Effects of Peroxide Initiator on the Structure and Properties of Ultralow Loss Thermosetting Polyphenylene Oxide-Based Composite

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Abstract: Although thermosetting polyphenylene oxide-(PPO) based composites with excellent dielectric properties have been widely accepted as superior resin matrices of high-performance copper clad laminate (CCL) for 5G network devices, there has been limited information regarding the composition–process–structure–property relationships of the systems. In this work, the effects of peroxide initiator concentration on the structure and dielectric properties of a free radical cured ultralow loss PPO/Triallyl isocyanate (TAIC) composite system were studied. As expected, the glass transition temperature ($T_g$) and storage modulus increased with the advancing of crosslinking, whereas the dielectric loss showed an “abnormal” rise with the increase in crosslink density. Extensive studies were carried out by varying the initiator contents and characterizing the structure with spectroscopy, thermal analysis, and positron annihilation lifetime spectrum (PALS) techniques. The results show that the competition of polarity, crosslink density, free volume, and free TAIC are the key factors determining the dielectric properties of the composites.

Keywords: copper clad; thermosetting PPO; dielectric properties; peroxide initiator

1. Introduction

The rapid development of 5G communication technology has initiated a large-scale, worldwide effort to develop more sophisticated electronic systems requiring high-performance dielectric materials for printed circuit boards (PCBs). The main development directions of PCBs in the 5G era are a high layer count design, high frequency, and high levels of signal integrity, as well as high wiring density, which have incurred more stringent demands on the PCB substrate materials, i.e., copper clad laminates (CCL). In addition to lead-free compatibility and reliability requirements, a low dielectric constant ($\varepsilon_r$) and extremely low dielectric loss ($\tan\delta$) under high frequency are the most challenging material properties which have attracted great attention in both industry and academic institutions [1–5].

The $\varepsilon_r$ and $\tan\delta$ of traditional thermosetting CCL matrix materials, such as epoxy resin and cyanate ester (CE) resin, are intrinsically too high to meet the requirements of high-frequency applications [6,7]. Therefore, new polymer systems with low $\varepsilon_r$ and $\tan\delta$ are being developed for high-frequency and high-speed applications. Among the available polymers, polyphenylene oxide (PPO) has great potential since it has a highly rigid skeleton structure with aromatic rings, without hydrolysable groups or polar side...
groups. These structural characteristics endow PPO with excellent hydrolytic stability, low moisture absorption, high thermal stability, and excellent electrical properties in a wide temperature range [8–11].

For PCB applications, thermosetting PPO systems with excellent performance have been developed by capping the ends of PPO oligomers with reactive groups and curing with chemicals containing multiple functional groups, such as triallyl isocyanurate (TAIC), divinyl benzene, and 1,2-polybutadiene (PB), etc. [12–18], or copolymerizing with epoxy, cyanate ester, and benzoxazine, etc. [19–29]. Among the most significant developments, Peters et al. [30] invented a methyl methacrylate capped bi-functional PPO oligomer (M-PPE-M) which was later commercialized by SABIC under the tradename of NORYLTM SA9000 and studied several cured systems with t-butyl styrene, divinyl benzene, TAIC, dibrominostyrene, etc., as crosslinking agents [31–33]. Outstanding dielectric, thermal and mechanical properties, as well as good water resistance, were achieved simultaneously in these systems. For example, for the TAIC cured systems at 1GHz, the dielectric constant slightly changed between 2.5 and 2.57, whereas the dielectric loss was always as low as about 0.001 over a wide composition range (M-PPE-M/TAIC = 0% ~ 80%) [32]. Such dielectric properties were much better than that of traditional PCB substrate materials, such as epoxy-based ($D_k \approx 2.85$, $D_f \approx 0.013$ at 1GHz) and cyanate ester-based systems ($D_k \approx 2.73$, $D_f \approx 0.0051 ~ 0.0117$ at 1GHz) [32,34].

In the past decade, several PPO-based substrate materials with similar formulations have been successfully commercialized and have started to play key roles in 5G devices [12–16] and a lot of research has been carried out regarding the combination of PPO oligomers with different crosslink agents, copolymers, or reinforcement components (DVB, PB, Epoxy, CE, etc.) [17–29]. However, little work has been reported regarding the characterization of chemical structures and the relationships between crosslinked structures and the dielectric properties of thermosetting PPO-based ultralow dielectric loss systems.

The dielectric properties of polymers are related to their polarizability. The relationship between the molar polarizability $P$, the dielectric constant, and the molecular polarization $\alpha$ can be expressed in Equation [35]:

$$P = \frac{\varepsilon_r - 1}{\varepsilon_r + 2} \times \frac{M}{\rho} = \frac{N_A \alpha}{3 \varepsilon_0}$$

where $P$ is the molar polarizability, $\varepsilon_r$ is the relative permittivity, $\varepsilon_0$ is the permittivity in vacuum, $M$ is molecular weight of a repeat unit, $\rho$ is density, $\alpha$ is polarizability, and $N_A$ is the Avogadro constant.

Molecular polarizability refers to the proportionality constant for the formation of dipole under the influence of an electric field [35]. Based on the relation, all the molecular and structural parameters that enhance polarizability and density, such as the higher polarity of functional groups or atoms, higher molecular movability, inclusion of polar or conductive impurities, and less free volume, etc., will lead to a higher dielectric constant, and vice versa [35–45]. Dielectric loss is related to the energy dissipated due to the oscillating electric field. It is also related to the dielectric relaxation time $\tau$, which is the time taken for the dipoles to return to their original position after the application of the electric field. When the rate of the electric field oscillate is much faster than the relaxation time, the polarization process in a molecule is unable to follow the rate of the change of the applied oscillating electric field and this results in energy absorption and dissipation as heat [35]. The molecular and structural parameters affecting polarizability usually have similar effects on dielectric loss. Synchronous changes of dielectric constant and dielectric loss have been observed in most studies.
For thermosetting polymer systems crosslinked through free radical mechanisms, the type and concentration of initiator are the critical factors determining the dielectric properties by influencing the crosslinking density and creating low-polarity functional groups. Wu et al. [46] studied the blends of 1,2-polybutadiene/styrene-butadiene-styrene triblock copolymer/ethylene-propylene-dicyclopentadiene (1,2-PB/SBS/EPDM) blends cured with two different organic peroxides, cumyl peroxide (DCP) and bis(1-(tert-butylperoxy)-1-methylethyl)-benzene (BIPb). The results showed that BIPb is more reactive and more effective in achieving high crosslink density and converting –C=C– double bonds with relatively high polarizability into –C-C– single bonds with low polarizability, which leads to low \( D_k \) and \( D_f \). Higher initiator concentration resulted in lower \( D_k \) and \( D_f \) regardless of initiator type through similar mechanisms, i.e., higher cross-link density that more effectively hindered the movement of polar groups, and further reduced the polarizability by replacing more –C=C– double bonds with new –C-C– bonds. In addition, the effects of the by-products from the decomposition of initiators on the dielectric properties cannot be ignored [47,48]. Walker [47] investigated the effects of initiator (DCP) content on the dielectric properties of crosslinked polyethylene (XLPE). The results showed that, with higher initial DCP concentration, more byproducts were produced which resulted in an increase in dielectric loss.

Thermosetting PPO-based systems using TAIC as the crosslink agent have been widely used as the matrix of high-performance CCLs [13–16,49–51]. Usually, the PPO/TAIC ratio ranges from 1.0:1.0 to 4.0:1.0 and more typically from 1.2:1.0 to 2.0:1.0, which ensures excellent dielectric properties and good processability. A low PPO/TAIC ratio gives lower \( D_f \) [32]; however, the prepreg starts to become sticky at room temperature which causes processing issues. In this work, a series of PPO/TAIC-based composites with ultralow dielectric loss were prepared with a fixed ratio of PPO/TAIC at 1.33:1.0 and a low loss glass fabric as the reinforcement. The effects of initiator concentration on the chemical structure, thermal-mechanical and dielectric properties of a thermosetting PPO/TAIC-based composite system were studied. The dielectric constant of the precured prepreg and extensively cured laminate samples decreased with the higher concentration of the free radical initiator, whereas the dielectric loss of the samples exhibited significantly different or even opposite trends. The mechanisms were discussed based on the competition of multiple factors that affect the polarizability of thermosetting systems.

2. Materials and Methods

2.1. Materials

Methyl methacrylate terminated PPO resin (NORYL\textsuperscript{TM} SA9000) with \( \text{Mn} = 2300 \) and \( T_g = 160 ^\circ C \) was purchased from SABIC (Riyadh, Saudi Arabia). Triallyl isocyanurate (TAIC, 98%, AR) was supplied by Shanghai Aladdin Bio-Chem Technology Co., LTD (Shanghai, China). The free radical initiator, BIPb-96 (mixture of bis(tert-butylidioxyisopropyl) benzene (40%) and di(tert-butylperoxyisopropyl) benzene (60%), 96%) was purchased from Hunan Enpai New Material Company (Changsha, China). Their chemical structures are shown in Scheme 1. The solvent, methyl ethyl ketone (MEK, 98%, AR), was purchased from Shanghai Aladdin Bio-Chem Technology Co., LTD. The L-glass fabrics (Style 2116) were provided by Asahi Kasei E-materials Corporation (Tokyo, Japan). All the solvents and reagents were used without further purification.

2.2. Sample Preparation

SA9000, TAIC, and MEK were mixed with the compositions shown in Tables 1 and 2 in a 3 L beaker by stirring the mixture for 120 min in a fume hood at room temperature. Then BIPb-96 was added to the beaker and stirred for a further 60 min to obtain the varnish. The solid content of the varnish was set at 70% by weight which ensured sufficient dissolution and dispersion of components and maintained the appropriate viscosities for impregnation.
methylethyl)-benzene (BIPb). The results showed that BIPb is more reactive and more effective in achieving high crosslink density and converting –C=C– double bonds with relatively high polarizability into –C-C– single bonds with low polarizability, which leads to low $D_k$ and $D_f$. Higher initiator concentration resulted in lower $D_k$ and $D_f$ regardless of initiator type through similar mechanisms, i.e., higher cross-link density that more effectively hindered the movement of polar groups, and further reduced the polarizability by replacing more –C=C– double bonds with new –C-C– bonds. In addition, the effects of the by-products from the decomposition of initiators on the dielectric properties cannot be ignored [47,48]. Walker [47] investigated the effects of initiator (DCP) content on the dielectric properties of crosslinked polyethylene (XLPE). The results showed that, with higher initial DCP concentration, more byproducts were produced which resulted in an increase in dielectric loss.

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![Scheme 1. Chemical structure of (a) NORYLTM SA9000; (b) TAIC; (c) bis(tert-butyldioxyisopropyl) benzene; and (d) di(tert-butylperoxyisopropyl) benzene.](image)

### Table 1. The composition of casting samples.

| Sample    | SA9000:TAIC:MEK (Mass Ratio) | BIPb (wt% of mSA9000 + TAIC) |
|-----------|-------------------------------|------------------------------|
| P/T-0.25  | 4:3:3                         | 0.25                         |
| P/T-0.50  | 4:3:3                         | 0.50                         |
| P/T-0.75  | 4:3:3                         | 0.75                         |
| P/T-1.00  | 4:3:3                         | 1.00                         |
| P/T-1.25  | 4:3:3                         | 1.25                         |
| P/T-1.50  | 4:3:3                         | 1.50                         |

### Table 2. The composition of prepreg and laminate samples.

| Sample   | SA9000:TAIC:MEK (Mass Ratio) | BIPb (wt% of mSA9000 + TAIC) |
|----------|-------------------------------|------------------------------|
| P/T-0.25-L | 4:3:3                      | 0.25                         |
| P/T-0.75-L | 4:3:3                      | 0.75                         |
| P/T-1.50-L | 4:3:3                      | 1.50                         |

Pieces of glass fabrics were fully immersed in the varnish and passed through a pair of steel bars with preset gaps to make sure the glass fabrics were uniformly coated with specific amount of varnish. The coated glass fabrics were placed in a fume hood to dry for 24 h and then baked in an explosion-proof oven at 150 $\degree C$ for 7 min to obtain the prepreg samples. Extensively cured laminate and cast samples were prepared by curing prepreg or resin powders taken from the prepreg samples, respectively. In addition, a P/T-0.00-L sample (without initiator) was air-dried at room temperature. The curing procedure is shown in Figure 1.
Scheme 1. Chemical structure of (a) NORYL™ SA9000; (b) TAIC; (c) bis(tert-butyldioxyisopropyl) benzene; and (d) di(tert-butylperoxyisopropyl) benzene.

2.2. Sample Preparation

SA9000, TAIC, and MEK were mixed with the compositions shown in Table 1 and Table 2 in a 3 L beaker by stirring the mixture for 120 min in a fume hood at room temperature. Then BIPb-96 was added to the beaker and stirred for a further 60 min to obtain the varnish. The solid content of the vanish was set at 70% by weight which ensured sufficient dissolution and dispersion of components and maintained the appropriate viscosities for impregnation.

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| P/T-0.50 | 4:3:3                         | 0.50                         |
| P/T-0.75 | 4:3:3                         | 0.75                         |
| P/T-1.00 | 4:3:3                         | 1.00                         |
| P/T-1.25 | 4:3:3                         | 1.25                         |
| P/T-1.50 | 4:3:3                         | 1.50                         |

Table 2. The composition of prepreg and laminate samples.

| Sample  | SA9000:TAIC:MEK (Mass Ratio) | BIPb (wt% of mSA9000 + TAIC) |
|---------|-------------------------------|------------------------------|
| P/T-0.25-L | 4:3:3                     | 0.25                         |
| P/T-0.75-L | 4:3:3                     | 0.75                         |
| P/T-1.50-L | 4:3:3                     | 1.50                         |

Figure 1. The curing procedure.

2.3. Measurements

The solid-state nuclear magnetic resonance (SSNMR) spectra were recorded on a Bruker III 400 M spectrometer (Rheinstetten, Germany) equipped with a solid-state high-resolution apparatus. The conventional cross-polarization/magic angle spinning (CP/MAS) method was used for high-resolution solid-state \(^{13}\)C measurements, and the spinning rate was 12 kHz.

Fourier transform infrared spectra were recorded on a PerkinElmer Fourier transform infrared reflection (FTIR) spectrometer Spotlight 400 N (Shelton, FL, USA) with a scanning step of 2 cm\(^{-1}\) and accumulation of 8 scans. Both the KBr tableting method and universal attenuated total reflectance accessory (UATR) were applied. For the KBr tableting method, the sample was mixed with KBr powder in a proportion of 1:100–1:200 and the total weight was controlled to 200 mg. For quantitative studies, the weight of samples and KBr powder were precisely weighed to 0.01 mg before mixing.

High-performance liquid chromatography (HPLC) was performed on a Waters e2695 (Milford, CT, USA) series liquid chromatograph system, which was equipped with a diode array detector (DAD) controlled by an Empower chromatographic workstation. A Sunfire C18 (150 cm × 4.6 cm, 5 µm) was adopted for the analysis. The mobile phase consisted of acetonitrile (chromatographically pure, 70%) and ultrapure water (30%) using an isocratic elution. The mobile phase was set at a flow rate of 1.0 mL/min and the column temperature was conditioned at 25 °C.

The Liquid-state nuclear magnetic resonance (NMR) spectra were measured on a Bruker Avance III 400 M with BBO probe (Rheinstetten, Germany). Samples were dissolved in deuterated chloroform (CDCl\(_3\)) using tetramethylsilane (TMS) as the internal reference at room temperature.

Dynamic mechanical analysis (DMA) was performed in a tension mode using NETZSCH DMA 242E (Selb, Germany) from room temperature to 300 °C at a heating rate of 2 °C min\(^{-1}\) and a frequency of 1 Hz in air. The specimen dimensions were about 15 mm × 5 mm × 0.4 mm.
Melt viscosity (MV) was measured on a Brookfield CAP2000+H (Toronto, ON, Canada) at 171 °C and with spindle rotating speed 23 rpm. The minimum readings within 60 s were recorded as the MV.

Dielectric constant and dielectric loss were tested using an Agilent Technologies ENA series network analyzer (E5071C, Santa Clara, CA, USA) with the split-post dielectric-resonator (SPDR) at 10 GHz, 50% relative humidity, and room temperature. Before testing, each sample with thickness about 0.4 mm was dried in a 110 °C oven for 1 h and then cooled to room temperature in a desiccator for 30 min before testing.

Differential scanning calorimetry (DSC) curves were obtained with a NETZSCH DSC 214 (Selb, Germany) from room temperature to 300 °C under a nitrogen atmosphere with a heating rate of 2 °C·min⁻¹.

Free volume was obtained by positron annihilation lifetime spectrum (PALS, DPLS3000, Anhui, China). There were two identical samples of the size at 1.5 mm × 10 mm × 10 mm in a 1.1 × 10⁶ Bq ²²Na positron source. And a 1.28 MeV γ-ray was emitted by the ²²Na nucleus and the positron within a few picoseconds. The positron lifetime was determined by the time delay between the emission of the birth gamma (1.28 MeV) and an annihilation photon of 0.511 MeV. PALS were obtained using the fast–fast coincidence system. The time resolution was 220 ps and the channel width was 12.6 ps. Program PATFIT was used to process the data and get the lifetime spectra. The positron source components (392 ps/16.40%, 2.05 ns/0.70%) were subtracted before analyzing each spectrum. The variance of the fits was about 1.1.

3. Results and Discussion
3.1. Chemical Structure of Cured Matrix

The SSNMR spectra of cured samples were shown in Figure 2. The peak positions were consistent for the two samples, indicating the same curing mechanisms and no changes in chemical structure were induced by the varied concentration of initiators. The signals at 126.0 ppm and 136.5 ppm coming from –C=C– of SA9000 [52] were not detected, showing that all the –C=C– of SA9000 had participated in the reaction.

Figure 2. The SSNMR spectra of cured samples.
The FTIR spectra of cured P/T-0.25 and P/T-1.50 together with that of SA9000 and TAIC were shown in Figure 3. The FTIR spectra of P/T-0.25 and P/T-1.50 were identical in terms of the number and position of the peaks, which agreed with SSNMR results that there are no differences in chemical structure caused by initiator concentrations.

The existence of the absorption peaks at 1645 and 3086 cm\(^{-1}\) suggested that some –C=\(-\)C– groups from TAIC did not participate in the crosslinking reaction [52]. To quantify the amount of unreacted –C=\(-\)C– groups from TAIC, the peak at 3086 cm\(^{-1}\) was employed for the calculation using Lambert–Beer law [53] with the following steps:

The powder from P/T-0.00-L was weighed out in five different amounts and thoroughly mixed with the appropriate amount of KBr powder to keep the total mass of the mixture at ~200 ± 0.5 mg. The mixture was pressed into a disc for the FTIR scanning. The obtained spectra were shown in Figure 4 and the calculated mole numbers of –C=\(-\)C–(n–C=C–) in TAIC in the samples were listed in Table 3. The peak areas were plotted against the n–C=C– in Figure 5 and a linear regression equation y = 74.79x was obtained with the R\(^2\) = 0.975. By calculating the peak areas obtained from the FTIR spectra of cured P/T-0.25 and P/T-1.50 (Figure 3b), the residual ratio of the –C=C– of TAIC in these two samples was 23.33% and 17.5%, respectively, revealing that more –C=C– in TAIC participated in the reaction as the mass fraction of the initiator increased.

### Table 3

| mTAIC (mg) | mKBr (mg) | n TAIC (mmol) | n –C=C– (mmol) | A (Peak area) |
|------------|-----------|---------------|----------------|-------------|
| 0.05       | 0.30      | 0.025         | 0.0125         | 0.9          |
| 0.10       | 0.30      | 0.050         | 0.0250         | 1.8          |
| 0.15       | 0.30      | 0.075         | 0.0375         | 2.8          |
| 0.20       | 0.30      | 0.100         | 0.0500         | 3.8          |
| 0.25       | 0.30      | 0.125         | 0.0625         | 4.8          |

### Figure 3.

(a) The FTIR spectra of SA9000, TAIC, cured P/T-0.25, and P/T-1.50; (b) locally enlarged FTIR spectra of cured P/T-0.25 and P/T-1.50.

### Figure 4.

FTIR spectra of P/T-0.00-L samples of different amounts.
Table 3. Calculation of fitting data of different amounts of P/T-0.00-L sample powders.

| P/T-0.00-L Powder (mg) | m_{TAIC} (mg) | n_{TAIC} (mmol) | n_{C=C-} (mmol) | A_{Peak area-3086} |
|------------------------|---------------|-----------------|-----------------|-------------------|
| 0                      | 0             | 0               | 0               | 0                 |
| 1.56                   | 0.6686        | 0.0027          | 0.0081          | 0.48              |
| 2.87                   | 1.2300        | 0.0049          | 0.0148          | 0.83              |
| 4.52                   | 1.9371        | 0.0078          | 0.0233          | 1.78              |
| 5.99                   | 2.5671        | 0.0103          | 0.0309          | 1.88              |
| 8.60                   | 3.6857        | 0.0148          | 0.0444          | 3.72              |

Figure 5. The calibration curve for \(-C=C-\) absorbance of TAIC at 3086 cm\(^{-1}\).

3.2. Unreacted TAIC in Laminates

Matsumoto et al. [54] studied the curing reaction of TAIC and pointed out that the unsaturated bonds of TAIC are difficult to be fully consumed because TAIC chains have bulkier isocyanurate side groups than the chains derived from the polymerizations of other multi-allyl compounds, and rigid TAIC chains lead to a reduced interaction between polar isocyanuric rings belonging to both the polymer chain and TAIC monomer. These dissociated \(-C=C-\) bonds either came from the partially reacted TAIC molecules or from the unreacted free TAIC molecules in the system (Scheme 2). To identify and quantify the soluble components of the cured samples, a certain amount of cured P/T-0.25 and P/T-1.50 samples were crushed and extracted for 12 h in boiled MEK and the extracts were studied by HPLC, \(^1\)H-NMR, and \(^{13}\)C-NMR.
other multi-allyl compounds, and rigid TAIC chains lead to a reduced interaction between polar isocyanuric rings belonging to both the polymer chain and TAIC monomer. These dissociated –C=C– bonds either came from the partially reacted TAIC molecules or from the unreacted free TAIC molecules in the system (Scheme 2). To identify and quantify the soluble components of the cured samples, a certain amount of cured P/T-0.25 and P/T-1.50 samples were crushed and extracted for 12 h in boiled MEK and the extracts were studied by HPLC, 1H-NMR, and 13C-NMR.

Scheme 2. Cross-linked network in the sample.

The standard curve for quantitative HPLC analysis was established by measuring several TAIC/MEK solutions with a given TAIC concentration, as shown in Figure 6a,b. Based on the linear regression equation obtained and the chromatograms of the extracts (Figure 6c), it was found that there was 276.72 mg of TAIC in the extract of P/T-0.25, which meant that 14.74% of the TAIC did not take part in the reaction and it occupied 6.24% of the total mass of the sample. Furthermore, in P/T-1.50, these values were 7.49 mg, 0.48%, and 0.20% (Table 4).

Similar phenomena were observed with NMR measurements. 1H-NMR and 13C-NMR spectra of the extracts were shown in Figure 7a,c. When Figure 7a,c were enlarged in equal proportions (Figure 7b,d), similar phenomena as the HPLC could be obtained. There was 286.29 mg of TAIC in the extract of P/T-0.25, but not in P/T-1.50. It was found that 15.26% of the TAIC in P/T-0.25 did not participate in the reaction, which was 6.46% of the total mass of the sample (Table 3).
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Figure 6. (a) Chromatograms of extracts from P/T-0.25 and P/T-1.50; (b) HPLC chromatograms of TAIC/MEK solution with different TAIC mass fractions; (c) TAIC calibration curve. The abscissa was the detected TAIC quality, and the ordinate was the area of TAIC peak.

Figure 7. (a,b) 1H-NMR spectra of different samples’ extractions; (c,d) 13C-NMR spectra of different samples’ extractions.

Table 4. Quantitative analysis results of different test methods.

| Sample   | Weight (mg) | HPLC (mg) | wt% (in TAIC) | wt% (in Sample) | 1H-NMR (mg) | wt% (in TAIC) | wt% (in Sample) |
|----------|-------------|-----------|---------------|-----------------|-------------|---------------|-----------------|
| P/T-0.25 | 4433.0      | 276.72    | 14.74%        | 6.24%           | 286.29      | 15.26%        | 6.46%           |
| P/T-1.50 | 3690.4      | 7.49      | 0.48%         | 0.20%           | 0           | 0             | 0               |
3.3. Thermal-Mechanical Properties and Free Volume

Dynamic mechanical analysis (DMA) was used to characterize the $T_g$ and the crosslink density ($v_e$) of the samples. The storage modulus ($E'$) curves and the tan$\delta$ curves as a function of temperature for samples with different initiator mass fractions were shown in Figure 8a,b. The peak temperature of the tan$\delta$ was denoted as $T_g$. The storage modulus ($E'$) of DMA in the rubber plateau area (40 °C above $T_g$) was used to calculate the $v_e$ based on the following formula [55–57]:

$$v_e = \frac{E'}{3RT}$$

where $E'$ was the storage modulus of the thermoset in the rubbery plateau region at $T_g + 4$ °C, R was the gas constant, and T was the absolute temperature at $T_g + 40$ °C.

The $T_g$ and $v_e$ of samples were shown in Table 5. As expected, with more initiator added into the system, the crosslinking reaction advanced more extensively, which resulted in a higher $T_g$ and higher crosslink density.

### Table 5. The thermal properties of different initiator mass fraction samples.

| BIPb% | $T_g$-tan$\delta$/DMA (°C) | $E'/T_g + 40$ °C (MPa) | $v_e$(mol/m$^3$) |
|-------|-----------------------------|------------------------|------------------|
| 0.25  | 172.0                       | 9.0                    | 814.53           |
| 0.50  | 200.5                       | 24.7                   | 2076.97          |
| 0.75  | 213.0                       | 37.7                   | 3097.26          |
| 1.00  | 229.9                       | 74.7                   | 5937.05          |
| 1.25  | 231.6                       | 77.3                   | 6134.94          |
| 1.50  | 236.0                       | 100.0                  | 7876.82          |

PALS was used to measure the free volume of different samples in the system. As shown in Figure 9 and Table 6, the free volume became larger with the increased initiator content.

3.4. Dielectric Properties

As shown in Figure 10a, at 10 GHz with a resin contents of about 52% the cured PPO/TAIC/L-glass composites exhibited dielectric constants from 3.24 to 3.10 and ultralow dissipation factors from 0.0023 to 0.0031, respectively. Before curing, the dielectric constants of prepreg samples decreased significantly with higher initiator concentration, while the dielectric loss stayed almost the same. After curing, the dielectric constant of the laminates kept the same trend as that of prepreg but moved to
higher levels compared to the corresponding prepreg samples. The dielectric loss, however, showed an ascending trend at the same time. This trend was further confirmed with the resin cast samples (Figure 10b), indicating that the variation of the dielectric properties of the composites was mainly caused by the intrinsic variation of the resin matrix.

Figure 8. (a) The storage modulus ($E'$) curves of different initiator mass fractions samples; (b) the tan $\delta$ curves of different initiator mass fractions samples.

Table 5. The thermal properties of different initiator mass fraction samples.

| BIPb% | $T_g$-tan $\delta$/DMA (°C) | $E'/T_g + 40$ °C (MPa) | $\nu e$ (mol/m$^3$) |
|-------|----------------------------|-------------------------|---------------------|
| 0.25  | 172.0                      | 9.0                     | 814.53              |
| 0.50  | 200.5                      | 24.7                    | 2076.97             |
| 0.75  | 213.0                      | 37.7                    | 3097.26             |
| 1.00  | 229.9                      | 74.7                    | 5937.05             |
| 1.25  | 231.6                      | 77.3                    | 6134.94             |
| 1.50  | 236.0                      | 100.0                   | 7876.82             |

Figure 9. Positron lifetime spectra measured for cured samples.

Table 6. Summary of the PALS analysis results of different samples.

| BIPb% | $\tau_3$ (ns) | $r_3$ (Å) | $V_3$ (Å$^3$) | $I_3$ (%) | FFV (%) |
|-------|---------------|-----------|---------------|-----------|---------|
| 0.25  | 2.19          | 3.02      | 115.03        | 40.1      | 6.9     |
| 0.50  | 2.29          | 3.10      | 125.03        | 38.5      | 7.2     |
| 0.75  | 2.33          | 3.14      | 129.06        | 38.4      | 7.4     |
| 1.00  | 2.44          | 3.22      | 140.62        | 36.8      | 7.6     |
| 1.25  | 2.51          | 3.28      | 147.27        | 33.0      | 8.0     |
| 1.50  | 2.57          | 3.32      | 153.42        | 31.5      | 8.5     |

For thermosetting polymer systems, it had been commonly agreed that, with the increase in crosslinking density, the orientation of the groups is progressively hindered and usually results in a lower dielectric constant and dielectric loss [43–45]. In the current PPO/TAIC system, however, it was obvious that more polarization and relaxation mechanisms were involved, and the significance of each mechanism varied depending on the advancement of curing, environment of the functional groups, and crosslinking structures.
Thus, the plasticization effect was weakened, which explained the decrease in the dielectric constant of different samples.

Table 7. Summary of the PALS analysis results of different samples.

| Prepreg Sample | Melt Viscosity (cP) |
|----------------|---------------------|
| P/T-0.25-L     | 10,739              |
| P/T-0.75-L     | 20,884              |
| P/T-1.50-L     | 23,454              |

When the samples were cured at elevated temperatures, several changes took place in the system. First, with the advancement of the curing reaction, more and more TAIC molecules reacted with SA9000 and either generated crosslink networks by bridging SA9000 molecules or formed bulky pendant groups by grafting onto SA9000 chains. Both factors caused decreased dipolar mobility, contributed to lower polarizability, and drove the dielectric constants to a lower level. At the same time, the lessening of the plasticizing effects of TAIC were also in favor of the reduction of the dielectric constant. On the other hand, however, there were several new factors generated during the curing which may have promoted an increase in dielectric constants. PPO and TAIC molecules are highly miscible at this composition [58]. It has been revealed that plasticizers in polymers can reduce intermolecular interaction, enhance the segmental mobility of polymer chains, and promote the dipole moments, thus increasing the dielectric constant [59,60]. Such mechanisms may play important roles in the current system.

While baked at 150 °C, some polymerization reactions were initiated which resulted in an increase in melt viscosity (Table 7). The DSC scan of P/T-0.25-L prepreg in Figure 11 showed that the curing reaction started at about 120 °C and had dual exothermic peaks at 160 °C and 180 °C, respectively, whereas that of pure TAIC started at 100 °C and reached an exothermic peak at 155 °C. The results revealed that multiple reaction processes are involved in the curing of PPO/TAIC systems. It is most likely that the peak at 160 °C can be mainly attributed to the curing of pure TAIC, and the reaction between PPO and TAIC occurred later at higher temperature. FTIR spectra in Figure 12 and the calculated results in Table 8 confirmed that, with the increase in initiator content in original compositions, more TAIC participated in homopolymerization and less plasticizer was left in the matrix. Thus, the plasticization effect was weakened, which explained the decrease in the dielectric constant of prepreg samples with higher initiator concentration. On the other hand, however, the dielectric loss at the prepreg stage did not change much with the variation of initiator content and the level of TAIC polymerization, which can be explained by the fact that the TAIC monomers have a highly symmetric structure and poly(TAIC) has a very low dielectric loss [32].

Table 7. Melt viscosity of different samples.

| Prepreg Sample | Melt Viscosity (cP) |
|----------------|---------------------|
| P/T-0.25-L     | 10,739              |
| P/T-0.75-L     | 20,884              |
| P/T-1.50-L     | 23,454              |

Figure 10. Dielectric properties of different initiator mass fractions: (a) laminate samples (the resin content was 52 ± 3%); (b) casting samples.
symmetrical, and the polarity was very low. When PPO and TAIC were copolymerized, the structural symmetry of both components was diminished and the dipole movements were enhanced, which resulted in significantly higher molecular and group polarity. Moreover, short TAIC branches can weaken the interaction force between polymer chains and increase the movability of polymer chains. Furthermore, chain transfer products from free radicals may also contribute to the overall polarizability of a system as polar impurities. For example, the primary radicals from the initiator may have been terminated by taking atoms from the other reactant and producing hydroxyl groups. Although the presence of alcoholic hydroxyl groups was not detected by ATR-FTIR or SSNMR [52], such byproducts from chain transfer reactions were inevitable and contribute to an increase in dielectric properties.

![DSC curves of P/T-0.25-L and TAIC homopolymerization system (the mass ratio of TAIC: BIPb was 99:1).](image1)

**Figure 11.** DSC curves of P/T-0.25-L and TAIC homopolymerization system (the mass ratio of TAIC: BIPb was 99:1).

![FTIR spectra of different prepreg samples.](image2)

**Figure 12.** FTIR spectra of different prepreg samples.
Table 8. Residual rate of –C=C– in TAIC of different prepreg samples.

| Prepreg Sample | Residual Rate of –C=C– in TAIC |
|----------------|--------------------------------|
| P/T-0.25-L     | 56.06%                         |
| P/T-0.75-L     | 52.62%                         |
| P/T-1.50-L     | 45.94%                         |

Therefore, the overall trends of dielectric constant and dielectric loss were the superposition of these competing factors. It was evident that the factors increasing polarizability prevailed after curing compared to that of the corresponding prepreg samples, which resulted in an increase in dielectric constant and loss. Moreover, the polarizabilities were enhanced with the higher concentration of the initiator which led to the significant ascending of dielectric loss, whereas increases in the dielectric constant were partially offset by the increased free volume and reduced plasticizing effect at higher initiator concentration (Figure 9 and Table 6).

4. Conclusions

The PPO/TAIC-based polymer composites exhibited excellent dielectric properties which are suitable for high-frequency applications. In this work, the composition–process–structure–property relationship of PPO/TAIC-based polymer composites with ultralow dielectric loss was studied. Special attention was focused on the effects of peroxide initiator concentration upon the dielectric properties. For prepreg samples, the plasticizing effect of TAIC played a key role which caused a decrease in the dielectric constant with the initiator content, while it did not affect dielectric loss much. When the samples were cured at high temperatures, more mechanisms affecting polarizability developed with the advancement of the reaction, including some factors that contributed to lower polarizability, such as increased crosslink density, free volume and quantities of bulky pendant groups, and reduced plasticizing effect, etc., as well as some factors that promoted polarizability such as enhanced polarity, diminished molecular symmetry, branching, and hydroxyl impurities, etc. As a result of competition, the latter mechanisms dominated the trends and both the dielectric constant and dielectric loss of the cured samples increased significantly with higher initiator concentration compared to the corresponding prepreg samples.

Author Contributions: Conceptualization, K.W., Z.L., Q.L. and X.Y.; methodology, X.Y.; validation, K.W. and Z.L.; formal analysis, X.Y., Z.F., D.L., Y.M. and C.L.; investigation, X.Y. and Z.F.; data curation, X.Y.; writing—original draft preparation, X.Y.; writing—review and editing, K.W.; visualization, X.Y.; supervision, K.W. and Z.L.; project administration, K.W. All authors have read and agreed to the published version of the manuscript.

Funding: This research was funded by the Department of Science and Technology of Guangdong Province (Project No.2020B010175001).

Institutional Review Board Statement: Not applicable.

Informed Consent Statement: Not applicable.

Data Availability Statement: The data presented in this study are available on request from the corresponding author.

Acknowledgments: The authors sincerely acknowledge the Department of Science and Technology of Guangdong Province (Project No. 2020B010175001) for the financial support. The authors would like to thank Yi Zhang from Sun Yat-Sen University for assistance with positron annihilation lifetime spectroscopy measurement.

Conflicts of Interest: The authors declare no conflict of interest.
References

1. Andrews, J.G.; Buzzi, S.; Choi, W.; Hanly, S.V.; Lozano, A.; Soong, A.C.K.; Zhang, J.C. What Will 5G Be? IEEE J. Sel. Areas Commun. 2014, 32, 1065–1082. [CrossRef]
2. Alsulami, M.M.; Akkari, N. The Role of 5G Wireless Networks in the Internet-of-Things (IoT). In Proceedings of the 2018 1st International Conference on Computer Applications Information Security (ICCAIS), Riyadh, Saudi Arabia, 4–6 April 2018; pp. 1–8. [CrossRef]
3. Tasaki, T.; Shiotsai, A.; Tsuji, M.; Nakamura, T.; Yamaguchi, T. The Low Dk / Df Adhesives for High Frequency Printed Circuit Board Using the Novel Solvent Soluble Polyimide. In Proceedings of the 2015 10th International Microsystems, Packaging, Assembly and Circuits Technology Conference (IMPACT), Taipei, Taiwan, 21–23 October 2015; pp. 221–224. [CrossRef]
4. Tseng, F.P.; Chuang, K.Y.; Liao, I.S.; Chiou, K.C. High Frequency Substrate Materials with Highly Thermal Resistance and Low Dielectric Properties. In Proceedings of the 2013 8th International Microsystems, Packaging, Assembly and Circuits Technology Conference (IMPACT), Taipei, Taiwan, 22–25 October 2013; pp. 109–112. [CrossRef]
5. Xu, M.; Chen, S.; Li, X.; Zhu, S.; Ren, D.; Liu, X. Fabrication of Phthalonitrile-Based Copper-Clad Laminates and Their Application Properties: Thermo-Stability and Dielectric Properties. Adv. Ind. Eng. Polym. Res. 2020, 3, 194–201. [CrossRef]
6. Maier, G. Low dielectric constant polymers for microelectronics. Prog. Polym. Sci. 2001, 26, 3–65. [CrossRef]
7. Zhang, Z.; Pei, J.; Liang, G.; Yuan, L. Methyl silsesquioxane/cyanate ester resin organic–inorganic hybrids with low dielectric constant. J. Appl. Polym. Sci. 2011, 121, 1004–1012. [CrossRef]
8. Chao, H.S.I.; Whalen, J.M. Poly(2,6-Dimethyl-1,4-Phenylene Ether) (PPO®) Capping and Its Significance in the Preparation of PPO®/Epoxy Laminates. J. Appl. Polym. Sci. 1996, 59, 473–481. [CrossRef]
9. Liang, G.; Meng, J.; Zhao, L. Synthesis of Styrene–Maleic Anhydride Random Copolymer and its Compatibilization to Poly(2,6-dimethyl-1,4-phenylene ether)/Brominated Epoxy Resin. Polym. Int. 2003, 52, 966–972. [CrossRef]
10. Seike, Y.; Okude, Y.; Iwakura, I.; Chiba, I.; Ikeno, T.; Yamada, T. Synthesis of Polyphenylene Ether Derivatives: Estimation of Their Dielectric Constants. Macromol. Chem. Phys. 2003, 204, 1876–1881. [CrossRef]
11. Krijgsman, J.; Feijen, J.; Gaymans, R.J. Segmented Copolymers of Uniform Tetra-Amide Units and Poly(Phenylene Oxide): 1. Synthesis and Thermal-Mechanical Properties. Polymer 2004, 45, 4677–4684. [CrossRef]
12. Su, M.S. Composite Material, High-Frequency Circuit Substrate Made Therefrom and Making Method Thereof. U.S. Patent 9890276B2, 13 February 2018.
13. Almen, G.R.; Wang, K.; Yuan, Q. Thermosetting Resin Composition Containing a Polyphenylene Ether and a Brominated Fire Retardant Compound. U.S. Patent 9243164B1, 26 January 2016.
14. Fung, D.R.; Liao, T.C.; Chen, H.S.; Chen, C.L. High-Frequency Copper Foil Covered Substrated and Compound Material Used Therein. U.S. Patent 9243132B2, 26 January 2016.
15. Almen, G.R. Thermosetting Resin Composition Containing a Polyphenylene Ether and a Brominated Fire Retardant Compound. U.S. Patent 9054651B1, 9 June 2015.
16. Tokiwa, T.; Utsumi, T.; Endo, M. Curable Resin Composition. U.S. Patent 8703277B2, 22 April 2014.
17. Yokoyama, D.; Saito, H.; Kita, Y.; Fujitwara, H.; Inagaki, K. Developing Thermosetting Resin System with Low Molecular Weight PPE to Achieve Lower Dielectric Property. J. Neto. Polym. 2012, 33, 336–340. [CrossRef]
18. Ishib, Y.; Kuroki, M.; Oda, H.; Arai, T.; Katayose, T. Novel Thermosetting Poly (Phenylene Ether). Synthesis, Blend, and Application to Copper Clad Laminates. Kobunshi Ronbunshu 1997, 54, 171–182. [CrossRef]
19. Chen, C.H.; Gu, Z.C.; Tai, Y.L.; Jeng, R.J.; Lin, C.H. Identification of the Reaction Mechanism between Phenyl Methacrylate and Epoxy and Its Application in Preparing Low-Dielectric Epoxy Thermosets with Flexibility. Polymer 2018, 140, 225–232. [CrossRef]
20. Chen, C.H.; Jhang, T.Y.; Abu-Omar, M.M.; Lin, C.H. Structure-Property Relationship of Vinyl-Terminated Oligo(2,6-Dimethyl-1,4-Phenylene Ether)s (OPEs): Seeking an OPE with Better Properties. Eur. Polym. J. 2019, 117, 94–104. [CrossRef]
21. Chen, C.H.; Lee, K.W.; Lin, C.H.; Jhang, T.Y. Low-Dissipation Thermosets Derived from Oligo(2,6-Dimethyl Phenylene Oxide)-Containing Benzoxazines. Polymers 2018, 10, 411. [CrossRef] [PubMed]
22. Weng, L.; Zhang, Y.; Zhang, X.; Liu, L.; Zhang, H. Synthesis and Properties of Cyanate Mixed Resin Systems Modified by Polyphenylene Oxide for Production of High-Frequency Copper Clad Laminates. J. Mater. Sci. Mater. Electron. 2018, 29, 2831–2840. [CrossRef]
23. Yang, M.C.; Higashihiara, T.; Su, H.W.; Ueda, M.; Chen, W.C. Crosslinked Copolymer with Low Dielectric Constant and Dissipation Factor Based on Poly(2,6-Dimethylphenol-Co-2,6-Diphenylphenol) and a Crosslinker. J. Polym. Sci. Part A Polym. Chem. 2016, 54, 3218–3223. [CrossRef]
24. Mo, J.; Xia, L.; Pan, P.; Shentu, B.; Weng, Z. Reactive Blend of Epoxy-Novolac Resin and Epoxide-Terminated Low-Molecular-Weight Poly(Phenylene Oxide). J. Appl. Polym. Sci. 2013, 127, 4879–4888. [CrossRef]
25. Gao, R.; Gu, A.; Liang, G.; Dai, S.; Yuan, L. Properties and Origins of High-Performance Poly(Phenylene Oxide)/Cyanate Ester Resins for High-Frequency Copper-Clad Laminates. J. Appl. Polym. Sci. 2011, 121, 1675–1684. [CrossRef]
26. Hwang, H.J.; Hsu, S.W.; Wang, C.S. Low Dielectric and Flame-Retardant Properties of Thermosetting Redistributed Poly(Phenylene Oxide). J. Vinyl Addit. Technol. 2009, 15, 54–59. [CrossRef]
27. Hwang, H.J.; Hsu, S.W.; Wang, C.S. Low Dielectric Thermoset from Redistributed Poly(Phenylene Oxide). J. Macromol. Sci. Part A 2008, 45, 1047–1054. [CrossRef]
28. Hwang, H.J.; Hsu, S.W.; Chung, C.L.; Wang, C.S. Low Dielectric Epoxy Resins from Dicyclopentadiene-Containing Poly(Phenylene Oxide) Novolac Cured with Dicyclopentadiene Containing Epoxy. React. Funct. Polym. 2008, 68, 1185–1193. [CrossRef]

29. Nunoshige, J.; Akahoshi, H.; Liao, Y.; Horiuchi, S.; Shibasaki, Y.; Ueda, M. Mechanical and Dielectric Properties of a New Polymer Blend Composed of 1,2-Bis(Vinylphenyl)Ethane and Thermosetting Poly(Phenylene Ether) Copolymer Obtained from 2,6-Dimethylphenol and 2-Allyl-6-Methylphenol. Polym. J. 2007, 39, 828–833. [CrossRef]

30. Peters, E.N. Polyphenylene Ether Macromonomers. I. Property Enhancements in Thermoset Resins via Novel Telechelic Oligomers. In Proceedings of the Annual Technical Conference Proceedings (ANTEC 2007), Cincinnati, OH, USA, 6–11 May 2007; pp. 2125–2128.

31. Peters, E.N.; Fisher, S.M.; Guo, H.; Howe, R.; Degenzague, C. Polyphenylene Ether Macromolecules. VII. Performance in t-Butyl Styrene/Divinyl Benzene Resin System. In Proceedings of the Annual Technical Conference Proceedings (ANTEC 2010), Orlando, FL, USA, 16–20 May 2010; pp. 1858–1861.

32. Peters, E.N.; Fisher, S.M.; Guo, H. PPE Macromonomers. IX. Performance Enhancements of Triallyl Isocyanurate Resins. In Proceedings of the Annual Technical Conference Proceedings (ANTEC 2011), Boston, MA, USA, 1–5 May 2011; Volume 3, pp. 2823–2826.

33. Peters, E.N.; Jeevanath, M.; Fisher, S.M.; Guo, H. Polyphenylene Ether Macromonomer: X. Vinyl Terminated Telechelic Macromers. In Proceedings of the Annual Technical Conference Proceedings (ANTEC 2011), Boston, MA, USA, 1–5 May 2011; pp. 2819–2822.

34. Shieh, J.Y.; Yang, S.P.; Wu, M.F.; Wang, C.S. Synthesis and characterization of novel low-dielectric cyanate esters. J. Polym. Sci. Part A Polym. Chem. 2004, 42, 2589–2600. [CrossRef]

35. Ahmad, Z. Polymer Dielectric Materials; IntechOpen: London, UK, 2012. [CrossRef]

36. Zhu, L.; Wang, Q. Novel Ferroelectric Polymers for High Energy Density and Low Loss Dielectrics. Macromolecules 2012, 45, 2937–2954. [CrossRef]

37. Zhou, D.L.; Li, J.H.; Guo, Q.Y.; Lin, X.; Zhang, Q.; Chen, C.; Han, D.; Fu, Q. Polyhedral Oligomeric Silsesquioxanes Based Ultralow-k Materials: The Effect of Cage Size. Adv. Funct. Mater. 2021, 31, 2102074. [CrossRef]

38. Luo, Y.; Jin, K.; He, C.; Wang, J.; Sun, J.; He, F.; Zhou, J.; Wang, Y.; Fang, Q. An Intrinsically Microporous Network Polymer with Good Dielectric Properties at High Frequency. Macromolecules 2016, 49, 7314–7321. [CrossRef]

39. Dlubek, G.; Saarinen, K.; Fretwell, H.M. The Temperature Dependence of the Local Free Volume in Polyethylene and Polytetrafluoroethylene: A Postion Lifetime Study. J. Polym. Sci. Part B Polym. Phys. 1998, 36, 1513–1528. [CrossRef]

40. Yu, J.; Huo, R.; Chao, W.; Wu, X.; Wang, G.; Jiang, P. Influence of Interface Structure on Dielectric Properties of Epoxy/Alumina Nanocomposites. Macromol. Res. 2012, 20, 816–826. [CrossRef]

41. Siddabattuni, S.; Schuman, T.P.; Dogan, F. Improved Polymer Nanocomposite Dielectric Breakdown Performance through Barium Titanate to Epoxy Interface Control. Mater. Sci. Eng. B 2011, 176, 1422–1429. [CrossRef]

42. Huang, X.; Iizuka, T.; Jiang, P.; Ohki, Y.; Tanaka, T. Role of Interface on the Thermal Conductivity of Highly Filled Dielectric Epoxy/AIN Composites. J. Phys. Chem. C 2012, 116, 13629–13639. [CrossRef]

43. Guo, H.; Zheng, J.; Gan, J.; Liang, L.; Wu, K.; Lu, M. Relationship between Crosslinking Structure and Low Dielectric Constant of Hydrophobic Epoxies Based on Substituted Biphenyl Mesogenic Units. RSC Adv. 2015, 5, 88014–88020. [CrossRef]

44. Yang, D.; Tian, M.; Kang, H.; Dong, Y.; Liu, H.; Yu, Y.; Zhang, L. New Polyester Dielectric Elastomer with Large Actuated Strain at Low Electric Field. Mater. Lett. 2012, 76, 229–232. [CrossRef]

45. Bakule, R.; Havranek, A. The Dependence of Dielectric Properties on Crosslink Density of Rubbers. Rubber Chem. Technol. 1978, 51, 72–80. [CrossRef]

46. Wu, B.; Mao, X.; Wang, C.; Deng, T.; Li, R.; Xu, Y.; Tang, X. Different Organic Peroxides that Cure Low-k 1, 2-PB/SBS/EPDM Composites for High-Frequency Substrate. J. Vinyl Addit. Technol. 2020, 26, 524–535. [CrossRef]

47. Walker, R.C.; Hamedi, H.; Woodward, W.H.H.; Rajagopalan, R.; Lanagan, M. Impacts of Crosslinking and Degassing on the Conductivity, Dielectric Loss, and Morphology of Low-Density Polyethylene and Crosslinked Polyethylene. In Broadband Dielectric Spectroscopy: A Modern Analytical Technique; American Chemical Society: Washington, DC, USA, 2021; pp. 239–260. [CrossRef]

48. El Harfi, J.; Kingman, S.W.; Dimitrakis, G.; Robinson, J.P.; Irvine, D.J. Dielectric Properties of Free Radical Initiators Investigation of Thermal Decomposition Products. Ind. Eng. Chem. Res. 2012, 51, 15811–15820. [CrossRef]

49. Yu, X.; Fang, Z.; Qin, Y.; Lu, D.; Liu, Q.; Luo, L.; Lin, Z.; Wang, K. FTIR and NMR Characterization of Thermosetting Methyl Methacrylate Terminated Poly(2,6-Dimethyl-1,4-Phenylene Oxide)—Triallyl Isocyanurate Copolymer. J. Polym. Res. 2021, 28, 272. [CrossRef]

50. Yuki, K.; Hiroaki, F.; Hirosuke, S. Thermosetting Resin Composition, Prepreg, Metal Clad Laminate Plate, and Printed Wiring Board. U.S. Patent 9708468B2, 18 July 2017.

51. Lin, H.; Wang, Z.; Zhong, D. Polyphenylene Ether Resin Composition. U.S. Patent 10167389B2, 1 January 2019.

52. Thomas, A.K. Circuit Materials and Articles Formed Therefrom. U.S. Patent 9809690B2, 7 November 2017.

53. Reig, F.B.; Adelantado, J.G.; Moreno, M.M. FTIR quantitative analysis of calcium carbonate (calcite) and silica (quartz) mixtures using the constant ratio method. Application to geological samples. Talanta 2002, 58, 811–821. [CrossRef]

54. Matsumoto, A.; Hirai, F.; Sumiyama, Y.; Aota, H.; Takayama, Y.; Kameyama, A.; Nakaniishi, T. Further Discussion of Steric Effect on the Radical Polymerization of Triallyl Isocyanurate as Compared with Its Isomer Triallyl Cyanurate: Polymerization and Copolymerization of Corresponding Trimethylallyl Compounds. Eur. Polym. J. 1999, 35, 195–199. [CrossRef]
55. Hill, L.W. Calculation of Crosslink Density in Short Chain Networks. *Prog. Org. Coat.* 1997, 31, 235–243. [CrossRef]
56. Pan, X.; Webster, D.C. Impact of Structure and Functionality of Core Polyol in Highly Functional Biobased Epoxy Resins. *Macromol. Rapid Commun.* 2011, 32, 1324–1330. [CrossRef]
57. Chattopadhyay, D.K.; Panda, S.S.; Raju, K.V.S.N. Thermal and Mechanical Properties of Epoxy Acrylate/Methacrylates UV Cured Coatings. *Prog. Org. Coat.* 2005, 54, 10–19. [CrossRef]
58. Rusli, A.; Cook, W.; Liang, G.G. Allylic Monomers as Reactive Plasticizers of Polyphenylene Oxide. Part I: Uncured Systems. *Eur. Polym. J.* 2011, 47, 1775–1784. [CrossRef]
59. Abaci, U.; Guney, H.Y.; Yilmazoglu, M. Plasticizer Effect on Dielectric Properties of Poly(Methyl Methacrylate)/Titanium Dioxide Composites. *Polym. Polym. Compos.* 2021, 29, 565–574. [CrossRef]
60. Shelly, M.; Mathew, M.; Pradyumn, P.P.; Francis, T. Dielectric and Thermal Stability Studies on High Density Polyethylene–Chitosan Composites Plasticized with Palm Oil. *Mater. Today Proc.* 2021, 46, 2742–2746. [CrossRef]