smart materials exhibiting LCST property and their miscibility/immiscibility with water triggered by temperature are entirely switchable and reversible.

Introduction

Materials research is focused on understanding the properties, structural optimization, and application of developed materials. Thermoresponsive materials are smart materials that are designed and synthesized to flexibly respond to changes in the surrounding temperature.^{1–4} Lower and upper critical solution temperature (LCST and UCST, respectively) systems are the two typical phase behaviors of thermoresponsive materials with water.⁴ In an LCST system, the solubility of the material in water decreases with an increase in temperature and a homogeneous solution is formed below its critical temperature (T_c); that is, its components are miscible at all concentrations below the LCST. In contrast, for UCST systems, two heterogeneous mixtures become homogeneously mixed upon heating above T_c . To date, many polymeric materials^{1,2} have been found to exhibit LCST phase transitions in water, whereas there are scarce reports on small-molecule materials exhibiting LCST properties.^{3,4} Because of the recent advances and significant progress in zwitterionic materials (ZMs) by Ohno and co-workers,³ we envisioned that small molecules such as sulfobetaines⁵ are zwitterionic, their structures are chemically tuneable, and in particular the sulfonate motif can be readily modified. Consequently, these sulfobetaine-based ZMs should be new candidate

smart materials, potentially exhibiting LCST thermoresponsive property (Fig. 1).

Herein, we report the concise two-step synthesis of a library of small-molecule ZMs (ZM 1a–e, ZM 2a–e, and ZM 3a–e), exploring their ability to undergo an LCST-type phase transition with water, structurally engineering their T_c values, and lastly presenting a proof-of-concept preliminary application for the analysis of biomolecules. Sulfobetaine ZMs are highly charged organic salts, with their cation and anion covalently tethered, sustaining their overall electroneutrality (Fig. 1). These ZMs carry unique advantages, including the absence of undesirable ion exchange and negligible vapor pressure. Furthermore, many ZMs are known to be biocompatible, non-fouling and resistant nonspecific protein adsorption in solution and on surfaces, mainly owing to their ultra-high dipole moments (18–30 D and 35–41 D for sulfobetaines and zwitterionic imidazolium sulfonates, respectively, while 1.9 D for water) and

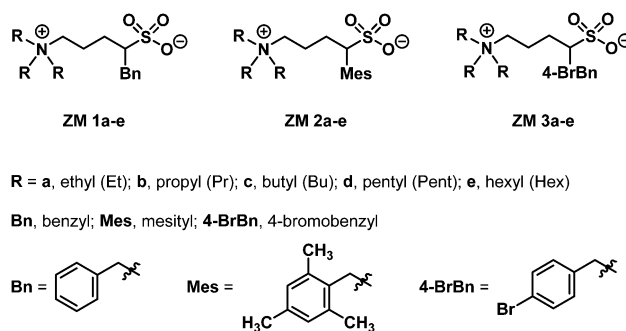


Fig. 1 Structures of small-molecule zwitterionic materials (ZM 1a–e, ZM 2a–e, and ZM 3a–e).

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† Electronic supplementary information (ESI) available: Experimental procedures, ¹H and ¹³C NMR, and high-resolution mass spectrometry (HRMS) spectra and data of all 16 ZMs (ZM 1a–e, ZM 2a–e, ZM 3a–e, and ZM 4c). See DOI: <https://doi.org/10.1039/d2ma01063a>

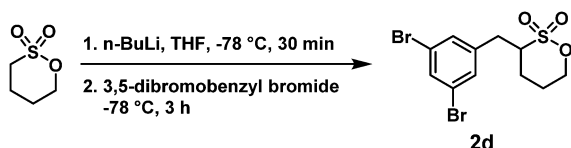
superhydrophilicity.^{5–9} In addition to their strong attraction with water,¹⁰ ZMs readily associate with themselves *via* interactions between their cationic and anionic groups. Accordingly, the intermolecular strong hydration and self-association ability of ZMs make them susceptible to external stimuli and phase responsive to temperature changes in water.^{7,8} These ZMs exhibit exceptional properties compared to common ionic liquids and have potential as valuable materials for studies of biomacromolecules with high selectivity and efficiency.

Experimental

Synthetic procedures

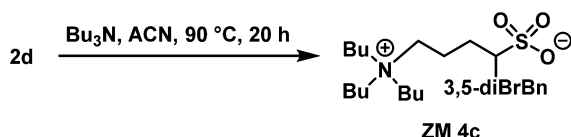
The detailed experimental procedures, ¹H and ¹³C NMR spectra, and high-resolution mass spectrometry (HRMS) data of all 16 ZMs (**ZM 1a–e**, **ZM 2a–e**, **ZM 3a–e**, and **ZM 4c**) are summarized in the (ESI[†]).

Preparation of α-(3,5-dibromobenzyl)-1,4-butanedisulfone (**2d**)



Briefly, *n*-BuLi (1.1 equiv., 2.5 M in hexane) was slowly added to a stirring solution of 1,4-butanedisulfone (0.35 mL, 1.0 equiv.) in THF (0.5 M) at -78 °C. This α-deprotonation reaction on 1,4-butanedisulfone was carried out at -78 °C for 30 min. Then, a solution of 3,5-dibromobenzyl bromide (1.2 equiv. in THF) was added at -78 °C. The reaction mixture was stirred for another 3 h, and finally quenched with water. The resulting mixture was extracted with ethyl acetate three times. The combined organic layers were concentrated and purified by column chromatography (silica gel; ethyl acetate/hexane = 1:10–1:2, v/v) to give α-(3,5-dibromobenzyl)-1,4-butanedisulfone (**2d**) as a colorless liquid with 63% isolated yield. ¹H NMR (400 MHz, CDCl₃) δ 1.83–2.07 (m, SOCH₂CH₂ + SCHCH₂CH₂, 4H), 2.68–2.75 (m, SCHCH₂Ar, 1H), 3.21–3.28 (m, SCH, 1H), 3.41–3.46 (m, SCHCH₂Ar, 1H), 4.46–4.61 (m, SOCH₂CH₂, 2H), 7.30 (d, *J* = 1.6 Hz, aryl H, 1H), 7.30 (d, *J* = 1.6 Hz, aryl H, 1H), 7.59 (dd, *J* = 1.6, 1.6 Hz, aryl H, 1H); ¹³C NMR (100 MHz, CDCl₃) δ 24.04, 27.77, 33.69, 60.18, 73.94, 123.44, 131.14, 133.21, 139.96; FAB-HRMS *m/z* [M + H]⁺ calculated for C₁₁H₁₃Br₂O₃S⁺ 382.8947, 384.8926, 386.8906, found 382.8953, 384.8943, 386.8925.

Synthesis of zwitterionic material **ZM 4c**



Tributylamine (1.5 equiv.) and acetonitrile (3.0 M) were added to a 4 mL sample bottle containing **2d** (0.27 g, 1.0 equiv.), and the corresponding α-benylation reaction was performed at 90 °C for 20 h. The reaction product was readily purified by column chromatography (silica gel; dichloromethane/

methanol = 30:1–6:1, v/v) to afford the desired **ZM 4c** as a white solid (mp 191 °C) with 62% isolated yield. ¹H NMR (400 MHz, CDCl₃) δ 1.00 (t, *J* = 7.3 Hz, N(CH₂)₃CH₃, 9H), 1.38–1.47 (m, N(CH₂)₂CH₂CH₃, 6H), 1.57–1.63 (m, NCH₂CH₂CH₂CH₃ + SCHCH₂CH₂, 7H), 1.79–1.85 (m, SCHCH₂CH₂, 1H), 1.97–1.99 (m, SCHCH₂CH₂, 2H), 2.63–2.69 (m, SCHCH₂Ar, 1H), 2.90–3.34 (m, SCH + NCH₂, 9H), 3.61–3.65 (m, SCHCH₂Ar, 1H), 7.35 (s, aryl H, 1H), 7.50 (s, aryl H, 2H); ¹³C NMR (100 MHz, CDCl₃) δ 13.78, 19.62, 19.90, 24.06, 26.54, 37.15, 59.03, 59.44, 59.82, 122.98, 131.25, 131.87, 144.89; ESI-HRMS *m/z* [M + H]⁺ calculated for C₂₃H₄₀Br₂NO₃S⁺ 568.1090, 570.1070, 572.1049, found 568.1082 ([M + H]⁺), 570.1062 ([M + H]⁺), 572.1040 ([M + H]⁺), 590.0901 ([M + Na]⁺), 592.0879 ([M + Na]⁺), 594.0859 ([M + Na]⁺).

Results and discussion

Library synthesis of zwitterionic materials

Using α-benzylated sulfonates as anions, we aimed at exploring their potential to participate in intermolecular π–π and hydrophobic interactions among ZMs; that is, for a ZM-in-H₂O system exhibiting an LCST phase transition, a homogeneous solution appeared at low temperatures (due to strong and favorable interactions such as hydrogen bonding between the sulfonyl groups in the ZM and water), and, upon heating above its *T_c*, two immiscible phases were formed, likely resulting from the rupture of the aforementioned hydrogen bonding and the enhancement of π–π interactions between ZMs. Accordingly, we synthesized a small-molecule ZM library with the aim to facilitate the discovery of LCST-type zwitterionic materials.

Although there are two examples of substituted sulfobetaines incorporated in zwitterionic polymers in the literature,^{11,12} small-molecule sulfobetaine ZMs featuring embedded functional substituents carrying thermoresponsive properties remain totally unexplored. Sulfobetaines can be readily prepared from tertiary amine reactions with inexpensive 1,3-propanedisulfone or 1,4-butanedisulfone. Sulfones and their derivatives are useful scaffolds of sulfonyl drugs in medicinal chemistry research and valuable heterocyclic intermediates in natural product synthesis.¹³ In this work, we describe a new strategy to synthesize sulfobetaine-based ZMs with various benzyl groups attached directly on the sulfone moiety. Fig. 2 illustrates our library synthesis of **ZM 1a–e**, **ZM 2a–e**, and **ZM 3a–e**, where the key element **2a–c** could be structurally engineered by initially metalating 1,4-butanedisulfone **1** using butyllithium,¹⁴ and in the same pot, mixing with the corresponding substituted benzyl bromides (benzyl bromide for **2a**,

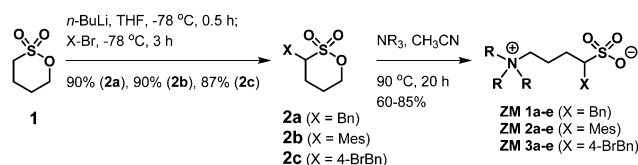


Fig. 2 Synthesis of **ZM 1a–e**, **ZM 2a–e**, and **ZM 3a–e**.



mesityl bromide for **2b**, and 4-bromobenzyl bromide for **2c**), resulting in the formation of α -alkylated sultones **2a–c**, followed by their ring opening with trialkylamines under heating conditions to finally afford the desired ZMs with 54–74% overall isolated yields in 2 steps. These syntheses proved straightforward on a multigram scale. The detailed experimental procedures, ^1H and ^{13}C NMR spectra, and high-resolution mass spectrometry (HRMS) data of all the ZMs (**ZM 1a–e**, **ZM 2a–e**, and **ZM 3a–e**) are summarized in the (ESI†).

Discovery of small-molecule zwitterionic materials exhibiting LCST thermoresponsive property

Our structural design of sulfobetaine-based ZMs was based on the chemical construction of a relatively hydrophilic, short-chain tetraalkylammonium cation with a substantially hydrophilic sulfonate anion carrying an engineered hydrophobic benzyl moiety. These benzylated sulfonates in ZMs should offer significant influence for phase separation with water, and structurally tune the ZMs to fine-balance their overall temperature-responsive property. Ultimately, the rational tuning of these ZMs enabled the discovery of thermoresponsive ZMs.

Fig. 3 shows a small library of 15 zwitterionic sulfobetaine salts (**ZM 1a–e**, **ZM 2a–e**, and **ZM 3a–e**) and their phase behaviors toward temperature changes in water. To discover ZMs carrying thermoresponsive property, each zwitterionic salt was mixed with water in a mass ratio of 1 : 3 (w/w), and then the mixture was placed in a water bath (4 °C) followed by gradual heating until it reached 90 °C. The phase transition temperature (T_c) for LCST was experimentally determined at a temperature point when the solution turned rigid during heating, as observed by the naked eye. To facilitate this determination, the organic dye Coomassie blue (0.006 wt% in water) was used to make the phase separation visible. Pleasantly, in the library of

15 ZMs investigated, two zwitterions **ZM 2c** and **ZM 3c** (labeled in green) were found to exhibit the LCST property with a T_c value of 84 °C and 60 °C, respectively. The library screening results for the ZMs labeled in red and blue indicate the formation of a homogeneous one-phase solution and a heterogeneous two-phase mixture between 0 °C and 90 °C, respectively (Fig. 3). In the case of the thermoresponsive ZMs discovered, because of the higher hydrophobicity on the benzyl group ($\log P$: Bn < Mes < 4-BrBn; computed by Xlog P 3.3.0),¹⁵ the structurally more hydrophobic **ZM 3c** (T_c = 60 °C) exhibited a lower T_c value than that of **ZM 2c** (T_c = 84 °C), as expected (Fig. 3). Furthermore, it has been reported that the phase property of a thermoresponsive material is a fine balance between the hydrophobicity and hydrophilicity of the ionic salts investigated,^{3,16} which is consistent with our results, as shown in Fig. 3, where both **ZM 2c** and **ZM 3c** (green) were identified to reside on the rim between being totally hydrophilic (red) and totally hydrophobic (blue).

Structural optimization of zwitterionic sulfobetaine materials

Because the T_c values of **ZM 2c** and **ZM 3c** obtained were not close to the body temperature (37 °C), both ZMs are less attractive for the direct analysis of biomolecules. Accordingly, we predicted that further structural optimization by incorporating two substituents on the benzyl group, such as 3,5-dibromobenzyl, carrying greater hydrophobicity should, in principle, result in a lower T_c value in water (Fig. 4). Accordingly, we employed α -3,5-dibromobenzylated 1,4-butanedisulfone (**2d**) and proceeded to synthesize **ZM 4c** from tributylamine and **2d** with the aim to discover new thermoresponsive ZMs carrying a lower T_c value. This newly engineered ZM was successfully prepared using the same synthetic route as depicted in Fig. 2. Finally, **ZM 4c** was afforded with 39% overall isolated yield in 2 steps (see Experimental section). The detailed experimental procedures, ^1H and ^{13}C NMR spectra, and HRMS data of **ZM 4c** are summarized in the Experimental section and ESI†.

We were very pleased to discover that **ZM 4c** indeed exhibited LCST thermoresponsive property, carrying a lower T_c value (38 °C) (Fig. 5). This structural modification of the ZMs (*i.e.*, from 4-BrBn in **ZM 3c** to 3,5-diBrBn in **ZM 4c**) and the successful engineering of **ZM 4c** clearly demonstrated the true value of the fine tunability of sulfobetaine zwitterion structures. The T_c value of this **ZM 4c** is at body temperature, and thus it was employed as our candidate ZM for the analysis of biomolecules.

Fig. 6A shows that **ZM 4c** forms a one-phase homogeneous solution with water at ambient temperature (25 °C), and upon heating above 38 °C, a two-phase system started to develop with

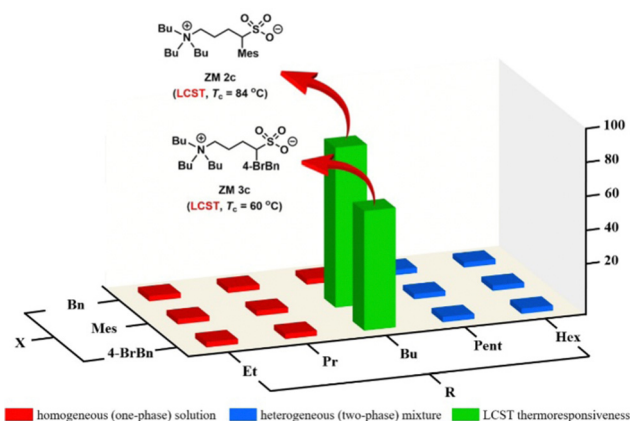


Fig. 3 Thermoresponsive screening of a small library of fifteen zwitterionic sulfobetaine salts (**ZM 1a–e**, **ZM 2a–e**, and **ZM 3a–e**) upon mixing with water (1 : 3, w/w) at temperatures between 4 °C and 90 °C. Phase transition results shown in red and blue indicate an entirely homogeneous one-phase solution and a heterogeneous two-phase mixture, respectively, between 4 °C and 90 °C. In the library, two ZMs (labeled in green) show LCST phase transitions: **ZM 2c** (T_c = 84 °C) and **ZM 3c** (T_c = 60 °C).

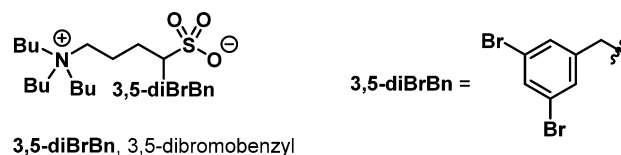


Fig. 4 Structure of zwitterionic sulfobetaine salt **ZM 4c**.



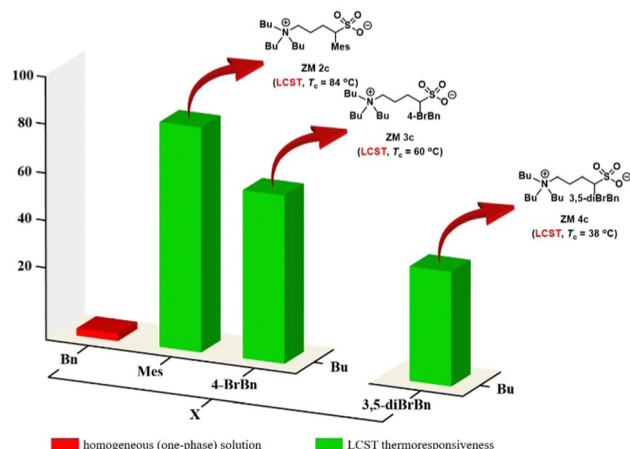


Fig. 5 Phase transitions of **ZM 1c**, **ZM 2c**, **ZM 3c** and **ZM 4c** upon mixing with water (1 : 3, w/w) at temperatures between 4 °C and 90 °C. Additional **ZM 4c** showed LCST-phase transition with attractive $T_c = 38$ °C.

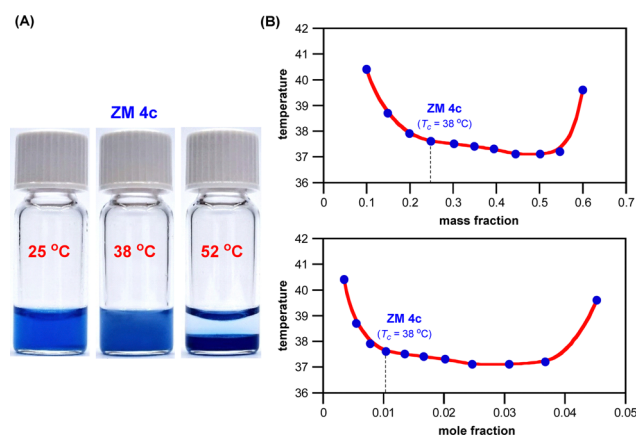


Fig. 6 (A) Temperature-dependent phase transitions of binary mixtures (3 : 1, w/w) of water with **ZM 4c** ($T_c = 38$ °C) at 25 °C, 38 °C and 52 °C, respectively. Coomassie blue dye (0.006 wt% in water) was added to make the phase separation more noticeable. (B) Phase diagrams (both T_c vs. mass fraction and T_c vs. mole fraction) of mixtures of water with **ZM 4c**. Solid red lines are used as the guide for the eye.

this ZM in the bottom layer, and ultimately a well separated two-layer solution mixture was formed at much higher temperatures (52 °C). The temperature-switchable mixing and demixing with water were entirely reversible, clearly illustrating that **ZM 4c** is a smart material exhibiting LCST phase transition with water.

For LCST systems, their phase transition temperature highly depends on the mass fraction and mole fraction of the thermoresponsive materials in water. Fig. 6B shows the phase diagrams of a mixture of water with **ZM 4c**, revealing the typical cup-shape concave curves with the lowest critical temperatures near its mass ratio of 1 : 1 and mole ratio of 1 : 32 (**ZM 4c**/H₂O).

This successful development of **ZM 4c** as an LCST-type thermoresponsive zwitterionic material clearly highlights the structural tunability of sulfobetaines, which is a fruitful ZM example using the combinatorial discovery platform developed in this work.

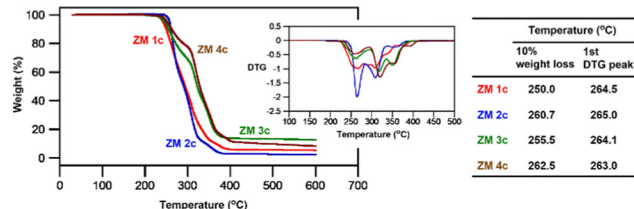


Fig. 7 Thermogravimetric analysis (TGA) of representative **ZM 1c**, **ZM 2c**, **ZM 3c**, and **ZM 4c** sulfobetaine materials measured under nitrogen at a heating rate of 20 °C min⁻¹. Inset: Derivative thermogravimetry (DTG) curves of **ZM 1c**, **ZM 2c**, **ZM 3c**, and **ZM 4c** with their first DTG peak at 265 °C, 265 °C, 264 °C, and 263 °C, respectively. Thermal analyses were carried out using a PerkinElmer STA 6000 thermogravimetric analyzer (PerkinElmer, Waltham, MA, USA).

Thermal stability of ZMs

Sulfobetaines are among most widely used ZMs, and many of them are studied and used at elevated temperatures (e.g., withstanding steam sterilization of medical ZM devices and oilfield antifouling ZMs).¹⁷ Accordingly, thermal stability is a prerequisite for the application of ZMs. Fig. 7 depicts the thermogravimetric behaviors and summarizes the key thermal stability data of the representative **ZM 1c**, **ZM 2c**, **ZM 3c**, and **ZM 4c**. The thermogravimetric analysis (TGA) indicates that, similarly for all ZMs, the major weight loss was observed after 250 °C and all the ZMs displayed multi-stage degradation processes with their first derivative thermogravimetry (DTG) peaks at 265 °C, 265 °C, 264 °C, and 263 °C (their decomposition temperatures T_d at 10% mass loss are 250 °C, 261 °C, 256 °C, and 263 °C) (table in Fig. 7), respectively. The results from the TGA thermograms (**ZM 3c** and **ZM 4c** vs. **ZM 1c**, **ZM 2c**) suggested that the bromo substituent enhanced the thermal stability of the ZMs. We also found that the greater the number of substituents on the benzyl group (**ZM 1c** vs. **ZM 2c**, **ZM 3c**, and **ZM 4c**; also, **ZM 3c** vs. **ZM 2c** and **ZM 4c**), the higher the degradation temperatures, resulting in more thermally stable ZMs. These thermal decomposition temperatures are similar to that of zwitterionic sulfonates reported in the literature.^{18–20}

A preliminary, proof-of-concept application

We developed novel zwitterionic materials considering their potential applications. Consequently, we successfully engineered α -benzylated sulfobetaines as thermoresponsive materials carrying LCST property in water, and then performed a preliminary study with the aim to prove that these ZMs are compatible with biomacromolecules for temperature-triggered phase separation; that is, the target biomolecules favorably partition in the ZM-rich layer, which will make future studies of biomolecular recognition and transformation possible.

Today, the separation and purification of biomolecules are routinely carried out using various chromatographic procedures, which are employed in almost all bioengineering processes. Therefore, it is necessary to design new technology platforms for concentrating dilute solutions or aqueous solutions containing only minute amounts of precious biomolecules. In this work, we explored the potential of ZMs for

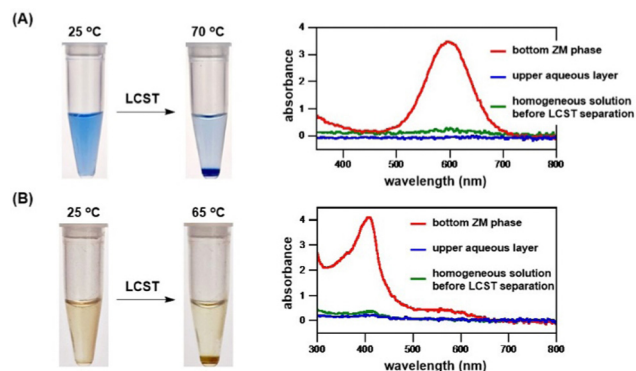


Fig. 8 Photos and spectra of temperature-dependent phase transition demonstrating biomolecule enrichment from its aqueous mixtures (1:20, w/w) of **ZM 4c** in (A) solution of Coomassie blue dye ($37.5 \mu\text{g mL}^{-1}$, $45 \mu\text{M}$) and (B) solution of human hemoglobin protein (1 mg mL^{-1} , $15 \mu\text{M}$). UV-vis spectra measured using a NanoDrop 2000 spectrophotometer (Thermo Scientific, Waltham, MA) for quantitatively determining the degree of enrichment in Coomassie blue dye (λ_{max} at 597 nm) and human hemoglobin protein (Soret band at 406 nm), respectively. In this case, only a minute amount of **ZM 4c** (4 mg in 80 μL water) was needed to achieve sample enrichment.

use as an immediate application to enrich dilute biomolecule solutions. Here, **ZM 4c** ($T_c = 38^\circ\text{C}$) was used as a candidate zwitterionic material, simply because it exhibits an LCST phase transition around body temperature in water. As a proof-of-concept application, we chose a high molecular-weight Coomassie blue dye (MW 826) and human hemoglobin protein (MW 64 500) for ease of visualization by the naked eye to test if its concentration enrichment of a highly diluted solution could be realized by **ZM 4c**. Pleasingly, as shown in Fig. 8, biomolecules were greatly enriched upon adding **ZM 4c** and mixing with dilute dye and protein solutions, respectively. Consequently, the Coomassie blue dye and hemoglobin protein were readily enriched by 16- and 11-fold, respectively. According to the spectra shown in Fig. 8A, the upper aqueous phases (blue spectra) were essentially free of the blue dye, that is, most, if not all the Coomassie dye was preferentially partitioned and concentrated in the ZM-rich bottom phase. This **ZM 4c** also readily enriched human hemoglobin protein (Fig. 8B). In our hands, no precipitation of hemoglobin was observed (e.g., its Soret band was preserved) during temperature-switchable mixing and demixing.

Conclusion

Ohno and co-workers previously reported the 3-step synthesis of a novel 3-(*N,N*-dihexyl-*N*-pentylammonio)propane-1-sulfonate zwitterionic salt ($\text{N}_{665}\text{C3S}$), which exhibited an intriguing LCST thermoresponsive property in water.²⁰ We reported in this work the 2-step synthesis of a small library of 16 ZMs, discovered 3 ZMs (**ZM 2c**, **ZM 3c**, and **ZM 4c**) carrying good thermal stability and LCST phase separation from water, and underlined the important role of ZMs for use in preliminary biomolecule analysis. Also, we demonstrated the successful structure optimization of **ZM 4c** to highlight its close

relationship with thermoresponsiveness. Recently, our group also reported *N*-triflimide (NTf)-based zwitterionic ionic liquids (ZILs) that exhibit LCST phase transitions in water.²¹ To the best of our knowledge, this is the first report on α -benzylated sulfobetaine ZMs as small-molecule smart materials exhibiting LCST property and that their miscibility/immiscibility with water triggered by temperature are entirely switchable and reversible. However, although our combinatorial approach for the discovery of smart materials is highly promising and effective, it is still being developed and more successful ZM examples are needed.

Author contributions

Yen-Ho Chu: conceived the idea, funding acquisition, designed experiments. Pin-Hsuan Chen, Hsin-Heng Huang: performed experiments. Yen-Ho Chu, Pin-Hsuan Chen: Interpreted the data. Yen-Ho Chu: wrote the manuscript and all authors discussed and approved the paper.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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