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The base-free multicomponent polymerization of elemental sulfur, difluoromethylene phosphobetaine and amines toward electron-deficient aromatic polythioureas†

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Polythioureas are emerging materials with fascinating properties, such as self-healing and adhesion, high refractive indices, high dielectric constants, and heavy metal ion adsorption. With the increasing requirement for polymer structures and properties suiting a wide range of potential applications, versatile synthetic approaches are demanded to access a great diversity of polythiourea structures efficiently and economically. In this work, a commercially available difluorocarbene precursor, difluoromethylene phosphobetaine (PDFA), was selected to react with sulfur and electron-deficient aromatic amines to enable the efficient syntheses of electron-deficient polythioureas from amine monomers with low reactivity based on the high reactivity of a difluorothiocarbonyl intermediate. A one-pot catalyst-free multicomponent reaction of sulfur, PDFA and an amine was designed, which could take place efficiently in DMAc at 60 °C under nitrogen and was applicable for aromatic amines with both electron-donating and electronwithdrawing groups, producing thioureas in high yields (up to 93%). Most importantly, catalyst-free multicomponent polymerizations of sulfur, PDFA, and diamines were also developed in DMAc at 60 °C with commercially available monomers, showing high efficiency (M_w s up to 65 900 g mol⁻¹ and yields up to 99%) and wide monomer applicability, providing an efficient synthetic approach for syntheses of polythioureas. Moreover, these polythiourea thin films showed high refractive indices (n_{633} nm up to 1.8133), suggesting their potential application in optical devices.

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Introduction

Polythioureas, with their unique hydrogen bonds, large dipole moment, large molar refractivity, metal-coordination ability, and others, have been developed into a group of promising polymer materials, and have attracted increasing attention owing to their potential applications as self-healing materials, ¹⁻³ light refraction materials, ⁴ dielectric materials, ⁵⁻⁸ dynamic polymer networks, ⁹ optoelectronic materials, ¹⁰ and precious metal recovery materials. ¹¹ Owing to the different requirements for various applications, polythioureas with different chemical structures and different thermal properties

and mechanical properties are required. Versatile synthetic approaches are hence demanded to access a great diversity of polythiourea structures.

The commonly adopted synthetic approaches for polythioureas have mainly been divided into two kinds. One is polycondensations between diamine monomers and thiophosgene, diimidazolethiocarbonyl, or thiourea. These sulfur-containing reagents are generally toxic and smelly, difficult to handle, release harmful small molecules, and some even required special synthetic conditions, such as microwave assistance. The other method is polyadditions between diisothiocyanates and diamine monomers. 13 While the reaction is efficient, the diisothiocyanate monomers were difficult to synthesize, and there was only limited commercially available diisocyanate compound. Recently, carbon disulfide was utilized for the synthesis of polythioureas; 14,15 however, an equivalent amount of sulfurcontaining small molecules such as H2S would be released as a byproduct, which was environmentally unfriendly. The limitations of the sulfur-containing monomers have vastly hindered the exploration of the chemical structures of polythioureas.

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Elemental sulfur, which abundantly exists in nature, is one of the major byproducts of the petroleum and natural gas industries, 16-19 and is a non-toxic, safe, odorless solid powder. It could serve as an ideal and economical monomer for the synthesis of sulfur-containing polymers. Great efforts have been made to utilize elemental sulfur to prepare polymeric materials, such as inverse vulcanization to synthesize chalcogenide hybrid inorganic/organic polymers.^{20–24} Polymerizations of sulfur were also reported for the synthesis of poly(benzothiazole)s, 16 polydisulfides, 25 poly(O-thiocarbamate)s, 26 polythioamides 27,28 and others. Among various sulfur-utilization approaches, multicomponent polymerization (MCP), with three or more monomers reacting together in a one-pot procedure to afford the targeted polymeric product, 29-33 offers an efficient approach to convert elemental sulfur to various sulfur-containing polymers, such as polythioamides, 34,35 polythiocarbamates, 26 and polythioureas.11 Additionally, it has shown fascinating advantages, including high efficiency, simple operation, mild conditions, robustness, and, most importantly, great polymer product structural diversity. We have reported the MCPs of elemental sulfur, aliphatic diamines, and diisocyanates to afford polythioureas with various aliphatic or semi-aromatic structures. 11,22,27 The MCPs of elemental sulfur, CH2Cl2 and aromatic diamines were also recently developed to afford a series of aromatic polythioureas with high efficiency.³⁶ MCPs of sulfur, chloroform and aromatic diamines were also reported to produce polythioureas with several repeating unit structures through the use of dichlorocarbene active species in the presence of a large amount of the strong base t-BuOK.³⁷ However, for aromatic amines with electron-withdrawing groups, which exhibit lower reactivity and weaker nucleophilicity compared to those with electron-donating groups, it was challenging to realize efficient transformations to thioureas. It is hence still difficult to access electron-deficient aromatic polythioureas, which are required and highly desired for potential applications in organic catalysis³⁸ and for other optical and electronic applications.39

To react efficiently with the less-reactive aromatic diamines, difluorocarbene as an active reaction intermediate could be a promising candidate; it was commonly used for fluoromethylations, $^{40-43}$ insertion reactions of X–H bonds (X = O, N, S), and

1] cycloaddition reactions involving bonds. 44 Xiao et al. developed a difluoromethylene phosphobetaine (PDFA) reagent to produce difluorocarbene in situ to participate N-trifluoromethylation, 45 further in C-trifluoromethylthiolation, 46,47 etc. In these reactions, PDFA was reported to react with sulfur to generate thiocarbonyl fluoride in situ, which then reacted with various amines to afford trifluoromethylated products, isothiocyanates, and HCF₂S-substituted heterocycles. 45 For example, PDFA, sulfur and an aromatic amine could react in 1,2-dimethoxyethane at 80 °C to afford the isothiocyanate in 85% yield, which could further react with dimethylamine to produce thiourea-containing chloromethiuron; additionally, a vicinal diamine with two neighboring -NH2 groups on a benzene ring was reported to react with PDFA and sulfur to afford cyclic thiourea in 75% yield at 80 °C in 1,2-dimethoxylethane (DME). The highly reactive difluorothiocarbonyl intermediate could hence undergo nucleophilic attack reactions with amines, including less-reactive electron-withdrawing-group-containing aromatic amines, to afford the corresponding polythioureas under suitable reaction conditions (Fig. 1).

In this work, to develop a polymerization approach for the synthesis of polythioureas, the one-pot multicomponent reaction of PDFA, sulfur, and two amine molecules was first investigated to directly produce a thiourea product. As solvents such as DME were not favored for the solubility of polythioureas with abundant intermolecular hydrogen bonds and strong polarity, polar solvents such as dimethylacetamide (DMAc) were adopted. Moreover, utilizing the highly reactive thiocarbonyl fluoride generated from PDFA and sulfur, electrondeficient aromatic amines might react efficiently to produce thiourea from sulfur. Herein, one-pot catalyst-free multicomponent reactions of sulfur, PDFA and electron-withdrawinggroup-containing aromatic amines were reported to afford thiourea products in high yields. Efficient and rapid polymerizations of sulfur, PDFA and various aromatic and benzyl diamines were developed without any catalyst under mild conditions, producing a series of polythioureas with high yields and high $M_{\rm w}$ s of up to 65 900 g mol⁻¹. In particular, aromatic polythioureas with electron-withdrawing carbonyl groups were successfully prepared in 85% yield and with a $M_{\rm w}$ of 18 600 g mol⁻¹, and they proved to possess the highest

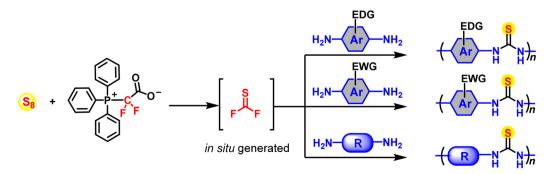


Fig. 1 A strategy for polythioureas synthesized from diamines containing electron-withdrawing groups

refractive index of 1.8133 at 633 nm among the tested aromatic polythioureas.

Results and discussion

Catalyst-free multicomponent reactions (MCRs) of sulfur, PDFA, and amines

To investigate the one-pot MCR of sulfur, PDFA, and an aromatic amine, an MCR was conducted in DME at 50 °C for 0.5 h, and 3-acetylaniline 1a was studied as an example of an electron-withdrawing-group-containing aromatic (Table 1). Luckily, the reaction went smoothly without any catalyst to afford the thiourea product 4a in 64% yield, calculated based on the ¹H NMR spectrum, suggesting that the conversion of sulfur, PDFA, and an amine to an isocyanate, and its further reaction with amine to generate a thiourea could be combined in a one-pot MCR. The MCR was then investigated in DME, THF, DMSO, DMF, and DMAc to screen a suitable solvent, and DMAc afforded the optimal yield of 80%. MCRs of sulfur, PDFA, and 1a in DMAc were then conducted at different temperatures ranging from 50 to 80 °C, suggesting that there was no obvious change in yield within this temperature range. The monomer loading ratio was then studied in DMAc at 60 °C, where the ratio of [1a] to [3] was changed from 2.0:1.2 to 2.0:1.4 while keeping the ratio of sulfur to PDFA fixed; the highest yield of 94% (91% isolated yield) was obtained with [1a]:[3] = 2.0:1.4.

With the high efficiency of the catalyst-free MCR, the monomer scope of amines was then studied, and a series of commercially available electron-withdrawing-group-containing aromatic amines **1a–1g** was selected to react with sulfur and PDFA in DMAc at 60 °C for 0.5 h (Fig. 2). Under the

optimized conditions, these electron-deficient aromatic amines generally reacted efficiently, affording the corresponding thiourea products 4a-4g in high yields. Aromatic amines with benzoyl, ester, -NO2, and sulfonyl groups generally worked well in the MCR, producing rarely reported aromatic thiourea structures in 84-90% isolated yields, demonstrating the good structural tolerance of the MCR. 3-Cyanoaniline 1e showed a lower yield, probably because of a side reaction involving the -CN group. Besides these electrondeficient aromatic amines, p-toluidine 1h with an electrondonating methyl group could also react efficiently under the same conditions, affording the thiourea 4h in 93% yield, showing a similarly high yield to the electron-deficient aromatic amines. Moreover, in addition to aromatic amines, the benzylamine 1i was also investigated, and the MCR of sulfur, PDFA and 1i also proceeded smoothly to afford the thiourea 4i in 62% yield. The catalyst-free MCR hence exhibited a broad amine substrate scope, accommodating electron-deficient and electron-rich aromatic amines as well as aliphatic amine, suggesting its robustness and wide applicability.

During the catalyst-free MCR of p-toluidine **1h**, sulfur and PDFA in DMAc, after reacting at 60 °C for 20 min, the reaction solution was cooled in an ice bath and the mixture was filtered for high-resolution mass spectrometry (HRMS) analysis. The characteristic peaks of thiocarbonyl fluoride (found 82.9770, calcd 82.9767) and isothiocyanate (found 150.0374, calcd 150.0377) were observed, indicating the key intermediates involved in the reaction mechanism (Fig. S1†). In this MCR, the difluorocarbene active species **A** was generated from PDFA upon heating, with the release of Ph₃P=S and CO₂; **A** then reacted with sulfur to produce thiocarbonyl fluoride **B** according to the literature. ⁴⁵ The nucleophilic substitution of the amine then took place to afford the intermediate **C**, and after

Table 1 Optimization of the reaction conditions for the MCRs of 1a, sulfur, and PDFA

| Entry | $[1a]: 1/8[S_8]: [3]$ | Solvent | T (°C) | $Yield^{a}$ (%) |
|-------|-----------------------|---------|--------|-----------------|
| 1 | 2:2.4:1.2 | DME | 50 | 64 |
| 2 | 2:2.4:1.2 | THF | 50 | 67 |
| 3 | 2:2.4:1.2 | DMSO | 50 | 54 |
| 4 | 2:2.4:1.2 | DMF | 50 | 72 |
| 5 | 2:2.4:1.2 | DMAc | 50 | 80 |
| 6 | 2:2.4:1.2 | DMAc | 60 | 83 |
| 7 | 2:2.4:1.2 | DMAc | 70 | 82 |
| 8 | 2:2.4:1.2 | DMAc | 80 | 82 |
| 9 | 2:2.6:1.3 | DMAc | 60 | 87 |
| 10 | 2:2.8:1.4 | DMAc | 60 | $94 (91)^b$ |
| 11 | 2:2.0:1.3 | DMAc | 60 | 88 |
| 12 | 2:3.0:1.3 | DMAc | 60 | 90 |

The reaction was carried out under nitrogen conditions at the corresponding temperature with 1 mL of solvent; the concentration of **1a** was 1.0 M, and the reaction time was 0.5 h. ^a Calculated based on ¹H NMR spectra, using 1,3,5-trimethylbenzene as the internal standard. ^b Isolated yield.

Fig. 2 MCRs of sulfur, PDFA and aromatic amines to form thioureas; the single-crystal-based ORTEP diagrams of compounds 4d-4i are given.

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 F^- was eliminated, the isothiocyanate intermediate **D** was generated, which then underwent nucleophilic addition with another amine molecule to afford the thiourea product **E** (Fig. S2 \dagger).

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Moreover, several single crystals of these thiourea compounds, including unreported thiourea compounds, were obtained via solvent evaporation or diffusion. The thioureas containing -CF₃, -CN, -NO₂, and sulfonyl groups, 4d-4g, generally adopted "W"-shaped molecular configurations in their single crystal structures (Fig. 2). For example, in the single crystal structure of the NO2-containing thiourea 4f, two THF solvent molecules were captured through N-H···O hydrogen bonds. The -NO2 groups also contributed to the formation of rich hydrogen bond networks in the crystal structure (Fig. S3†). Different from that, the single crystal of the CH₃-containing thiourea 4h was monoclinic with a space group of Pbcn, adopting a "V"-shaped molecular conformation, which was the same as reported crystals prepared via other synthetic approaches.⁴⁸ A dense hydrogen bond network was formed through -N-H...S bonds with a length of 2.496 Å (Fig. S3†). Different from the two above-mentioned types of thioureas, 4i prepared from a benzylamine possessed an asymmetric conformation. Different types of thiourea structures hence adopted different molecular conformations because the hybridization of C–N bonds imparts *cis-trans* isomeric properties on the C–N bonds of the thiourea moiety, influencing the spatial arrangement of associated N–H bonds. The presence of multiple and diverse hydrogen bonds endows the polythioureas with various hydrogen bond networks.

Catalyst-free multicomponent polymerizations of elemental sulfur, PDFA, and diamines

To explore the multicomponent polymerizations of sulfur, PDFA and diamines, commercially available diphenylmethane-diamine 5a was selected as an example to polymerize with sulfur and PDFA. The MCP was conducted in DMF at 40 °C for 3 h under nitrogen with $[5a]:1/8[S_8]:[PDFA]=1.0:2.6:1.3$, and a polymer product with a M_w of 10 200 g mol⁻¹ was obtained in 60% yield. It should be noted that besides the thiourea moieties in the polymer product, sulfur atoms were also transformed to Ph₃P=S as a byproduct. In solvents with higher polarity, such as DMAc and DMSO, the polymerization yields improved, and the best result was obtained in DMAc, with the polymeric product with a M_w of 16 800 g mol⁻¹ obtained in 84% yield (Table S1†). The temperature was found

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Table 2 Optimization of the MCP conditions using diamines, elemental sulfur, and PDFA^a

| Entry | T (°C) | $[5a]: 1/8[S_8]: [3]$ | $M_{\rm w}^{\ b} \left({\rm g \ mol}^{-1} \right)$ | $M_{ m w}/M_{ m n}$ | Yield (%) |
|-------|--------|-----------------------|---|---------------------|-----------|
| 1 | 40 | 1.0:2.6:1.3 | 16 800 | 1.34 | 84 |
| 2 | 50 | 1.0:2.6:1.3 | 20 900 | 1.26 | 82 |
| 3 | 60 | 1.0:2.6:1.3 | 53 900 | 1.82 | 94 |
| 4 | 70 | 1.0:2.6:1.3 | 36 900 | 1.51 | 98 |
| 5 | 80 | 1.0:2.6:1.3 | 31 700 | 1.50 | 93 |
| 6 | 60 | 1.0:2.0:1.0 | 13 300 | 1.24 | 65 |
| 7 | 60 | 1.0:2.4:1.2 | 22 900 | 1.35 | 79 |
| 8 | 60 | 1.0:2.8:1.4 | 37 500 | 1.73 | 97 |
| 9 | 60 | 1.0:3.0:1.5 | 18 500 | 1.47 | 88 |
| 10 | 60 | 1.0:2.4:1.3 | 49 500 | 1.73 | 90 |
| 11 | 60 | 1.0:2.8:1.3 | 51 400 | 1.77 | 94 |

^a The substrate 5a (1.0 mmol) was combined with Ph₃P⁺CF₂CO₂ and S₈ in 1 mL of DMAc and the mixture was stirred for 3 h at the corresponding temperature under N₂. ^b The data were obtained from GPC testing in DMF, with monodisperse PMMA as the standard sample.

to be crucial for polymerization, and the MCP was conducted at a range of temperatures from 40 to 80 °C (Table 2). The polymerization results obtained upon increasing the temperature from 40 to 60 °C suggested that the $M_{\rm w}$ of the polymer gradually increased, achieving 53 900 g mol⁻¹ at 60 °C, whereas further heating resulted in a decrease of the $M_{\rm w}$. Considering the temperature required for the efficient decomposition of PDFA to form a carbene intermediate and that a high temperature might induce side reactions, such as generating byproduct with N-SCF₃ to terminate the polymerization reaction, 45 60 °C was found to be the optimal temperature. The monomer feed ratio for polymerization was then optimized at 60 °C with a fixed ratio of PDFA to sulfur (Table 2). Upon increasing the amount of PDFA, both the yield and $M_{\rm w}$ first increased and then decreased, and when [5a]: 1/8 $[S_8]:[PDFA] = 1:2.6:1.3$, the highest M_w was obtained, suggesting a balancing of the effects from excess PDFA and sulfur.

Surprisingly, unlike other sulfur- and aromatic-aminebased polymerizations, the addition of base was not necessary or even beneficial to the MCP, and the incorporation of K₂CO₃, Na₂CO₃, NaOH, KF, or triethylamine did not improve the polymerization results (Table S2†), suggesting that simple base-free conditions were optimal, probably due to the reduction of base-promoted side reactions such as the generation of trifluoromethylthio-containing byproducts. 42 The progression of the MCP over time was also investigated; it could afford the polymer product with a $M_{\rm w}$ of 42 000 g mol⁻¹ in 87% yield within 10 min, reaching 65 900 g mol⁻¹ in 93% yield after 1 h, suggesting fast and efficient polymerization (Table S3†). Last but not least, the influence of the concentration of the monomer 5a was then investigated while fixing the monomer feed ratio. A low concentration of 0.25 M resulted in relatively low $M_{\rm w}$ s, while the monomer could not be completely dissolved at a high concentration of 1.0 M; 0.5 M was found to be the optimal concentration (Table S4†).

A kinetic study of the MCP

The real-time polymerization process during the MCP of sulfur, PDFA and 5a was then monitored in DMSO at 60 °C using in situ IR measurements (Fig. 3). Due to the interference caused by the coincidence of the C=O stretching vibration peak of DMAc with the C=S bond, DMSO was chosen as the reaction solvent. In the FT-IR spectra of the model compounds **4h** and **P1** (Fig. S4†), the double peaks at \sim 3400 cm⁻¹ associated with the stretching vibration of the $-NH_2$ group of p-toluidine disappeared, while the stretching vibration peak of -NHappeared at ~3200 cm⁻¹. Meanwhile, the stretching vibration peaks of the C=S bonds of 4h and P1 were located at 1555 cm⁻¹ and 1537 cm⁻¹, respectively. Through a comparison of the in situ IR spectra of 5a, PDFA and P1, the peak at 1537 cm⁻¹ was attributed to the stretching vibration of C=S bonds, and the peak at 1702 cm⁻¹ was attributed to the stretching vibration of C=O bonds in PDFA (Fig. S5†). The stacked time-dependent IR spectra of the polymerization solution of sulfur, PDFA and 5a in DMSO showed that a new increasing peak appeared at 1537 cm⁻¹, and the peak intensity at 1702 cm⁻¹ continuously decreased (Fig. 3A). The timedependent peak intensities at 1537 cm⁻¹ suggested that the reaction could be completed in 15 min (Fig. 3B and C), proving the rapid and efficient polymerization.

To explore the monomer scope of the catalyst-free MCP, a series of commercially available diamines 5a-5h was selected to be used as the monomers; these were polymerized with sulfur and PDFA under the optimal conditions in DMAc at 60 °C under nitrogen (Fig. 4 and Fig. S6†). Polythiourea P1 with a high $M_{\rm w}$ of 65 900 g mol⁻¹ was obtained in 93% yield, and good polymerization results were also obtained for the ether- and thioether-containing polythioureas P2 and P3. While these aromatic polythioureas could generally be dissolved in DMF, DMSO, and DMAc, P4 and P5 showed limited solubility; during their synthesis, large amounts of insoluble

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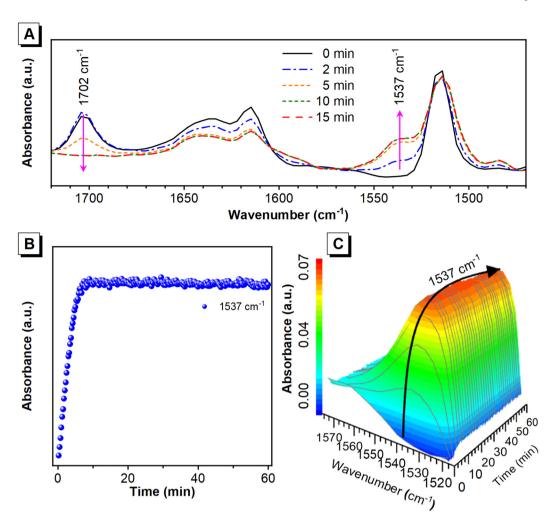


Fig. 3 (A) Stacked time-dependent in situ FT-IR spectra of a polymerization solution of 5a, S₈ and PDFA in DMSO. (B) The time-dependent peak intensities at 1537 cm $^{-1}$ during the MCP. (C) The three-dimensional in situ FT-IR profile of the peaks in the range of 1580-1520 cm $^{-1}$. The monitored polymerization was carried out at 60 °C in 2 mL of DMSO, with a feed ratio of [5a]: 1/8[S₈]: PDFA = 1.0: 2.6: 1.3, using 0.5 mmol of 5a as a standard.

precipitate were formed, and only the soluble part was used for characterization. The diamine 5f with large steric hindrance might twist the polymer chain and avoid strong interchain stacking, which might produce a polythiourea with improved solubility, while sacrificing yield. Furthermore, the MCPs of aliphatic diamines were not as efficient as those with aromatic diamines, and polythioureas P7 and P8 possessed decreased yields and $M_{\rm w}$ s. The polydispersity of the molecular weights of these polymers was generally low for a step-growth polymerization approach, which was probably associated with the special solubility of these polythioureas.

Most importantly, considering challenging electron-withdrawing-group-containing aromatic amine monomers, the sulfonyl- and carbonyl-group-containing electron-deficient aromatic diamines 5i and 5j were studied in the catalyst-free MCP (Fig. 5), and they were polymerized with sulfur and PDFA in DMAc at 60 °C for 1 h. The electron-deficient aromatic polythioureas P9 and P10 were obtained in 76% and 85% yields, respectively, with $M_{\rm w}$ s of 7900 g mol⁻¹ (**P9**) and 18 600 g mol⁻¹

(P10). These rarely reported results suggested the successful polymerization of electron-deficient aromatic diamine monomers.

Structural characterization

The chemical structures of these polythioureas were confirmed based on their ¹H and ¹³C NMR spectra (Fig. 6 and S7†). The resonance of -NH₂ protons seen in the spectrum of 5i at δ 5.87 disappeared in the ¹H NMR spectra of **4b** and **P10**; meanwhile, the -NH- protons of the thiourea groups of 4b and P10 emerged at δ 10.45 and δ 10.44, respectively (Fig. 6A). Similarly, in the ¹³C NMR spectra of 5j, 4b and P10 (Fig. 6B), the C=O peak appeared at δ 192.49, δ 199.83 and δ 193.31, respectively. Most importantly, the C=S peak emerged at δ 184.33 and δ 178.98 in the spectra of **4b** and **P10**, respectively. The ¹H and ¹³C NMR spectra of P1-P9 are also provided in Fig. S8-S11.† The -NH- proton resonances of the thiourea groups of 4h and P1 are located at δ 9.58 and δ 9.67, respectively, in the ¹H NMR spectra, while the C=S resonances of 4h

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Fig. 4 (A) The MCPs of elemental sulfur, PDFA and diamine monomers, (B) the structures of the electron-donating-group-containing aromatic diamines 5a-5h, and (C) the chemical structures and polymerization results for P1-P8. M_w s are determined using GPC in DMF based on a PMMA standard sample.

Fig. 5 MCPs of the electron-withdrawing-group-containing diamines 5i-5j. Mws are determined using GPC in DMF based on a PMMA standard sample.

and **P1** are at δ 179.63 and δ 179.49, respectively, in the ¹³C NMR spectra. Compared with those of P1, the -NH- proton resonance peaks of P9 and P10 appeared at a lower field value of $\sim \delta$ 10.5, and the resonance peaks for the aliphatic polythioureas P7 and P8 appeared at a higher field value of δ 8.0. The characteristic C=S resonance peaks generally emerged at $\sim \delta$ 185. Moreover, in the XPS spectra, the binding energies for the S 2p orbitals of 4h and P1 are closely matched at 163.18/ 161.89 eV and 163.40/162.10 eV, respectively, proving the similar chemical state of sulfur, and also indicating that there was no elemental sulfur residue in the polymer product (Fig. S12†). In the FT-IR spectra of P2-P10, the C=S bond

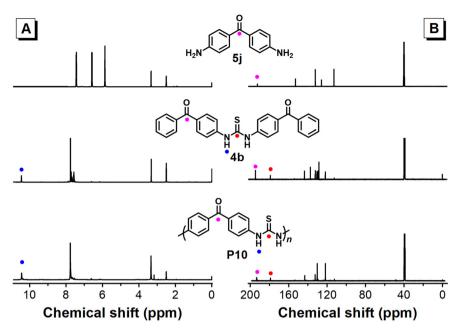


Fig. 6 (A) 1 H NMR and (B) 13 C NMR spectra of 5j, the model compound 4b, and P10 in DMSO- d_{6} .

peaks generally emerged at $1530-1594~{\rm cm}^{-1}$ (Fig. S13 and S14†).

The thermal stability and refractivity of these polythioureas

Thermogravimetric analysis (TGA) indicated that these polythioureas enjoyed good thermal resistance, with their decomposition temperatures ($T_{\rm d}$) at 5 wt% weight loss ranging from 191 °C to 246 °C; the intermolecular hydrogen bonds among the abundant thiourea moieties and the large number of aromatic rings might contribute to the good thermal stability (Fig. 7A). Differential scanning calorimetry (DSC) analysis of these polythioureas was also conducted. The glass transition

temperature was observed for P10 to be 127 °C (Fig. S15†), and no obvious thermal transition temperatures were observed for the other polymers.

Most of the polymers possess satisfying solubility, enabling facile thin film preparation via a spin-coating method. The wavelength-dependent refractive indices of thin films of **P1-P3**, **P6-P7**, and **P9-P10** were measured within the wavelength range of 400–1700 nm (Fig. 7B). The aromatic polythioureas generally showed higher refractive indices compared with the polythiourea **P7** prepared from 1,4-bis(aminomethyl)benzene; and **P10** with electron-withdrawing carbonyl groups possessed the highest $n_{633 \text{ nm}}$ value of 1.8133, which was exceptionally

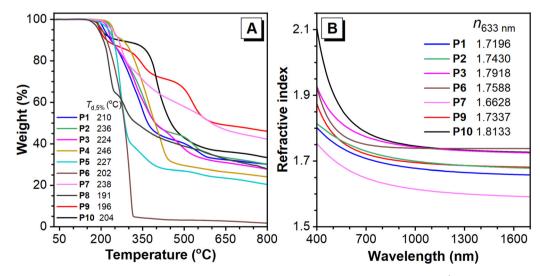


Fig. 7 (A) TGA thermograms of P1–P10, upon heating from 50 °C to 800 °C with a heating rate of 10 °C min⁻¹ under N_2 , and (B) the wavelength-dependent refractive indices of thin films from 400–1700 nm.

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high for organic polymer materials (Table S5†). Moreover, several polymers possessed high n values above 1.7 at 1700 nm, which have rarely been reported. The transmittance spectra of these polymers were also investigated, and their spin-coated thin films on quartz plates were measured. As shown in Fig. S16,† among the tested polythiourea samples of uniform thin films on quartz plates, P1, P2, and P10 showed excellent light transmittance of above 96% beyond 400 nm. Photos of solid samples and thin films on quartz plates of P10 also suggested its high transmittance. These aromatic polythioureas, especially P10 with electron-withdrawing groups, possessed excellent optical properties, and they could be promising optical materials for optical lenses and CMOS systems.

Conclusions

In this work, catalyst-free multicomponent reactions of sulfur, PDFA and amines were developed in DMAc, which proved to be applicable and efficient for both electron-rich and electron-deficient amines, as well as benzylamine. PDFA was then used as a monomer for polymerization with sulfur and diamine without any catalyst; this could react efficiently in DMAc at 60 °C under mild conditions, affording aromatic polythioureas. Notably, aromatic amines containing electron-withdrawing groups could also proceed efficiently, which greatly expanded the amine monomer scope and range of obtained polythiourea structures. Tests of the optical properties of these polythiourea films showed that the electron-deficient polythioureas possessed high refractive indices, reaching 1.8133 at 633 nm, and they could find promising applications in optical sensors, semiconductors, and catalysis.

Data availability

The data supporting this article have been included as part of the ESI.† Crystallographic data for compounds **4d–4i** have been deposited at the Cambridge Crystallographic Data Centre under the CCDC 2407294–2407296, 2407300, 2245406 and 2407301† and can be obtained from **https://www.ccdc.cam.ac.uk**.

Conflicts of interest

The authors declare no competing financial interests.

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