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Facile one-pot synthesis and self-healing properties of tetrazole-based metallopolymers in the presence of iron salts†

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As promising intelligent materials, self-healing materials have the capability of healing or repairing inflicted damage. Simply, inexpensive and commercial synthetic process is one of the hard issues. By using a facile one-pot two-step methodology, polyacrylonitrile-*r*-poly(butyl acrylate) (PAN-*r*-PnBA) was successfully prepared and modified to Fe³⁺/tetrazole-based metallopolymers (MPs) with significant self-healing properties. FeCl₃·6H₂O was used as a catalyst to modify cyano groups with sodium azide (NaN₃) and generate tetrazole group for coordinating with FeCl₃·6H₂O itself. The results indicated that the prepared MPs without any stimulus, can rapidly and autonomously heal inflicted damage at ambient temperature. These self-healing materials which were rapidly prepared with commercial and inexpensive substrates could facilitate the potential for application in a wide range.

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Introduction

Self-healing^{1–5} is one of the most fascinating and challenging abilities to heal or repair inflicted damage. Nowadays, self-healing materials^{6–12} are under rapid development to improve the lifetime, safety, energy efficiency and environmental impact of synthetic materials. The past few decades have witnessed a rapid development in exerting such features into artificial systems where polymers are used for self-healing applications. Several self-healing strategies, such as the utilization of healing agents,¹³ reversible covalent^{14,15} and noncovalent bonds,¹⁶ have been reported in polymer systems. Different dynamic non-covalent bonding interactions, such as hydrogen-bonding,^{17,18} ionic interactions,^{19,20} π–π stacking²¹ and metal–ligand interaction,²² have been used to improve the properties of materials with self-healing ability.

Metallopolymers (MPs) or metal–ligand (M–L) complexes^{23,24} can be designed as functional materials with high mechanical properties based on a wide range of thermodynamic and kinetic parameters. Some prominent examples of self-healing materials have been reported by using reversible M–L interactions. Furthermore, the MPs prepared by the interaction of M–L have excellent mechanical properties. For example, a series of

functional materials, based on the coordination of dopamine^{24,25} and metal ions, are reported with notable self-healing properties. Harrington M. J. *et al.* reported²⁶ that self-healing functional materials like mussel byssal threads were developed through the coordination between Fe and catechol ligands. A novel polymer with bidentate triazole pyridine ligands, which cross-linked with Fe²⁺ or Co²⁺ salts,²⁷ was developed with notable self-healing performance. As mentioned above, these methods have the advantages of providing a favorable selectivity and design of MPs, but most of the polymers are achieved by designing complex monomer polymerization and difficult synthetic process.

Tetrazoles^{28,29} which have a five membered ring containing one carbon and four nitrogen atoms are heterocyclic compounds. Polyvinyltetrazoles (PVTs)^{30–32} and its derivatives have attracted broad attention because of their interesting properties (high thermostability, biological activity and good solubility) and wide applications (polyelectrolytes, distinctive complex, dynamite and biocompatible material). However, the vinyltetrazole³³ is not a common monomer for commercial source because of the difficult synthesis. To the best of our knowledge, there have been no reports and research on self-healing materials by using coordination with tetrazoles groups and metal ions from PVTs or its derivatives.

Polyacrylonitrile (PAN)^{34–36} and its copolymers are commercial polymers for preparing fibers, membranes and so on. By using simple and efficient synthesis, the cyano groups in PAN can react with sodium azide (NaN₃) to prepare PVT^{30–32,37,38} with various molecular weights. Matyjaszewski *et al.*³⁹ reported well-defined PAN and PAN-*co*-PSt prepared by atom transfer radical polymerization (ATRP). Then, the prepared polymers were

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successfully modified by the “cyano-azide” type of click chemistry with NaN_3 as a reactant and zinc chloride as a catalyst. After the click chemistry of cyano-azide under microwave radiation, functional PVT-grafted resins⁴¹ with high capacity for the adsorption of heavy metal ions were successfully prepared by Wei and coworkers.

Herein, we designed a PAN-*r*-PnBA random copolymer with the M-L complexes strategically embedded in the soft matrix having low glass transition temperature (T_g). By using a facile one-pot two-step methodology, polyacrylonitrile-*r*-poly(butyl acrylate) (PAN-*r*-PnBA) was successfully prepared and modified to MPs with significant self-healing properties. Iron(III) chloride hexahydrate ($\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$) was used as a catalyst to modify cyano groups with sodium azide (NaN_3) and generate tetrazole group for coordinating with $\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$ itself. The prepared MPs without any stimulus have rapid and autonomous self-healing capability with good mechanical strength under ambient conditions.

Experimental section

Materials

Acrylonitrile (AN, >99%) and *n*-butylacrylate (nBA, >99%) were purchased from Tianjin Damao Chemical Reagents Factory (Tianjin, China). AN and nBA were distilled under reduced pressure. Sodium azide (NaN_3 , >97%), iron(III) chloride hexahydrate ($\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$, >99%), were purchased from Zhengzhou Alfa Chemical Company (Zhengzhou Alfa, China). *N,N*-Dimethylformamide (DMF, 99.9%), tetrahydrofuran (THF, 99.9%), dimethyl sulfoxide (DMSO, 99.9%) and methanol (99.9%) were purchased from Tianjin Ruijin Chemistry Company (Zhengzhou Alfa, China) and used as received. All other chemicals were obtained from Shanghai Chemical Reagents and used as received unless mentioned. Deionized water was obtained from a UPH-I-40L super-pure water machine (Ludong University Deionized Water).

Typical procedure for preparation of Fe(III)-tetrazole-based metallopolymers

The molar ratio of $[\text{AN}]_0/[\text{nBA}]_0/[\text{AIBN}]_0 = 100/100/0.2$ was shown as a typical radical polymerization procedure. After adding AN (15 mL, 0.23 mol), nBA (33 mL, 0.23 mol), AIBN (80 mg, 0.46 mmol) and DMF (15 mL), the dried single-necked flask flame sealed and then transferred into an oil bath at 65 °C with a stir bar. After the desired time, the single-necked flask continued to react with NaN_3 (0.3 g, 4.6 mmol), $\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$ (1.2 g, 4.6 mmol) and a small amount of DMF in an oil bath at 120 °C for about 2 h. Afterward, the flask was cooled to ambient temperature, then the contents were dissolved in DMF. The polymer solution was precipitated into methanol (~300 mL) with stirring. After filtration, the obtained polymer was dried until constant weight at 30 °C under vacuum oven.

Characterization

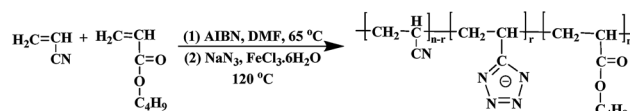
The FT-IR spectra measurements were performed on a MAGN-AIR-550 FTIR. The UV-vis spectra were performed with

a Beijing Purkinje UV-visible spectrophotometer T6 New Century equipped with Persee/UV Probe software. Differential scanning calorimetry (DSC) of copolymer and MPs was characterized by using a Netzsch DSC 204 F1 Phoenix instrument. The same test condition with a heating rate of 10 °C min^{-1} was used from -120 to 50 °C under a nitrogen atmosphere. Energy dispersive spectrometer (EDS) was measured on Bruker XFlash 6160 under a nitrogen atmosphere. The rheology data were performed on a DFR-2 Rheometer from TA Instruments. Stainless steel parallel plates with a diameter of 25 mm were carried for the test of all MPs and the gap between plates was set at 1 mm. For the convenience of tests, all MPs were cut into pellets with a diameter of 25 mm and about 1 mm thickness. Tensile testing was measured *via* LDW-5 Microcomputer control electronic universal material testing machine from Shanghai Songdun Machine Equipment Co., Ltd. The images of healing process were performed on a DMM-300C metallurgical microscope (Shanghai Caikon Optical Instrument Co., Ltd.).

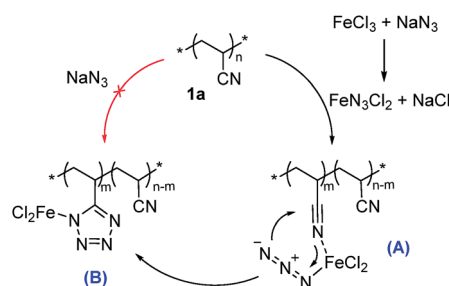
Results and discussion

Preparation of Fe³⁺/tetrazole-based metallopolymers (MPs)

The synthesis of the tetrazole-based metallopolymers (MPs) is described in Scheme 1 through one-pot two-step. The PAN-*r*-PnBA was synthesized *via* radical polymerization (Scheme 1a). In one-pot, the different proportion of NaN_3 and $\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$ was added directly to proceed click reaction after 2 hours. Initially, $\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$ reacted⁴⁰ with NaN_3 to produce the FeN_3Cl_2 catalytic species. The [3 + 2] cycloaddition between the $\text{C}\equiv\text{N}$ bond of cyano **1a** and FeN_3Cl_2 took place readily to form the intermediate (A); pre-coordination of the nitrogen atom of the CN group of **1a** with iron azide to form complex (B) would accelerate this cyclization step (Scheme 2). After washing by methanol, MPs which tetrazole moieties coordinated with ferric ion were simply and easily obtained with the self-healing capability.



Scheme 1 The scheme for the prepared of MPs using one pot two step.



Scheme 2 Proposed mechanism for the modification of PAN.



The FT-IR spectra for MPs with different amounts of $\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$ and NaN_3 are shown in Fig. S1.† For PAN-*r*-PnBA, a typical peak at 2250 cm^{-1} is the characteristic absorption of cyano ($\text{C}\equiv\text{N}$) stretching vibration. A typical peak at 1726 cm^{-1} attributes to the $\text{C}=\text{O}$ stretching vibration of the ester groups in the nBA. For MPs, based on modified PAN-*r*-PnBA (MPAN-*r*-PnBA), the peak at 2250 cm^{-1} becomes weak due to that the $\text{C}\equiv\text{N}$ in PAN is partially modified to tetrazole groups. The broad peak at 3400 cm^{-1} represents the skeleton stretching vibration absorption peak of the tetrazole ring. In addition, the new peak at 1671 cm^{-1} attributes to the characteristic absorption of the Cl-Fe stretching vibrations. These results indicate that the chemical modification of MPs is obtained. Energy dispersive spectrometer (EDS) spectrum test of the MPs is also shown in Fig. 1(a). From the results, it can be seen that the metal element Fe is contained in addition to the basic elements C, N, O of the copolymer. The elemental proportions of C, N, O, Cl and Fe are 60.4%, 17.2%, 15.8%, 3.6% and 3.0%, respectively, which is consistent with the results of coordination with the tetrazole group. Fig. 1(b) shows the UV-vis spectra of PAN-*r*-PnBA and MPs. The concentration of absolute absorbance of the three polymers is 2.0 mg mL^{-1} . From 800 nm to 250 nm, the relatively stronger absorption at about 270 nm for M-L complexation than that of PAN-*r*-PnBA indicates the presence of coordination between Fe^{3+} and tetrazole ring,²² which are prepared through the reaction of cyano group with NaN_3 catalyzed by $\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$. By using differential scanning calorimetry (DSC) in the heating rate $10\text{ }^\circ\text{C min}^{-1}$, the glass transition temperature (T_g) of copolymer and MPs is also measured. From Fig. 1(c), the results show that the T_g of PAN-*r*-PnBA is about $8.14\text{ }^\circ\text{C}$. Copolymer PAN-*r*-PnBA exhibits a T_g which is parently lower than that of the homopolymer PAN. This proves that the presence of nBA units in PAN-*r*-PnBA could dramatically decrease its glass transition temperature. It is also found that MPs, in which the cyano groups of copolymer are partly replaced to tetrazole groups, could dramatically decrease its glass transition temperature. The T_g of MPs with 25 : 1, 50 : 1, 75 : 1 are $2.05\text{ }^\circ\text{C}$, $-1.60\text{ }^\circ\text{C}$, $-2.34\text{ }^\circ\text{C}$ and $-8.50\text{ }^\circ\text{C}$, respectively. The increasing T_g from $-8.50\text{ }^\circ\text{C}$ to $2.05\text{ }^\circ\text{C}$ is caused by the increasing amount of Fe^{3+} and the coordination between the tetrazole groups and metal ions. These results demonstrate that coordination between the tetrazole groups and metal ions can enhance the intermolecular interactions between the polymer chains more drastically, thus significantly affecting the self-healing properties.

Rheological analysis of MPs

The self-healing properties of copolymers and their MPs can be provided insight into the rheological characterization. By using a parallel plate rheometer, the rheological properties of copolymers and MPs are investigated at ambient temperature. For the self-healing materials, the storage modulus (G' , correspond to the energy storage after materials received deformation) demonstrates the energy storage of material after straining, while loss modulus (G'' , correspond to the energy loss after the materials received deformation) denotes energy loss of material

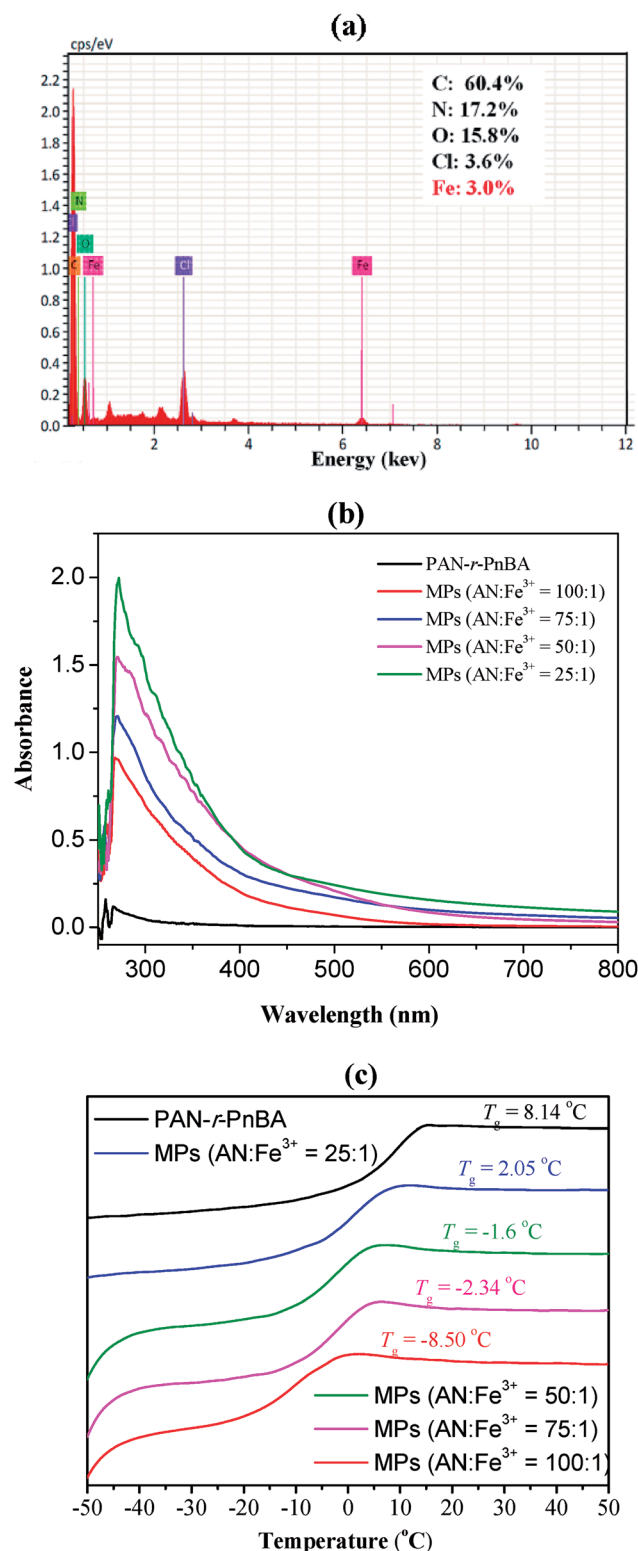


Fig. 1 (a) Element distribution spectra (EDS) of the MPs; (b) UV-vis spectra of PAN-*r*-PnBA and the different ratio of MPs in DMF solution (2.0 mg mL^{-1}); (c) DSC curves of the PAN-*r*-PnBA and different ratio of MPs with the heating rate $10\text{ }^\circ\text{C min}^{-1}$.

during the generation of deformation. G' and G'' are critical to characterizing the mechanical and self-healing properties of materials. Strain sweep measurement of MPs is performed at



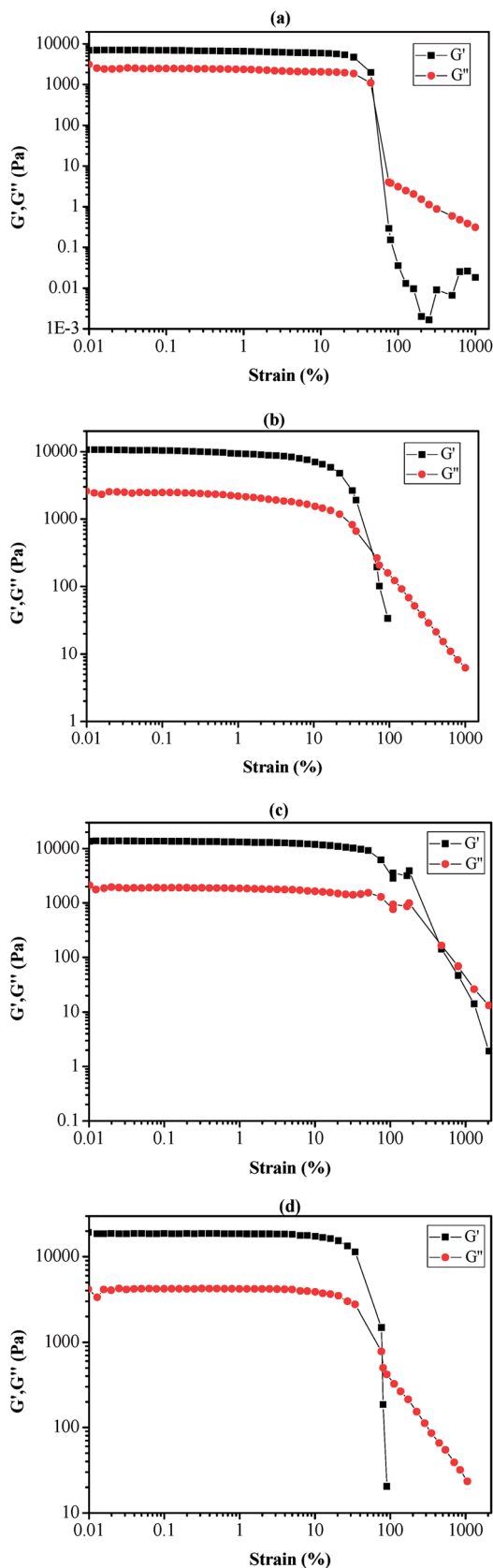


Fig. 2 The values of storage modulus (G') and loss modulus (G'') are investigated through dynamic strain sweep. Polymerization conditions: $[AN]_0/[nBA]_0/[AIBN]_0/[FeCl_3 \cdot 6H_2O]_0/[NaN_3]_0 = 100/100/0.2/1/1$ (a), $75/75/0.15/1/1$ (b), $50/50/0.1/1/1$ (c), $25/25/0.05/1/1$ (d), respectively.

25 °C, which shows a typical solid state rheological property (Fig. 2 and S2†). For PAN-*r*-PnBA, the G' is probably closed to G'' in Fig. S2,† which suggests that it present poor viscoelasticity. In Fig. 2, the G' is larger than G'' , which proves that the MPs with different molar ratio boast of excellent self-healing potential.

In the initial state with the 0.01% strain, it is worth noting that the G'/G'' is 6986.8 Pa/3139.8 Pa, 10 717.5 Pa/2592.2 Pa, 13 686.9 Pa/2592.2 Pa and 19 354.6 Pa/4154.6 Pa with the molar ratio of $[polymer]_0/[NaN_3]_0/[FeCl_3 \cdot 6H_2O]_0$ of 100/1/1, 75/1/1, 50/1/1 and 25/1/1, respectively. It is notable that G' increases with the increasing molar ratio and is all bigger than G'' for MPs. The increase of G' may be caused by the interactions of metal–ligand bonding by 4VT functional group with ferric ion. In addition, it is clear that G' is larger than G'' , when the strain is in the range of 0.01% to 10% (Fig. S2†) to 45% (Fig. 2(a)), 60% (Fig. 2(b)), 470% (Fig. 2(c)), 80% (Fig. 2(d)), with the increased amount of $[NaN_3]_0$ and $[FeCl_3 \cdot 6H_2O]_0$, respectively. This results show the prepared MPs materials of 50/1/1 could withstand relatively large deformation damage. Moreover, it indicates that MPs show excellent adhesion behavior, which was the result of the modified cyano group to 4VT group at ambient temperature. However, the value of G' is less than G'' with further increasing the strain.

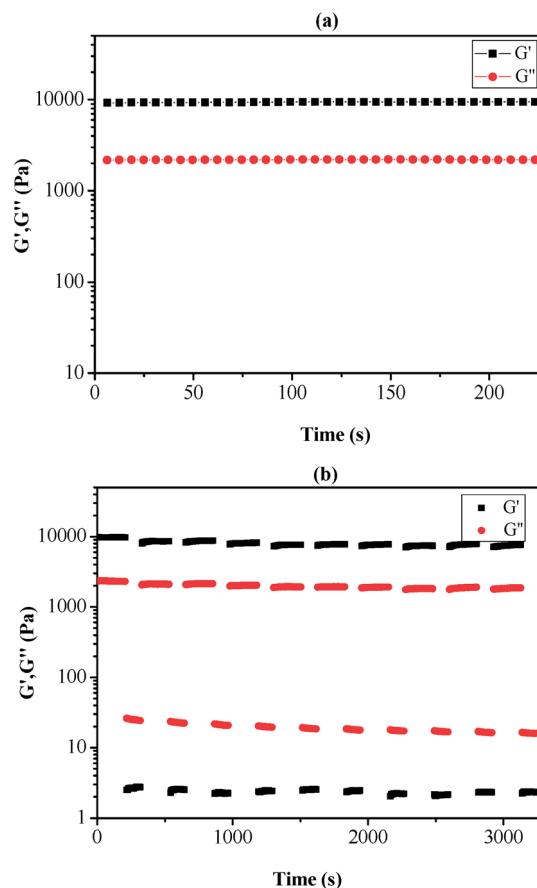


Fig. 3 The values of storage modulus (G') and loss modulus (G'') with MPs (AN : $Fe^{3+} = 75 : 1$) are investigated by (a) an immediate recovery after the 1000% strain deformation ($t = 200$ s, $\gamma = 1\%$), (b) repeated dynamic strain amplitude cyclic test ($\gamma = 1\%$ or 400%).



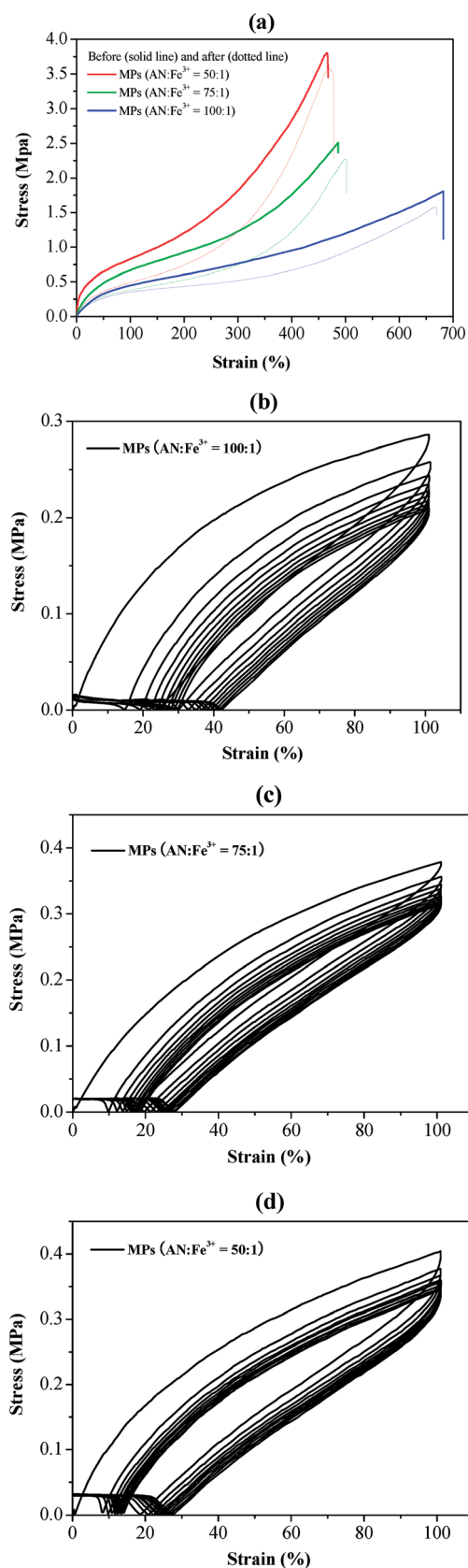


Fig. 4 Tensile stress–strain curves of self-healing MPs (a), mechanical recovery curves of MPs with the molar ratio of 100 : 1 (b), 75 : 1 (c) and 50 : 1 (d) (AN : Fe³⁺) under 10 cyclic tensile tests.

After large strain deformation ($\gamma = 1000\%$), a 1% of strain is immediately applied for a period of time and it is found that the mechanical property of MPs could fully recover without any loss (Fig. 3(a) and S3†). Repeated dynamic strain step tests are carried out on MPs to investigate the self-healing property, in which γ is alternated with 1% (200 s) and 400% (200 s) for ten cycles. As shown in Fig. 3(b), when a lower strain ($\gamma = 1\%$) is applied to the 50 : 1 MPs, the value of G' and G'' keep constant (G' is larger than G''). However, when subjected to a large strain ($\gamma = 400\%$), G' drop dramatically within a few seconds and become lower than G'' in value. After large strain deformation ($\gamma = 400\%$), a 1% of strain is immediately applied for a period of time and it is found that the mechanical property of MPs can recover to its initial value instantaneously without any loss (Fig. 5(b)). After ten cycles test, the G' and G'' can still recover to initial value without significant decrease. This phenomenon indicates that all the fabricated MPs are totally reversible and endowed with outstanding self-healing property.

Study on self-healing performance

The stretchability of the MPs of different ratio (100 : 1, 75 : 1 and 50 : 1), is characterized with the same gauge length and stretching speed. The stretch process is shown as Fig. S4.† The mechanical performance of tensile strength can be increased by the increasing percent of the incorporation of PVT and the amounts of Fe³⁺. For example, By increasing the ratio from 100 : 1, 75 : 1 to 50 : 1, tensile strength is remarkably increased from 1.80 MPa, 2.5 MPa to 3.8 MPa, respectively. In our self-healing experiment, MPs films are typically cut into two pieces and then put together to heal itself about 4 hours. The self-healing efficiency of MPs films, which is closed to the ratio of the original to self-healing tensile strength, are also illustrated in Fig. 4(a). Without any external stimuli, the tensile modulus can reach the original levels after about 4 hours at ambient temperature. The self-healing efficiency of 100 : 1, 75 : 1 and 50 : 1, is about 85.7%, 90.8% and 94.7%, respectively. The results show that different molar ratio of MPs films have excellent self-healing properties. Additionally, by using loading-unloading cyclic tensile tests, the mechanical properties of self-healed MPs are discussed in depth to study the effects of coordination of 4VT and Fe³⁺. Self-healed MPs are recycled to 10

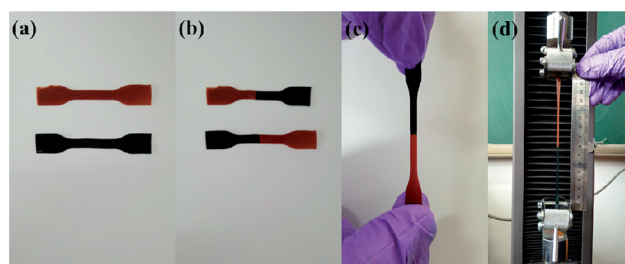


Fig. 5 (a) The stretch strips added with or without dyes (malachite green); (b) cut two different colors of the stretch strips and contact them together; (c) MPs were healed at room temperature for about 30 min; (d) tensile testing were measured via Microcomputer control electronic universal material testing machine.



cyclic tensile tests between the strain 0% and 100%. These MPs are not broken and returned to their original length. For instance, the stress of these MPs is 72.8%, 83.2% and 85.5% of its original values about 10 strain cycles (Fig. 4(b–d)) in the molar ratio of 100 : 1, 75 : 1 and 50 : 1, respectively. It indicates that all of the MPs can reversibly break and reform in the process of loading-unloading cycles. Therefore, the coordination of Fe^{3+} and tetrazole group deliver remarkable self-healing performance.

In order to clearly demonstrate the self-healing ability of MPs, the healing experiments were carried out by two different

colored strips (added with or without malachite green). Fig. 5(a) shows that the two strips have different colors. Afterwards, the two strips were cut from the middle, and the different colored strips were intertwined and healed at ambient temperature for 30 min, as shown in Fig. 5(b). Fig. 5(c) shows that the two color strips have been healed together. The stretched strip was subjected to a tensile test in Fig. 5(d). The results show that MPs exhibits excellent self-healing properties at ambient temperature.

To observe the self-healing process, PAN-*r*-PnBA and different molar ratio of MPs films are sealed at ambient

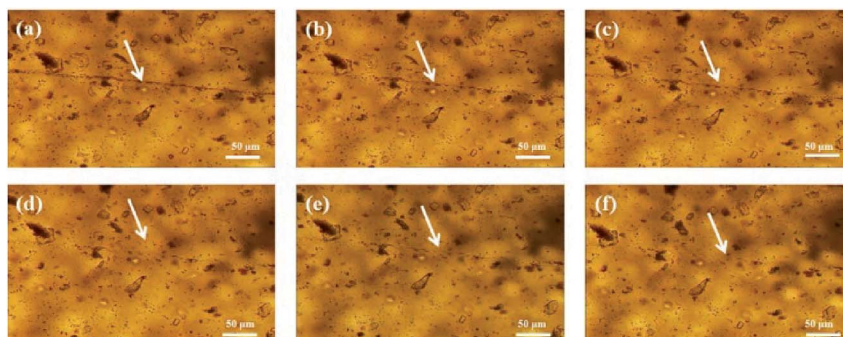


Fig. 6 Self-healing processes of MPs at ambient temperature (a) 0 min, (b) 10 min, (c) 30 min, (d) 1 hours, (e) 2 hours and (f) 4 hours, respectively. Polymerization conditions: $[\text{AN}]_0/[\text{nBA}]_0/[\text{AIBN}]_0/[\text{FeCl}_3 \cdot 6\text{H}_2\text{O}]_0/[\text{NaN}_3]_0 = 100/100/0.2/1/1$.

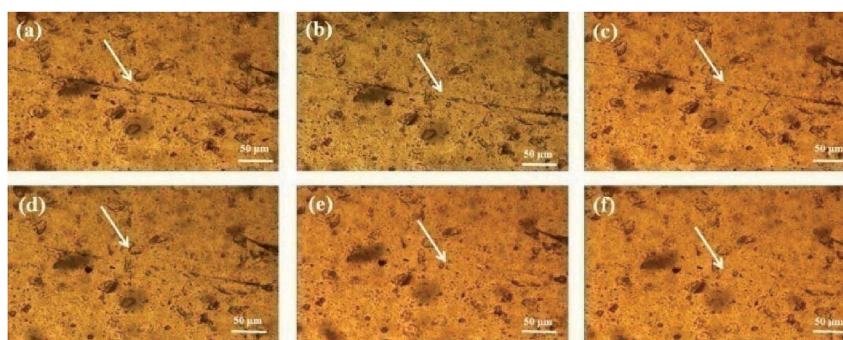


Fig. 7 Self-healing processes of MPs at ambient temperature (a) 0 min, (b) 10 min, (c) 30 min, (d) 1 hours, (e) 2 hours and (f) 4 hours, respectively. Polymerization conditions: $[\text{AN}]_0/[\text{nBA}]_0/[\text{AIBN}]_0/[\text{FeCl}_3 \cdot 6\text{H}_2\text{O}]_0/[\text{NaN}_3]_0 = 75/75/0.15/1/1$.

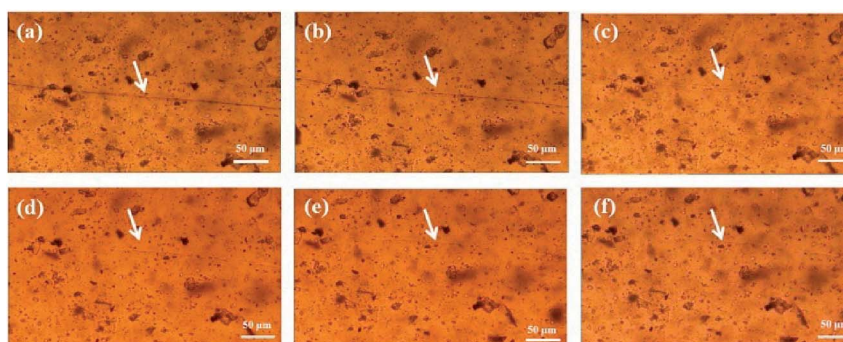


Fig. 8 Self-healing processes of MPs at ambient temperature (a) 0 min, (b) 10 min, (c) 30 min, (d) 1 hour, (e) 2 hours and (f) 4 hours, respectively. Polymerization conditions: $[\text{AN}]_0/[\text{nBA}]_0/[\text{AIBN}]_0/[\text{FeCl}_3 \cdot 6\text{H}_2\text{O}]_0/[\text{NaN}_3]_0 = 50/50/0.1/1/1$.



temperature and observed in metallurgical microscope. Then the films are cutted off with a scalpel and placed on glass substrates. An optical microscope is used to observe the self-healing process of these films. Fig. 6–8 and S5† show the variation of cracks at different times, which show self-healing properties of MPs. In Fig. S5,† for the PAN-*r*-PnBA copolymer without the coordination between tetrazole group and Fe³⁺, the crack of PAN-*r*-PnBA copolymer dose not heal even after 12 hours. After the modification of PAN-*r*-PnBA, the MPs films show different levels of self-healing property. As shown in Fig. 6, it gives the result that the crack is completely repaired after 4 hours at ambient temperature. The 75 : 1 MPs film can rapidly heal the crack after 1 hour without additional treatment (Fig. 7). Moreover, the self-healing films of 50 : 1 is almost completed the self-healing process within 30 min (Fig. 8) at ambient environment. In addition, the self-healing process of MPs film is taken a video. In the video, the MPs is completely cut off from the middle and healed at room temperature. After two minutes, the MPs can be healed together. All of these results indicates that MPs with highly tetrazole groups have excellent self-healing properties.

Conclusions

In conclusion, we provided a novel inspiration for the preparation of functionself-healing MPs by using one-pot two-step. After the modification of PAN's copolymers with FeCl₃·6H₂O as a catalyst, MPs used tetrazole group for coordinating with FeCl₃·6H₂O itself. The prepared Fe³⁺/tetrazole-based MPs showed significant self-healing ability. These MPs could rapidly achieve the self-healing of crack and could show the potential for application in a wide range. Furthermore, the simple, inexpensive, biocompatible and commercial nature of the MPs has potential application in replacing existing materials.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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