Cross-linking modification of ammonium polyphosphate via ionic exchange and self-assembly for enhancing fire safety properties of polypropylene

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Abstract: Modified ammonium polyphosphate (MAPP) as a novel mono-component intumescent flame retardant (IFR) was prepared via the ionic exchange between ammonium polyphosphate (APP) and piperazine sulfonate, which is synthesized by self-assembly using 1-(2-Amioethyl) piperazine (AEP) and p-amino benzene sulfonic acid (ASC) as raw materials. This all-in-one IFR integrating three functional elements (carbon, acid, and gas source) showed more efficient flame retardancy and excellent smoke suppression as well as better mechanical properties than the conventional APP. The incorporation of 22.5 wt.% MAPP into polypropylene (PP) eliminated the melt dripping phenomenon and passed the UL-94 V-0 rating. The results of the cone calorimetry test (CCT) revealed that the release of heat, smoke, and CO is significantly decreased, demonstrating that this novel IFR endows PP with excellent fire safety more effectively. For PP/MAPP composites, a possible IFR mechanism was proposed based on the analysis of the pyrolysis gas and char residues.

Keywords: Ionic exchange; self-assembly; Modified ammonium polyphosphate; Polypropylene; Combustion behavior
1. Introduction

With the wide application of polymers in construction, electronics, and other fields, the possibility of polymeric materials being exposed to flames has significantly increased [1]. However, most polymeric materials are liable to combust, possessing the characteristics of a high heat release rate [2]. Therefore, improving the fire resistance of polymeric materials is of great significance for diminishing fire hazards and protecting the safety of people's lives.

Polypropylene, one of the commercially available general-purpose plastics, has been applied to agriculture, industry automobile manufacturing, and film packaging, owing to its low toxicity, excellent mechanical properties, electrical insulation, and chemical resistance [3,4]. However, its further application and development are restricted by its disadvantages including the inherent flammability, severe dripping behavior, and the release of smoke and toxic gases during combustion [5]. Therefore, it is particularly significant to improve both its flame retardancy and smoke suppression performance simultaneously. Recently, the IFR system has completely replaced the halogen-containing flame-retardant system due to its high efficiency, low smoke, and low toxicity [6,7]. It is well known that the conventional IFR system is made of the acid source, blowing agent, and carbonizing agent [8]. Although APP can act as both an acid source and a blowing agent, it still fails to achieve a good IFR effect in case of the usage of APP alone [9]. Many strategies have been developed to improve its flame retardancy. The most important choice is to microencapsulate APP with silane coupling agents [10], polyurethane (PU) [11], melamine-containing polyphosphazene (PZMA) [12], or melamine-formaldehyde (MFT) [13] via in situ polymerization. The microencapsulation modification of APP has improved the flame retardant efficiency of PP more or less, but it is still needed to be compounded with the charring agent which preparation procedures were complicated and time-consuming. Therefore, the preparation of a mono-component flame retardant which integrates all three sources of IFR has become a research hotspot. Shao et al. tried to modify APP with various organic amines, such as ethylenediamine (EDA) [14], ethanolamine (ETA) [15], diethylenetriamine (DETA) [16], and piperazine (PA) [17] via ionic exchange to prepare three-in-one flame retardant. This method was facile whereas the flame retardancy was still needed to be improved. Thus, a novel modification technology should be developed to overcome this problem.

Self-assembly technology has been applied to produce substances with specific structures by exploiting the interaction between structural units, such as hydrogen bonding and ionic interactions [4,18-20]. Recently, self-assembly technology has been adopted to prepare flame retardants due to its facile preparation method and environmentally friendly. Su et al. [21] modified APP with melamine-formaldehyde and phytic acid (MF-PA) as building blocks by self-assembly reaction. The results demonstrated that the dispersivity of the modified APP in the matrix has been dramatically improved, and the LOI value of the IFR-PP system reached 35% and the UL-94 V-0 rating was achieved at a loading level of 25 wt.% of modified APP/ CFA (4:1, weight ratio). Jin et al. [22] prepared a novel macromolecular IFR (AM-APP) via supramolecular reactions between melamine and p-amino benzene sulfonic acid, followed by an ionic exchange with APP. With the incorporation of 22 wt.% AM-APP and 3 wt.% TiO$_2$, polyamide 11 composites showed high LOI value, upgraded UL-94 rating, and an 81.2 % reduction on pHRR value. Although some progress had been achieved in the modification of APP via self-assembly, it still needs to be matched with charring agents or metal oxides to pass the UL-94 V-0 rating. In addition, few pieces of literature have been reported on the synthesis of mono-component flame retardant...
through self-assembly technology and ionic exchange.

The current work reported the synthesis and characterization of a mono-component flame retardant prepared by self-assembly reactions between 1-(2-Amioethyl) piperazine and \( p \)-amino benzene sulfonic acid, followed by an ionic exchange with APP. The facile preparation method was considered environmentally friendly, using anhydrous ethanol and deionized water as the solvent. The flame retardancy, combustion behaviors, and flame retardant mechanism of the IFR-PP system were investigated in detail.

2. Materials and Methods

2.1 Materials

PP (F401), a granulated product with a melt flow index of 2.5 g (10 min)\(^{-1}\) (230 °C, 2.16 kg) was provided by Yangzi Petroleum Chemical Company (Nanjing, China). Ammonium polyphosphate (TY-432, APP crystalline form II, polymerization degree > 1000) was purchased from Yunnan Tianyao Chemical Co., Ltd (Kunming, China). \( p \)-amino benzene sulfonic acid (AR, 99.5 %) and 1-(2-Amioethyl) piperazine were supplied by Aladdin Industrial Corporation (Shanghai, China). Absolute ethyl ethanol (AR, 99.5 %) was obtained from Sinopharm Group Co., Ltd (Shanghai, China). Deionized water was self-made. All of the marketing materials were used directly without further purification.

2.2 Modification of APP

Figure 1 illustrated the modification process of ammonium polyphosphate involving two stages.

Firstly, the piperazine sulfonate (intermediate) was prepared via self-assembly as shown in stage I. Under stirring, ASC (17.3 g, 0.1 mol) was dissolved in deionized water at 90 °C. Hereafter, AEP (13.1 ml, 0.1 mol) was dropped into the ASC solution at a constant titration rate (2 s / drop). The reaction system was kept 95 °C for 3 h. After vacuum distillation and suction filtration, the straw yellow powder (AEP-ASC) was obtained and then dried at 80 °C in the vacuum oven for 24 h.

Secondly, the modified APP (MAPP) was synthesized via the ionic exchange between intermediate and APP as shown in stage II. To a three-neck round-bottom flask equipped with a stirrer, a thermometer and a reflux condenser were added APP (10 g), 50 mL ethanol, and 20 mL water. The mixture was refluxed under stirring for 12 h. Afterward, the intermediate (10 g) was introduced into the mixture, and the reaction continued for 2 h. Then, the light yellow and viscous solid was obtained from vacuum filtration and washed by deionized water (removing unreacted intermediate) three times. After drying under a high vacuum at 80 °C overnight, the final product (light yellow powder) was obtained and defined as modified APP (MAPP).
2.3 Sample preparation

The pure PP, APP, and MAPP were dried in a vacuum oven at 80 °C for 12 h. Then the PP blends containing flame retardants were mixed at 190 °C for 5 min in an internal mixer (US-70C, Changzhou Suyan Technology Co., Ltd. China) according to sample formulations in Table 1. The samples for fire behavior characterization were compression-molded under a plate vulcanizer (ZHY-W, Chengde Testing Machine Factory, China) at 195 °C for 5 min under 10 MPa of the pressure. The samples for mechanical measurements were injection-molded in a miniature injection molding instrument (WZS-10D, Xinshuo Precision Machinery Co., Ltd. China). The melting temperature was 200 °C, the mold temperature was 40 °C, the holding pressure was 0.6 MPa, and the holding time was 10 s.

2.4 Characterization

The Fourier transform infrared (FTIR) spectra were investigated by a Perkin Elmer instrument (Waltham, Massachusetts, USA) at room temperature. The samples were mixed with KBr pellets and scanned 32 times over a spectral range of 4000 - 450 cm⁻¹ with a resolution of 4 cm⁻¹.

X-ray diffraction (XRD) test was carried out by a power D/MAX2500 diffraction (Rigaku Corporation, Tokyo, Japan) using Cu Ka radiation under a scanning rate of 3°/s from 5° to 60° (2θ).

X-ray photoelectron spectroscopy (XPS) was determined by an ESCALAB 250XI system (Thermo Fischer, Massachusetts, USA).

The microstructures of flame retardants, char residues, and sections were measured using an SEM instrument (Zeiss SUPRA55, Jena, Germany). The specimens were sputter-coated with a conductive gold layer before observation.

The UL-94 vertical burning test was measured by a CZF-5 instrument (Shine Ray Instrument Co. Ltd., Nanjing, China) according to ASTM D3801 using the sample with the dimension of 130 mm×13 mm×3.2 mm.

Figure 1. The synthetic route of MAPP
Limiting oxygen index (LOI) was measured by an LOI analyzer (JF-3, Jiang Ning Co. Ltd., Nanjing, China) with sheet dimensions of 130 mm×6.5 mm×3.2 mm according to GB/T 2406-93 standard.

Thermogravimetric analysis (TGA) was conducted on a Perkin-Elmer TGA 4000 (Waltham, Massachusetts, USA) with a heating rate of 10 °C/min under nitrogen and oxygen atmosphere at temperatures ranging from 30 °C to 850 °C. TG-FTIR analysis was carried out under a nitrogen atmosphere at a heating rate of 10 °C/min from 30 °C to 850 °C. The flame retardant samples used in the tests were all fine solid powders with a mass of 8 ± 0.5 mg; the samples of pure PP and its blends used were block solids with a mass of 8 ± 0.5 mg.

The flammability of pure PP and PP blends were tested by a cone calorimeter device (Fire Testing Technology, East Grinstead, UK) according to ISO 5660-1. The dimension of the square sample was 100 mm×100 mm×3 mm, and the irradiation power was 35 kW/m².

Raman spectroscopy measurement was carried out with a DXR laser Raman spectrometer (Thermo Scientific, Massachusetts, USA) using a 532 nm helium-neon laser line at room temperature.

Tensile measurements were conducted on a universal tensile testing machine (WDT-5, Shenzhen Kai Qiang Experimental Instrument Co. Ltd, Guangzhou, China) according to GB/T 1040-2006 at a tensile speed of 50 mm/min. The Izod notched impact strength of the specimens was measured with an impact tester (XJU-22, Chengde Testing Machine Co. Ltd, Sichuan, China) according to GB/T 1843-2008. The values of all the mechanical properties were calculated as averages over five specimens.

3. Results and Discussion

3.1 Characterization of MAPP

3.1.1 FTIR Analysis

Figure 2a illustrates the FTIR spectra of AEP, ASC, and AEP-ASC. The characteristic peaks at 2939 and 2813 cm⁻¹ are attributed to the symmetrical and antisymmetric -CH₂- stretching vibration of AEP, and the peaks at 1315 and 1120 cm⁻¹ represent the -S=O stretching vibration of ASC [23]. Noticeably, the peak at 2647 cm⁻¹ (S-OH) [24] in the spectrum of ASC disappears and two new peaks at 2446 and 3265 cm⁻¹ (NH₂⁺ and NH₃⁺ bending vibration) are observed in the spectra of AEP-ASC, which indicates the formation of -NH₂⁺-O- and -NH₃⁺-O- [25, 26]. Furthermore, the spectrum of AEP-ASC shows sharp peaks at 3469, 3347 and 3232 cm⁻¹ owing to amino groups [27]. These results demonstrate that intermediate (AEP-ASC) is obtained by self-assembly. The chemical structures of the APP and MAPP are also explored by FTIR spectra (Figure 2b). Both APP and MAPP display the typical peaks at 1263 cm⁻¹ (P=O), 1087 cm⁻¹ (P-O symmetric stretching), and 882 cm⁻¹ (P-O asymmetric vibration) [28]. As compared with the spectra of APP, however, that of the MAPP displays some new peaks which are ascribed to the absorption of the benzene ring (1600 and 689 cm⁻¹, etc.), the stretching vibration of O=S=O (1320 and 1180 cm⁻¹) and especially the NH₂⁺ and NH₃⁺ bending vibration (2481 and 3248 cm⁻¹). The results initially demonstrate that the APP was successfully cross-linking modified with AEP-ASC by ionic exchange.
3.1.2 XRD Analysis

XRD patterns of APP and MAPP are presented in Figure 2c. Obviously, the position of the diffraction peaks of APP and MAPP are almost the same, indicating that the crystalline structure of APP is not influenced by cross-linking modification.

3.1.3 XPS Analysis

The XPS spectra of APP and MAPP are shown in Figure 2d. As can be seen, MAPP shows much higher carbon content (44%), lower contents of oxygen (32%), and phosphorus (8%) than APP. Furthermore, a new peak appears at 168.1 eV for MAPP, which is ascribed to the S\(_{2p}\) from the intermediate (AEP-ASC). The fitted N\(_{1S}\) spectra of APP and MAPP are presented in Figure 2e, f. For APP, the binding energy at around 401.2 eV is assigned to NH\(_{4}^+\), and the peak at 399.3 eV might correspond to the N (-P-NH-P-). This result of APP spectra agrees with Wang’s report\(^{[17]}\). Although the NH\(_{4}^+\) peak still appears for MAPP, three new peaks at 399.7 eV, 400.3 eV, and 400.7 eV are observed\(^{[16, 17]}\). It indicates that NH\(_2^+\) and NH\(_3^+\) are formed and take place of NH\(_{4}^+\) partially. This result further confirms the successful ionic interactions between intermediate and APP.

3.1.4 Surface Morphology and EDS test

SEM is used to investigate the micro-morphology of the APP and MAPP, and the corresponding images are shown in Figure 3. Apparent changes in the surface morphology between APP and MAPP are easily observed. The pure APP particle shows a smooth surface, and its particle size is about 10 µm (Figure 3a, b). However, the MAPP particle looks like a cross-linked aggregate of APP particles and displays a rough surface (Figure 3c).
surface (Figure 3c, d). It could be interpreted as the H-bonding and ion-dipole forces that contributed to particle aggregation and fusion\cite{16}. The changes in the surface morphology of MAPP are consistent with the proposed model as shown in Figure 1. Moreover, the elemental composition of the APP and MAPP is evaluated from the EDS test (Figure 3e). By comparison with APP, the MAPP presents higher content of C (24.27 %) and lower content of P (11.53 %) as well as the appearance of S element. Both the changes in the surface morphology and element composition between APP and MAPP demonstrate the successful cross-linking modification of APP evidently.

![Figure 3. SEM images of (a, b) APP and (c, d) MAPP; (e) EDS image of APP and MAPP](image)

3.2. Fire behavior

3.2.1 Reaction to small flame (UL 94 vertical burning and LOI)

The flame retardant data of the neat PP and IFR-PP blends are summarized in Table 1. The neat PP shows serious dripping with no rating level during the UL-94 test, and its value of LOI is merely 17 %, indicating its extreme flammability. In the case of the incorporation of APP alone, the PP/APP blend fails to pass the UL-94 V-0 test and gives of LOI value of 19 % even at its loading level up to 25 wt.%, revealing that APP is not very practical to improve the flame retardancy of PP. However, the LOI values and flame retardant rating of PP/MAPP blends are increased significantly with the incorporation of MAPP. For example, the IFR-PP blend containing 22.5 wt.% MAPP gives 30 % of LOI value and UL-94 V-0 rating. It demonstrates that the flame retardant efficiency of MAPP is much higher than that of APP for IFR-PP blends.

| Sample ID    | PP wt.% | APP wt.% | MAPP wt.% | Dripping or not | UL-94 rating 3.2mm | LOI % |
|--------------|---------|----------|-----------|-----------------|---------------------|-------|
| Pure PP      | 100     | 0        | 0         | Yes             | No rate             | 17    |
| PP/20%APP    | 80      | 20       | 0         | Yes             | No rate             | 17    |
| PP/22.5%APP  | 77.5    | 22.5     | 0         | Yes             | No rate             | 18    |
| PP/25%APP    | 75      | 25       | 0         | Yes             | V-2                 | 19    |
3.2.2 Combustion behavior under forced-flaming scenario (Cone calorimetry test)

Cone calorimetry test (CCT) is currently the ideal test to assess the burning behavior of polymeric materials under an ongoing fire or forced combustion conditions. The characteristic curves including the heat release rate (HRR), total heat release (THR), smoke production rate (SPR), total smoke production (TSP), CO, and CO₂ production (COP, CO₂P) are illustrated in Figure 4, and some important parameters such as time to ignition (TTI), the peak of heat release rate (PHRR), THR, and average mass loss rates (AMLR) are listed in Table 2.

HRR is generally the most critical performance parameter to characterize the fire intensity, and PHRR indicates the maximum degree of heat release during combustion. As presented in Table 2, the pure PP gives 718.3 kW/m² of PHRR value and 57.3 MJ/m² of THR, which are reduced to 354.7 kW/m² and 47.9 MJ/m² with the incorporation of 25 wt.% APP, respectively. The corresponding values are further reduced significantly with the substitution of APP with MAPP. For example, the IFR-PP containing 25 wt.% MAPP presents the lowest PHRR (155.9 kW/m²) and THR values (44.9 MJ/m²), which are reduced by 78.3 % and 21.6 %, respectively, as compared to those of the neat PP. The sharp decline of the PHRR and THR values indicates that MAPP has a more significant role in fire safety than APP.

Fire growth index (FGI) and fire performance index (FPI) are used to assess the fire hazard of polymers, which can be calculated according to the following equation [29] based on the HRR curves and TTI values,

\[ FGI = \frac{PHRR}{T_{PHRR}} \quad FPI = \frac{TTI}{PHRR} \]

The obtained FGI and FPI values of the neat PP and IFR-PP blends are also listed in Table 2. Generally, the lower FGI and higher FPI values imply higher fire safety properties [30]. As can be seen, the IFR-PP blends display much lower FGI values whilst greater FPI values than that of the pure PP. Noticeably, the FPI value (0.22 m²/s/kW) of PP/25 %MAPP is increased by 57.1 % whilst the FGI value (0.72 kW/m²) is decreased by 61.5 % as compared with these corresponding values of PP/25%APP (0.14 m²/s/kW of FPI and 1.87 kW/m² of FGI, respectively). Furthermore, taking the much lower average mass loss rate (AMLR, 0.027 g/s) of PP/25%MAPP than that (0.043 g/s) of PP/25%APP into consideration, it can be concluded that MAPP is superior to APP in fire safety.

It is well known that most victims suffocate to death by inhalation of smoke and CO in fire accidents. As such, smoke suppression is very significant for flame retardant materials. As can be seen in Table 2, the total smoke production (TSP) is reduced from 48.8 m² for pure PP to 6.4 m² for PP/APP while their peaks of the smoke production rate (PSPR) are almost the same. Fascinatingly, both the PSPR and TSP values of PP/25%MAPP are reduced drastically as compared with that of the pure PP or PP/25%MAPP, indicating the superior smoke suppression of MAPP. Moreover, the CO production (0.15 kg/kg) of PP/25%MAPP specimen is half of that of pure PP (0.32 kg/kg), implying the possibility of suffocation can be significantly inhibited during evacuation.
Table 2. Data of the pure PP and its flame-retardant blends during Cone calorimeter combustion

| Sample     | Pure PP | PP/25%APP | PP/20%MAPP | PP/22.5%MAPP | PP/25%MAPP |
|------------|---------|-----------|------------|--------------|------------|
| TTI (s)    | 47      | 49        | 40         | 38           | 35         |
| PHRR (kW/m²) | 718.3   | 354.7     | 201.0      | 192.4        | 155.9      |
| TₚHRR (s)  | 105     | 190       | 255        | 235          | 215        |
| THR (MJ/m²)| 57.3    | 47.9      | 50.7       | 47.2         | 44.9       |
| TSP (m²)   | 48.8    | 6.4       | 6.5        | 5.2          | 4.2        |
| Mean COY (kg/kg) | 0.32 | 0.11      | 0.17       | 0.17         | 0.15       |
| PSPR (m²/s) | 0.5     | 0.045     | 0.026      | 0.024        | 0.016      |
| AMLR (g/s) | 0.059   | 0.043     | 0.033      | 0.032        | 0.027      |
| FPI (m²/s/kW) | 0.07 | 0.14      | 0.20       | 0.20         | 0.22       |
| FGI (kW/m²) | 6.84    | 1.87      | 0.79       | 0.82         | 0.72       |

Figure 4. HRR (a), THR (b), SPR (c), TSP (d), CO₂P (e), and COP (f) curves of samples during the combustion in CCT

3.3. Thermal stability analysis

Figure 5 and Figure 6 presents the TG and DTG curves of the flame retardants and IFR-PP samples under nitrogen and air atmosphere, respectively. Corresponding data, such as the initial thermal decomposition temperature (Tₜ₅wt%), the temperature at maximal degradation (Tₘₐₓ), Rate of Tₘₐₓ, and char residue at 800 °C, are listed in Table 3 and Table 4. Obviously, the pure PP shows a single degradation process with no char residue left at 800 °C no matter whether under nitrogen or air atmosphere, indicating pure PP fails to form char alone. On the contrary, two-stage degradation is observed for all IFR-PP blends. The first-stage at low temperature is mainly ascribed to the degradation of the IFR and PP, and the second-stage at high temperature is attributed to the decomposition products (polyphosphoric acids, pyrophosphate, and metaphosphate acid) of APP or MAPP [26]. Furthermore, all samples display a lower degradation temperature under the air atmosphere than that under the nitrogen atmosphere owing to
thermo-oxidative degradation \[^{[24]}\]. As compared with the pure PP, all PP/APP samples display greater \(T_{5\text{wt\%}}\) values, which is ascribed to the protection of the thick polyphosphoric acid fluid produced by the decomposition of APP \[^{[25]}\]. In the case of substitution of APP with MAPP, the \(T_{5\text{wt\%}}\) values of PP/MAPP samples move to lower temperatures since the \(T_{5\text{wt\%}}\) of MAPP is lower than that of APP. It is noteworthy that the char residue of PP/25\%MAPP is up to 5.4 % at 800 °C under the air atmosphere, which is higher than that of pure PP and PP/APP samples. Such an observation demonstrates that MAPP can form a stable carbon layer, which acts as a barrier between oxygen and PP matrix \[^{[27]}\].

![Figure 5](https://doi.org/10.20944/preprints202010.0561.v1)

**Figure 5.** (a) TG and (b) DTG curves of APP and MAPP. (c, e) TG and (d, f) DTG curves of PP and its flame-retardant blends under the nitrogen atmosphere.

**Table 3.** TGA data of the IFR-PP samples under nitrogen atmosphere

| Sample       | \(T_{5\text{wt\%}}\) (°C) | \(T_{\text{max1}}\) (°C) | \(T_{\text{max2}}\) (°C) | Rate of \(T_{\text{max1}}\) (wt.%/min) | 800°C Char residues (%) |
|--------------|---------------------------|---------------------------|---------------------------|----------------------------------------|-------------------------|
| APP          | 345.1                     | 364.4                     | 623.9                     | 7.3                                    | 20.5                    |
| MAPP         | 303.1                     | 351.7                     | 588.4                     | 4.2                                    | 20.1                    |
| Pure PP      | 373.5                     | 455.4                     | /                         | 17.7                                   | 0                       |
| PP/20%APP    | 387.4                     | 448.8                     | 687.9                     | 8.4                                    | 8.1                     |
| PP/22.5%APP  | 395.9                     | 456.6                     | 719.1                     | 8.2                                    | 10.9                    |
| PP/25%APP    | 407.9                     | 463.7                     | 747.4                     | 8.1                                    | 12.5                    |
| PP/20%MAPP   | 383.0                     | 454.6                     | 704.6                     | 8.8                                    | 8.3                     |
| PP/22.5%MAPP | 368.0                     | 446.1                     | 694.8                     | 8.0                                    | 8.5                     |
| PP/25%MAPP   | 361.2                     | 437.6                     | 670.9                     | 7.6                                    | 8.6                     |
Figure 6. (a) TG and (b) DTG curves of APP and MAPP. (c, e) TG and (d, f) DTG curves of PP and its flame-retardant blends under the air atmosphere

Table 4. Data of TGA for PP and its flame-retardant blends under air atmosphere

| Sample          | $T_{5wt}$ (°C) | $T_{max1}$ (°C) | $T_{max2}$ (°C) | Rate of $T_{max1}$ (wt.%/min) | 800°C Char residues (%) |
|-----------------|----------------|-----------------|-----------------|-------------------------------|-------------------------|
| APP             | 346.6          | 359.5           | 626.8           | 7.4                           | 22.9                    |
| MAPP            | 270.9          | 320.7           | 615.0           | 6.7                           | 5.7                     |
| Pure PP         | 246.4          | 334.8           | /               | 8.5                           | 0                       |
| PP/20%APP       | 287.9          | 381.1           | 561.3           | 6.0                           | 3.7                     |
| PP/22.5%APP     | 284.2          | 374.1           | 583.7           | 5.7                           | 3.9                     |
| PP/25%APP       | 275.0          | 366.0           | 589.9           | 5.3                           | 5.3                     |
| PP/20%MAPP      | 279.3          | 362.1           | 557.1           | 6.4                           | 3.3                     |
| PP/22.5%MAPP    | 277.3          | 354.1           | 591.0           | 5.9                           | 4.1                     |
| PP/25%MAPP      | 273.8          | 346.0           | 596.6           | 5.6                           | 5.4                     |

3.4. Flame-retardant mechanism analysis

3.4.1 Gas phase analysis

The TG-FTIR test is performed to analyze the volatile gases released from IFR-PP samples, which helps to explore the flame-retardant mechanism of the gas phase. Figure 7 illustrates the FTIR curves and 3D spectra of the evolved gaseous products during the pyrolysis of pure PP, PP/25%APP, and PP/25%MAPP. All the samples display the following characteristic peaks: such as alkane (2960, 2920 cm$^{-1}$), alkene (1647, 1465, and 1371 cm$^{-1}$), and a diene (890 cm$^{-1}$) [10]. However, the intensity of the above-mentioned peaks in the spectra of PP/25%MAPP is weaker than that of the pure PP and PP/25%APP, suggesting that volatile components are suppressed effectively during the degradation process of PP/25%MAPP specimen [21]. As compared with the pure PP, both PP/25%APP and PP/25%MAPP specimens show characteristic peaks ascribed to P=O (1258 cm$^{-1}$), P-O (1087 cm$^{-1}$), and NH$_3$ (966 and 937 cm$^{-1}$), which are contributed to the thermal degradation of APP and scissions of polyphosphoric acids [32]. Furthermore, it is noticeable that a new peak at 1499 cm$^{-1}$ (SO$_2$) is found merely in
the spectrum of PP/25%MAPP specimen. The appearance of these non-combustible gas (NH$_3$ and SO$_2$) can dilute the released combustible gas (hydrocarbons, etc.).

![Figure 7. FTIR spectra and 3D images of the pyrolysis products of pure PP (a, a’), PP/25%APP (b, b’), and PP/25%MAPP (c, c’) at different temperatures.]

3.4.2 Analysis of char residue

The digital photographs of the residues after CCT are shown in Figure 8. Obviously, there is no residue left for pure PP and a minute quantity of residues for PP/25%APP. It indicates that the incorporation of the unmodified APP alone cannot promote the formation of an expanded char layer. However, more intumescent and expanded char residues with expansion heights of 1.7 cm (Figure 8c’) and 2.5 cm (Figure 8d’) are observed for PP/22.5%MAPP and PP/25%MAPP, respectively. The expansion height is usually adopted to assess the swelling degree or quality of the char layer. Accordingly, it is concluded that MAPP has a greater effect on the formation of the intumescent char layer than the unmodified APP, which can obstruct the heat and mass transfer between the gas and condensed phase.

![Figure 8. Digital photographs of residues of pure PP (a), PP/25%APP (b, b’), PP/22.5%MAPP (c, c’), and PP/25%MAPP (d, d’) samples after CCT.]

The microstructure of the residues after CCT was observed further via SEM, and
the resultant images are presented in Figure 9. Clearly, numerous large holes are observed both in the outer and inner char residues of PP/25%APP whilst compact and continuous char layer is observed on the surface of PP/MAPP blends. Especially, the appearance of some char bubbles in the PP/25%MAPP surface residues indicates that the incorporation of MAPP enhances the compactness of char residues and thus forms a much excellent IFR system [5]. However, the honeycomb structure still exists in the inner char residue of PP/MAPP samples, which is ascribed to the diffusion of pyrolysis gas.

Figure 9. SEM images of char residues of PP/25%APP (a, outer, a’, inner), PP/22.5%MAPP (b, outer, b’, inner), and PP/25%MAPP (c, outer, c’, inner) after CCT

The elemental compositions of the char residue of the PP/25%MAPP sample after CCT were investigated by the XPS analysis (Figure 10). As can be seen, the main elements in the residue are C, O, P, and N. In N$_{1S}$ spectra (Figure 10b), the peaks at 400.5 eV, 401.2 eV, and 402.3 eV are attributed to the P-N-P, NH$_4^+$, and P-C-N, respectively. [21, 23]. The O$_{1S}$ peaks (Figure 10c) at 531.1 eV and 532.2 eV are assigned to =O (including P=O and C=O) and -O- (including C-O-C and P-O-C [10]), respectively. In the case of P$_{2P}$ spectra (Figure 10d), the peaks around 133.9 eV and 134.5 eV correspond to P=O and P-O-C structure, respectively [22, 33]. The results above demonstrate that the existence of P-C-N and P-O-C structure plays a vital role in forming a dense and continuous char layer.
Raman spectroscopy is a useful tool to characterize the graphitization degree of char residues. The Raman spectra are mainly divided into G bands (about 1590 cm\(^{-1}\), showing the graphitic structure) and D bands (about 1365 cm\(^{-1}\), representing lattice defect of carbon atom). Furthermore, the graphitization degree of the char residue can be assessed by the area ratio of two peaks (\(A_D/A_G\)). Generally, the lower ratio of \(A_D/A_G\) means the higher graphitization degree and better thermal-oxidative stability of the char\[^{[34,35]}\]. The \(A_D/A_G\) values are calculated to be 2.57 for PP/25%MAPP, 2.24 for PP/22.5%MAPP, and 2.12 for PP/25%MAPP, respectively. As such, it is concluded that the incorporation of MAPP facilitates the graphitization to form much dense and continuous surface char layers, which is in agreement with the results of SEM observation.
3.4.3 Possible flame-retardant mechanism

Based on the analysis of the volatile gas and char residues, the possible flame-retardant mechanism is proposed and illustrated in Figure 12. In the initial stage of the combustion, the dehydration of MAPP and the cleavage of ionic bonds result to release of some incombustible gases (such as NH$_3$, SO$_2$, and water vapor) which can fully dilute the oxygen and heat. With the temperature rising, MAPP is thermally decomposed into polyphosphoric acid, pyrophosphoric acid, or metaphosphoric, all of which can esterify and crosslink with the char precursor (piperazine ring) [10]. Moreover, the phosphate ester radicals and pyrophosphate radicals released from the decomposition of MAPP readily quench O$_2$, H$_2$, and OH radicals [36]. Additionally, the benzene ring in the intermediate acts as both the cross-linking agent and crystal core [23]. As such, the char layer containing P-O-C and P-C-N structure is formed and its quality is improved owing to the crosslinking networks. It is noteworthy that although the initially formed char layer did not swell, its continuous and compact structure hinders the gas overflowing and the accumulation of the incombustible gases keeps the char layer expanding. As a result, an intumescent and dense char layer is formed, which protects the underlying substrate from further thermal decomposition. The physical (expansion and morphology) and chemical (thermal stability) properties of the char layer were enhanced after introducing MAPP and the possible IFR mechanism is shown in Figure 12 [21].

![Figure 11: Raman spectra of char residues of PP/25%APP, PP/22.5%MAPP, PP/25%MAPP after CCT](image)
Figure 12. Schematic diagram of the flame-retardant mechanism

3.5 Mechanical properties

Figure 13 illustrates the typical stress-strain curves of pure PP and IFR-PP blends, and the mechanical properties data are provided in Table 5. The pure PP gives 39.1 MPa of tensile strength, 147.1 % of elongation at break, and 4.6 kJ·m⁻² of impact strength, respectively. Although the mechanical properties of IFR-PP blends are decreased to a certain extent by comparison with the pure PP, the PP/25%MAPP sample shows better mechanical properties than the PP/25%APP. Namely, the tensile strength, elongation at break, and impact strength of PP/25%MAPP are increased by 12.3 %, 110.5 %, and 37.9 % as compared with PP/25%APP, respectively. This difference might be related to interface interaction between the dispersion phase (MAPP or APP) and the PP matrix. Figure 14 presents the SEM images of the impact fractured surface of PP, PP/25%APP, and PP/25%MAPP. APP particles fail to adhere to the PP matrix, and the interface between the APP particles and PP matrix is clearly visible (Figure 14b). Conversely, the MAPP particles are uniformly embedded in the PP matrix (Figure 14c), which leads to an improvement of the elongation at break and impact strength. This observation is probably ascribed to that the modification of APP improved the interfacial adhesion, and the addition of MAPP had little effect on the mechanical properties of IFR-PP samples.
Figure 13. Typical stress-strain curves of pure PP and its flame-retardant blends

Table 5. Mechanical properties data of pure PP and its flame-retardant blends

| Sample       | Tensile strength (MPa) | Elongation at break (%) | Impact strength (kJ m$^{-2}$) |
|--------------|------------------------|-------------------------|-----------------------------|
| Pure PP      | 39.1±0.4               | 147.1±3.0               | 4.6±0.5                     |
| PP/25%APP    | 25.3±2.5               | 59.8±6.8                | 2.9±0.2                     |
| PP/25%MAPP   | 28.4±1.1               | 125.9±4.8               | 4.0±0.2                     |

Figure 14. SEM images of the impact fractured surface of pure PP (a), PP/25%APP (b), and PP/25%MAPP (c)

4. Conclusion

A novel mono-component flame retardant (MAPP) was prepared by the modification of conventional APP with piperazine sulfonate. The MAPP showed more effective flame retardancy and smoke suppression than the traditional APP. The PP/MAPP composite passed the UL-94 V-0 rating and achieved 30 % of LOI value at 22.5 wt% loading level of MAPP. The HRR, PHRR, and TSP values of PP/22.5%MAPP were reduced significantly as compared with those of the pure PP or PP/25%MAPP. The results of TGA-FTIR and char residue analysis revealed that some incombustible gases were released and the intumescent, dense, and continuous char residues were produced with the incorporation of MAPP. The PP/25%MAPP sample also displayed better mechanical properties than the PP/25%APP.
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