A Highly Effective, UV-Curable, Intumescent, Flame-Retardant Coating Containing Phosphorus, Nitrogen, and Sulfur, Based on Thiol-Ene Click Reaction

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Abstract: In this paper, a flame-retardant, UV-cured coating was prepared on the fiber composites' (FC) surface via a thiol-ene click reaction using pentaerythritol tetra(3-mercaptopropionate) (PETMP), triallyl cyanurate (TAC), and 2-hydroxyethyl methacrylate phosphate (PM-2). The synergistic effectiveness of phosphorus (P), nitrogen (N), and sulfur (S) was studied in detail by changing the proportion of these reactants. Sample S_{4}(N_{3}P_{2})_{6}, with a molar ratio of N and P elements of 3:2, and the thiol and vinyl groups of 4:6 had the highest LOI value (28.6%) and was self-extinguishing in the horizontal combustion test. It had the lowest peak heat release rate (PHRR) value (279.25 kW/m^2) and total smoke production (2.18 m^2). Moreover, the thermogravimetric analysis (TG) showed that the decomposition process of the coated composites was delayed. The conversion rate of the double bond and the thiol of S_{4}(N_{3}P_{2})_{6} was 100% and 92.0%, respectively, which showed that the cross-linked network structure was successfully formed. The tensile strength and the flexural strength of coated composites improved, and the transparency of the coating can reach 90%. These characteristics showed that the UV-cured coatings could be used in industrial production to effectively prevent fires.

Keywords: UV-curable coating; thiol-ene click reaction; flame retardancy; natural fiber

1. Introduction

A UV-curable coating can be used on different substrates [1–4] to enhance the materials with various properties, such as fire resistance [5,6], anti-fouling [7], and good optical transparency [8]. Compared with ordinary polymer coatings, UV-curable coatings have the advantages of short reaction time, low energy consumption, and simple operation [9]. The functional UV-curable coating is based on the modification of unsaturated resin—unsaturated polyester, acrylic resin, etc. These systems have problems that need to be solved urgently, such as high viscosity, large volume shrinkage after curing, oxygen inhibition, and a non-uniform crosslinking network [10,11]. Among them, oxygen inhibition is the main obstacle in the UV-cured process [12,13]. Oxygen reacts with the free radicals initiated by the photoinitiator in the chain growth step and finally generates peroxy free radicals with lower activity, which leads to the failure of the curing of the coating surface and the low conversion rate of the UV-cured film. So far, the most commonly used method in the industry is curing under an N_2 atmosphere, using more photoinitiators or using multiple initiators to form a photoinitiating system [12,14]. However, these methods lead to a sharp increase in curing costs and greater volume shrinkage of the coating, which results in a decrease of the adhesion of UV-cured coatings, an increase of internal stress, and the deterioration of mechanical properties [11].

The thiol-ene click reaction uses vinyl monomers and thiol monomers as prepolymers. The polymerization reaction is a free-radical step-growth reaction with the participation of
photoinitiators. The polymer molecular weight gradually increases, which greatly improves the double bond and thiol conversion rate. In this curing process, the thiol group and the peroxo radical undergo a hydrogen extraction reaction to produce a highly reactive thiol group, which effectively eliminates the problem of oxygen inhibition [1,11,15]. Moreover, the gradual growth mechanism of the thiol-ene click reaction leads to a delay in the gel point of the cured product, which can effectively release the shrinkage stress. Besides, the thiol-ene click reaction has the advantages of fast reaction speed and high selectivity. It has a non-negligible application prospect in the field of UV-cured coatings [16,17], and it is considered to be an effective method for preparing cross-linked network coatings [18,19].

To develop a green halogen-free flame-retardant coating on the surface of the natural fiber composites, the prepolymerer containing N, P, and S are cross-linked via the thiol-ene click reaction under UV irradiation in this paper. Pentaerythritol Tetra(3-mercaptopropionate) (PETMP), a kind of tetramercaptan monomer, was used as a film-forming agent, due to its good reactivity with the double bonds. Meanwhile, the S element in PETMP and the N element in triallyl cyanurate (TAC) could be used as the blowing agent to build an intumescent flame-retardant system, together with the phosphorous-containing monomer 2-hydroxyethyl methacrylate phosphate (PM-2). The effect of the relative ratio between N, P, and S elements on the flame-retardant effect of the coating was studied more precisely. At the same time, the effect of the relative ratio of double bonds and sulfhydryl groups on the conversion rate of the coating and the enhancement of the mechanical properties of the composite by the presence of the coating was also investigated.

In this paper, this coating was applied on the natural fiber composites, which not only improved the flame-retardant properties of the composites but also had good adhesion and did not have a negative impact on the mechanical properties. The curing reaction kinetics and the functional group changes of the coating before and after curing were evaluated by Fourier transform infrared spectroscopy (FTIR). The flame retardancy of the coated composites was investigated by limit oxygen index (LOI), horizon burning rate (HBR), and the cone calorimetry tests (CCT), respectively. The thermal properties of the coated composites were studied by thermogravimetric analysis (TGA). The mechanical properties of the coated composites were characterized by the tensile strength, the flexural strength, and the adhesion test. Finally, the scanning electron microscope/energy-dispersive spectrometer (SEM/EDS) was used to observe the surface morphology of the char layer structure of the samples after burning and acquired the content of C, O, N, P, and S of the char layers. To sum up, this work offered an efficient, quick, convenient, and simple method for preparing crosslinked flame-retardant coatings.

2. Experiment

2.1. Materials

Pentaerythritol tetra(3-mercaptopropionate) (PETMP, 98.0%) was purchased from Shanghai Yuhong Chemical Technology Co., Ltd. (Shanghai, China); triallyl cyanurate (TAC, 98%) was obtained from Shanghai Aladdin biochemical technology Co., Ltd. (Shanghai, China); 2-hydroxyethyl methacrylate phosphate (PM-2, 99%) was received from Guangzhou Lihou Trading Co., Ltd. (Guangzhou, China); and phenylbis(2,4,6-trimethylbenzoyl)phosphine oxide (BAPO, 98%) was supplied by Changshu Hengyao New Material Co., Ltd. (Suzhou, China). The UV absorption peaks of the photoinitiator BAPO are 250 nm and 370 nm. All reagents were analytical grade, and they were used as received without further purification. The chemical structures of raw materials are shown in Figure 1.
2.2. Preparation of Samples

The flax fiber, kenaf fiber, and polypropylene fiber were blended in the mass ratio of 1:1:2 to prepare the needle-punched nonwoven felts. The non-woven felts were hot-pressed into fiber composites (FC) at 180 °C and 8 MPa. The formulation of UV curable coatings is shown in Table 1. The coatings were obtained by mixing four materials in different proportions, which included a prepolymer containing thiol groups (PETMP), a prepolymer TAC containing nitrogen, a phosphate functional monomer (PM-2), and the photoinitiator BAPO. There is one C=C bond in PM-2 and three C=C bonds in TAC; the thiol monomer contains four thiol groups, and the ratio of monomers meets C=C: SH=1:1. From this, the best ratio of nitrogen and phosphorus can be explored. On this basis, the relationship between the ratio and the conversion rate can also be studied by changing the ratio of C=C and SH. Thereafter, the prepolymer solution was mixed thoroughly with the photoinitiator BAPO (3 wt% of the total weight of TAC, PM-2, and PETMP) at 60 °C for 30 min to obtain a clear liquid. The homogeneous mixture was coated on the fiber composites with a 200 µm film-applicator to form uniform films. Samples were cured under a 395 nm light from a UV lamp (Zhongshan Guzhen Yanxizhao Lighting Appliance Factory, AC90V-240V 395 nm 35W, Zhongshan, China) at a distance of 10 cm for 1 min or more to ensure an adequate click reaction. The structure of the prepolymer and a brief description of preparation for an idealized cross-linked network is shown in Figure 1.

Table 1. The formulation of the Ultraviolet (UV)-curing films.

| Coating  | PETMP (mol) | TAC (mol) | PM-2 (mol) | [SH]: [C=C] (mol) | N:P (mol) |
|----------|-------------|-----------|------------|-------------------|-----------|
| S\textsubscript{5}N\textsubscript{5}P\textsubscript{0} | 3           | 4         | 0         | 5:5               | 5:0       |
| S\textsubscript{5}N\textsubscript{4}P\textsubscript{1} | 3           | 3.2       | 2.4       | 5:5               | 4:1       |
| S\textsubscript{5}N\textsubscript{3}P\textsubscript{2} | 3           | 2.4       | 4.8       | 5:5               | 3:2       |
| S\textsubscript{5}N\textsubscript{2}P\textsubscript{3} | 3           | 1.6       | 7.2       | 5:5               | 2:3       |
| S\textsubscript{5}N\textsubscript{1}P\textsubscript{4} | 3           | 0.8       | 9.6       | 5:5               | 1:4       |
Table 1. Cont.

| Coating        | PETMP (mol) | TAC (mol) | PM-2 (mol) | [SH]: [C=C] (mol) | N:P (mol) |
|----------------|-------------|-----------|------------|-------------------|-----------|
| S₂N₀P₅         | 3           | 0         | 12         | 5:5               | 0:5       |
| S₄(N₃P₂)₆      | 1.5         | 1.8       | 3.6        | 4:6               | 3:2       |
| S₃(N₃P₂)₇      | 1.5         | 2.8       | 5.6        | 3:7               | 3:2       |
| S₂(N₃P₂)₈      | 1.5         | 4.8       | 9.6        | 2:8               | 3:2       |
| S₁(N₃P₂)₉      | 1.5         | 10.8      | 21.6       | 1:9               | 3:2       |

2.3. Characterization

2.3.1. Fourier Transform Infrared Spectroscopy (FTIR)

The photopolymerization kinetics were studied by the TENSOR 27 FTIR spectrometer (Germany Bruker, Berlin) in the range of 4000–400 cm⁻¹. The double bond conversion was monitored by measuring the area under the peak of 1637 cm⁻¹, and the peak at 2572 cm⁻¹ was used to measure the thiol group conversion. The peak at 1730 cm⁻¹ represented the stretching vibration peak of C=O, and the intensity of the peak did not change significantly during the FTIR tests. As a result, the peak at 1730 cm⁻¹ was used as an internal standard peak to neutralize the variation in the coatings caused by rapid curing [2,5,20]. The double bond conversion rate (DC) could be calculated according to Equation (1):

\[
DC (%) = \left(1 - \frac{A_t(1637 \text{ cm}^{-1})}{A_0(1637 \text{ cm}^{-1})} \right) \times 100\% \quad (1)
\]

The thiol group conversion (TC) could be calculated according to Equation (2):

\[
TC (%) = \left(1 - \frac{A_t(2572 \text{ cm}^{-1})}{A_0(2572 \text{ cm}^{-1})} \right) \times 100\% \quad (2)
\]

where \(A_0(1637 \text{ cm}^{-1})\), \(A_0(1730 \text{ cm}^{-1})\), and \(A_0(2572 \text{ cm}^{-1})\) represent the integral peak area of C=C, C=O, and -SH, before curing, respectively; the \(t\) at \(A_t\) represents the UV-curing time.

2.3.2. Limiting Oxygen Index (LOI)

The LOI of the coated composites was obtained by an LOI instrument (JF-3, Seoul, Korea FESTEC Company) according to GB/T 2406.2-2009 (China) oxygen index method. The sample size used for the test was 100 × 10 × 3 mm².

2.3.3. Horizontal Burning Rate (HBR)

The HBR test was performed using a burning test instrument (H1011D, Changchun-shiHosly apply technique graduate school, Changchun, China). The test was measured according to the Flammability of automotive interior materials (GB 8410-2006). The composites were ignited for 15 s, and then the fire was removed. Burning length and time were recorded to calculate the burning rate. The sample size used for HBR tests was 150 × 75 × 3 mm³.

2.3.4. Thermo-Gravimetric Analysis (TG)

The thermogravimetric (TG) analysis results were obtained on a Q500 thermal analyzer (TA Instruments, New Castle, DE, USA) under a nitrogen atmosphere (50 mL/min), and under an air atmosphere with a heating rate of 10 °C/min. About 5–10 mg of the samples was placed in an alumina crucible and heated from 25–800 °C.

2.3.5. Cone Calorimetry Test (CCT)

The CCT is one of the best methods to evaluate the combustion characteristics of the samples. The CCT was performed with a cone calorimeter apparatus (FTT0242, West
Sussex, UK) according to ISO 5660 (2002). In the test, the coated composites were exposed to a radiation cone with a heat flux of 50.0 kW/m². The sample size was 100 × 100 × 3 mm³.

2.3.6. Mechanical Properties

The mechanical tests, including the tensile strength and the bending strength, were performed using a universal testing machine WSM-5KN (Changchun intelligent instrument equipment Co., Ltd., Changchun, China) according to the TL52448. The dimensions of each specimen were 100 × 25 × 3 mm³, and the strain rate was 3 mm/min. All mechanical data had an effective average of 5.

2.3.7. Adhesion Test

The adhesion test was measured using the single blade pocket knife in accordance with ISO 2409:2013. The number of cuts in each direction was 6, and the cross-cut spacing was 3 mm.

2.3.8. Optical Transparency

The optical transparency of the UV-cured coatings was recorded using a UV-6100s Scanning UV-Vis Spectrophotometer (Shanghai Mepda Instrument Co., Ltd., Shanghai, China) in the range of 200–1100 nm.

2.3.9. Morphology Observation

The char residues after HBR tests were observed via scanning electron microscopy (SEM, 30 kv/Magellan400, FEI NanoPorts, Hillsboro, Oregon, USA). The energy-dispersive spectrometer (EDS, Magellan400, FEI NanoPorts, Hillsboro, Oregon, USA) on the SEM was used to observe the surface of the coated samples after HBR tests and compare the content of C, O, N, P, and S of the coatings on the surface of coated composites before and after the HBR test.

3. Results and Discussion

3.1. Curing Kinetics and Double Bond Conversion

In order to study the effect of the relative changes of ratio of -SH to C=C and the TAC to PM-2 on the curing performance of the coating, the FTIR technique was employed to monitor the chemical structure changes of the S₅₋ₓₐ₋ₓₐ₋ₓₐ₋ₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋ₓₐ₋xDC and S₅₋ₓₐ₋ₓₐ₋ₓₐ₋xDC series coatings before and after curing. As shown in Figure 2, the FTIR spectra of sample S₅₋ₓₐ₋xDC was recorded at different UV-irradiation times (5 s, 10 s, 20 s, 30 s, 60 s, and 300 s) [2,4,20,21]. The conversion rate of the thiol and vinyl groups gradually stabilized after 20 s. Besides, the C=C and -SH peak intensity remained almost unchanged at 60–300 s, which indicated that the curing process was complete and the double bond and the thiol had formed the cross-linked network structure. Thus, the coating formed when the curing time was 60 s.

Figure 2c showed that the relative change of PM-2 and TAC content had little effect on the double bond conversion rate. However, when the -SH and C=C molar ratio was 1:1, as the content of PM-2 increased, the -SH conversion rate gradually decreased as in Figure 2d. This may be because there were acrylate groups in PM-2; the double bond tended to undergo homopolymerization and does not copolymerize with thiol during the UV-curing process, which led to a higher double bond conversion rate than the thiol conversion rate (Figure 2c,d). The final conversion rate of thiol and double bond is shown in Table 2; the conversion rate of the double bond of coating S₅₋xDC reached 100%, and the conversion rate of thiol reached 96.2% and 92%, respectively. The double bond and thiol had an optimal ratio of homopolymerization and copolymerization, and the coatings had good film-forming properties. However, excessive C=C content led to homopolymerization and lower thiol conversion rate (Figure 2e,f), and the -SH conversion rates of samples S₅₋xDC and S₅₋xDC decreased to 87.6% and 81.8%, respectively. At this time, the oxygen inhibition effect in the homopolymerization reaction gradually appeared, so the double bond conversion rate also decreased to a certain extent. However, the double bond
conversion rate of sample $S_1(N_3P_2)_9$ remained at 92.8%. This showed that the presence of -SH can effectively counter oxygen inhibition and ensure the coating has good film-forming properties. To sum up, the entire coatings system had good curing efficiency, good film-forming performance, and could form a complete cross-linked network structure.

**Figure 2.** The Fourier Transform Infrared Spectroscopy (FTIR) spectra of sample $S_1(N_3P_2)_6$, which were recorded during the UV-curing (a); the enlarged FTIR spectra of the thiol group (2572 cm$^{-1}$) and the acrylate double bond (1637 cm$^{-1}$) of the coating $S_1(N_3P_2)_6$, after being subjected to UV-irradiation for various durations (b); the double bond conversion rate versus the irradiation time (c,e); and the thiol group conversion versus the irradiation time (d,f).
Table 2. The properties of the UV-cured, flame-retardant coatings.

| Coating   | LOI (%) | HBR (mm/min) or Self-Extinguishing Time (s) | C=O Conversion (%) | -SH Conversion (%) |
|-----------|---------|------------------------------------------|-------------------|-------------------|
| FC        | 22.8    | 14.3 mm/min                              | -                 | -                 |
| S₅N₅P₀   | 25.6    | 5.8 mm/min                               | 98.6              | 81.1              |
| S₅N₅P₁   | 26.7    | 125 s                                    | 99.7              | 79.0              |
| S₅N₅P₂   | 27.2    | 79 s                                     | 98.8              | 70.9              |
| S₅N₅P₃   | 26.9    | 94 s                                     | 98.2              | 65.4              |
| S₅N₅P₄   | 26.5    | 94 s                                     | 99.7              | 65.2              |
| S₅N₅P₅   | 26.3    | 91 s                                     | 99.3              | 62.0              |
| S₅(N₃P₂)₆| 28.6    | 85 s                                     | 100               | 92.0              |
| S₅(N₃P₂)₇| 27.0    | 140 s                                    | 100               | 96.2              |
| S₅(N₃P₂)₈| 25.9    | 214 s                                    | 97.1              | 87.6              |
| S₅(N₃P₂)₉| 25.5    | 220 s                                    | 92.8              | 81.8              |

3.2. Flame Retardancy of Fiber Composites

The formulation of the coating is the main factor affecting the flame-retardant performance. From Table 2, the proportion of flame-retardant elements was appropriate when the molar ratio of N and P was 3:2, and it played a synergistic flame-retardant role. The sample S₅N₅P₂ had the best flame-retardant performance among the S₅N₅P₆ series. The LOI value reached 27.2%, and was extinguished after 79 s. In addition, when the molar ratio of SH and C=O was 4:6, sample S₅(N₃P₂)₆ had the highest LOI value, up to 28.6%. In the horizontal combustion test, it quickly extinguished after being ignited for 85 s, which showed that S could also be used as a flame-retardant element and synergistically a flame-retardant with N and P elements. Figure 3 showed the carbon residue morphology of some samples after the horizontal combustion test. It can be seen that samples FC and S₅N₅P₀ were not self-extinguishing during the horizontal combustion test. The FC carbon layer was an amorphous and loose coke residue, and no dense carbon layer was formed. Sample S₅N₅P₀ formed a part of the carbon layer on the surface, but the carbon layer was disconnected and broken, and thus did not effectively prevent the material from burning. The samples containing P elements all self-extinguished, indicating that P elements have obvious solid-phase flame-retardant effects. In general, when the N, P, and S elements coexisted in the coating, it showed an obvious synergistic effect of the gas-phase flame retardant and a solid-phase flame retardant, and it obtained better flame retardancy.

Figure 3. The residual carbon layer sample after a horizontal combustion test.

3.3. Combustion Behavior

The cone calorimetry test can obtain the ignition time (TTI), the heat release rate (HRR), the peak heat release rate (PHRR), the total heat release rate (THR), the effective heat of combustion (EHC), the smoke production rate (SPR), the total smoke release (TSP), the CO and CO₂ production, and other important parameters that can evaluate the potential fire hazard of materials [4,22]. Samples S₅N₅P₀, S₅N₅P₂, S₅N₅P₅, S₅(N₃P₂)₆, and S₅(N₃P₂)₉ were selected for cone calorimetry analysis to further study the flame-retardant properties of the coating. The test results are shown in Figure 4 and Table 3. It is worth noting that
due to the presence of flame-retardant elements such as N, P, and S, the coating formed an expanded carbon layer on the surface of the composites in the combustion process. The carbon layer could be used as a good thermal insulation layer, so HRR had a downward trend in 40–70 s, resulting in the first peak of the HRR curve. Subsequently, the matrix composites gradually burned, degraded, and released flammable products from the cracks of the carbon layer under the action of heat. The existence of the fireproof layer prevented the spread of flame and the combustion of the matrix materials, which resulted in a second peak in the HRR curve. In the HRR curve, sample S₄(N₃P₂)₆ exhibited the lowest PHRR value, indicating that sample S₄(N₃P₂)₆ had a good flame-retardant effect. As shown in Figure 4b and Table 3, the THR values of all the coated composites were not much different because all samples were completely consumed during the cone calorimetry. The types of flame-retardant elements contained in the coating could slightly affect the overall flammability of the fiber composites. When the coating contained only S and N (sample S₅N₅P₀) or only S and P (sample S₅N₀P₅), the HRR of the composites was 52.44 MJ/m² and 50.89 MJ/m², which was higher than the coatings containing N, P, and S elements (around 47 MJ/m²). This could indicate that the three flame-retardant elements of N, P, and S had a synergistic flame-retardant effect. However, the three elements’ molar ratio had little effect on the HRR values of combustion. In addition, it must be mentioned that the TTI value of the coating samples with more phosphorus content decreased slightly, which might be attributed to the rapid rupture of the P-O-C bond and promoted the formation of the flame-retardant carbon layer.

Generally speaking, smoke and toxic gases will threaten people’s health by poisoning and suffocating them in a real-life fire scene. The curves of the SPR and TSP of the samples are shown in Figure 4c,d, and the relevant data are summarized in Table 3. The trend of the SPR curve was similar to the HRR curve. As shown in Figure 4c, the composites reached the first peak in 20 s and then decreased rapidly in 20–60 s. Sample S₄(N₃P₂)₆ had a higher initial peak value, which was mainly attributed to the violent combustion of the phosphorous-contained coatings, and then the phosphate groups formed a dense carbon layer during the combustion process. This caused the second peak of the SPR curve of sample S₄(N₃P₂)₆ to be lower than other samples. Therefore, sample S₄(N₃P₂)₆ had the lowest TSP value, reaching 2.18 m². This showed that the synergistic effect of N, S, and P elements could reduce smoke production in an appropriate proportion. CO and CO₂ production (Figure 4e,f) were also obtained to study the toxicity of burning products [22]. An increase in the CO production value of the coated composite was observed in the first 30 s, which might have been caused by rapid degradation and incomplete combustion of the coating. In the first 100 s since the formed carbon layer effectively prevented the escape of combustible gas CO, the CO production was significantly reduced within 100–200 s. However, there was a slow increase in CO production at the end of the combustion. This might have been because the early combustion consumed a lot of oxygen, resulting in insufficient oxygen content and incomplete combustion in the later stage of the combustion. Meanwhile, the TSP and CO production of the coatings with the N element were significantly reduced, indicating that the N element had an obvious gas-phase flame-retardant effect, which could effectively reduce the smoke emission. In addition, compared with other coating composites, sample S₄(N₃P₂)₆ had significantly lower CO₂ production, which could effectively reduce the risk of suffocation in a real fire.
Figure 4. (a) heat release rate (HRR), (b) total heat release rate (THR), (c) the smoke production rate (SPR), (d) the total smoke release (TSP), (e) CO production, and (f) CO$_2$ production curves as functions of the combustion time for the coated fiber composites.

Table 3. The cone calorimetry data of the coated composites.

| Samples          | TTI | PHRR  | Mean EHC | THR  | TSR  | TSP  |
|------------------|-----|-------|----------|------|------|------|
| $S_5N_5P_0$      | 20  | 384.42| 21.33    | 52.44| 243.55| 2.36 |
| $S_5N_3P_2$      | 11  | 294.55| 21.37    | 47.57| 254.18| 2.32 |
| $S_5N_0P_5$      | 10  | 296.37| 19.84    | 50.89| 434.56| 3.96 |
| $S_4(N_3P_2)_6$  | 18  | 279.25| 23.73    | 47.80| 217.03| 2.18 |
| $S_4(N_5P_2)_9$  | 7   | 362.53| 20.26    | 46.41| 321.83| 2.93 |
3.4. Thermogravimetric Analysis

The influence of the changes of S, N, and P elements in the coating on the thermal stability of the composite was studied by TG and DTG curves. The TG and DTG curves of UV-coated composites are shown in Figures 5 and 6. Table 4 showed the thermogravimetric parameters, including the initial degradation temperature ($T_{5\%}$), the temperature corresponding to the maximum degradation rate ($T_{max1}$, $T_{max2}$), and the carbon residue $Y_c$ at 700 °C. The sample FC had a high initial degradation temperature, and $Y_c$ can only reach 6.55% and 0.79% in N$_2$ and Air, respectively. With the increase of PM-2 content, the initial degradation temperature of the coated composites was gradually decreasing. This was mainly due to the lower bond energy of the P-O-C bond, which was not as stable as the C-C bond [23,24]. A small amount of the degradation of the coated composites below 200 °C was mainly due to the volatilization and degradation of a small part of the small-molecule impurities that were not fixed in the cross-linked network. In the DTG curve, the degradation process of FC corresponds to the thermal decomposition of hemicellulose (250–300 °C), cellulose (320–390 °C), and polypropylene (400–500 °C). The degradation peak (240–400 °C) in the first stage of the coated composites mainly corresponded to the breaking of the C-S bond in the cross-linking network, the volatilization of volatiles containing C and S (SO$_2$, SO$_3$, and CO$_2$), the decomposition of the phosphate group in the PM-2 structure, the scission of long chain branches in the solidified layer, and the preliminary formation of the carbon layer. The degradation of the cellulose and hemicellulose of the matrix composites was also included in this degradation peak. The degradation peak in the second stage was mainly the rupture of the cross-linked network and the main chain of the polypropylene product in the matrix material. Then, the entire matrix of the composites was almost completely degraded, leaving only part of the carbon residue [25,26]. In air atmosphere, the first degradation peak is at 200–300 °C and the second degradation peak ranges from 300–450 °C. Both decomposition stages are advanced, which indicates that the presence of oxygen accelerates the degradation process of the composite. The third degradation peak (at 500–600 °C) occurs for the thermal degradation in an air atmosphere, which is mainly attributed to the degradation of unstable carbon layers at high temperatures. It can be seen from Table 4 that, compared with $S_1(N_3P_2)_y$, $S_4(N_3P_2)_6$ and $S_5N_3P_2$ exhibited a good synergistic flame-retardant effect due to the appropriate relative proportions of N, P, and S, and $T_{max2}$ was also higher than other coating composites. The gas-phase and the solid-phase flame-retardant mechanism worked together and delayed the degradation process to a certain extent. In the $S_xN_yP_z$ series, sample $S_8N_0P_3$ had the highest $Y_c$ at 700 °C. However, in the samples of the $S_x(N_3P_2)_y$ series, even if the PM-2 content increased, $Y_c$ did not change significantly. This was mainly due to the decrease in the content of PETMP, which was the carbon chain provider. In addition, it can be observed from the DTG curves that due to the presence of the coating, the peak degradation rate of all coated composites was much lower than that of the uncoated fiber composite materials, which shows that the coating effectively reduced the degradation rate.
rate of all coated composites was much lower than that of the uncoated fiber composite materials, which shows that the coating effectively reduced the degradation rate.

Figure 5. The thermogravimetric (TG) analysis (a) curves and differential thermogravimetric (DTG) curves (b) of the series of $S_N^xP_y$, and the TG (c) curves and DTG curves (d) of the series of $S_x(N_3P_2)_y$ (in an $N_2$ atmosphere).

Figure 6. The TG (a) curves and DTG curves (b) of the series of $S_N^xP_y$, and the TG (c) curves and DTG curves (d) of the series of $S_x(N_3P_2)_y$ (in an air atmosphere).
Table 4. The thermal properties of the UV-cured composites.

| Samples   | $T_{5\%}$ (°C) | $T_{\text{max1}}$ (°C) | $T_{\text{max2}}$ (°C) | $Y_c$ (wt%) |
|-----------|----------------|-------------------------|-------------------------|-------------|
|           | $N_2$ | Air | $N_2$ | Air | $N_2$ | Air | $N_2$ | Air |
| FC        | 296    | 251 | 355   | -    | 435    | 349 | 6.55    | 0.79 |
| S$_2$N$_3$P$_0$ | 291   | 271 | 378   | 285  | 450    | 390 | 9.59    | 0.69 |
| S$_2$N$_4$P$_1$ | 265   | 259 | 376   | 291  | 458    | 368 | 11.14   | 1.51 |
| S$_2$N$_3$P$_2$ | 252   | 251 | 350   | 271  | 456    | 366 | 14.49   | 3.50 |
| S$_2$N$_2$P$_3$ | 251   | 243 | 291   | 290  | 464    | 365 | 16.53   | 4.64 |
| S$_2$N$_1$P$_4$ | 244   | 240 | 277   | 291  | 449    | 357 | 18.41   | 6.68 |
| S$_2$N$_0$P$_5$ | 237   | 239 | 276   | 269  | 422    | 366 | 20.78   | 5.82 |
| S$_4$(N$_3$P$_2$)$_6$ | 258   | 252 | 342   | 280  | 465    | 365 | 13.76   | 8.46 |
| S$_2$(N$_3$P$_2$)$_7$ | 238   | 244 | 285   | 271  | 459    | 357 | 14.10   | 4.88 |
| S$_2$(N$_3$P$_2$)$_8$ | 219   | 224 | 272   | 264  | 463    | 345 | 14.94   | 4.63 |
| S$_1$(N$_3$P$_2$)$_9$ | 225   | 200 | 271   | 264  | 456    | 388 | 14.49   | 4.37 |

3.5. Char Residues Surface Morphologies Analysis and Elements Analysis

After the horizontal combustion test, all samples were imaged by SEM to evaluate the surface morphology and chemical composition of the residual carbon (Figure 7). According to the previous literature [27], the dense carbon layer could isolate thermal oxygen and other volatile combustible gases well, thereby improving the flame retardancy [28]. The coating with a high content of PM-2 had a complete carbon layer structure. When the content of the PETMP groups was reduced (Figure 7f), the sufficient carbon layer could not form completely on the surface of the composites. There were cracks and through holes on the carbon layer. Meanwhile, the gas-phase flame-retardant did not have a sufficient effect to achieve a relative balance with the solid-phase flame-retardant mechanism, so the flame-retardant effect gradually decreased. In contrast, in the S$_2$N$_3$P$_2$ and S$_4$(N$_3$P$_2$)$_6$ coatings (Figure 7c,e), pores and closed bubbles appear in the carbon layer, which are caused by the release of incombustible gas during the combustion process. The carbon layer showed a honeycomb structure, which formed a temperature gradient on the carbon residue layer, inhibiting heat transfer, preventing oxygen diffusion, and providing better heat insulation performance for the base material [24,27,29,30].

Figure 7. The micro-morphologies of the char layer structure of the uncoated composite FC (a) and the coated composites samples of S$_2$N$_3$P$_0$ (b), S$_2$N$_3$P$_2$ (c), S$_2$N$_0$P$_5$ (d), S$_2$(N$_3$P$_2$)$_6$ (e), and S$_1$(N$_3$P$_2$)$_9$ (f), after horizontal burning rate (HBR) tests.
According to EDS analysis (Figure 8), the N element in samples completely disappeared, which indicated that the N element completely generated non-flammable gas during the combustion process (Table 5). The carbon residue on the surface of the $S_4(N_3P_2)_6$ carbon layer contained 69.53 wt% of C, 6.55 wt% of P, and 1.42 wt% of S. Compared with the element content before the combustion (Table 6), the content of S element was reduced, which showed the element S participated in the flame-retardant reaction through the gas phase flame-retardant mechanism. The phosphate group in the coatings generated polyphosphoric acid and its derivatives to promote the formation of the carbon layer during combustion. The P element in the coating was almost completely deposited in the carbon layer. In summary, the coating formed a carbonaceous layer rich in phosphorus on the surface of the composites in the combustion process, which delayed the pyrolysis reaction during the combustion process and had a good flame-retardant effect.

![Figure 8](image-url)

**Figure 8.** The energy-dispersive spectrometer (EDS) elements petra of sample $S_4(N_3P_2)_6$.

**Table 5.** The element percentage content of the char layer surface of $S_5N_0P_5$, $S_5N_3P_2$, $S_5N_5P_0$, $S_4(N_3P_2)_6$, and $S_1(N_3P_2)_9$, according to EDS tests.

| Coating     | C (wt%) | O (wt%) | N (wt%) | P (wt%) | S (wt%) |
|-------------|---------|---------|---------|---------|---------|
| $S_5N_0P_5$ | 83.73   | 11.89   | 0       | 0.5     | 3.88    |
| $S_5N_3P_2$ | 77.37   | 14.91   | 0       | 4.97    | 2.75    |
| $S_5N_5P_0$ | 59.63   | 26.25   | 0       | 12.72   | 1.41    |
| $S_4(N_3P_2)_6$ | 69.53 | 22.51   | 0       | 6.55    | 1.42    |
| $S_1(N_3P_2)_9$ | 68.79 | 20.71   | 0       | 10.5    | 0       |
Table 6. The mass fractions of N, P, and S contained in films of $S_5N_0P_5$, $S_5N_3P_2$, $S_5N_5P_0$, $S_4(N_3P_2)_6$, and $S_1(N_3P_2)_9$.

| Coating       | C-Content (wt%) | O-Content (wt%) | N-Content (wt%) | P-Content (wt%) | S-Content (wt%) |
|---------------|----------------|----------------|----------------|----------------|----------------|
| $S_5N_5P_0$   | 48.23          | 23.39          | 6.82           | 0              | 15.59          |
| $S_5N_3P_2$   | 42.41          | 31.27          | 3.28           | 4.85           | 12.51          |
| $S_5N_0P_5$   | 37.05          | 38.55          | 0              | 9.34           | 9.64           |
| $S_1(N_3P_2)_6$ | 42.58        | 32.23          | 3.90           | 5.76           | 9.92           |
| $S_1(N_3P_2)_9$ | 42.93        | 34.99          | 5.70           | 8.42           | 2.41           |

3.6. Adhesion and Mechanical Testing

Table 7 shows the tensile strength, flexural strength, and adhesion of the coated composites. When the molar ratio of -SH and C=C was 1:1, the change of TAC and PM-2 content had relatively little effect on adhesion. The coating did not fall off at all after the tests, and the adhesion grade was 0. This was mainly due to the fact that the thiol and the double bond underwent a click reaction via a free-radical step-by-step copolymerization [5,7,31,32]. The cross-linked network structure formed in this way had low shrinkage and high monomer conversion. In addition, PM-2 formed the anchor effect with PP on the surface of the composites through the penetration of its phosphate ester, so the coatings had better adhesion. Because of the formation of flexible thioether bonds, the lack of rigid structure led to the poor mechanical properties of the coating [33], so the tensile strength of the $S_5N_xP_y$ samples did not obviously increase. However, due to the better toughness of the coating produced by this method, the bending strength fluctuated about 42.6 Mpa, which was an increase of 24.3% compared with the uncoated composites.

Table 7. The mechanical properties of the coated composites and the adhesion grade of the coatings.

| Samples | Tensile Strength (MPa) | Flexural Strength (MPa) | Adhesion (Grade) |
|---------|------------------------|-------------------------|------------------|
|         |                        |                         | Coating Loss Rate (%) | Adhesion Grade |
| FC      | 26.05 ± 0.85           | 34.27 ± 2.50            | -                 | -              |
| $S_5N_5P_0$ | 24.88 ± 2.55         | 44.65 ± 1.45           | 0                 | 0              |
| $S_5N_3P_2$ | 25.87 ± 3.57         | 44.39 ± 2.62           | 0                 | 0              |
| $S_5N_0P_5$ | 25.76 ± 3.82         | 42.52 ± 2.56           | 0                 | 0              |
| $S_1(N_3P_2)_6$ | 24.38 ± 2.60       | 39.96 ± 3.35           | 0                 | 0              |
| $S_1(N_3P_2)_9$ | 25.24 ± 2.54       | 45.15 ± 2.81           | 0                 | 0              |
| $S_0(N_3P_2)_5$ | 24.15 ± 2.61       | 40.90 ± 2.36           | 0                 | 0              |
| $S_4(N_3P_2)_6$ | 26.11 ± 1.83        | 42.02 ± 2.74           | 3                 | 1              |
| $S_7(N_3P_2)_7$ | 28.34 ± 2.52        | 48.11 ± 1.64           | 5                 | 1              |
| $S_8(N_3P_2)_8$ | 28.95 ± 0.53        | 50.29 ± 2.29           | 14                | 2              |
| $S_9(N_3P_2)_9$ | 30.67 ± 2.63        | 52.31 ± 1.98           | 48                | 4              |

By fixing the relative content of TAC and PM-2 and changing the ratio of thiol to a double bond, the following dynamic emerged: the lower the thiol content, the worse the adhesion of the coating. The peeling rate of sample $S_1(N_3P_2)_9$ after the adhesion test even reached 48%. As the content of double bonds increases, the cross-linking of the coating tends to homopolymerize between the double bonds. The internal stress was not released due to the rapid curing of the coating, and the formed coating had a large volume shrinkage rate and high surface tension [7,34], thereby reducing the adhesion between the coating and the substrate. Additionally, the influence of the coating on the mechanical properties was related to the crosslinking method of the coating. The cross-linking density of the coating formed by the double-bond homopolymerization reaction became larger, which increased the hardness of the coating. Therefore, the tensile strength and flexural strength increased with the increase of the double bond content.
3.7. Optical Property

Figure 9 is the optical transparency spectrum of the UV-cured coatings. The thickness of the coatings was 200 µm. It was obvious from the digital photos of Figure 9b that the crosslinked coating had good transparency. The UV-vis spectrum showed that the transparency of these coatings at visible light wavelengths (400–800 nm) was about 90%, which was consistent with the performance of digital photos. The transmittance of the coating was close to 0 under 300 nm UV light, which indicated that the coating had a good blocking effect on short-wave UV light [20]. The relative content changes of PETMP, PM-2, and TAC in the film had little effect on the optical transparency of the UV curable film [5]. This kind of film with high transparency showed obvious application potential.

![Figure 9. The UV-vis transmittance spectra of the UV-cured coatings (a), and a digital photo of sample S4(N3P2)b (b).](image)

4. Conclusions

The UV-cured, flame-retardant coatings containing N/P/S were successfully prepared on the surface of the composites via a thiol-ene click reaction. The effect of the element ratio on the flame-retardant properties of the coating was investigated. Among the PETMP/TAC/PM-2 crosslinked networks, sample S4(N3P2)b showed the best flame retardancy, which was determined by LOI, HBR, and CCT. The LOI value of sample S4(N3P2)b reached 28.6%, and its self-extinguishing time was 85 s in the horizontal combustion test. Meanwhile, it had the lowest THR value, the lowest PHRR value, and the lowest TSP value. All these results confirmed the synergistic effectiveness of flame retardants between PETMP, TAC, and the PM-2. The thermal degradation results of the UV-cured coatings revealed that the P-containing prepolymers could promote the carbon chain into a compact and stable char layer in a timely fashion, which demonstrated that P and C exert a synergistic flame retardancy in the condensed phase. Moreover, the coatings consumed oxygen and generated CO2 and SO2, and N2 and NH3, which could dilute combustible gas produced during combustion around the fiber composites. This indicated that N and S exert flame retardancy in the gas phase, leading to the formation of the honeycomb porous carbon layer. This carbon layer was monitored in the SEM test. In addition, the UV-curable coating S4(N3P2)b also had a high double bond conversion rate and thiol conversion rate (100% and 92%, respectively) and had good mechanical properties. The flexural strength of the composites was increased to 42.02 MPa. The coating had good adhesion, and the shedding rate was within 5%. The transparency of this coating was about 90%, and it had good optical properties. The aforementioned results indicated that the PETMP/TAC/PM-2 cross-linked network had potential in the flame-retardant coating field. Therefore, this research provided new ideas for the development of flame-retardant coatings that meet the needs of practical applications.
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References

1. Wang, T.; Liu, T.; Ma, T.; Li, L.; Wang, Q.; Guo, C. Study on degradation of phosphorus and nitrogen composite UV-cured flame retardant coating on wood surface. Prog. Org. Coat. 2018, 124, 240–248. [CrossRef]
2. Wang, T.S.; Li, L.P.; Wang, Q.W.; Xie, G.J.; Guo, C.G. Castor oil based UV-cured coatings using thiol-ene click reaction for thermal degradation with flame retardance. Ind. Prod. Prog. 2019, 141, 111798. [CrossRef]
3. Wang, T.; Li, L.; Cao, Y.; Wang, Q.; Guo, C. Preparation and flame retardancy of castor oil based UV-cured flame retardant coating containing P/Si on wood surface. Ind. Crops Prod. 2019, 130, 562–570. [CrossRef]
4. Huang, Y.B.; Jiang, S.H.; Liang, R.C.; Liao, Z.W.; You, G.X. A green highly-effective surface flame-retardant strategy for rigid polyurethane foam: Transforming UV-cured coating into intumescent self-extinguishing layer. Compos. Part A Appl. Sci. Manuf. 2019, 125, 125. [CrossRef]
5. Xiao, Z.F.; Li, S.L.; Zhang, Z.J.; Mai, C.; Xie, Y.J.; Wang, Q.W. Fire retardancy of an aqueous, intumescent, and translucent wood varnish based on guanylurea phosphate and melamine-urea-formaldehyde resin. Prog. Org. Coat. 2018, 121, 64–72. [CrossRef]
6. Lokhande, G.; Chambhare, S.; Jagtap, R. Synthesis and properties of phosphate-based diacrylate reactive diluent applied to UV-curable flame-retardant wood coating. J. Coat. Technol. Res. 2017, 14, 255–266. [CrossRef]
7. Banerjee, J.; Pangule, R.C.; Kane, R.S. Antifouling Coatings: Recent Developments in the Design of Surfaces That Prevent Fouling by Proteins, Bacteria, and Marine Organisms. Adv. Mater. 2011, 23, 690–718. [CrossRef]
8. Wei, D.D.; Liao, B.; Huang, J.H.; Zhang, M.; Pang, H. Fabrication of castor oil-based hyperbranched urethane acrylate UV-curable coatings via thiol-ene click reactions. Prog. Org. Coat. 2019, 135, 114–122. [CrossRef]
9. Contreras, PP.; Agarwal, S. Photo-polymerizable, low shrinking modular construction kit with high efficiency based on vinylcyclopropanes. Polym. Chem. 2016, 7, 3100–3106. [CrossRef]
10. Alvankarian, J.; Majlis, B.Y. Exploiting the Oxygen Inhibitory Effect on UV Curing in Microfabrication: A Modified Lithography Technique. PLoS ONE 2015, 10, e0119658. [CrossRef]
11. Corcione, C.E.; Striani, R.; Frigione, M. UV-cured methacrylic-silica hybrids: Effect of oxygen inhibition on photo-curing kinetics. Thermochim. Acta. 2014, 576, 47–55. [CrossRef]
12. Husar, B.; Ligon, S.C.; Wutzel, H.; Hoffmann, H.; Liska, R. The formulator’s guide to anti-oxygen inhibition additives. Prog. Org. Coat. 2014, 77, 1789–1798. [CrossRef]
13. Chen, W.B.; Wang, L.; Xie, Y.J.; Zhang, M.; Zhang, H.; Tang, G.F. Synthesis and preliminary photopolymerization evaluation of novel photoinitiators containing phototrigger to overcome oxygen inhibition in the UV-curing system. J. Photochem. Photobiol. A Chem. 2020, 388, 112187. [CrossRef]
14. Wei, J.; Liu, F. Novel Highly Efficient Macrophotoinitiator Comprising Benzophenone, Cointiator Amine, and Thiouroles for Photopolymerization. Macromolecules 2009, 42, 5486–5491. [CrossRef]
15. Roper, T.; Kwee, T.; Lee, T.; Guymon, C.A.; Hoyle, C.E. Photopolymerization of pigmented thiol ene systems. Polymer 2004, 45, 2921–2929. [CrossRef]
16. Wang, Q.; Chen, G.X.; Cui, Y.Y.; Tian, J.F.; He, M.H.; Yang, J.W. Castor Oil Based Biocatalyst as a Highly Stable and Self-Initiated Oligomer for Photoinitiator-Free UV Coatings. ACS Sustain. Chem. Eng. 2017, 5, 376–381. [CrossRef]
17. Liu, C.H.; Li, T.C.; Zhang, J.H.; Chen, S.F.; Xu, Z.C.; Zhang, A.Q.; Zhang, D.H. Preparation and properties of phosphorous-nitrogen containing UV-curable polymeric coatings based on thiol-ene click reaction. Prog. Org. Coat. 2016, 90, 21–27. [CrossRef]
18. Wang, Z.Y.; Liang, H.B.; Yang, H.T.; Xiong, L.; Zhou, J.P.; Huang, S.M.; Zhao, C.H.; Zhong, J.; Fan, X.F. UV-curable self-healing polyurethane coating based on thiol-ene and Diels-Alder double click reactions. Prog. Org. Coatings 2019, 137, 105282. [CrossRef]
20. Miao, J.T.; Yuan, L.; Guan, Q.B.; Liang, G.Z.; Gu, A.J. Water-Phase Synthesis of a Biobased Allyl Compound for Building UV-Curable Flexible Thiol Ene Polymer Networks with High Mechanical Strength and Transparency. *Acs Sustain. Chem. Eng.* 2018, 6, 2902–2909. [CrossRef]

21. Wu, C.D.; Wu, J.L.; Ma, K.K.; Zhang, D.L.; Xiong, S.Y.; Zhang, J.; Luo, G.Q.; Chen, F.; Shen, Q.; Zhang, L.M.; et al. Synthesis of AA7075-AA7075/B4C bilayer composite with enhanced mechanical strength via plasma activated sintering. *J. Alloys Compd.* 2017, 701, 416–424. [CrossRef]

22. Huang, Y.B.; Jiang, S.H.; Liang, R.C.; Sun, P.; Hai, Y.; Zhang, L. Thermal-triggered insulating fireproof layers: A novel fire-extinguishing MXene composites coating. *Chem. Eng. J.* 2020, 391, 123621. [CrossRef]

23. Wang, X.; Song, L.; Xing, W.Y.; Lu, H.D.; Hu, Y.A. A effective flame retardant for epoxy resins based on poly(DOPO substituted dihydroxyl phenyl pentaerythritol diphosphonate). *Mater. Chem. Phys.* 2011, 125, 536–541. [CrossRef]

24. Phalak, G.; Patil, D.; Patil, A.; Mhaske, S. Synthesis of acrylated cardanol diphenyl phosphate for UV curable flame-retardant coating application. *Eur. Polym. J.* 2019, 121, 109320. [CrossRef]

25. Xing, W.Y.; Hu, Y.; Song, L.; Chen, X.L.; Zhang, P.; Ni, J.X. Thermal degradation and combustion of a novel UV curable coating containing phosphorus. *Polym. Degrad. Stab.* 2009, 94, 1176–1182. [CrossRef]

26. Liu, R.; Zhu, G.G.; Li, Z.Q.; Liu, X.Y.; Chen, Z.G.; Ariyasivam, S. Cardanol-based oligomers with “hard core, flexible shell” structures: From synthesis to UV curing applications. *Green Chem.* 2015, 17, 3319–3325. [CrossRef]

27. Qu, L. Research on Properties of Phosphorus/Nitrogen/Silicon/Graphene-Containing Synergic Flame-Retardant Epoxy resin. Ph.D. Thesis, Jilin University, Changchun, China, 2020.

28. Ma, T.T.; Guo, C.G. Synergistic effect between melamine cyanurate and a novel flame retardant curing agent containing a caged bicyclic phosphate on flame retardancy and thermal behavior of epoxy resins. *J. Anal. Appl. Pyrolysis* 2017, 124, 239–246. [CrossRef]

29. Bai, Z.M.; Song, L.; Hu, Y.; Yuen, R.K.K. Preparation, Flame Retardancy, and Thermal Degradation of Unsaturated Polyester Resin Modified with a Novel Phosphorus Containing Acrylate. *Ind. Eng. Chem. Res.* 2013, 52, 12855–12864. [CrossRef]

30. Sun, J.; Wang, X.D.; Wu, D.Z. Novel Spirocyclic Phosphazene-Based Epoxy Resin for Halogen-Free Fire Resistance: Synthesis, Curing Behaviors, and Flammability Characteristics. *Acs Appl. Mater. Interfaces* 2012, 4, 4047–4061. [CrossRef]

31. Carioscia, J.A.; Lu, H.; Stanbury, J.W.; Bowman, C.N. Thiol-ene oligomers as dental restorative materials. *Dent. Mater.* 2005, 21, 1137–1143. [CrossRef]

32. Cramer, N.B.; Scott, J.P.; Bowman, C.N. Photopolymerizations of thiol-ene polymers without photoinitiators. *Macromolecules* 2002, 35, 5361–5365. [CrossRef]

33. Duan, W.; Li, M.; Sun, H.; Cheng, J.; Zhang, J. Preparation and charactrization of hybrid thiol-ene/epoxy UV-thermal dual-curing systems. *Adhesion* 2017, 38, 16–22.

34. Chiang, T.H.; Hsieh, T.E. A study of monomer’s effect on adhesion strength of UV-curable resins. *Int. J. Adhes. Adhes.* 2006, 26, 520–531. [CrossRef]