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Tandem diaza-Cope rearrangement polymerization: turning intramolecular reaction into powerful polymerization to give enantiopure materials for Zn²⁺ sensors†

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[3,3]-Sigmatropic rearrangement is a powerful reaction to form C–C bonds stereospecifically; however, owing to intrinsic simultaneous bond formation and breakage, this versatile method has not been utilized in polymerization. Herein, we report a new tandem diaza-Cope rearrangement polymerization (DCRP) that can synthesize polymers with defect-free C–C bond formation from easy and efficient imine formation. A mechanistic investigation by *in situ* ¹H NMR experiments suggests that this polymerization proceeds by a rapid DCR process, forming an enantiospecific C–C bond that occurs almost simultaneously with imine formation. This polymerization produces not only highly stable polymers against hydrolysis due to resonance-assisted hydrogen bonds (RAHBs) but also chiral polymers containing enantiopure salen moieties, which lead to high-performance Zn²⁺-selective turn-on chemosensors with up to 73-fold amplification. We also found that their optical activities and sensing performances are heavily dependent on the reaction temperature, which significantly affects the stereoselectivity of DCR.

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Introduction

Pericyclic reaction, the reorganization of π -bonds in a concerted manner, is one of the most widely used transformations in synthetic organic chemistry. Among the various pericyclic reactions, highly efficient cycloadditions, such as Diels–Alder and Cu-catalyzed azide–alkyne cycloaddition reactions, have been employed as powerful tools to prepare polymers^{1–10} because of their high efficiency and orthogonality (Scheme 1A(a)). [3,3]-Sigmatropic rearrangement is another powerful and reliable pericyclic reaction, allowing for the stereoselective construction of C–C bonds.¹¹ However, this rearrangement cannot be applied to polymerization other than the post-modification of side-chains^{12–16} because it is an intramolecular reaction, which intrinsically forms and breaks the bond simultaneously (Scheme 1A(b)).

Meanwhile, imine formation between an amine and an aldehyde is an efficient and straightforward reaction and occurs reversibly under equilibrium control. Taking advantage of this dynamic nature, imine formation is widely employed for synthesizing polyimine vitrimers,^{17–22} rotaxanes, and catenanes.^{23,24} Ironically, extending this method to obtain high-

molecular-weight linear polymers is still quite challenging due to its inherent reversible imine formation (Scheme 1B).^{25–38} Therefore, to solve the depolymerization issue, an

(A) Pericyclic Reactions



(B) Linear Polymers via Imine Bond Formation



(C) This Work: Diaza-Cope Rearrangement Polymerization



Scheme 1 Polymerization via pericyclic reactions and imine formation.

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Fig. 1 ^1H NMR and ^{13}C NMR spectra of (R,R) -P1.



Fig. 2 (a) Monitoring the polymerization using (S,S) -1 and 2a by *in situ* ^1H NMR spectroscopy in $\text{DMF-}d_7$. (b) Plots showing consumption of 2a (black), formation of the imidazolidine intermediate (green), and phenolic O–H groups (blue and red). (c) Detailed polymerization mechanism.

intermediate *via* imine formation was rapidly formed up to 55% (see Fig. S1† for details).^{46,47} However, (S,S) -3, the second intermediate, comprising the normal O–H signal, was hardly observed, whereas only the RAHB O–H signal corresponding to (R,R) -4 or (R,R) -P1 gradually increased, implying clean conversion (Fig. 2a and b). This suggests that a DCR is rapid, occurring almost simultaneously with imine formation (Fig. 2c). In other words, (R,R) -P1 was predominantly produced by C–C bond polymerization rather than imine polymerization (if so, the

subsequent DCR process would have slowly generated C–C bonds on the pre-formed polymer). On the other hand, in the case of (R,R) -P2 containing electron-donating groups, the activation energy for DCR increased,⁴² thereby decelerating the rearrangement. As a result, both normal and RAHB O–H (δ 10.7 ppm and δ 13.7 ppm, respectively) were observed at the beginning of the reaction, but the polymerization progressed to show only the enthalpically favored RAHB O–H (Fig. S2†). In short, because of the defect-free DCR signal regardless of the electronic characters, this tandem DCRP becomes a novel strategy to form more challenging C–C bonds from easier imine formation.

DCR polymers contain C–C bonds in the main backbone, while the resulting imine bonds at the side-chains are also stabilized by stronger RAHBs. Therefore, we could test their stability against hydrolysis by comparing with analogous poly-imine. To conduct a control experiment, we synthesized **P4** by using a diamine monomer ((S,S) -7) without –OH groups, ensuring that the RAHB that drove the completion of DCR was now absent.³⁹ Therefore, **P4** (M_n of 14.1 kDa) contained both the initial imine ((S,S) -8) and the rearranged C–C bond ((R,R) -9) in the main chain (Fig. 3a). As expected, **P4** in the THF solution containing 1% H_2O underwent depolymerization, lowering M_n to 5.4 kDa within a day. After three days, **P4** was further hydrolyzed to 3.6 kDa, which is 1/4 of the initial molecular weight (Fig. 3b). However, as shown by SEC analysis, the molecular weight of (R,R) -P1 from the complete DCRP did not change under identical conditions even after seven days (Fig. 3c). Moreover, (R,R) -P1 remained stable in CDCl_3 in an NMR tube for three days, but even a small amount of residual acid in CDCl_3 was sufficient to depolymerize **P4** within 11 h, as seen by the reduction in the imine signal (δ 8.3 ppm) and the significant increase in the aldehyde signal (δ 10.0 ppm) (Fig. S3†).

The DCR of small molecules undergoes complete stereo-specific inversion of stereochemistry at rt because the corresponding transition state (TS) bears all the aryl substituents in pseudo-equatorial positions making this pathway most



Fig. 3 Control experiment to compare stability against hydrolysis. (a) Scheme for **P4**. (b) SEC traces of **P4** measured at the initial and measured one/three days after dissolving in $\text{THF} : \text{H}_2\text{O}$ (99 : 1). (c) SEC traces of (R,R) -P1 measured at the initial and seven days after dissolving in $\text{THF} : \text{H}_2\text{O}$ (99 : 1).



Fig. 4 (a) Table for optical rotation values showing how temperature affects the stereospecificity of tandem DCRP. (b) Circular dichroism spectra of the resulting polymers (2.5 mg mL⁻¹ in CHCl₃ at rt, 0.2 mm cell).

kinetically preferred.⁴² Likewise, to investigate the stereochemistry of the resulting polymers, we prepared **P1** at various temperatures and compared their optical rotations ($[\alpha]_D^{24}$) (Fig. 4a and Table S4[†]). While **P1** synthesized from (S,S)-1 at rt exhibited the highest $[\alpha]_D^{24}$ of +135.95, the values significantly decreased from +82.82 to +25.92 as the polymerization temperature increased from 50 °C to 140 °C (entries 1–6 in Fig. 4a). In addition, (S,S)-**P1** obtained from the enantiomer (R,R)-1 at rt (entry 8 in Table 1) showed an expected $[\alpha]_D^{24}$ of similar absolute value but the opposite sign (–134.71) (entry 7 in Fig. 4a). Lastly, **P1** prepared from the racemic mixture or *meso*-isomer of **1** resulted in $[\alpha]_D^{24}$ close to 0 (entries 8 and 9 in Fig. 4a). To further support this trend, we measured circular dichroism spectra, which showed a decrease in the amplitudes of the two Cotton effects with an increase in the reaction temperature (Fig. 4b). Furthermore, (R,R)-**P1** and (S,S)-**P1** synthesized at rt showed Cotton effects of the same amplitude but of the opposite sign. According to the computational studies, the next alternating pathway *via* the second lowest TS (7.7 kcal mol⁻¹ higher) would produce a *meso* product, an achiral diastereomer.⁴² Therefore, it is suspected that the higher reaction temperature led to more *meso* products in **P1**, thereby lowering $[\alpha]_D^{24}$ and the amplitude of Cotton effects accordingly.

To confirm the effect of temperature on *meso*-isomer formation, we thoroughly analyzed the ¹H-NMR spectra of **P1** from (S,S)-1 and *meso*-**P1** synthesized at various temperatures (Fig. 5a and S4[†]). As the polymerization temperature increased, another peak labeled as *g'* appeared and gradually increased at δ 13.38 ppm (Fig. 5a and S4a–f[†]). Notably, apart from the *g* and



Fig. 5 (a) ¹H NMR spectra (in CD₂Cl₂) showing RAHB O–H signals of various **P1** (in Table S4[†]) synthesized from chiral and *meso* diamines at various reaction temperatures. (b) Plot showing the relationship between $\ln([\alpha]_D^{24})$ and stereospecificity.

g' peaks, an additional RAHB O–H signal (*g''* at δ 13.05 ppm) started to grow for **P1** synthesized at higher temperatures. Fortunately, the last *g''* peak was easily identified as the *meso* diastereomer as it perfectly matched with the authentic sample of *meso*-**P1** prepared from *meso*-1 at rt (Fig. 5a and S4g[†]). From these data, we came to a plausible conclusion that the three types of RAHB O–H signals are due to diad tacticities of the salen side-chains in the following cases: (i) *g* for the homo diad (cm³) from the chiral-salen-rich units; (ii) *g''* for the homo diad (mm) from the achiral *meso*-salen-rich units; (iii) *g'* for the hetero diad (cm) from the chiral and *meso*-salen-rich units adjacent to each other. Based on this assignment, the content of the chiral salen unit depending on the temperature was estimated from the integration of all RAHB O–H signals (Fig. S5a[†]). Consistent with the trend observed in $[\alpha]_D^{24}$, the chiral salen content also decreased from 100% to 60.7% as the temperature increased from rt to 140 °C. Moreover, one could plot $[\alpha]_D^{24}$ to the chiral content, and they showed an exponential relationship (Fig. S5b[†]). Taking the natural logarithm of $[\alpha]_D^{24}$ provided a good linear relationship with the chiral content, which should allow for some predictability (Fig. 5b). In short, increasing the reaction temperature lowered the stereospecificity as well as $[\alpha]_D^{24}$ due to the formation of the *meso*-isomer.

One useful application of salen moieties is their potential as a metal chemosensor.^{48–50} Since (R,R)-**P1**–**P3** contain salen side-chains, their UV/vis absorption and fluorescence responses to various metal cations in THF/H₂O (9 : 1 v/v) were investigated (Fig. 6 and S8–S12[†]). Interestingly, all the metal cations showed insignificant fluorescence except for Zn²⁺, which showed obvious turn-on fluorescence enhancement ($I - I_0$) (Fig. 6a, b and S8[†]). On the other hand, no sensing ability was observed



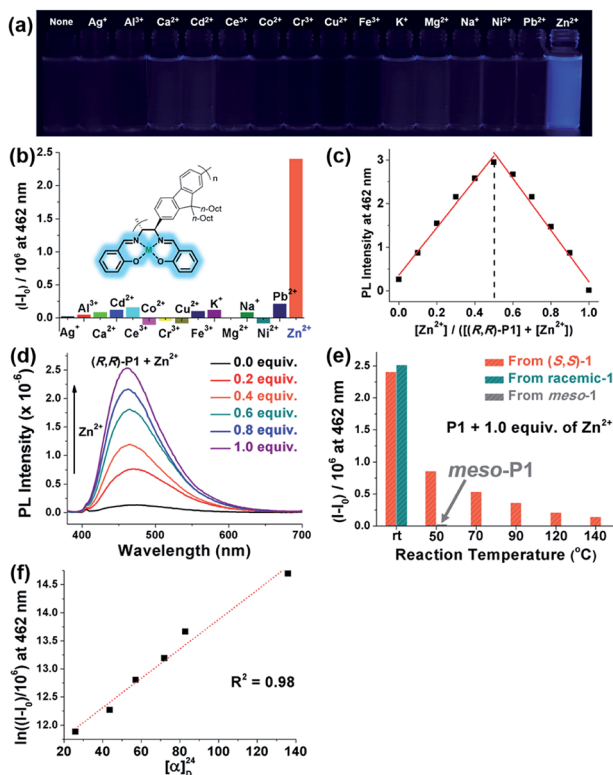


Fig. 6 Metal sensing experiments using **P1** in THF : H₂O = 9 : 1 (v/v) solution (10 μM) at 298 K. (a) Photoluminescence of (*R,R*)-**P1** solution in the absence or presence of 1.0 equiv. of various metal cations under 365 nm UV irradiation. (b) Emission changes of (*R,R*)-**P1** in the presence of 1.0 equiv. of various metal cations. (c) Job plot of (*R,R*)-**P1** vs. Zn²⁺ (total concentration = 20 μM). (d) Emission spectra of (*R,R*)-**P1** with increasing amounts of Zn²⁺ (0.0–1.0 equiv.). (e) Effect of polymerization temperature on the Zn²⁺ sensing performance of **P1**. (f) Correlation between fluorescence enhancement ($I - I_0$) and optical rotation ($[\alpha]_{\text{D}}^{24}$) of **P1**.

for **P4**, which only contains diimine side-chains without –OH groups (Fig. S9†). The Job plot using (*R,R*)-**P1** showed the maximum fluorescence intensity at 0.5 mole fraction, indicating 1 : 1 complexation of Zn²⁺ to the salen group (Fig. 6c). Increasing the concentration of Zn²⁺ from 0.0 to 1.0 equiv. resulted in a gradual increase in the fluorescence intensities. In particular, (*R,R*)-**P3** was an excellent Zn²⁺ sensor showing a high amplification of fluorescence (I/I_0) up to 73-fold and a limit of detection value of 0.577 μM (37.7 ppb) (Fig. 6d, S10 and S11†). Notably, **P1** prepared at various temperatures showed different sensing performances (Fig. 5e and S12†). **P1** synthesized at rt using enantiopure (*S,S*)-**1** or the racemic mixture of chiral **1** exhibited the highest I/I_0 of 19-fold, whereas this I/I_0 decreased significantly with the temperature to 4.8-fold (25% compared to that at rt) at 50 °C, 2.6-fold (14%) at 90 °C, 1.9-fold (10%) at 120 °C, and even further down to 1.5-fold (8%) at 140 °C. The $I - I_0$ and $[\alpha]_{\text{D}}^{24}$ also showed an exponential relationship and applying the natural logarithm to $I - I_0$ provided a good linear relationship with $[\alpha]_{\text{D}}^{24}$, allowing for prediction of the sensor performances (Fig. 6f and S12c†). Furthermore, to examine the effect of the *meso*-salen unit, we conducted a sensing

experiment using *meso*-**P1** prepared at 50 °C to achieve comparable conversion (entry 9 in Table 1 and Fig. S4h†). Interestingly, *meso*-**P1** showed no sensing ability for Zn²⁺ at all, indicating that the stereochemistry of the salen moiety was crucial for Zn²⁺ detection (Fig. 6e). Therefore, we concluded that the high sensing performances of (*R,R*)-**P1**–**P3** synthesized at rt were due to the exclusive formation of the chiral salens, which formed a tetrahedral geometry suitable for selective Zn²⁺ binding.⁵⁰ However, as the reaction temperature increased, the degree of DCR to the undesired *meso*-isomer increased, and their sensing performance decreased due to the resulting square-planar geometry from the *meso* configuration in which selective Zn²⁺ detection is impossible.

Conclusions

In conclusion, we developed a new tandem diaza-Cope rearrangement polymerization (DCRP) to synthesize high-molecular-weight polymers up to 33.3 kDa with excellent stability against hydrolysis. The key factor for the success was the introduction of a defect-free, rapid, and thermodynamically favored DCR process into the polymerization mechanism by forming C–C bonds almost simultaneously with efficient imine formation. Taking advantage of the highly stereospecific DCR, we prepared several highly enantiopure polymers having either (*R,R*)- or (*S,S*)-salen moieties. Lastly, these polymers showed excellent performance as turn-on chemosensors selective for Zn²⁺ detection.

Conflicts of interest

There are no conflicts to declare.

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