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Recent advances in the development and applications of nonconventional luminescent polymers

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Traditional luminescent polymers typically contain extended π - π conjugated structures that suffer from associated aggregation-caused quenching (ACQ) and high cytotoxicity issues. Therefore, there is a huge demand for stand-alone luminescent polymers that can circumvent these drawbacks and can be employed in diverse applications. In view of that, nonconventional luminescent polymers (NLPs) are being developed and studied extensively for their distinctive characteristics and potential applications in wide-ranging areas, and for an in-depth understanding of photophysical processes in science. These polymers do not contain classical chromophores, but often involve isolated benzene rings or only electron-rich moieties (e.g. amine, C=O, -OH, ether, imide, and heteroatoms N, O, P, S, etc.). This review article highlights the recent progress in the field of NLPs, including their synthesis, underlying emission mechanisms, and applications in materials and biological sciences. Besides, it is identified to be the foremost review that extensively emphasizes on both the fluorescence and phosphorescence features of NLPs. Finally, some prospects have been suggested for further exploration in this flourishing field of research.

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1. Introduction

Conventional luminescent polymers are classified as polymers integrated with large π -aromatic building blocks. On account of their strong emission properties, these materials have been the center of great attention for a variety of applications in organic lasers,^{1,2} organic light-emitting diodes (OLEDs),^{3,4} solar cells,⁵ sensors,^{6,7} bioimaging,⁸ etc. Despite their wide utility, complex synthesis and toxicity of large conjugated systems, poor miscibility in water medium and low resistance to photobleaching restrict further improvement and applications of such luminogens. Additionally, they are eminently luminescent in dilute solutions but exhibit weaker/no emission in their concentrated or aggregated state due to the aggregation-caused quenching (ACQ) effect.⁹ The aforesaid bottlenecks have prompted researchers to contrive a new class of photoluminescent materials devoid of large π - π conjugated structures, referred to as nonconventional luminescent polymers (NLPs).¹⁰ This new category of luminescent polymers often consists of isolated benzene rings or involves different

subgroups (-OH, -NH₂, C=C, and C≡N) and electron-rich heteroatoms (N, O, P, and S). Unlike traditional luminogens, their emission phenomenon shows certain unique characteristics including the following: they (i) do not require any excitation of delocalized electrons and (ii) act as collective molecular entities rather than a single or independent unit leading to substantial emission in their concentrated/aggregated states.¹¹ Moreover, the advantages of these photoluminescent polymers over conventional ones are easy preparation, low cost, environmental friendliness, good processability, excellent biocompatibility, and low toxicity or even complete non-toxicity, thereby enabling them to be molded into potential materials for desired applications.^{12,13} Because of such intriguing properties, hitherto diverse luminogens have been demonstrated aiming at the identification of emission mechanisms, and unfortunately, it remains disputed. Clustering-triggered emission (CTE) and/or aggregation-induced emission (AIE) have been proposed to rationalize the emission properties of these nonconventional luminogens. Nevertheless, the emergence of such materials in the areas of luminescent polymers has led to immense leap-frogging from the perspective of applications based on their inexplicable luminescence properties. Eventually, the past decade realization of rational synthetic strategies have offered numerous nonconventional luminescent polymers and actuated their implementation in a myriad of applications namely light-emitting diodes (LEDs), light-harvesting systems, chemical sensing, drug delivery, intracellular temperature measurements, cell imaging, ink-free anti-counterfeiting printing, data encryp-

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Fig. 2 (A) BSA structure with nonaromatic (purple and yellow) and aromatic (green, cyan and blue) residues. (B) Photographic images of the solid powder and tablet of BSA taken under 312 nm UV excitation in air or under vacuum. Reprinted with permission from ref. 22. Copyright (2019), WILEY-VCH.

pure organic luminophores, amorphous SA solids possess the p-RTP feature. Cross-linking with Ca^{2+} boosted the film emission because of the existence of strong chain-chain association and junction zone formation between G residues and Ca^{2+} . Rheological measurements proved that the entanglement of chains and their overlap in concentrated counterparts are invariant with the enhanced emissions, thus certifying the CTE mechanism. Moreover, SA was utilized for endosome specific intracellular imaging of HeLa cells, which has a great significance in biomedical research and clinical diagnostic applications.¹⁸

Amino acids are the constituent components of peptides and proteins.¹⁹ Earlier it was generally believed that protein emission occurred primarily because of three aromatic amino acids, namely phenylalanine, tyrosine, and tryptophan. But, Ye *et al.* showed that oligopeptides and polypeptides based on nonaromatic amino acids emitted fluorescence in the solid state.²⁰ It was observed that those oligopeptides adopted ordered conformations such as β -sheets, α -helices and turns stabilized through inter/intramolecular hydrogen bonds. It was speculated that these abundant hydrogen bonds bring the highly polarized amide group into close proximity, leading to the through-space interaction of amide groups' electrons resembling those of the aromatic structure based on the through-space π electron interaction. In 2018, Zhang and co-workers found that peptides could not only emit but also their basic building blocks aliphatic amino acids could generate noticeable visible light emission in concentrated solutions, and particularly in the solid state.²¹ In addition to prompt fluorescence, persistent RTP was observed in powders of poly (amino acids) of ϵ -poly-L-lysine (ϵ -PLL) after ceasing UV illumination. In concentrated solutions, amino acid molecules may form diverse nanoaggregates approaching each other with the aid of intermolecular hydrogen bonding interactions; however, in dilute solutions they exist as molecularly dissolved individuals. Furthermore, NH_2 , $\text{C}=\text{O}$, and OH subunits afforded through space electronic communications among different π and n (lone pair) electrons, resulting in electron cloud overlap and thus extended electron delocalization together with simul-

taneously rigidified conformations. These clustered chromospheres are responsible for the unexpected emission.

Recently, Yuan and coworkers reinvestigated the photoluminescence of natural protein taking BSA as the model protein under different conditions (Fig. 2A). Like polysaccharide SA, BSA showed AIE and concentration enhanced emission characteristics. Intriguingly, BSA exhibited distinct p-RTP in the form of amorphous powder or a tablet form under vacuum even under ambient conditions (Fig. 2B). This fact may be ascribed to the conformation rigidification caused by the cluster formation among the peptide backbone and pendant groups, which resulted in through-space electronic communications. Some other natural proteins, notably human serum albumin (HAS) and ovalbumin (OVA), showed similar emission characteristics to BSA.²²

2.2. Synthetic macromolecules

2.2.1. Poly(amidoamine). Poly(amidoamine) (PAMAM) dendrimers consist of amido, amine, and alkyl groups. Although the first starburst-dendritic PAMAM was synthesized by Tomalia *et al.* in 1985, the serendipitous intrinsic fluorescence from PAMAM was noted by Larson and Tucker in 2001.²³ They have examined the intrinsic luminescence of carboxylate-terminated PAMAM dendrimers utilizing two different techniques: excitation-emission matrices (EEMs) and lifetimes. This unanticipated fluorescence was assumed plausibly because of an $n-\pi^*$ electronic transition from the amido groups throughout the dendritic structure. The overall relative fluorescence intensity was found to be enhanced on increasing dendrimer generation. Again, lifetimes were also susceptible to the dendritic size and lengthened with generation, most likely due to an increase in the number of emissive species and structural rigidity (decrease in interior mobility) of the dendritic interior. PAMAM dendrimers and hyperbranched PAMAM (HPAMAM) are the foremost examples and most investigated NLPs comprising nonconventional chromophores. Notably, there is continued curiosity in this theme without interruption to the present.

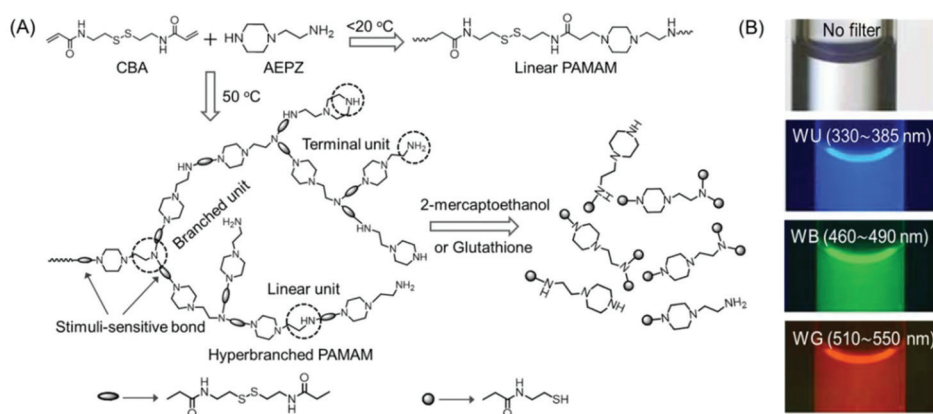


Fig. 3 (A) Synthetic strategy adopted for the preparation of linear and hyperbranched PAMAMs, and stimuli-induced degradation of the hyperbranched analog. (B) Fluorescence microscopy images of aqueous solutions of hyperbranched PAMAMs in a capillary acquired with 40x magnification. Reprinted with permission from ref. 24. Copyright (2009), WILEY-VCH.

In 2009, Pan's research group synthesized a novel disulfide-functionalized biodegradable HPAMAM by the Michael addition polymerization of *N,N'*-cystaminebisacrylamide (CBA) and 1-(2-aminoethyl) piperazine (AEPZ) (Fig. 3A). The resultant HPAMAMs displayed an absorption band at 360 nm in the UV-Vis spectrum and excitation wavelength-dependent bright wide-band fluorescence (Fig. 3B). In contrast, an aqueous solution of the linear PAMAMs did not display the UV-Vis absorption band, and only a weak emission was found. It was believed that the tertiary nitrogen with lone pair electrons inside the hyperbranched structure acted as the emissive species. This was verified by converting the tertiary amine into a quaternary ammonium group upon the reaction of the polymers with CH_3I . The absorption band at 360 nm disappeared almost completely, and a drastic decrease in the photoluminescence intensity at around 460 nm was observed in the case of quaternized HPAMAM. Such disulfide-functionalized polymers were easily cleaved under mild conditions by several stimulus reagents, such as 2-mercaptoethanol or glutathione *via* disulfide/thiol chemistry.²⁴ Chu *et al.* demonstrated the use of intrinsically fluorescent PAMAM dendrimers as an efficient nonviral gene carrier that could perform multiple functions such as bioimaging, delivery and transfection at the same time. To this end, NH_2 -terminated generation 4 PAMAM dendrimers were readily treated with an oxidizing agent to yield blue fluorescence with a moderate quantum yield (QY). Three negatively charged antisense oligonucleotides NGF2, NGF1 and (AS-ODN) p75 were electrostatically associated with the positively charged PAMAM dendrimer to produce fluorescent complexes at various nitrogen-to-phosphorus (N/P) ratios. Similar to the hyperbranched polyethyleneimine-mediated gene transfection, these complexes were also found to be capable of knocking down the specific protein expression to a certain level, confirmed by western blot analysis. The intrinsic fluorescence properties of PAMAM dendrimers enabled direct analysis of cellular internalization behavior by fluorescence microscopy and flow cytometry.²⁵

In another report, biodegradable less toxic β -cyclodextrin (β -CD) was incorporated into HPAMAM by a simple and efficient one-step Michael addition copolymerization reaction to improve biocompatibility and enhance photoluminescence. The amplified fluorescence of the resultant β -CD containing HPAMAM (HPAMAM-CD) was assumed to be generated from the restricted rotational motion of terminal chains due to the presence of large β -CD units, which suppressed collisional relaxation and a self-quenching phenomenon. The HPAMAM-CD could condense plasmid DNA (pDNA) very efficiently. The intrinsic luminescence of HPAMAM-CD aided in the study of the cellular uptake and gene transfection processes of the complexes through confocal laser scanning microscopy (CLSM) and flow cytometry, without any additional fluorescence labeling.²⁶

Wang and coworkers developed a novel conventional chromophore-free photoluminescent supramolecular hyperbranched polymer (SHP) *via* inclusion complexation between diethylenetriamine (guest) and α -cyclodextrin (host). The SHP displayed excitation wavelength-dependent wide-band fluorescence. The inclusion complex prevented the chain movement and increased the rigidity of the SHP, which eventually suppressed the non-radiative decay and amplified the fluorescence.²⁷

A series of photoluminescent HPAMAMs with dual thermal and pH sensitivity, and reducible degradability were successfully prepared by the Michael addition mediated copolymerization of tris(2-aminoethyl) amine and two bisacrylamide monomers CBA and *N,N'*-hexamethylenebisacrylamide, and the subsequent modification with isobutyric anhydride at room temperature. The fluorescence intensity was enhanced by lowering the pH and oxidizing the polymer by air. The tertiary amine oxide was assumed to be the fluorescent species.²⁸

Shi and coworkers reported the synthesis of a label-free green fluorescent and biocompatible PAMAM dendrimer derivative *via* surface modification of pristine PAMAM with acetaldehyde for traceable and controlled drug delivery.

Further PEGylation enables the fluorescent PAMAM to demonstrate excellent intracellular tracking in human melanoma SKMEL28 cells. The encapsulation of a model anticancer drug, doxorubicin (DOX), into the new nanocarriers helps this nanocomposite achieve higher drug loading capability and better therapeutic efficiency compared to the hydrophobic DOX and the original DOX loaded PAMAM nanocomposite.²⁹

Hao's research group unveiled HPAMAMs as novel mechano-responsive fluorescent polymers where the luminescence intensity of HPAMAM films increased upon stretching the sample and the fluorescence intensity quickly returned to the nearly original value automatically once the strain was relaxed. HPAMAMs exhibited sensitivity for the entire strain range from 0 to 250% or even up to 350%. Besides, they showed good recyclability in terms of fluorescence responses.³⁰

2.2.2. Poly(amino esters). Poly(amino esters) are another class of NLPs lacking any traditional chromophore groups in their structure but rich in aliphatic tertiary amino groups. In 2005, Liu and coworkers reported the synthesis of a family of hyperbranched poly(amino esters) (HPAE) with a variety of terminal groups by a $2A_2 + BB'B''$ strategy *via* Michael addition reaction utilizing 2-aminoethyl piperazine and 1,4-butane diacrylate (BDA) which were protected with hydroxyl (OH), amino and vinyl groups (Fig. 4). All the hyperbranched polymers exhibited blue luminescence in water, much like perfect PAMAM dendrimers, suggesting that amidoamine interior compositions and perfect dendrimer architectures were not essential for exhibition of NTIL emissive behaviour.³¹

The first report of degradable dual pH- and temperature-responsive dendrimers displaying photoluminescence was provided by Shen *et al.* This dual-stimuli responsive dendrimeric polymer was proved to be a perfect drug-delivery carrier. DOX was incorporated into the dendrimers at low temperatures without using organic solvents. The loaded DOX release was very slow under physiological conditions, but very fast at acidic pH, such as the lysosomal pH (pH 4–5) which caused quenching of the autofluorescence of the dendrimer. Moreover, the

fluorescence intensity decreased linearly when the DOX content was increased from the range of 1–16 wt%. Hence, the encapsulation and release of DOX can be monitored from the dendrimer's autofluorescence activity.³²

Yan *et al.* reported melt polycondensation of triethanolamine and itaconic anhydride for facile preparation of water-soluble poly(amino ester) polyols. The synthesized polymers displayed unexpected bright blue emission which originated due to the clusters of carbonyl (C=O) and oxidized tertiary amine functionalities. These fluorescent polymers have been employed for consecutive detection of unlabeled Fe^{3+} ions and L-cysteine *via* “on-off-on” fluorescence switching.³³

An unprecedented multicolor photoluminescent HPAE containing non-conventional chromophores was synthesized *via* one-step $A_2 + B_3$ melt polycondensation using *N*-methyldiethanolamine and citric acid. The as-synthesized polymers exhibited a full-spectrum of bright photoluminescence including blue, cyan, green and red fluorescence under different excitation wavelengths. This unique emissive characteristic of a non-conjugated polymer is caused by the heterogeneity and wide size distribution of the polymer aggregates. Besides, its fluorescence was very sensitive to Fe^{3+} ions.³⁴

Recently, another HPAE has been reported from the polycondensation reaction between maleic acid and triethanolamine by Yan *et al.* The as-prepared HPAE did not display multicolor fluorescence; rather it emitted only blue light on excitation with different wavelengths, which was assigned to the homogeneous electron delocalization. Intriguingly, the HPAE exhibited a multi-stimuli-responsive photoluminescence feature against solvent, pH, temperature and Fe^{3+} , enabling it to act as a multifunctional macromolecular probe towards the sensing of solvent polarity, temperature, pH and Fe^{3+} .³⁵

Water-soluble fluorescent hyperbranched poly(amido acids) with terminal $-CO_2H$ functionalities were synthesized using straightforward self-condensation of *N*-(3-aminopropyl) diethanosuccinate amine (AB_2 monomer). The amphiphilic poly(amido acids) formed nanoscopic globular micelle structures with a size around 30–50 nm in water and showed strong blue emissions with a QY of 23%. Moreover, their photoluminescence intensity was highly influenced by solution pH.³⁶

2.2.3. Polyethyleneimines. Another family of polymers devoid of traditional chromophores with intrinsic photoluminescence behavior is polyethyleneimines. Stiriba and coworkers reported unprecedented intrinsic fluorescence properties of hyperbranched polyethyleneimines (HPEIs) and linear polyethyleneimines (LPEIs) without the presence of traditional luminophores. In both the polymer architectures, the emission intensity was found to be affected by the oxidation of the PEI's amine groups and specific chemical modification (*i.e.*, methylation) and substantially strengthened by lowering the polymer solution pH towards the more acidic region. In contrast to the HPEI, the less compact linear analog LPEI was shown to produce strong intrinsic emissive species with long excited lifetimes. This observation clearly ruled out the state-



Fig. 4 Synthetic strategy for the preparation of HPAE with diverse terminal functional groups: monohydroxy, diol and amine. Reproduced with permission from ref. 31. Copyright (2005), American Chemical Society.

ment that an NLP should possess a highly branched architecture.³⁷

A series of hyperbranched graft copolymers (HPEI-*g*-HPG) were synthesized through the 'grafting from' method. These copolymers contain HPEI as the core and hyperbranched polyglycerol (HPG) as a shell with different numbers of glycidol units. HPEI-*g*-HPGs emitted blue fluorescence centered at *ca.* 470 nm albeit it did not contain any traditional fluorophores. The fluorescence intensity was found to be enhanced by a combination of factors such as increasing the shell thickness, oxidation of the polymer by air at 95 °C, adding tetrahydrofuran (THF) into the solvent in an appropriate ratio, and lowering the pH of the solution below 4. It was believed that tertiary amine oxide acted as a primary fluorescence center.³⁸

A water-soluble amphiphilic polyethyleneimine-poly(D,L-lactide) (PEI-PDLLA) copolymer was synthesized which emitted intrinsic blue light in both the solid-state and concentrated solution under a 365 nm UV lamp (Fig. 5A). Further transformation of the amphiphilic copolymer into polymeric nanoprobe (PEI-PDLLA NPs) was attained by a facile nanoprecipitation method. Interestingly, the synthesized NPs exhibited concentration enhanced fluorescence properties and a significant excitation-dependent multi-wavelength emission characteristic (Fig. 5B). The *in vitro* and *in vivo* imaging techniques corroborated tumor-targeted imaging and lysosome-specific staining capability of the PEI-PDLLA NPs with good biocompatibility and excellent photostability (Fig. 5C).³⁹

Self-monitoring and triple-collaborative nanotheranostics (NPICS) was successfully constructed from the positively charged amphiphilic autofluorescent copolymer PEI-poly(lactide) (PEI-PLA) which can simultaneously load NIR dye IR825 and hydrophobic antiangiogenesis agent combretastatin A4

(CA4), and interact with the negatively charged heat shock protein 70 (HSP70) inhibitor (siRNA against HSP70). The NPICS was applied for dual-modal imaging-guided cancer anti-angiogenesis as well as gene silencing enhanced photothermal therapy. Importantly, the inherent luminescence of PEI-PLA assisted it in not only serving as the drug carrier but also self-monitoring of the biodistribution and tumor accumulation in a real-time manner by virtue of fluorescence microscopy imaging.⁴⁰

2.2.4. Polysiloxanes. Feng and coworkers synthesized several generations of siloxane-PAMAM (Si-PAMAM) dendrimers by alternating aza-Michael and amidation reactions. Strong blue photoluminescence was noted from the low generation of Si-PAMAM even without the addition of an extra oxidizing reagent. Interestingly, the fluorescence emission intensity of the Si-PAMAM was found to enhance rapidly with the increase of generation. In addition, the AIE behavior of the Si-PAMAM was observed in a water-methanol mixture. Such unusual emission was attributed to the aggregation of its C=O groups caused by N → Si coordination bonds.⁴¹ This emission intensity was found to decrease with increasing temperature and quench rapidly in the presence of Fe³⁺ and Hg²⁺ ions.⁴²

In this context, Yan *et al.* introduced two novel water-soluble hyperbranched polysiloxanes (WHPs) simultaneously carrying primary amine and -OH groups *via* melt polycondensation, that is, transesterification reaction between the ethoxyl groups of (3-aminopropyl)triethoxysilane and -OH groups of dihydric alcohols such as 2-methyl-1,3-propanediol (MPD) and neopentylglycol (NPG) using a one-step process under solvent-free and catalyst-free conditions. Both the WHPs exhibited bright blue luminescence. However, the emission intensity was much intense for the NPG moiety containing compound compared to that of the other, plausibly due to the more easily restricted intramolecular rotations of the former than that of the later. They showed that the terminal OH functionalities instead of primary amine groups in WHPs were mainly responsible for producing the blue-luminescent species. Moreover, the photoluminescence intensities of the two unhydrolyzed polymers were enhanced after hydrolysis.⁴³

The same group has also synthesized fluorescent hyperbranched polysiloxanes (HBPSi) comprising epoxy and -OH groups, and observed that the aggregation of oxygen-rich heteroatoms from both the -OH and epoxy groups in the polymers played a vital role in the formation of fluorescent species.⁴⁴ It is noteworthy that these hyperbranched polymers displayed reinforced photoluminescence in water after polyetherification by 20 wt% polyether amine. However, the fluorescence intensity of the modified polymer in water diminished on further increase of the polyether content from 20 to 40 wt%, owing to its increased dissolubility that obstructs the aggregation of the polymer and as a result an incompact texture was formed.⁴⁵

Using a similar synthetic strategy they constructed luminescent HBPSi, composed of plenty of -OH groups and non-conjugated C=C bonds. With increasing molecular weight and concentration of the polymer, the fluorescence intensity was

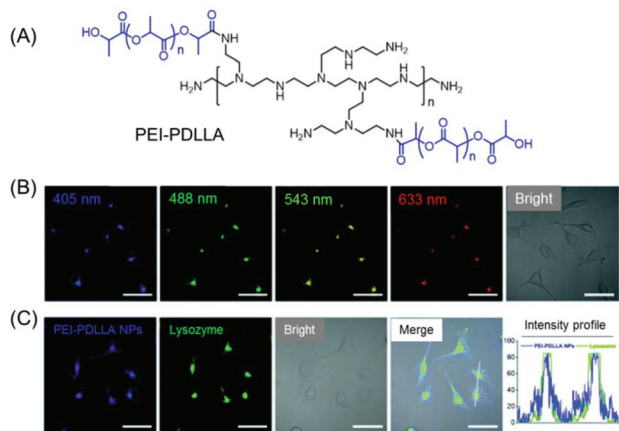


Fig. 5 (A) Molecular structure of an intrinsically fluorescent PEI-PDLLA polymer. (B) Excitation wavelength-dependent multiemission cellular imaging of MCF-7 cells incubated with PEI-PDLLA NPs. Scale bars = 50 μ m. (C) CLSM images of PEI-PDLLA (blue fluorescence) treated MCF-7 cells followed by staining with LysoTracker Red (green fluorescence). The intensity profiles indicated the polymer's ability to function as an alternative dye for lysosome imaging. Scale bar is 50 μ m. Reprinted with permission from ref. 39. Copyright (2019), Royal Society of Chemistry.

found to be dramatically strengthened, and even the brightest light emission was observed in the solid-state. The authors concluded that both the $-OH$ groups and $C=C$ bonds play a pivotal role in creating the emissive centers, and their persistent aggregation was responsible for the glow.⁴⁶

For the first time, carbosiloxane dendrimers bearing $C=C$ bonds were fabricated from a monomer containing two kinds of $C=C$ and a dithiol *via* orthogonal thiol-ene click chemistry. Generation 1 dendrimers showed a stronger blue glow than the monomer containing two types of $C=C$ bonds, whereas polydimethylsiloxane (PDMS) showed no glow at all under similar conditions. In comparison with the non-luminescent PDMS, the authors speculated that the introduction of sulfur is accountable for such unusual emission; however, they mentioned that more experimental data are required to understand the mechanism.⁴⁷ Again, OH -terminated HBPSi containing aliphatic tertiary amines have been reported to show blue fluorescence both in solution (water) and in the bulk state, and the emission intensities in water increased progressively with concentration. It was revealed that the $-OH$ group, rather than the aliphatic tertiary amine, is crucial for the formation of blue-luminescent species. Additionally, Fe^{3+} ions acted as a highly selective fluorescence quencher of the polymers.⁴⁸

Zhang and coworkers unprecedentedly constructed hydrophilic diamond-structured organosilica nanocrystals (*ca.* 2–6 nm) with finely tunable fluorescence (460–625 nm) from nonfluorescent precursors utilizing the classic Stöber strategy. The synthesized nanocrystals exhibited size-dependent fluorescence emission originating from the π - π interaction of orderly stacked vinyl groups throughout the particle.⁴⁹

A series of water-soluble intrinsically luminescent comb polysiloxanes containing various ratios of mercaptopropyl groups and polyether were synthesized *via* thiol-ene click reaction by Feng's research group. It was presumed that the unprecedented fluorescence is generated because of the formation of $S \rightarrow Si$ coordination bonds that facilitate the through-space charge transfer of the lone pair electron in sulfur to the 3d

empty orbital of silicon. However, in response to Hg^{2+} and Fe^{3+} , the $S-Si$ interaction was broken and eventually quenching of fluorescence occurred.⁵⁰

Novel imidazole functionalized polysiloxanes were synthesized for the consecutive detection of Fe^{3+} and cysteine by fluorescence quenching and a lighting mechanism. Here, imidazole groups and the whole polysiloxane structure contributed to the fluorescence. The present system could be useful for monitoring the Fe^{3+} /cysteine cycle *in situ* in living HeLa cells.⁵¹

The HBPSi or common unconventional luminescent polymers discussed above suffer from low fluorescence intensity and efficiency; in particular the QY is low in comparison with those of the traditional conjugated luminescent materials. Recently, Feng *et al.* showed how the fluorescence QYs of HBPSi could be improved. They synthesized HBPSi with both $C=O$ and vinyl groups, which exhibited a nontraditional intrinsic luminescence QY up to 43.9%, the highest value obtained to date among the reported silica-containing hyperbranched fluorescent polymers. To elucidate the fluorescence mechanism, the results were compared with those of two reference oligomers, which contain either $C=O$ groups or vinyl groups (Fig. 6A). The absolute fluorescence QYs for only $C=O$ groups and only vinyl groups bearing polymers have been reported to be 16.3% and 10.5%, respectively. Thus, the high fluorescence QY of the designed hyperbranched polymer was attributed to the synergism of vinyl and $C=O$ groups along with the $Si-O$ groups which facilitated the through-space conjugation (Fig. 6B).⁵²

Although the QYs of such nontraditional luminescent polysiloxanes substantially improved, their bioapplications are still associated with a major shortcoming *viz.* low water-solubility.

Accordingly, to enhance the fluorescence intensity and water solubility of HBPSi, β -CD, a rigid biodegradable molecule with plenty of hydroxyl groups was decorated onto the surface of the polymer architecture. The β -CD decorated HBPSi showed promise not only as a drug delivery vehicle that can

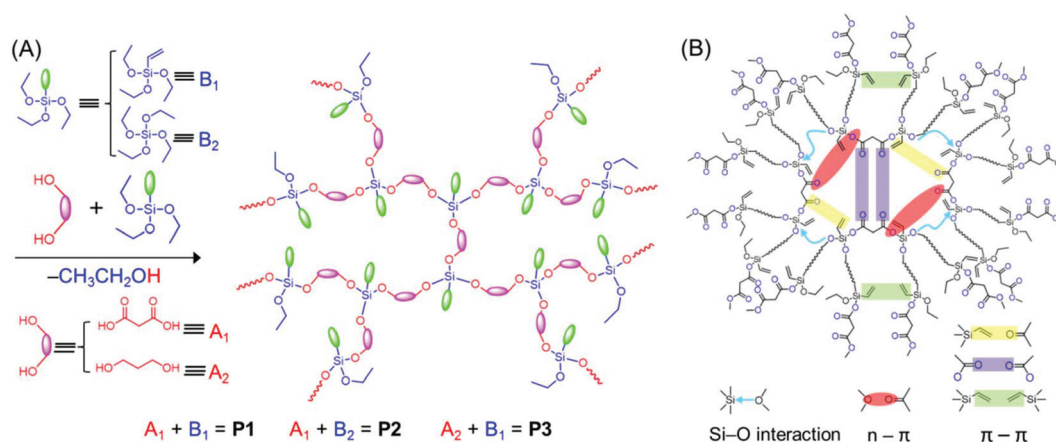


Fig. 6 (A) One-step synthetic method for the preparation of P1–P3. (B) Schematic diagram of different interactions between the functional groups, responsible for the bright-blue emission of P1. Reprinted with permission from ref. 52. Copyright (2019), American Chemical Society.

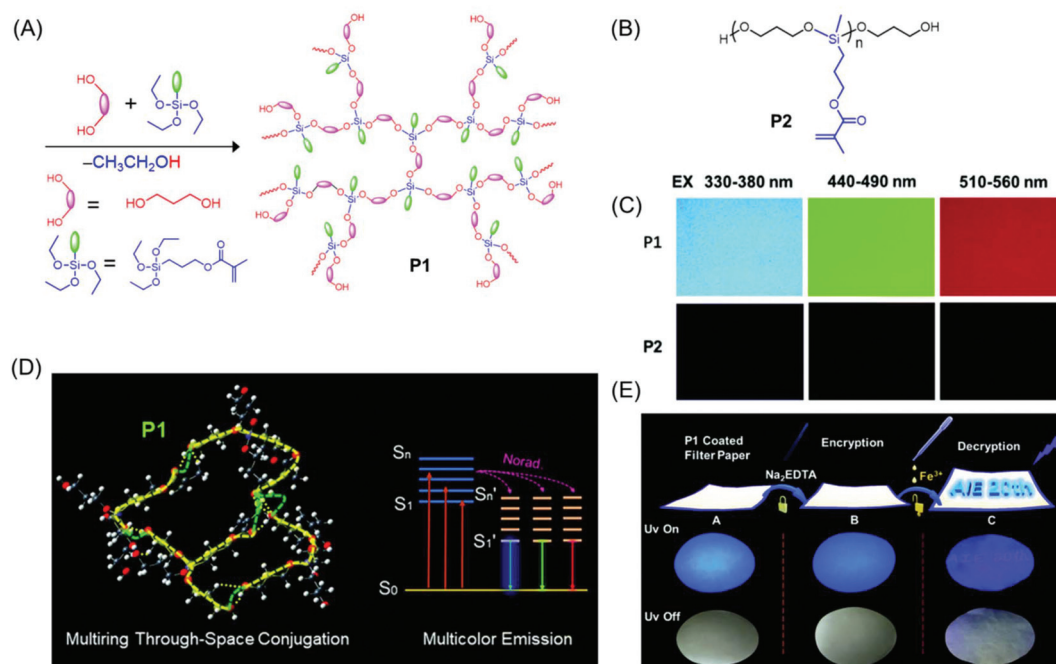


Fig. 7 (A) Synthetic route employed for the preparation of **P1**. (B) Chemical structure of **P2**. (C) Fluorescence microscopy images of hyperbranched (**P1**) and linear (**P2**) polysiloxanes acquired under different UV filters. (D) Occurrence of multiring through-space conjugation through a silicon-bridge in **P1** and schematic representation of its luminescence mechanism. (E) Chronological steps involved in the data encryption process with **P1** coated security paper and their photographs under both natural and UV light ($\lambda = 365$ nm). Reprinted with permission from ref. 54. Copyright (2020), Royal Society of Chemistry.

achieve high loading capacity and pH-responsive drug release performance, but also in simultaneous cell imaging.⁵³

Very recently, a novel multicolor emissive HBPSi with adjacent C=O and C=C groups was designed and developed by Tang *et al.* (Fig. 7A). To explore the multicolor emission mechanism, a reference linear polymer with the same chemical components was synthesized (Fig. 7B), and its photophysical properties were compared with a more compact hyperbranched one. The hyperbranched analog showed higher QY (7.71%) and luminescence lifetime (1.0 ns) compared to the linear one (QY = 1.12%, lifetime = 0.57 ns). Notably, the hyperbranched polymer displayed multicolor luminescence at different excitation wavelengths (Fig. 7C) in contrast to the linear polymer which emitted only weak blue fluorescence. An innovative luminescence mechanism namely “multiring-induced multicolor emission” was proposed based on the results obtained from density functional theory (DFT) calculations (Fig. 7D). Furthermore, on account of their quenching effect in response to Fe^{3+} , the application of the hyperbranched polymer in data encryption was accomplished by using disodium ethylenediaminetetraacetate dehydrate (Na_2EDTA) as the “ink” and iron(III) chloride (FeCl_3) as a “key” to decode the encrypted information on the security paper (Fig. 7E).⁵⁴

2.2.5. Polyacrylonitrile. Zhang and coworkers were the first to report the intrinsic emission from polyacrylonitrile (PAN), a semicrystalline non-conjugated polymer which contains a simple polyethylene backbone and directly attached cyano pen-

dants (Fig. 8A). PAN did not emit in dilute solutions, but became highly fluorescent when concentrated (Fig. 8A) or aggregated as solid powders, films or nanosuspensions, with quantum efficiency (ϕ) values up to 16.9%, revealing a distinct AIE phenomenon (Fig. 8B and C). Along with the prompt fluorescence response, PAN powders were capable of exhibiting delayed fluorescence (DF) and RTP generated from a long-lived triplet manifold, which is rarely observed in other unorthodox luminogens. Such unique emission of non-conjugated PAN was attributed to the clustering of nitrile groups in solid aggregates and condensed solutions that behaved as the exact chromophores and are liable for the observed light emission. In such clusters, the overlap of π and lone pair (n) electrons among nitrile groups extended the delocalization and simultaneously rigidified the conformation, thus causing the emission (Fig. 8D). Again, triplet emission is the result of the $n\text{-}\pi^*$ transition of lone pairs of electrons from the nitrogen atoms of nitrile groups.⁵⁵

Surface-grafted PAN brushes were synthesized by copper (Cu)-mediated photoinduced atom transfer radical polymerization (ATRP) using α -bromophenyl acetic acid-based initiator-modified silicon wafers. The AIE behavior of PAN was validated by making polymer brushes with various film thicknesses, and it depends on the molecular weight of the grafted chains and grafting density. It was observed that, as the film thickness increases the emission is more pronounced, which is attributed to the formation of larger fluorescent clusters.⁵⁶



Fig. 8 (A) Chemical structure of intrinsically emissive PAN. Photographs of PAN in only DMF and (B) a DCM/DMF mixture with various f_{DCM} obtained under a 365 nm UV lamp. (C) Luminescence images of (i) solid powders and (ii) a film of PAN on exposure to 365 nm UV excitation. (D) Schematic illustration of plausible intra- and inter-molecular interactions present within nitrile clusters in the aggregated state. (E) Photoluminescence image of SR6 ENRs taken under 365 nm UV irradiation. (F) Photographic images of ENRs at 77 K in UV lamp-on/lamp-off mode at different time courses. Figures A, B, C and D are reprinted with permission from ref. 55. Copyright (2016), John Wiley and Sons. Figures E and F are reprinted with permission from ref. 57. Copyright (2020), Royal Society of Chemistry.

In a recent work, the non-conjugated PAN was implemented for the fabrication of electrospun nanofiber ribbons (ENRs) by electrospinning and high heat stretching above the glass transition temperature (T_g). The resultant mesoscopically ordered ENRs were light-weight and found to exhibit tremendous mechanical strength and toughness. It is noteworthy that upon excitation at 340 nm, the ribbons showed polarized deep blue luminescence with anisotropy and QY values of 0.37 and 31%, respectively. Along with the prompt luminescence (Fig. 8E), delayed deep blue and green phosphorescence was also observed at room temperature (Fig. 8F). In a nutshell, the authors have exemplified a very fascinating material with the combination of the unique properties of emission, anisotropy and mechanical behavior.⁵⁷

2.2.6. Polyurethane. A series of nonaromatic polyurethanes (PUs) were outlined and developed by the polyaddition reaction between diisocyanates and simple diols. The as-synthesized PUs were nonfluorescent in dilute solutions, but turned out to be highly emissive in their aggregated state as films/powder or concentrated solutions, featuring AIE and concentration enhanced emission characteristics. Moreover, the polymer was used for PA detection and cellular imaging.⁵⁸

An unprecedented aggregation-induced long-persistent phosphorescence (>1 s) in a non-conjugated PU system at 77 K was reported for the first time by Bryce *et al.* It was proposed that the appearance of such unusual behavior is because of the formation of intra- and/or inter-molecular C=O clusters at low temperature that facilitates spatial electronic overlap between π electrons and lone pairs. Interestingly, the incorporation of conjugated aromatic units into the non-conjugated PU backbone has been shown to increase the lifetime of long-persistent phosphorescence, revealed that the intersystem crossing rate was enhanced through the inter- or intra-molecular $n-\pi^*$ transition from the electron-rich C=O groups to the con-

jugated aromatic units.⁵⁹ In the follow-up work, they prepared a series of fluorescent non-conjugated PU derivatives by incorporating classical π -aromatic chromophores, and applied them for highly selective and sensitive recognition of PA both in solution and in the solid state. The sensing mechanism was ascribed to the combined effect of Förster resonance energy transfer (FRET), photoinduced electron transfer (PET) and strong IFE.⁶⁰

Zhao and coworkers reported NLPs based on dynamic covalently cross-linked polyhydroxyurethane (PHU2), synthesized from amines and aliphatic multifunctional cyclic carbonates avoiding the use of toxic isocyanates (Fig. 9A). It was observed that the fluorescence intensity of PHU2 progressively became stronger without any sign of quenching on increasing the curing time from 0 to 8 h (Fig. 9B). It is noteworthy that it displayed long persistent phosphorescence under cryogenic conditions. In addition to the unique luminescence properties, the new NLP possessed excellent mechanical strength along with smart characteristics including shape memory (Fig. 9C) and self-healing (Fig. 9D and E). The efficient transcarbamoylation dynamic exchange reaction between carbamate and -OH groups was responsible for the good shape memory property and self-healing behavior of PHU2. Utilizing these alluring characteristics of the synthesized NLP, "light-assisted ink-free screen printing" was established to prepare anti-counterfeiting materials, which eliminated the use of expensive security inks.⁶¹

Very recently, a series of novel hyperbranched polyurethane (HPDU) elastomers have been developed by Zhang *et al.* using a simple polycondensation reaction among commercially available poly(tetramethylene) glycol, methylene-bis(4-cyclohexyl isocyanate) and diazolidinyl urea by following the $A_2 + B_4$ strategy (Fig. 9F). The resultant PUs were transparent (Fig. 9G) and showed super-tough, puncture-resistant (Fig. 9H) and photo-



Fig. 9 (A) Chemical structure of PHU2 and (B) its fluorescence images taken under daylight and UV excitation ($\lambda = 365$ nm) at various reaction time periods at 90 °C. (C) The shape memory properties of a PHU2 sheet and its self-healing behaviour monitored at 160 °C using (D) a polarizing microscope and (E) a SEM. (F) Chemical structure of HPDU. (G) Transparent photograph of a HPDU4 film. (H) Illustration of puncture resistant properties of the HPDU10 film. (I) Schematic view of a crack shaped as XJTU with bright-blue fluorescence on the film surface. (J) Photographs of crack XJTU on the HPDU4 film surface when exposed to UV excitation after being healed for different time intervals at 90 °C. Figures A, B, C, D and E are reprinted with permission from ref. 58. Copyright (2020), American Chemical Society. Figures F, G, H, I and J are reprinted with permission from ref. 59. Copyright (2020), American Chemical Society.

luminescence elastomeric behavior with crack self-diagnosis and healing tracking ability. Upon irradiation with 365 nm UV light, the fracture surface furnished strongest emission intensity as compared to the entire film, which was found to diminish gradually with self-healing. Consequently, the change in the emission intensity on the crack guides in the diagnosis of the crack location and concomitant tracking of the progress of the self-healing process (Fig. 9I and J).⁶²

Carbon dioxide (CO₂)-derived siloxane (Si–O–Si)-containing linear PHU was designed and synthesized and it possessed strong photoluminescence (QY = 23.6%), high photostability, and broad absorption and emission characteristics both in solution and bulk states. The hydrophobicity and flexibility of the siloxane linkages in PHU led to strong fluorescence through the significant aggregation of hydroxyurethane chromophores along with H-bonding interactions. To demonstrate a practical application, a low voltage cool white LED was fabricated with a color rendering index (CRI) of 83 by combining the polymer as a single phosphor with a commercially available UV chip.⁶³

Recently, a white LED with a high CRI of 95 has been reported from a non-conjugated PHU microsphere (PHUM) prepared *via* the crosslinking of 1,6-hexanediamine and trimethylolpropane tri(cyclic carbonate) ether. The high degree of crosslinking as well as various H-bonding structures prompted the construction of carbamate clusters with different sizes and broad distributions, enabling a wide-range emission spectrum (white light emission).⁶⁴ Another report used a similar synthetic strategy to prepare non-traditional fluorescent PHU. The collective results of the methylthiazolyldiphenyl-tetrazolium bromide (MTT), hemolysis and cytokine release assays for PHU certified its much lower cyto-

toxicity and immunotoxicity along with good blood compatibility with a <5% hemolysis rate in comparison with the traditional isocyanate-derived PU. This polymer was applied for fluorometric detection of Fe³⁺, and the LOD was calculated to be 4.56 μ M.⁶⁵

2.2.7. Polyesters and polycarbonates. Yan *et al.* reported the preparation of fully bio-based hyperbranched polyesters by adopting a one-pot A₂ + B₃ step-growth polymerization reaction between succinic or adipic acid and glycerol. Upon irradiation with a 365 nm UV lamp, a strong blue light emission from synthetic polyesters was observed. The fluorescence intensity strengthened with the increment of the molecular weights and solution concentrations of the polyesters. Notably, an absolute QY of 16.75% was achieved for the resulting polyesters. This unanticipated fluorescence behavior of the aliphatic hyperbranched polyesters could be attributed to the aggregation of C=O groups.⁶⁶ They also described the unprecedented strong blue photoluminescence characteristics of non-conjugated hyperbranched polycarbonates (HBPCs). The hyperbranched polycarbonates were synthesized *via* catalyst or solvent-free one-pot A₂ + B₃ transesterification reaction between diethyl carbonate and triol (glycerol⁶⁷ or trimethylolpropane⁶⁸). The resultant HBPC exhibited excitation wavelength-dependent and concentration-dependent emission behaviors in ethanol. The authors have envisioned that the intermolecular H-bond probably formed between the oxygen atom of the carbonate group and the unreacted –OH group suppressed the non-radiative decay processes through rigidifying the polymer conformation and thus furnished bright emission.

2.2.8. Polymers containing isolated benzene rings. Poly (ethylene terephthalate) (PET) comprises isolated, small aromatic systems with merely short-range electron delocalization.



Fig. 10 Overall emission properties of PET at different states. Reprinted with permission from ref. 69. Copyright (2018), American Chemical Society.

Despite PET emitting weak UV light (emission at 330 nm for individual terephthalate units) in dilute solutions, it became highly emissive in concentrated solutions or in the aggregated (solid) state (Fig. 10). Moreover, in solid films, PET displayed noticeable fluorescence and RTP dual emissions with crystallization-enhanced emission (CIDE) characteristics.⁶⁹

Li *et al.* demonstrated a new class of polymers based on a non-conjugated methacrylate backbone appended with a single switchable boron chromophore (Fig. 11A). Impressively, the homopolymers displayed rare multicolor emission in solution which was found to be dependent on the degree of polymerization (DP). Moreover, these homopolymers exhibit solvent and temperature-tunable fluorescence color changes (Fig. 11B and C).

The color-switching mechanism is associated with the structural transition from an open form (tricoordinate boron

chromophore, blue-emitting) to a closed-form (tetracoordinate boron chromophore, red-emitting). In a nutshell, the solvent, temperature and DP (molecular weight of the polymer) have immense effects on the open-to-closed form ratios of the attached pendant boron chromophore (Fig. 11D). Notably, a pure white light emission was attained in solution by finely adjusting the homopolymer molecular weight, and also in a thin-film/solid-state by copolymerizing the luminescent boron monomer with non-fluorescent monomers in an appropriate ratio.⁷⁰

Long-lived room temperature phosphorescence (LRTP) has emerged as an appealing optical feature in the area of organic photonics and electronics.⁷¹ Amorphous polymer poly(styrene sulfonic acid) (PSS) showed green phosphorescence in air (dry solid-state) with a maximum lifetime up to 1.22 s. The RTP lifetime was greatly influenced by the molecular weight of PSS and percentage of sulfonic acid groups in PSS incorporated through sulfonation reaction. It was presumed that the inter-/intra-chain H-bonding network amongst the pendant sulfonic acid groups suppressed non-radiative transition processes *via* restricting the motion of the chromophores, enabling ultra-long RTP lifetimes. However, exposure to water caused RTP quenching of solid-state PSS, and restoration occurred by removal of water. This reversible “off-on” RTP switching of PSS powder upon exposure and expulsion of water was realized in lifetime-encoding application simply by injecting water vapor within the PSS film using a mask.⁷²

Huang *et al.* reported the achievement of the longest lifetime (RTP lifetime is around 2.1 s) of organic phosphorescence in polymer luminogens by a facile ionic bonding cross-linking strategy. Instead of PSS, poly(sodium 4-styrenesulfonate)

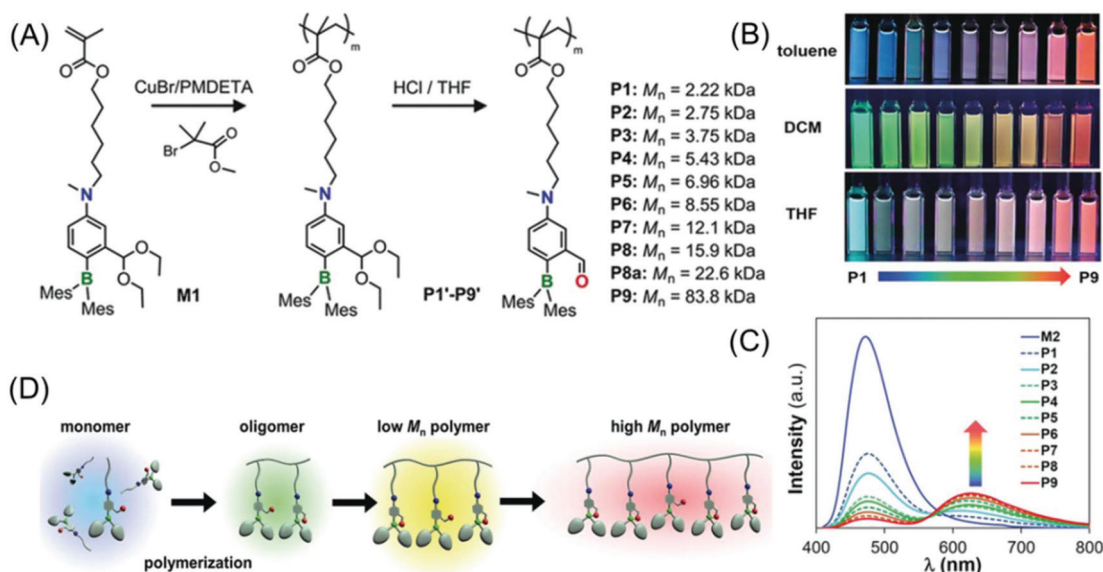


Fig. 11 (A) Synthesis of methacrylate-based polymers containing pendant switchable boron chromophores. (B) Photographs of P1–P9 showing multicolour emission in toluene, DCM and THF acquired under UV illumination of 365 nm. (C) Emission spectra of P1–P9 in toluene. (D) Schematic view of the DP-dependent boron chromophore structural change and fluorescent colors of the methacrylate polymers. Reprinted with permission from ref. 70. Copyright (2019), WILEY-VCH.

(PSSNa) as an ionic polymer model was taken to prove the hypothesis. Similar steady-state fluorescence and phosphorescence spectra were observed after the replacement of Na^+ with different cations (Li^+ , K^+ , Rb^+ and NH_4^+). The universality of the ionic interlocking for L RTP was verified by introducing the ionic bonding cross-linking into nonaromatic polymers. So a set of totally four ionic polymers, namely PSSNa, poly(acrylic acid sodium salt) (PAANA), poly(maleic acid sodium salt) (PMANA), and poly(maleic acid-co-4-styrenesulfonic acid sodium salt) (PSSNa-co-PMANA), were designed and synthesized. On chromophore variations, the L RTP emission colors were successfully transformed from yellow to blue. It is noteworthy that both aromatic and nonaromatic polymers exhibited ultralong phosphorescence on varying the excitation wavelength under ambient conditions.⁷³ The delay time-dependent RTP emission was first reported in the case of poly(4-styrenesulfonic acid-co-maleic acid) salt by Gong *et al.* The existence of the free volume that provided space for altering the conformation in order to relax the excited states was primarily responsible for the observed phenomenon.⁷⁴

On the other hand, amorphous purely organic phosphorescent polymer materials with color-tunable and long-lived emission by a concise chemical ionization strategy have been reported. Poly(4-vinylpyridine) (PVP) was chosen as an ionization model. Upon ionization with 1,4-butanedisulfone, the PVP-based ionization product poly(4-vinylpyridine)butane-1-sulfonate (PVP-S) (Fig. 12A) showed distinct phosphorescence behavior with an emission lifetime of 578.36 ms, which is 525 times longer than that of the PVP polymer under ambient conditions. Intriguingly, the phosphorescence emission color could be precisely tuned over the full-spectrum of the visible region from blue to red, by changing the excitation wavelength (Fig. 12B), whereas, the PVP polymer did not show any excitation-dependent phosphorescence at room temperature, even when cooled to 77 K. It was proposed that the strong ionic interactions between PVP and sulfonic acid help in generating ultralong phosphorescence in the PVP-S polymer by improving

the local rigidity and suppressing the non-radiative decay of the excited triplet state through the restriction of the vibration and rotation of the molecules (Fig. 12C).⁷⁵

The development of the foremost instance of the amorphous RTP polymer containing nonmetal dative bonds has been reported. In the copolymer of vinylphenylboronic acid and acrylamide, nonmetal dative bonds formed between boron and nitrogen/oxygen atoms enhanced the probability of an intersystem crossing (ISC) process to populate the triplet excitons; the charge transfer and subsequent inhibition of the motions of the phosphors resulted in an effective RTP lifetime in air (Fig. 12D). Additionally, the dynamic dative bonds and H-bonds endowed the material with self-healing and outstanding anti-counterfeiting capabilities.⁷⁶

Jin's research group recently reported non-conjugated RTP phosphoramidic acid oligomers consisting of a phosphoryl group having $\text{d}\pi\text{-p}\pi$ bonds and a diamino group as a H-bond donor with ultralong phosphorescence lifetimes and phosphorescence QY up to 776.9 ms and 4.6–10.5%, respectively. The system was easily synthesized by the polycondensation reaction of H_3PO_4 (phosphoric acid) and variable diamines including 1,3-bis(aminomethyl)cyclohexane (BAC), norbornane dimethylamine (ND) and *m*-xylylenediamine (XY). The ultralong phosphorescence lifetime was attributed to the H-bond trapping excited electrons, albeit it was not fully understood.⁷⁷

2.2.9. Maleimide and maleic anhydride containing polymers. Park's research group developed a novel ratiometric pH sensor with tricolor (green, yellow and orange) fluorescence based on a poly(*N*-phenylmaleimide) (PPMI)-containing block copolymer (Fig. 13A). In addition to the pH-tunable reversible color-switching behavior (Fig. 13B), the PPMI derivatives exhibited strong solvatochromism (Fig. 13C). Based on theoretical calculations they inferred that the keto–enol tautomerism in succinimides played an important role in the observed multicolor fluorescence, as the tautomeric equilibria are greatly influenced by the solution pH and solvent polarity (Fig. 13D).⁷⁸



Fig. 12 (A) Synthetic route to an amorphous PVP-S polymer. (B) Phosphorescence spectra of PVP-S acquired at 77 K at various excitation wavelengths. Inset shows the afterglow multicolor phosphorescence emission on excitation with a wide range of wavelengths from 280 to 500 nm at 77 K. (C) Schematic illustration of ionization-induced ultralong phosphorescence of PVP-S. Reprinted with permission from ref. 75. Copyright (2019), WILEY-VCH. (D) Graphical representation of the RTP polymer involving boron–nitrogen dative bonds and their fluorescence images under sunlight and UV “on–off” conditions. Reprinted with permission from ref. 76. Copyright (2020), American Chemical Society.



Fig. 13 (A) Chemical structure of poly(ethylene oxide-*b*-*N*-phenylmaleimide) (PEO-*b*-PPMI). (B) Ratiometric fluorescence changes and multiple emission colors of PEO-*b*-PPMI at different pH values in water on excitation at 445 nm. (C) Emission profiles ($\lambda_{\text{ex}} = 445$ nm) of PPMI in different solvents. (D) Tautomeric structures of PPMI, responsible for pH-tunable emission and solvatochromism phenomena. Reprinted with permission from ref. 78. Copyright (2015), American Chemical Society.

Tang and coworkers showed how a pure oxygenic non-conjugated polymer based on poly[(maleic anhydride)-*alt*-(vinyl acetate)] (PMV) could emit strong blue-light upon UV illumination (Fig. 14A). Moreover, its photophysical behavior could be manipulated by varying the solvent (Fig. 14B). Because of the rigid conformation of succinimide-based polymers, a heterodox cluster by the collection of many C=O groups is generated. It was believed that the clusters formed within the locked

C=O groups by interacting in a through-space manner generated the actual chromophore, leading to bright emission upon UV light irradiation. Again, the existence of the polymer/solvent complex is accountable for the solvatochromism.⁷⁹

Qiao and coworkers have demonstrated how the emission intensity/QY of PMV could be enhanced, and how the emission color could be altered from common blue to red. Hydrolysis and esterification reaction of PMV with a strongly alkaline sodium hydroxide solution afforded PMV-Na, which emitted high-intensity blue-green light in solution and solid-states. The calculated QYs of the PMV-Na solution and solid were found to be as high as 35% and 66%, respectively. By thermally treating at around 170 °C for 5 min, blue-emissive PMV-Na solids could be converted into a red emission agent. It was recommended that the construction of isolated C=C bonds is responsible for the unusual red emission from such non-conjugated PMV polymers (Fig. 14C and D).⁸⁰ In the subsequent year, a new kind of PMV derivative was synthesized. The new red-emissive PMVs were synthesized by utilizing large alcohol molecules such as diacetone alcohol produced from acetone under basic catalysis. Two types of C=O chromophores were present in the newly synthesized PMV derivatives: a blue light-emitting sodium carboxylate group and a red light-emitting bulky carboxylic ester group, and both exhibited excitation-dependent luminescence. Also, such non-conjugated photoluminescence polymers having dual chromophores showed AIE characteristics.⁸¹

Novel multicolored AIE-active photoluminescent polymers were developed by mixing the PMV solid with basic aqueous solutions of variable pH in suitable ratios at room temperature. The PMV/alkaline sodium hydroxide ratio played a key role in tuning the emission color from cyan to red.⁸² The mechanism behind the above phenomenon was not fully elucidated. However, it has been assumed that the addition of NaOH altered the clustering state of the carbonyl chromophores present in PMV.

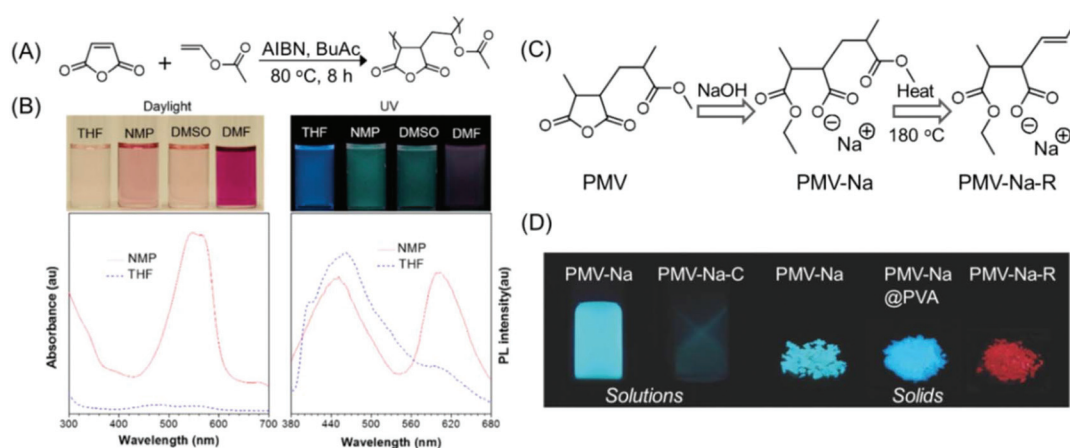


Fig. 14 (A) Synthesis of PMV. (B) Photographs of PMV solutions under daylight/365 nm UV illumination in different organic solvents and corresponding UV and fluorescence spectra ($\lambda_{\text{ex}} = 330$ nm) in NMP and THF. (C) The synthetic route to PMV-Na and PMV-Na-R. (D) Photoluminescence images of PMV-derived polymers in solution and in the solid state upon UV excitation at 365 nm. Figures A and B are reprinted with permission from ref. 79. Copyright (2015), American Chemical Society. Figures C and D are reprinted with permission from ref. 80. Copyright (2017), WILEY-VCH.

Wang's research group utilized a conventional free radical precipitation copolymerization technique to synthesize a series of poly(maleic anhydride-*alt*-vinyl pyrrolidone) (PMVP) copolymers with variable molecular weights. The resultant PMVPs emitted strong light in both DMSO solvent and solid-state, demonstrating their AIE characteristics. Besides, the PMVPs exhibited distinct excitation-dependent fluorescence (EDF) and molecular weight-dependent fluorescence (MWDF). More importantly, their emissions were significantly red-shifted in comparison with those of the homopolymers poly(maleic anhydride) (PMAh) and poly(*N*-vinyl-2-pyrrolidone). They suggested that the copolymerization of *N*-vinyl-2-pyrrolidone with MAh increased the flexibility of polymer chains and as a result the rigidity of the polymer chain was compromised, which facilitated the stronger and more extended intra- and/or inter-chain $n-\pi^*$ and $\pi-\pi^*$ interactions between the anhydride and pyrrolidone groups in the aggregation state, leading to the strong and red-shifted emission of PMVPs.⁸³

To prove the previous hypothesis that the polymer chain flexibility plays an imperative role in enhancing the photoluminescence of NLPs, they synthesized poly(itaconic anhydride) (PITA) and poly[(1-octene)-*co*-(itaconic anhydride)] (POITA) by using 1-octene and itaconic anhydride (ITA) as the comonomers.⁸⁴ The photophysical properties of these as-synthesized copolymers were compared with those of PMAh and poly[(maleic anhydride)-*alt*-(2,4,4-trimethyl-1-pentene)] (PMP) to elucidate the mechanism in terms of chain flexibility. In comparison with PITA, POITA exhibited red-shifted and stronger luminescence both in solution and in the solid state. The finding is completely different from the result of PMP, where the bulky 2,4,4-trimethyl-1-pentene (TMP) segregated the MAh moieties and thus obstructed their intra-chain interactions, leading to very weak emission. Thus, the increased chain flexibility enabled the functional groups to adopt proper conformations to brighten up the emissions.

The photoluminescence characteristics of poly(itaconic anhydride-*co*-vinyl caprolactam) (PIVC) and poly(itaconic anhydride-*co*-vinyl pyrrolidone) (PIVP) were recently highlighted keeping in mind that in some cases the chain flexibility is beneficial for boosting the fluorescence of NLPs.⁸⁵ ITA was chosen, as it has a cyclic acid anhydride ring that is attached to the polymeric structure *via* only one carbon atom; therefore these structures possess much higher flexibility in comparison with PMAh. It was found that the emissions of both the copolymers are significantly red-shifted (emission maxima in the orange-red region) in solutions. Interestingly, PIVP and PIVC displayed bright white and orange-red emission in the solid-state upon 365 nm UV illumination, respectively. Both the polymers emitted blue fluorescence at a low concentration in DMSO, but surprisingly the emissions changed to orange as the concentration was increased. It was speculated that two distinct chromophores were present causing the blue and orange emissions, and the number of chromophores corresponding to the orange emission increased with the increase of the polymer concentration.

The reversible addition-fragmentation chain transfer (RAFT) method was employed to prepare alternating copolymers based on the poly(styrene-*alt*-maleimide) skeleton by the sequence-defined copolymerization of *para*-methoxydiethylene glycol-substituted styrenes and *N*-(2-salicylaldehyde-aminoethyl)maleimides. The as-synthesized alternating copolymers exhibited a fast and highly sensitive "off-on" fluorometric response toward Zn^{2+} with a LOD value of 0.25 μM by inhibiting the PET process. Apart from this, a notable green fluorescence "off-on" switching (230-fold) was arrested while the pH changed from 11 to 12.⁸⁶

Polyhedral oligomeric silsesquioxane (POSS)-containing homopolymer poly(maleimide isobutyl POSS) and various alternating copolymers namely poly(styrene-*alt*-maleimide isobutyl POSS), poly(4-acetoxystyrene-*alt*-maleimide isobutyl POSS) and poly(4-hydroxystyrene-*alt*-maleimide isobutyl POSS) were synthesized through free radical polymerization and subsequent post-polymerization modification reactions. Although the POSS-containing polymer does not contain any common fluorescent moiety, it exhibited emission at high concentration and in the solid-state. Moreover, the POSS-appending homopolymer displayed stronger photoluminescence emission in the bulk state with a much higher ϕ compared to those of the POSS-integrated alternating copolymers, presumably due to the former's crystallinity and aggregation of locked C=O groups of five-membered rings.⁸⁷

Qiao and coworkers developed a new family of photoluminescent alternating copolymers by the copolymerization of MAh and diverse olefin derivatives. These fluorescent polymers were totally transparent and exhibited thermoplastic behavior. Additionally, the copolymers prepared from styrene or its derivatives and MAh were found to be phosphorescent.⁸⁸

Concurrently, we reported the development of dual pH- and thermo-responsive luminescent alternating copolymers based on the poly(styrene-*alt*-maleimide) skeleton (Fig. 15A). To this end, an *N*-substituted maleimide monomer bearing a diethylene oxide side-chain, (*N*-(methoxydiethyleneglycol) maleimide) and a *tert*-butylcarbamate (Boc)-protected leucine appended styrenic monomer was copolymerized at an equimolar feed ratio to obtain well-defined copolymers with perfectly alternating sequences of these two monomers. The as-prepared Boc-protected copolymers showed bright-blue emission in organic solvents, whereas the deprotected copolymers retained their luminescence behavior in organic solvents and also exhibited dual pH/temperature-tunable fluorescence activity in water.⁸⁹ This conventional fluorophore-free water-soluble luminescent copolymer was applied as a turn-off fluorescent sensor for selective, sensitive and instantaneous detection of the nitro explosive PA in 100% water. Besides, this polymeric sensor showed the ability to detect PA in both solid and vapor phases (Fig. 15B). The ground-state electrostatic complex formation with concomitant ground-state electron transfer from the analyte (picrate anion) to the polymeric probe was proposed as the fluorescence quenching mechanism.⁹⁰

In another report, we engineered an alternating polyampholyte based on the same polymeric building block by copoly-



Fig. 15 General structure of a poly(styrene-*alt*-maleimide) copolymer with variable pendants R1 and R2. (A) Solution properties and unusual photo-luminescence behaviour of a dual pH and thermo-responsive copolymer and (B) its PA sensing application. (C) Chemical structure of leucine-derived alternating 1:1 polyampholyte luminogen and its implementation in CS₂ sensing in both solution and vapour phases. (D) Schematic representation demonstrating pH-induced drug delivery application using such an alternating copolymer. Figure A reprinted with permission from ref. 89. Copyright (2016), Royal Society of Chemistry. Figure B reprinted with permission from ref. 90. Copyright (2017), Royal Society of Chemistry. Figure C reprinted with permission from ref. 91. Copyright (2019), Royal Society of Chemistry. Figure D reprinted with permission from ref. 92. Copyright (2019), American Chemical Society.

merization of two rationally designed monomers: L-leucine-derived functional group protected maleimide and styrene-based monomers *via* RAFT polymerization. The expulsion of protection groups resulted in the formation of a charge-neutral polyampholyte having an anionic and a cationic functional entity placed in a perfectly alternating fashion. Interestingly, the as-synthesized polyampholyte displayed emission behavior in aqueous media and was applied for the specific recognition of carbon disulfide (CS₂) amongst volatile organic compounds (VOCs) in pure water (Fig. 15C). In addition, a colorimetric “naked-eye” detection of CS₂ vapor with significant sensitivity and selectivity under ambient conditions was accomplished with this polymeric probe.⁹¹

Next, to explore the biomedical applications of such intrinsically emissive alternating copolymers, an autofluorescent amphiphilic brush copolymer was fabricated as a pH-sensitive drug delivery nanocarrier for the intracellular administration of the fluorescent inactive aromatic nitrogen mustard chlorambucil (CBL) drug into the cancer cells. To serve the purpose, a CBL functionalized *N*-substituted maleimide monomer was copolymerized with a poly(ethylene glycol) (PEG) attached styrenic macromonomer, which provided a well-defined luminescent alternating copolymer. These copolymers were self-assembled into nanoparticles in aqueous medium where the CBL drug was enclosed into the core of the self-assembled nanostructures. At endolysosomal pH 5.0, 90% of the CBL was released from the nanocarrier by the cleavage of pH-responsive aliphatic ester linkages, connecting CBL to the polymeric chain. The unique luminescence property assisted intracellular monitoring of CBL-incorporated nanocarriers by virtue of fluorescence microscope imaging is noteworthy (Fig. 15D).⁹²

The generation of fully controlled 1:1 alternating D- and L-alanine containing sequence-regulated polymers based on the poly(styrene-*alt*-maleimide) backbone using an *N*-substituted maleimide monomer bearing Boc-L-alanine and a styrene conjugated Boc-D-alanine monomer *via* the RAFT polymerization process has been reported. It is noteworthy that these copolymers displayed fluorescence in various organic solvents and retained their fluorescence activity even after deprotection as observed in organic and aqueous media.⁹³ Recently, we employed an alanyl-alanine tethered maleimide monomer and a glycyl-glycine functionalized styrenic monomer to make these two dipeptides containing 1:1 alternating copolymers on the same polymer skeleton. Unusual luminescence characteristics in solution and in the solid state were observed, where only blue emission occurred in solution but in the solid-state tri-color emission (blue, green and red) was observed.⁹⁴

Yang *et al.* observed that although succinimides act as common fluorescence quenchers of proteins, and common succinimide monomers are nonfluorescent but succinimide-based compounds and also succinimide-containing polymers became fluorescent only after functionalization with dithiol/amino substituents without introducing any conjugated fluorescent entity. The unexpected fluorescence of dithio/amino-succinimides originated from the spatial separation between the highest occupied molecular orbital (HOMO) and the lowest unoccupied molecular orbital (LUMO), realized from DFT calculations. Compared with monomers, the 2-amino-succinimide-based polymers showed considerable enhancement in emission intensity (up to ~200-fold) owing to the isolation effect where the non-radiative decay occurring among succinimide units was significantly suppressed.⁹⁵

Poly(amide-imide)s (PAIs) belong to a class of outstanding engineering materials which express the combined properties of both polyamides (PAs) and polyimides (PIs) *viz.* superior thermal stabilities, high mechanical strength, chemical resistance and facile processability.^{96,97} Broadly, PAIs are nonemissive, and do not exhibit fluorescence except when they are totally aromatic or semi-aromatically connected with emissive species. Therefore, the construction of fluorescent aliphatic PAIs has been challenging and has rarely been reported. Meanwhile, PAIs mostly suffer from their solubility issue in common organic solvents and water, restricting their potential roles in biological applications. However, Yang *et al.* demonstrated the synthesis of luminescent aliphatic PAI with excellent solubility in both organic and aqueous media. To this end, initially, a functional thiolactone-maleimide monomer was synthesized using a Cu(I)-catalyzed click reaction. An efficient polycondensation between the thiolactone-maleimide and an aliphatic diamine afforded fluorescent aliphatic PAIs containing pendant thiols and 2-aminosuccinimide fluorophores through the combination of the ring-opening of thiolactones and amine-maleimide Michael addition reaction. Furthermore, the released thiol was employed for PEGylation *via* a thiol-methacrylate Michael addition that significantly resolved the solubility issues of PAIs in both common organic solvents and aqueous media.⁹⁸ Using similar types of chemical reactions, they also fabricated hyperbranched aliphatic PAI with fluorescence behavior. The as-prepared polymer was found to be soluble in various solvents and showed solvent-dependent emission with moderate QYs.⁹⁹

Again, by using a similar synthetic route, non-conjugated PEGylated PAI was prepared with isolated benzene rings. The authors serendipitously discovered that this amphiphilic PEGylated PAI self-assembled into nanovesicles with unexpected red fluorescence in water. It was believed that the synergistic effects of π - π stacking interactions and H-bonding endowed non-conjugated PAI with a significant bathochromic shift (>100 nm) from the blue-green fluorescence of the amino-succinimide fluorophore to red emission. Moreover, the self-assembled structure shielded PAIs from the interaction with water and suppressed the H-bond formation between amino-succinimide fluorophores and water, prohibiting the quenching of fluorescence in protic solvents and as a consequence the PAIs were found to be emissive in water. Finally, the graft copolymer showed excellent biostability and photostability and a promising cell imaging ability.¹⁰⁰

Tang *et al.* developed an easy and effective multicomponent tandem polymerization method to afford a series of NTIL poly(aminomaleimide)s (PAMIs) with well-defined structures and moderate molecular weights in excellent yields from alkynes and diamines. The as-synthesized PAMIs were employed as macromonomers to carry out further polymerization which resulted in PAMIs that barely contained simple and single maleimide moieties in the polymer main-chain. However, they exhibited strong blue-green emission in DMSO/ethyl acetate (EA) mixtures. Interestingly, the emission color and intensity did not change with the addition of different EA fractions (f_{EA})

in the DMSO/EA solvent mixture, although EA is a bad solvent for PAMI and it aggregated in solvent mixtures with a high f_{EA} . PAMIs showed outstanding biocompatibility, and were therefore advantageous for bioimaging applications.¹⁰¹

Although Tang's group proposed that the unexpected photoluminescence behavior of maleimide or MAH-based homooligomers and alternating copolymers is a result of through-space electronic communication among the locked C=O groups of MAH or maleimide units, the mode of interaction of these non-aromatic groups was not known; and therefore further investigation was required. To prove the hypothesis they synthesized the homooligomer of oligo(maleic anhydride)s (OMAHs) and an alternating polymer of PMP as well as their copolymers, and their systematic structure-property relationships were investigated. Instead of a small vinyl acetate comonomer, they have used more bulky TMP as a comonomer partner to isolate the MAH units, thereby inhibiting their intra-chain through-space interactions, and this assists in the examination of the effect of the in-between distance on the fluorescence characteristics in condensed phases. OMAHs showed distinct absorption at *ca.* 365 and 458 nm, whereas PMP did not absorb above 300 nm. Furthermore, under UV irradiation (365 nm), OMAH4 showed intense emissions, but PMP emitted very weakly both in solution and in the solid state (Fig. 16). It was anticipated that bulky *t*-butyl groups segregated MAH repeating units in PMP. Again simulation results revealed that the distance between two adjacent MAH units of OMAHs is in the range of 2.84–3.18 Å, whereas for PMP, the distance was 4.90–5.37 Å. It was further unveiled that the existence of the intra- and inter-chain π - π^* interactions of the C=O groups of MAH units present in OMAHs is responsible for unexpected light emission. Detailed experimental results and theoretical simulation study led to the coining of the term CTE or clusteroluminescence mechanism for rationalizing the inexplicable luminescence of such non-aromatic polymers.¹⁰²

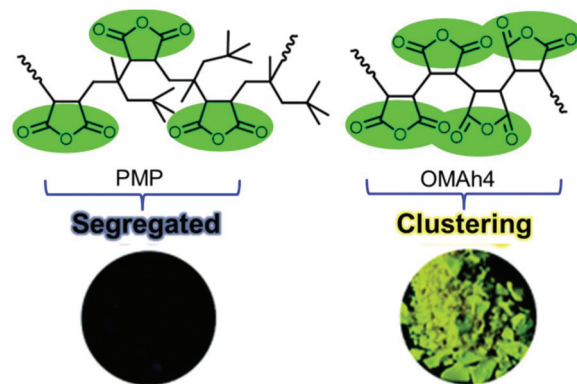


Fig. 16 Chemical structures of PMP and OMAH4 and corresponding luminescence photographs of their solids when exposed to 365 nm UV illumination. Reprinted with permission from ref. 102. Copyright (2017), Royal Society of Chemistry.

Turner's research group synthesized *tert*-butyl carboxylate functionalized four alternating copolymers: two poly(stilbene-*alt*-maleic anhydride) and two poly(styrene-*alt*-maleic anhydride), and their fluorescence properties were studied in detail. The stilbene-containing copolymers I and II exhibited strong fluorescence with high QYs of 71.2% and 65.9%, respectively, whereas both of the styrene-containing copolymers III and IV (Fig. 17) exhibited weak fluorescence with low QYs of 2.6% and 3.1%, respectively. The fluorescence remained unaltered even after the removal of *tert*-butyl groups which makes the resulting copolymer completely water-soluble. The origin of the fluorescent band with high QY was assigned to the extra phenyl groups and "through space" π - π interactions between the phenyl rings of stilbene and C=O groups from the succinimides.¹⁰³

2.2.10. Other polymers. You *et al.* reported one of the earliest examples of NTIL polymers comprising non-fluorescent monomers containing amide bonds. They prepared a simple linear poly(*N*-isopropyl acrylamide) (PNIPAM) using a trithiocarbonate chain transfer agent (CTA) *via* RAFT polymerization. Unexpectedly, the resultant homopolymer was able to exhibit luminescence characteristics (excitation = 380 nm; emission = 410 nm) with a QY of 63.6%. Consequently, PNIPAM displayed a strong blue luminescence when exposed to a UV lamp of 366 nm. Meanwhile, a remarkable enhancement in the steady-state fluorescence intensity was observed as a function of molecular weight. DFT calculations suggested that this inexplicable fluorescence may be due to the π - π electronic interaction between the C=O unit of NIPAM and the benzene ring of CTA present in the polymeric pendant and chain end, respectively. Furthermore, the investigators synthesized a well-defined luminescent multiblock copolymer comprising both PNIPAM and poly(oligoethylene glycol acrylate) (POEGA) using the same polymerization technique and used it for cell imaging purposes.¹⁰⁴

Tang and coworkers attempted to understand the chromophore responsible for the intrinsic fluorescence of non-aromatic polymers such as PNIPAM and poly(*N*-*tert*-butylacrylamide) (PNTBA). By further investigating the emission features of a group of non-aromatic bio-based and synthetic peptides,

they proposed that the fluorescence in the aggregation state/condensed phase in such non-conventional fluorescent polymers is attributed to the electronic interaction amongst the amide groups, where H-bonds play a decisive role in bringing these functionalities in the immediate vicinity.¹⁰⁵

Autoluminescent micelles were fabricated through the self-assembly of amphiphilic polymer molecules having an intrinsic unconventional fluorophore and AIE-active polyetheramide (PEA). A one-pot synthetic route was employed to prepare the PEA molecule consisting of a hydrophilic poly(ethylene oxide) (PEO) backbone and two hydrophobic chains on both the ends from palmitic acid and commercially obtainable polyetheramide (Jeffamine ED2003). The AIE behavior of PEA emerged from the structural rigidification caused by the restriction of intramolecular rotation by virtue of multiple H-bonding interactions within the amide groups rather than from the clusters of oxygen atoms with lone pair electrons of PEO segments as the aggregation of the PEO-PPO-PEO triblock copolymer (Pluronic F127) was found to be non-emissive.¹⁰⁶

Wang *et al.* identified a new series of nonconventional fluorescent polynorbornenes bearing aminosuccinimide side groups. Under UV illumination, these polymeric materials showed bright-blue emission (470–478 nm) with large Stokes shifts of over 110 nm.¹⁰⁷ The same research group also discovered the intrinsic fluorescence of a common non-conjugated compound, poly(*N*-vinylpyrrolidone) (PVP), a linear polymer with side-chain pyrrolidone rings (a lactam) connected to a simple polyethylene backbone. Although the monomer *N*-vinyl-2-pyrrolidone (NVP) was very weakly emissive, its polymer version PVP exhibited strong intrinsic fluorescence. Surprisingly, the oxidized hydrolysate of PVP showed much stronger fluorescence, about 1000 times stronger than those of unhydrolyzed compounds (Fig. 18). The increased fluorescence intensity was attributed to the formation of secondary amine oxides.¹⁰⁸

We unveiled the unprecedented luminescence, AIE activity and heat-induced AIE characteristics of well-known non-conjugated thermoresponsive poly(*N*-vinylcaprolactam) (PNVCL), a higher homolog of PVP. The PNVCL film displayed multicolor emission (blue, green and red) upon illumination with

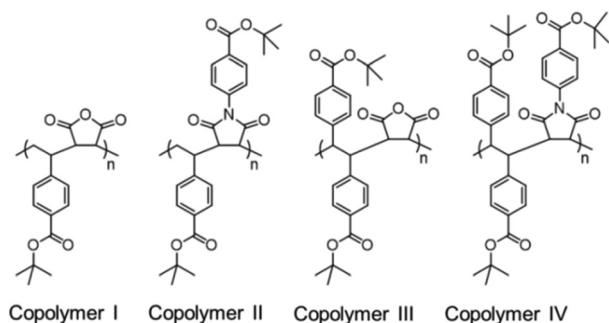


Fig. 17 Molecular structures of stilbene-containing luminescent alternating copolymers. Reprinted with permission from ref. 103. Copyright (2018), WILEY-VCH.

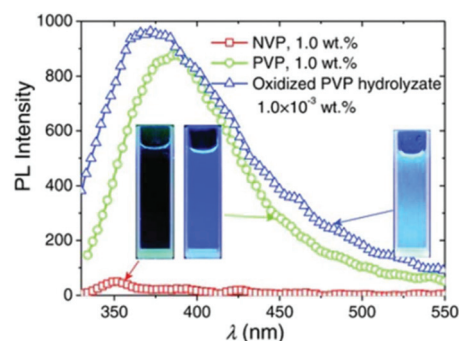


Fig. 18 Fluorescence spectra of NVP, PVP and oxidized PVP hydrolysate in water. Reprinted with permission from ref. 108. Copyright (2015), WILEY-VCH.

different excitation wavelengths, suggesting the coexistence of multiple emissive species with different energy levels. Impressively, the PNVL (concentration = 0.1 wt% in water) exhibited a bright blue emission upon heating above its lower critical solution temperature (LCST) (37.5 °C), while it was nonfluorescent below this temperature (Fig. 19A). This fact was ascribed to the heat-assisted AIE phenomenon. Subsequently, these fascinating properties, temperature-assisted aggregation and AIE characteristics, were ingeniously utilized to apply PNVL as a suitable fluorescent polymeric thermometer to determine the intracellular temperature (Fig. 19B).¹⁰⁹

Yuan *et al.* showed that the CTE hypothesis to rationalize the inexplicable fluorescence properties of nonconventional luminogens is also applicable to those compounds with merely oxygen moieties (*i.e.*, ether and -OH). Impressively, in solid-state xylitol, PEG and F127 displayed cryogenic persistent phosphorescence sustaining for almost 12 s, and even persistent RTP emission was observed from xylitol crystals.¹¹⁰

Wang and coworkers observed autofluorescence of hydrogels made of polyacrylamide (PAM) or polyacrylic acid (PAA) and their derivatives devoid of any traditional fluorophore. This autofluorescence was successfully applied for self-monitoring the gelation process, as the internal structural changes of the gels strongly correlated with the fluorescence response. It was found that with the decrease of water content in the as-prepared gels, the fluorescence intensity gradually increased. When the water content was >40%, a very weak fluorescence was observed, while the emission intensity sharply increased

once the water content decreased from 40% to 20%. Moreover, the gel exhibited dazzling fluorescence on further diminishing the water content to 10%. However, the quenching of fluorescence takes place with some particular metal ion namely Fe^{3+} , the erasure. Accordingly, information rewriting in the presence of visible and UV light, respectively, was successfully realized by using an ethanol solution containing Fe^{3+} as a “writer” and ethanol as a “remover”.¹¹¹

Yuan *et al.* reported the preparation of new sulfur-containing polythioether *via* a Michael polyaddition between 1,4-butanedithiol and 1,4-butanedioldiacrylate. The synthesized polythioether was found to be emissive under UV irradiation in aggregates, despite the absence of remarkable conjugations. Furthermore, boosting of the emission efficiency was realized by the oxidation of polythioether to polysulfoxide and polysulfone both in the solid state and in concentrated solutions. The clustering of sulfur-containing functional groups ($\text{O}=\text{S}=\text{O}$ or $\text{S}=\text{O}$, S) and ester moieties (COO) assisted in the “through-space” electronic interaction and concurrently rigidified the molecular conformations, making the clusters fluorescent active under UV light irradiation.¹¹²

They also reported the unique intrinsic luminescence and additionally p-RTP from a group of non-conjugated and aromatic-ring-free amorphous polymers PAA, PAM and PNIPAM (Fig. 20A). The polymers exhibited no emission in dilute solutions, but became highly luminescent in solid powders/films, nanosuspensions and concentrated solutions. Interestingly, PAA and PAM solids showed distinct p-RTP in air, while PNIPAM solids did so under vacuum or nitrogen (Fig. 20B and C). Furthermore, ionization-induced increased efficiency and prolonged p-RTP were realized in the case of PAANA and PAMHCl powders resulting from the neutralization of PAA and PAM with NaOH or fuming HCl, respectively. The p-RTP of PNIPAM was also modulated by pressurization (Fig. 20D).¹¹³

Tang and coworkers reported the synthesis of luminescent well-defined polythioamide *via* catalyst-free atom economical multicomponent polymerization (MCP) reaction using elemental sulphur, aliphatic diamines, and aromatic diynes as monomers. The synthesized polythioamides were found to be emissive both in solution and in the powder state upon UV irradiation. They envisioned that the H-bonding between the secondary amine and thiocarbonyl group along with the $n \rightarrow \pi^*$ interaction among thioamides causes the formation of “heterodox clusters” which may behave as an actual luminogen.¹¹⁴

Recently, Ling *et al.* achieved colorful ultralong clusterization-triggered phosphorescent (CTP) PAMs by controlling the polymerization requisites (Fig. 21A). Two reaction parameters, specifically initiators and polymerization temperatures, impose immense effects on the emission intensity and color of the CTP PAMs. It was found that water-soluble initiators ($(\text{NH}_4)_2\text{S}_2\text{O}_8$, and $\text{Na}_2\text{SO}_3/(\text{NH}_4)_2\text{S}_2\text{O}_8$) can cause quite high molecular weights of PAMs and strong persistent CTP than oil-soluble 2,2'-azobis(2-methylpropionitrile) (AIBN) initiated PAM (PAM1) and there was a correlation between the afterglow color and polymerization temperature. PAMs initiated at high temperature (PAM2 and PAM3) or aged at 100 °C (PAM5) displayed



Fig. 19 (A) Synthetic route to PNVL and its thermo-induced fluorescence photographs on exposure to a 365 nm UV lamp in an aqueous medium. (B) CLSM images of PNVL treated MCF-7 cells at 25, 35 and 38 °C. Scale bar is 10 μm . Reprinted with permission from ref. 109. Copyright (2020), Royal Society of Chemistry.



Fig. 20 (A) Molecular structures of PAA, PAM and PNIPAM. (B) Schematic diagram of the variation of p-RTP features of the mentioned polymers. (C and D) Luminescence pictures of different films and powders before and after ceasing the UV excitation at 312 nm and room temperature. PNIPAM/p means PNIPAM powders are pressed at 2000 kg cm⁻² pressure for one min. Reprinted with permission from ref. 113. Copyright (2019), Royal Society of Chemistry.



Fig. 21 (A) Prompt fluorescence ($\lambda_{\text{ex}} = 347$ nm) response of polyacrylamides (PAM2–PAM5). The inset shows their afterglow images while the 365 nm UV source was switched off. (B) Schematic representation demonstrating two different orientations of the polymer chain of PAMs: small and large clusters were formed by intra- and inter-molecular hydrogen bonding, respectively. Reprinted with permission from ref. 115. Copyright (2020), Royal Society of Chemistry.

green ultralong RTP by the formation of side-by-side small clusters through intra-chain H-bonding among acrylamide (AM) units, and that PAM4 obtained at low polymerization temperature without heat treatment exhibited a yellow color afterglow *via* the formation of head-to-head large clusters mainly through inter-chain hydrogen bonding (Fig. 21B).¹¹⁵

3. Conclusions and outlook

Intrinsic luminescence was serendipitously discovered from a few nonaromatic biomacromolecules and subsequently further systematic investigation was performed in some other natural polymers. These biopolymers include a few polysaccharides such as starch, cellulose, chitosan, dextran, glycogen, sodium alginate, *etc.*, and some proteins such as BSA, OVA and HSA.

Apart from natural polymers, other synthetic nonaromatic polymers, for example, PAMAMs, PAE, PEIs, PEA, PU, polyester, polycarbonate, PAN, PEG, PAA, PAM, PNIPAM, PNVP, and PNVCL lacking traditional aromatic chromophores also furnish intrinsic fluorescence. So the above libraries of luminescent polymers strongly indicate that the aromatic entities are not a key requirement to acquire light emission behavior from such polymers. In addition, a few polymers with isolated benzene ring structures, such as PET, PSS, PVP-S and PVP, showed exceptional fluorescence and phosphorescence behavior.

The CTE mechanism was envisaged to justify the unorthodox photophysical characteristics of nonconventional luminogens. That is, the electron cloud overlap of nonconventional chromophores bearing lone-paired (n) electrons of O, N, P and S atoms and π orbitals of double and/or triple bonds or C=O

groups in the cluster with concurrent conformational rigidification can account for the unprecedented luminescence behavior of nontraditional luminophores. However, most of the reported NLPs normally emit excitation-independent blue luminescence upon irradiation with UV light and excitation-dependent multicolor emission with relatively low QYs at longer wavelength ranges. The aforementioned drawbacks would limit their application possibilities, particularly in the area of bioimaging, due to strong biological autofluorescence in the blue region and potential tissue damage on exposure to UV light.¹¹⁶ As the underlying photoluminescence mechanism of NLPs is not yet crystal clear, only very few reports could be made on the successful preparation of NLPs with long-wavelength emissions. Therefore, achieving intrinsic long-wavelength emission from NLPs upon excitation with visible light of higher QYs is highly desirable for unequivocal understanding of their luminescence mechanism/sources and for widening their practical enforcement.

Researchers before 2017 mostly focused on the fluorescence processes of the NLPs, but their fascinating phosphorescence behavior, particularly room temperature phosphorescence (RTP), was almost overlooked. However, in the last 3 years, special attention has been paid to glorious developments of RTP materials. Various strategies such as hydrogen bonding, ionic cross-linking, strong ionic interactions, nonmetal dative bond formation, *etc.* have been employed for designing many fascinating polymer-based RTP materials and strengthening their emission characteristics. It is noteworthy that another review article highlighting the phosphorescence properties of NLPs was published by Bryce *et al.*¹¹⁷ when this manuscript was in the evaluation stage. Generally, nonconventional polymers exhibited cryogenic phosphorescence or RTP mostly in the solid-state, so it is a great challenge to generate phosphorescence in the solution phase especially in aqueous medium.

The discovery of the cryogenic phosphorescence and RTP phenomenon in NLPs is a huge breakthrough and one of the most recent advancements. This will include another dimension within the field of bioimaging in the foreseeable future due to the longer excitation lifetime of phosphorescence relative to fluorescence, and therefore more sophisticated instruments could be potentially used or invented to look inside the cells and subsequently perceive the dynamics of life processes. To date, phosphorescent NLPs have been utilized mainly for information encryption. But owing to their high biocompatibility and less toxicity, these RTP polymers could be beneficial as *in vitro* and *in vivo* bioprobes to improve the detection sensitivity by diminishing the signal-to-noise ratio. Furthermore, organelle targeted (such as the nuclei, lysosomes, and mitochondria) bioimaging could be a research hotspot for low cytotoxicity NLPs. By highlighting the fluorescence and phosphorescence behavior of NLPs in this article, we firmly believe that this manuscript will help new and experienced researchers from broad research areas to advance their knowledge and to come up with new possibilities for expanding the horizon of these NLPs.

Conflicts of interest

There are no conflicts to declare.

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properties of the NLPs as they have been developed very recently. Hence, in the present review article, we have provided a comprehensive and extensive discussion on the newly discovered photophysical properties including dual-emission (fluorescence and phosphorescence) of NLPs along with their underlying mechanisms and advanced applications which are transpiring at a rapid pace nowadays.

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