

Article

Cerium Salts: An Efficient Curing Catalyst for Benzoxazine Based Coatings

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Abstract: The effect of three different cerium salts ($\text{Ce}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$, $\text{CeCl}_3 \cdot 7\text{H}_2\text{O}$ and $\text{Ce}(\text{OOCCH}_3)_3 \cdot 5\text{H}_2\text{O}$) on the ring-opening polymerization (ROP) of a model diamine-based benzoxazine (4EP-pPDA) was investigated. With the incorporation of the cerium salts, the curing temperature of 4EP-pPDA is reduced substantially, and the glass transition temperatures of the resulting networks are increased significantly. The three cerium salts exhibit different catalytic activities, which were analyzed by FT-IR, NMR, and energy-dispersive X-ray (EDX). $\text{Ce}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$ was found to exhibit the best catalytic effect, which seems to be related to its better dispersibility within 4EP-pPDA benzoxazine precursors.

Keywords: benzoxazine; cerium salt; ROP; catalyst; low curing temperature

1. Introduction

Benzoxazine resins are relatively new incomers in the field of thermosetting materials [1,2] and they are obtained from the ring-opening polymerization of 1,3-benzoxazine precursors [3]. The latter can be straightforwardly synthesized in high yield (>90wt%) by a Mannich-like condensation of an amine, a phenol, and formaldehyde, either in solution or in bulk [4–6]. Moreover, the diverse and large number of commercially available phenolic derivatives and amine derivatives allows the preparation of a very wide range of monomers including the incorporation of additional substituents which can bring new functionalities to materials such as self-healing [7], self-cleaning [8], photo-sensing [9] properties, etc. Benzoxazine resins have attracted substantial attention these last two decades in the first place, it is mainly because they exhibit a combination of excellent properties [10,11] such as a high glass transition temperature, near-zero volumetric change upon curing, low water absorption, good electrical resistance, low surface free energy, and high char yield. Many endeavors have been made to explore the possible applications in the automotive, aerospace, and construction industries in the form of matrix resin or surface coating [8,12,13]. However, for most benzoxazine precursors, their thermal ring-opening polymerization usually occurs at a high curing temperature (i.e., over 180 °C) not compatible with several manufacturing processes of materials. This drawback could limit the use of benzoxazine, and it particularly cannot withstand such high temperature heating as surface coatings

for several substrates [14–16]. Therefore, the search for reaction paths reducing the curing temperature has become one of the main focuses to overcome in this field.

Although it is not fully elucidated, it is commonly accepted that the thermal ring-opening polymerization of benzoxazine monomers occurs via cationic polymerization and the mechanism includes mainly three steps: coordination ring-opening of the oxazine ring, an electrophilic attack, and, finally, rearrangement [17]. In this context, several strategies have been proposed and reported in the literature to lower the polymerization temperature of benzoxazine precursors.

First, thanks to the great flexibility in the design of the molecular structure of benzoxazine monomers, several studies have focused on the understanding of the effects of structural parameters—including the nature, position, and hindrance that chemical groups bear on the aromatic backbone—on the reactivity of the corresponding monomers. For instance, the comparison of structurally different phenylenediamine-based benzoxazines with resorcinol/hydroquinone-based benzoxazines has indicated that the curing exotherm seems to be influenced to a large extent by the position of the oxazine ring on the aromatic units [18]. The influence of bridging groups on thermal ring-opening polymerization was also studied on diphenol-based benzoxazine monomers [19]. The authors observed that electron-withdrawing groups promoted thermal activated polymerization and that the curing temperature decreased by lowering the bond energy of C–O on oxazine rings. The distance separating the oxazine groups also seems an important parameter in the case of linear aliphatic diamine-based benzoxazine monomers. Indeed, it appears that the polymerization exotherm position decreases to a lower temperature range with the shortening of the length of the aliphatic diamine chain [20]. Moreover, the synthesis of multifunctional benzoxazines is also reported to undergo polymerization at lower temperatures when the number of oxazine functionalities increases [21,22].

In addition to these structural considerations, it also appears that the introduction of certain appropriate moieties like hydroxyl [23,24], imidazole [25], and amine [26] groups onto the backbone structure of benzoxazine monomers lowers the temperature of the ring-opening of the oxazine. In fact, it is now well known that the introduction of electron-withdrawing or electron-donating groups on the structural skeleton of benzoxazine monomers can greatly affect their reactivity by activating the ring-opening or stabilizing the intermediates [27]. Indeed, the catalyst effect of neighboring groups accelerating benzoxazine polymerization has been clearly shown in the case of carboxylic acid- [28], hydroxyl- [29], and amide [30] containing benzoxazine monomers. Moreover, the substitution on the ortho-position was found to further stimulate and activate the ring-opening polymerization when compared to its para-substituted counterpart as well in the case of a methylol functional benzoxazine [23,31] as in the case of an ortho-amide-imide functional benzoxazine [30].

Another strategy uses the reactivity of benzoxazine functions towards other functional groups such as amines. This approach is a rather new polymerization method via copolymerization than a direct acceleration of benzoxazine polymerization. Nevertheless, it is worth noting that in the case of copolymerization of benzoxazines with amines, in addition to the significant lowering of the polymerization temperature, an enhancement of the resulting mechanical and thermal properties of the networks has been achieved [32]. Another study reported the preparation and use of a polyamine-based oligomer to accelerate benzoxazine polymerization. Again, in this case, the polyamine was able to promote both the curing and, simultaneously, enhance the glass transition temperature as well as the thermal stability of the cured network [33].

A third strategy to reduce the curing temperature of benzoxazines consists of using initiators of cationic ring-opening polymerization. Some researchers have employed Lewis acids and nucleophilic catalysts to promote polymerization of benzoxazine, consequently reducing the curing temperature [34–36]. Indeed, several Lewis acids such as PCl_5 , PCl_3 , and POCl_3 , and transition metal salts such as TiCl_4 , AlCl_3 , CuCl_2 , AgCl , ZnCl_2 , NiCl_2 , FeCl_3 , and LiI have been tested successfully. Moreover, Liu et al. observed that LiI was especially efficient [37]. Recently, Akkus et al. highlighted that for amine salts, the nucleophilicity of the counterion had high impact, and they observed that the anion I^- exhibited the best catalytic behavior [38]. Organic-based catalysts such as acetylacetonato

complexes with iron, manganese, or cobalt, sulfonates such as para-toluene sulfonate, or organic oniums such as sulfonium hexafluoroantimonates were also found to be highly efficient for promoting the polymerization of benzoxazine precursors at a lower temperature [39–41].

Although many approaches have already been considered, they do not always allow retaining or improving the properties of the pristine polybenzoxazine network. Therefore, there is still a need for developing alternative solutions, which, in addition, should also consider environmental and toxicological issues for practical use, especially in the coating industry.

In this paper, we report the synthesis of a new diamine-based benzoxazine (4EP-pPDA) synthesized by the reaction of bio-basable reactants (4-ethylphenol, 1,4-phenylenediamine, and paraformaldehyde) and its thermal ring-opening polymerization in the presence of different kinds of cerium salts. The structure chosen for the benzoxazine monomers is based on both environmental and theoretical criteria. Indeed, 4-ethylphenol is a biomass-derived phenolic compound which can be obtained following an enzymatic process from natural and widely spread *p*-coumaric acid [42]. In addition to its bio-based character, the interest of 4-ethylphenol structures also lies in the absence of additional reactive functions on its structural backbone, thus allowing to use it as a model molecule to focus exclusively on the effect of cerium salts on the polymerization of benzoxazine functions. To the best of our knowledge, the ability of these metallic salts to accelerate benzoxazine polymerization has never been tested.

2. Materials and Methods

2.1. Materials

4-ethylphenol ($\geq 97.0\%$) and $\text{CeCl}_3 \cdot 7\text{H}_2\text{O}$ ($\geq 99.0\%$) were purchased from Alfa Aesar (Karlsruhe, Germany). 1,4-phenylenediamine ($\geq 99.0\%$) and $\text{Ce}(\text{OOCCH}_3)_3 \cdot 5\text{H}_2\text{O}$ ($\geq 99.9\%$) were purchased from Sigma-Aldrich (Darmstadt, Germany). $\text{Ce}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$ was purchased from Honeywell Fluka (Seelze, Germany). Paraformaldehyde ($\geq 95.0\%$) and absolute ethanol ($\geq 99.5\%$) were purchased from VWR (Leuven, Belgium). The starting reagents were used without further purification.

2.2. Preparation of 4EP-pPDA

The synthesis of 4EP-pPDA was adapted from a procedure in bulk reported elsewhere by Ishida et al. [4]. 4-ethylphenol 23.21 g (1.9×10^{-1} mol) and 1,4-phenylenediamine 10.00 g (9.25×10^{-2} mol) were mixed in a beaker with a mechanical stirrer at 120 °C until a homogeneous liquid was obtained. Then, paraformaldehyde in excess of 12.22 g (4.07×10^{-1} mol) was rapidly added under vigorous stirring to prevent bubbling from the rapid decomposition of paraformaldehyde into formaldehyde. The resulting mixture was reacted for seven additional minutes under continuous stirring. The crude reaction product was dissolved in refluxing ethanol (about 500 mL) and the precursors of the resin were precipitated upon cooling. The resulting precipitate was collected, filtered and abundantly washed with cold ethanol. Then it was dried under vacuum. A vitrified yellow resin was obtained (weight yield about 65%).

2.3. Preparation of 4EP-pPDA/Cerium Salt Mixtures

4EP-pPDA and cerium salts were first added into the dichloromethane and ethanol mixture solvents (dichloromethane/ethanol=90/10 v/v). After stirring for 2 h at room temperature, the solvent was removed by volatilizing and drying under vacuum. The content of each cerium salt in the 4EP-pPDA substrate was 5 mmol/100 g. In this work, we denote briefly the samples made from $\text{Ce}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$, $\text{CeCl}_3 \cdot 7\text{H}_2\text{O}$, and $\text{Ce}(\text{OOCCH}_3)_3 \cdot 5\text{H}_2\text{O}$ as 5mM $\text{Ce}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$, 5mM $\text{CeCl}_3 \cdot 7\text{H}_2\text{O}$, and 5mM $\text{Ce}(\text{OOCCH}_3)_3 \cdot 5\text{H}_2\text{O}$, respectively.

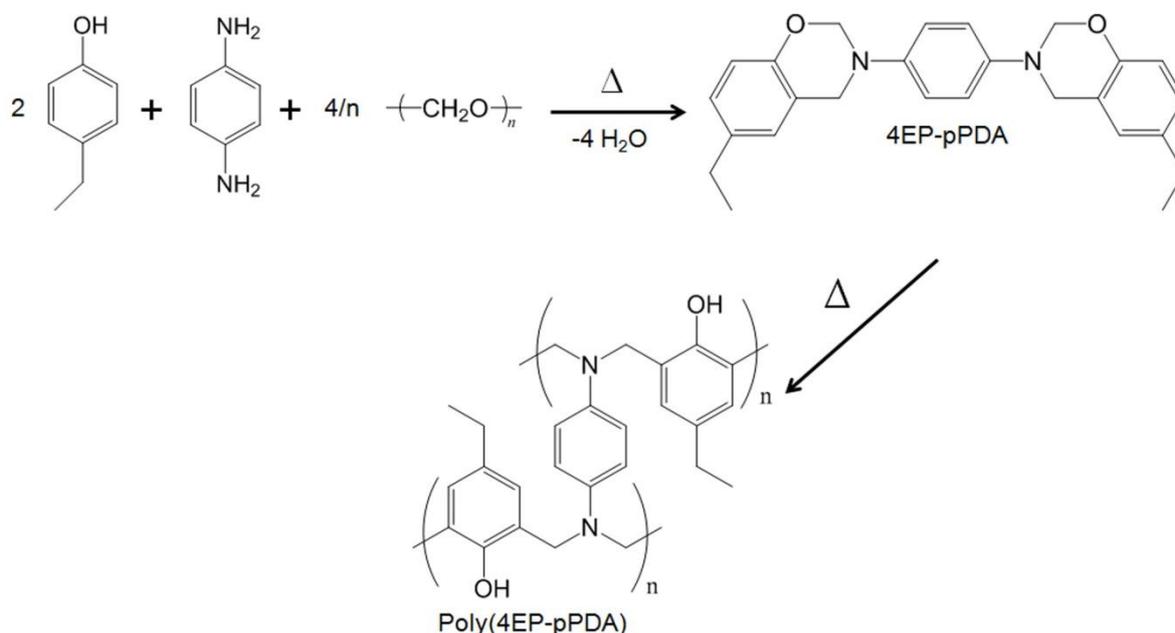
2.4. Measurements and Characterization

Fourier transform infrared (FT-IR) spectra were recorded using a BRUKER IFS 66v/S spectrometer (Bruker Instruments, Billerica, MA, USA) from 400 to 4000 cm^{-1} with a resolution of 4 cm^{-1} . $^1\text{H-NMR}$

spectra were obtained by using a BRUKER Avance DMX-500 MHz spectrometer (Bruker Instruments, Billerica, MA, USA) in CDCl_3 with TMS as the internal standard. Differential scanning calorimetry (DSC) was performed on a TA DSC Q200 calorimeter (TA Instruments, New Castle, DE, USA) from room temperature up to 300 °C with a heating rate of 10 °C/min and a nitrogen flow rate of 50 mL/min. Field-emission scanning electron microscopy (FE-SEM) and energy-dispersive X-ray (EDX) analysis were performed with a Hitachi S-4800 analyzer (Hitachi, Tokyo, Japan) operated at 20 kV. Thermogravimetric analysis (TGA) was performed on a TA TGA Q50 thermogravimetric analyzer (TA Instruments, New Castle, DE, USA) from room temperature to 550 °C with a heating rate of 10 °C/min under a nitrogen atmosphere UV-visible spectral measurements were performed on the elaborated powders with a Labsphere RSA-PE-19 reflectance accessory (Labsphere, N. Sutton, NY, USA).

3. Results and Discussion

A new diamine-based benzoxazine (4EP-pPDA) was synthesized by the reaction of 4-ethylphenol, 1,4-phenylenediamine, and paraformaldehyde as shown in Scheme 1.



Scheme 1. Synthesis of 4EP-pPDA and corresponding cured network of Poly(4EP-pPDA).

The chemical structure was investigated by FT-IR and $^1\text{H-NMR}$. The results are gathered in Figure 1. More specifically, the FT-IR spectrum of 4EP-pPDA shows absorption peaks at 1516, 1223, 1034, and 945 cm^{-1} . Based on similar benzoxazine derivatives reported in the literature [43–45], they are ascribed to the trisubstituted benzene ring stretching, asymmetric stretching vibration of C-O-C, symmetric stretching vibration of C-O-C, and out-of-plane C-H, respectively. In the $^1\text{H-NMR}$ spectrum of 4EP-pPDA, the characteristic protons appear at around 4.51 (Ar- CH_2 -N) and 5.24 (O- CH_2 -N) ppm, confirming the formation of the 4EP-pPDA molecular structure [45–47]. Furthermore, the integration ratios of the peaks agree well with the proposed chemical structure. These results further confirm the successful synthesis of 4EP-pPDA precursors.

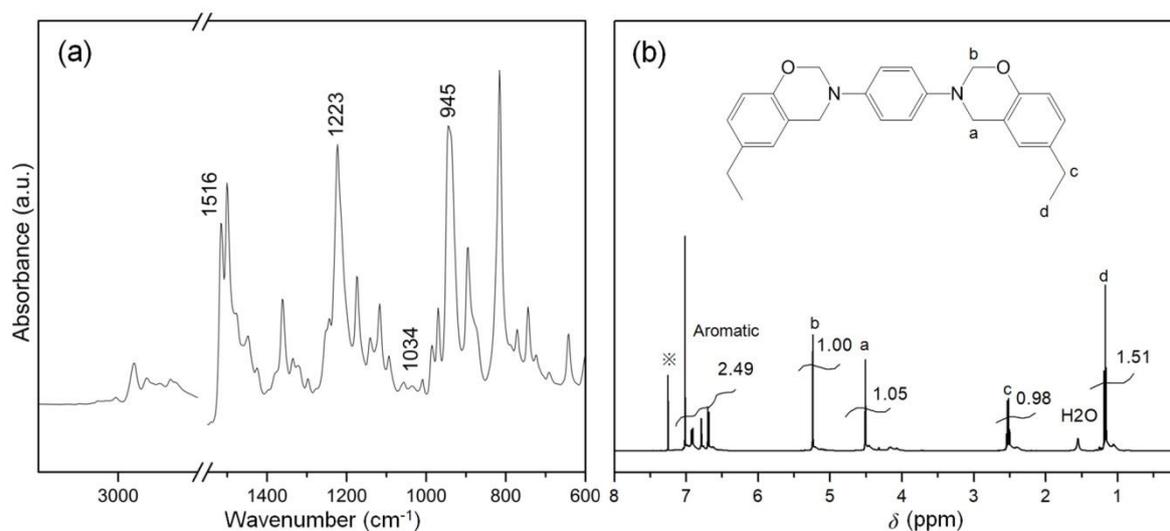
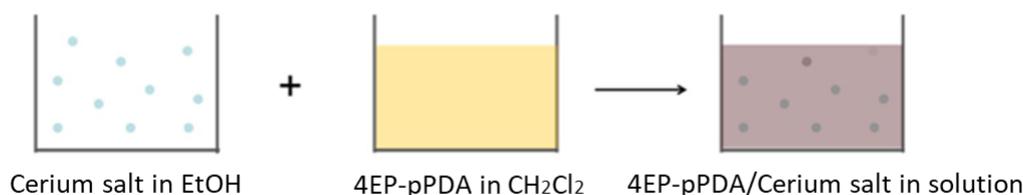


Figure 1. (a) FT-IR and (b) $^1\text{H-NMR}$ spectrum of 4EP-pPDA.

Thereafter, the cerium salts were incorporated within the 4EP-pPDA precursors via a solvent mixture of dichloromethane/ethanol (90/10 v/v) as shown in Scheme 2.



Scheme 2. Illustration of the preparation process of 4EP-pPDA/cerium salt mixtures.

Interestingly, the three salts (i.e., $\text{Ce}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$, $\text{CeCl}_3 \cdot 7\text{H}_2\text{O}$, and $\text{Ce}(\text{OOCCH}_3)_3 \cdot 5\text{H}_2\text{O}$) seem to dissolve well within the 4EP-pPDA-based solution and color changes are observed for the three systems. Indeed, the solutions containing 5 mM $\text{Ce}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$, 5 mM $\text{CeCl}_3 \cdot 7\text{H}_2\text{O}$, and 5 mM $\text{Ce}(\text{OOCCH}_3)_3 \cdot 5\text{H}_2\text{O}$ all become dark red (see Figure 2). This color change phenomenon is mostly due to the complexation between a 4EP-pPDA monomer with Ce ions via an oxygen or nitrogen atom in the oxazine ring as coordination atoms [48,49].

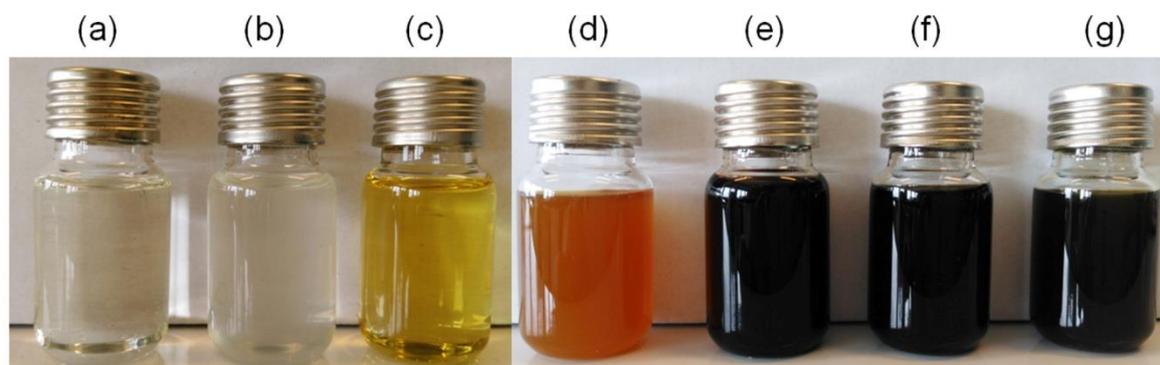


Figure 2. Photos of (a) $\text{Ce}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$, (b) $\text{CeCl}_3 \cdot 7\text{H}_2\text{O}$, (c) $\text{Ce}(\text{OOCCH}_3)_3 \cdot 5\text{H}_2\text{O}$, (d) 4EP-pPDA, (e) 5 mM $\text{Ce}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$, (f) 5 mM $\text{CeCl}_3 \cdot 7\text{H}_2\text{O}$, and (g) 5 mM $\text{Ce}(\text{OOCCH}_3)_3 \cdot 5\text{H}_2\text{O}$ in the solvents of dichloromethane/ethanol (90/10 v/v).

After removal of the solvent mixture, the color of the samples is modified and different shades between the three samples appear, as shown in Figure 3. Interestingly, the color of all samples looks uniform and homogeneous, suggesting a fine dispersion of all the cerium salts within the 4EP-pPDA precursors.



Figure 3. Photos of powders of (a) 4EP-pPDA, (b) 5 mM $\text{Ce}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$, (c) 5 mM $\text{CeCl}_3 \cdot 7\text{H}_2\text{O}$, and (d) 5 mM $\text{Ce}(\text{OOCCH}_3)_3 \cdot 5\text{H}_2\text{O}$.

The complexation between 4EP-pPDA monomers with Ce ions was further confirmed by UV-VIS spectroscopy. Indeed, as shown in Figure 4, a broadening and shifting of the band towards 500 nm attributed to the formation of cerium complexes with 4EP-pPDA monomers are observed in the case of systems 5 mM $\text{Ce}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$, 5 mM $\text{CeCl}_3 \cdot 7\text{H}_2\text{O}$, and 5 mM $\text{Ce}(\text{OOCCH}_3)_3 \cdot 5\text{H}_2\text{O}$.

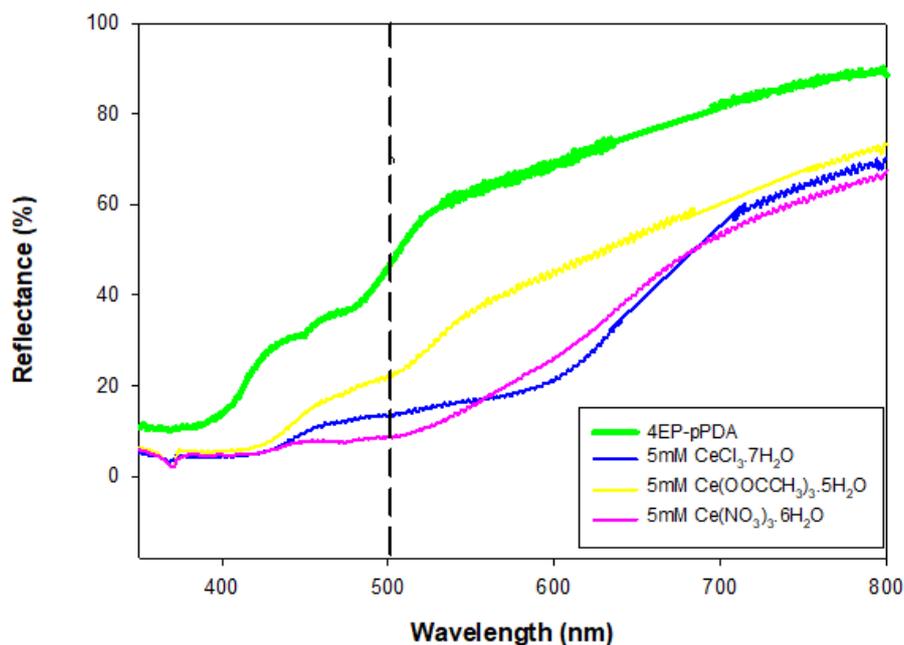


Figure 4. UV-vis spectra of 4EP-pPDA, 5 mM $\text{Ce}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$, 5 mM $\text{CeCl}_3 \cdot 7\text{H}_2\text{O}$, and 5 mM $\text{Ce}(\text{OOCCH}_3)_3 \cdot 5\text{H}_2\text{O}$.

The effect of cerium salts on the curing behavior of 4EP-pPDA precursors was studied by DSC, as shown in Figure 5a. The neat 4EP-pPDA precursors synthesized for this study exhibit an exothermic peak with a maximum temperature (T_p) at about 232 °C, corresponding to the ring-opening polymerization of benzoxazine. For all 4EP-pPDA systems containing cerium salt, one exotherm is also observed, but the T_p of these systems is shifted to lower temperatures. The presence of only one exotherm further confirms the good dispersion of cerium salts within the 4EP-pPDA matrix. It also shows the ability of cerium salts to accelerate the ring-opening polymerization of 4EP-pPDA

benzoxazine precursors. Although all three systems contain the same amount of cerium, the decrease in T_p appears not to be identical. Indeed, the following order is observed: 5 mM $\text{Ce}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$ (36°C) > 5 mM $\text{CeCl}_3 \cdot 7\text{H}_2\text{O}$ (20°C) > 5 mM $\text{Ce}(\text{OOCCH}_3)_3 \cdot 5\text{H}_2\text{O}$ (15°C), indicating differences in the catalytic activities for the three cerium salts. On the other hand, an obvious decrease in the enthalpy of the curing reaction (ΔH) is observed with the introduction of cerium salt. It also should be noted that $\text{Ce}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$, $\text{CeCl}_3 \cdot 7\text{H}_2\text{O}$, and $\text{Ce}(\text{OOCCH}_3)_3 \cdot 5\text{H}_2\text{O}$ all contain structural water which is released in the curing region of 4EP-pPDA, as observed by DSC in Figure 6. Hence, the reason for the reduction of ΔH can also be explained by the superposition of two antagonist phenomena, i.e., the benzoxazine polymerization, which is exothermic, and the releasing of crystal water, which is endothermic.

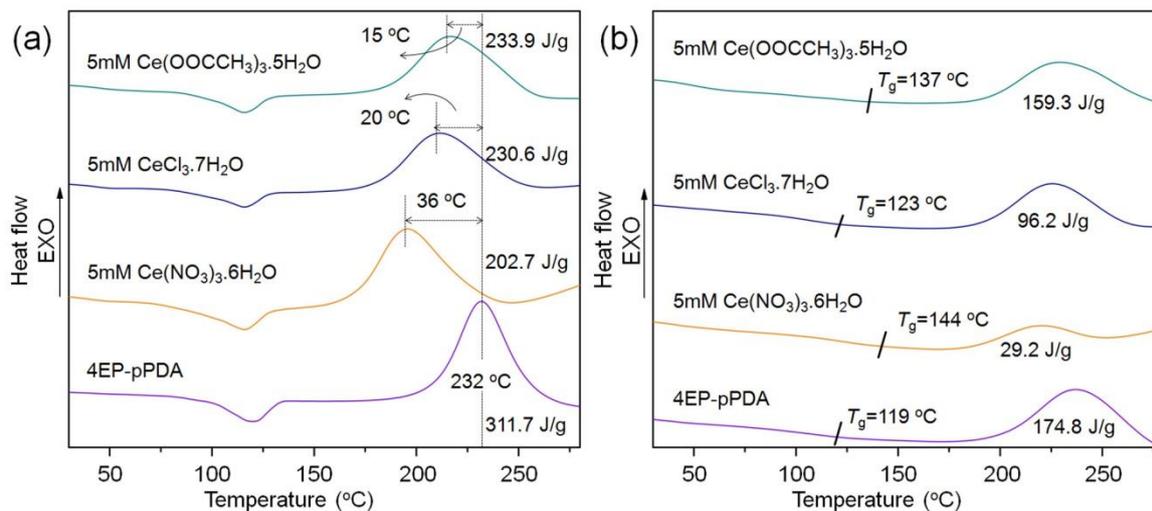


Figure 5. Differential scanning calorimetry (DSC) curves of 5 mM $\text{Ce}(\text{OOCCH}_3)_3 \cdot 5\text{H}_2\text{O}$, 5 mM $\text{CeCl}_3 \cdot 7\text{H}_2\text{O}$, 5 mM $\text{Ce}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$, and 4EP-pPDA (a) before and (b) after curing at 150°C for 4 h.

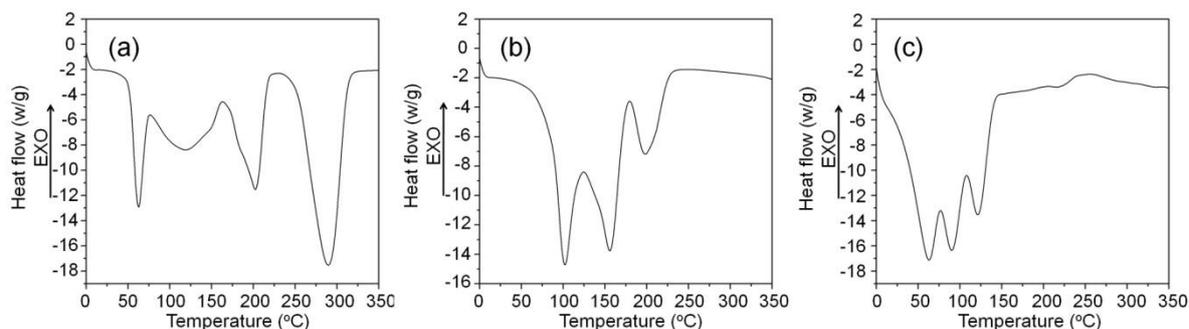


Figure 6. DSC curves of (a) $\text{Ce}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$, (b) $\text{CeCl}_3 \cdot 7\text{H}_2\text{O}$, and (c) $\text{Ce}(\text{OOCCH}_3)_3 \cdot 5\text{H}_2\text{O}$.

To try and clarify the activities of cerium salts, the 4EP-pPDA systems containing cerium salts were further monitored by DSC after isothermal curing at 150°C for 4h, as shown in Figure 5b. The polymerization enthalpy ΔH of a neat 4EP-pPDA system is found to decrease by 44%. By contrast, the enthalpies of 5 mM $\text{Ce}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$, 5 mM $\text{CeCl}_3 \cdot 7\text{H}_2\text{O}$, and 5 mM $\text{Ce}(\text{OOCCH}_3)_3 \cdot 5\text{H}_2\text{O}$ are found to decrease by 86%, 58%, and 32%, respectively. This result clearly indicates that among the three cerium salts, $\text{Ce}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$ is the one exhibiting the best catalytic efficiency, which can significantly shorten the curing time. Surprisingly, $\text{Ce}(\text{OOCCH}_3)_3 \cdot 5\text{H}_2\text{O}$ exhibits the worst performance whereas the activity of $\text{CeCl}_3 \cdot 7\text{H}_2\text{O}$ stands in between that of $\text{Ce}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$ and $\text{Ce}(\text{OOCCH}_3)_3 \cdot 5\text{H}_2\text{O}$. Furthermore, the order of glass transition temperatures (T_g) observed is 5 mM $\text{Ce}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$ (144°C) > 5 mM $\text{Ce}(\text{OOCCH}_3)_3 \cdot 5\text{H}_2\text{O}$ (137°C) > 5 mM $\text{CeCl}_3 \cdot 7\text{H}_2\text{O}$ (123°C) > 4EP-pPDA (119°C). The highest T_g of the 5 mM $\text{Ce}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$ system can be explained by the highest curing degree and the formation of the corresponding polybenzoxazine network.

To better understand the above results, representative samples of 4EP-pPDA and 4EP-pPDA modified by cerium salts after curing at 150 °C for 1h were analyzed by FT-IR and $^1\text{H-NMR}$. As shown in Figure 7a, by selecting the peak at 1516 cm^{-1} as the reference, the relative absorption intensities of the benzoxazine ring for 4EP-pPDA/cerium salt mixtures are found to be weaker than those of 4EP-pPDA. In addition, as presented in Figure 7b, the peaks at 5.24 and 4.51 ppm are split into two signals, and the integrations at 5.24 ppm are lower than those of the samples before curing, indicating the breakage of O-CH₂-N groups. The order of relative content for the O-CH₂-N group is 5 mM Ce(NO₃)₃·6H₂O < 5 mM Ce(OOCCH₃)₃·5H₂O < 5 mM CeCl₃·7H₂O < 4EP-pPDA. Furthermore, new signals at about 9.9 and 8.5 ppm can be observed, revealing the presence of an Ar-OH group [48]. The order of relative content for the Ar-OH group is 5 mM Ce(NO₃)₃·6H₂O > 5 mM Ce(OOCCH₃)₃·5H₂O > 5 mM CeCl₃·7H₂O > 4EP-pPDA. Therefore, from these results it appears clearly that during the curing process the cerium salts promote the breaking of the O-CH₂-N group and the corresponding cross-linking reaction.

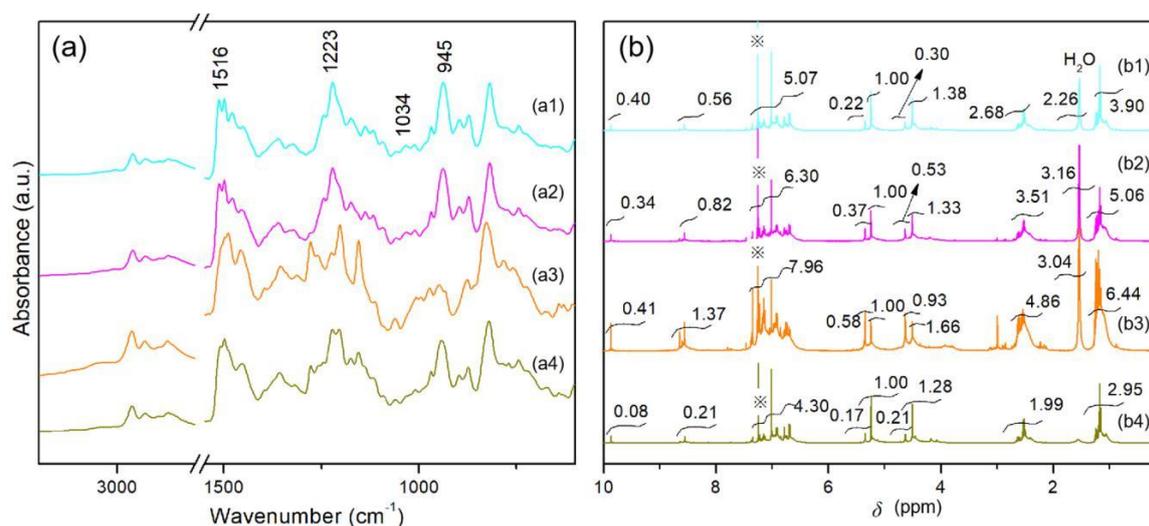


Figure 7. (a) FT-IR spectra and (b) $^1\text{H-NMR}$ spectra of the 5mM Ce(OOCCH₃)₃·5H₂O (a1, b1), 5mM CeCl₃·7H₂O (a2, b2), 5mM Ce(NO₃)₃·6H₂O (a3, b3), and neat 4EP-pPDA (a4, b4) after curing at 150 °C for 1 h.

Figure 8a–d presents the top view SEM images of 4EP-pPDA (a), 5 mM Ce(NO₃)₃·6H₂O (b), 5 mM CeCl₃·7H₂O (c), and 5 mM Ce(OOCCH₃)₃·5H₂O (d) after curing at 150 °C for 4 h. Clearly, some white particles can be seen in Figure 8c,d, which are attributed to the cerium salts. The elemental composition of the cured samples was also analyzed as shown in Figure 8e–h. The order of weight content of Ce is 5 mM Ce(NO₃)₃·6H₂O > 5 mM CeCl₃·7H₂O > 5 mM Ce(OOCCH₃)₃·5H₂O, which is consistent with the catalytic activity previously determined. The lowest cerium content observed for the sample 5 mM Ce(OOCCH₃)₃·5H₂O is mostly due to the lowest dispersibility of Ce(OOCCH₃)₃·5H₂O. When removing the solvent, most of the Ce(OOCCH₃)₃·5H₂O possibly precipitates out of 4EP-pPDA matrix. Hence, the differences in the catalytic activity are mostly due to the dispersibility of the cerium salt within the 4EP-pPDA matrix. If it is dispersed in the matrix at a molecular level, the complexation between the 4EP-pPDA monomer with Ce ion via the oxygen or nitrogen atom in the oxazine ring may more easily promote the breaking of a O-CH₂-N group, so the catalytic effect is obvious.

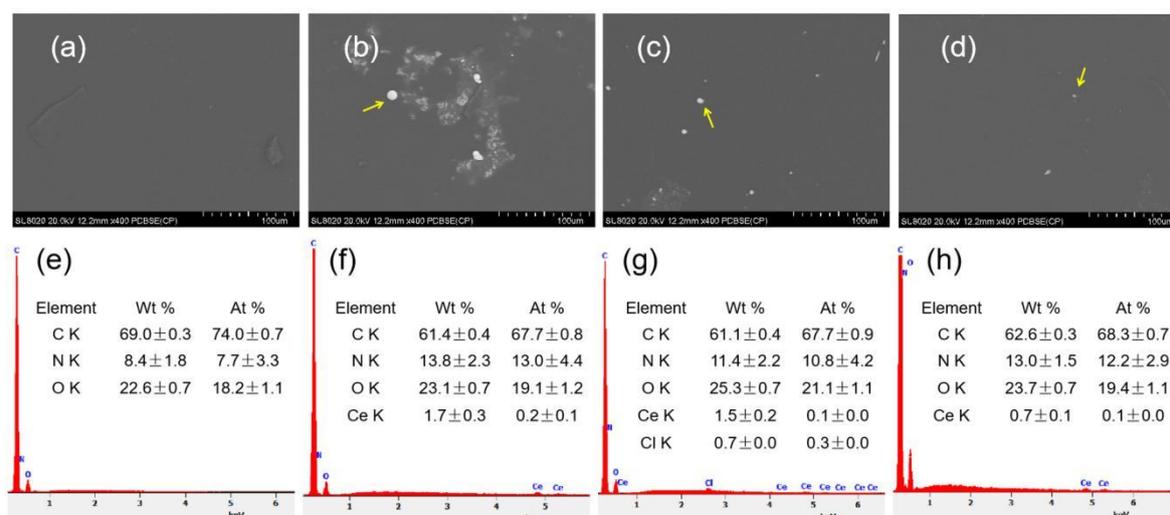


Figure 8. View of SEM images (first row) and EDX spectra (second row) of (a,e) 4EP-pPDA, (b,f) 5 mM $\text{Ce}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$, (c,g) 5 mM $\text{CeCl}_3 \cdot 7\text{H}_2\text{O}$, and (d,h) 5 mM $\text{Ce}(\text{OOCCH}_3)_3 \cdot 5\text{H}_2\text{O}$ after curing at 150 °C for 4 h.

In addition, the effect of the above three cerium salts on the thermal stability of 4EP-pPDA after curing at 150 °C for 4 h was investigated, as shown in Figure 9. As for the 5 wt % weight loss temperature ($T_{5\%}$), the $\text{Ce}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$ shows almost no influence, but the $\text{CeCl}_3 \cdot 7\text{H}_2\text{O}$ and $\text{Ce}(\text{OOCCH}_3)_3 \cdot 5\text{H}_2\text{O}$ present obvious improvement. As for the residue at 550 °C, the $\text{CeCl}_3 \cdot 7\text{H}_2\text{O}$ shows no influence, and the $\text{Ce}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$ and $\text{Ce}(\text{OOCCH}_3)_3 \cdot 5\text{H}_2\text{O}$ present little improvement. Therefore, the three cerium salts reported herein can accelerate the ring-opening polymerization of 4EP-pPDA benzoxazine precursors without sacrificing their thermal stability.

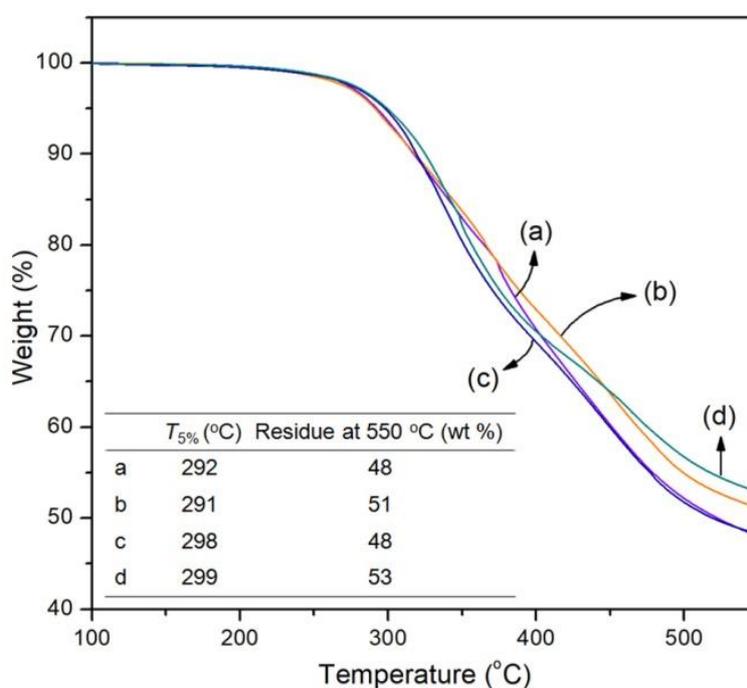


Figure 9. TGA curves of (a) 4EP-pPDA, (b) 5 mM $\text{Ce}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$, (c) 5 mM $\text{CeCl}_3 \cdot 7\text{H}_2\text{O}$, and (d) 5 mM $\text{Ce}(\text{OOCCH}_3)_3 \cdot 5\text{H}_2\text{O}$ after curing at 150 °C for 4 h.

4. Conclusions

Model benzoxazine precursors (4EP-pPDA) were synthesized from bio-basable 4-ethylphenol, 1,4-phenylenediamine, and paraformaldehyde and used to test the ability of three different cerium salts (i.e., $\text{Ce}(\text{OOCCH}_3)_3 \cdot 5\text{H}_2\text{O}$, $\text{CeCl}_3 \cdot 7\text{H}_2\text{O}$, and $\text{Ce}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$) to catalyze its ring-opening polymerization. Although the three cerium salts studied were found to accelerate the opening of a O-CH₂-N group and promote the corresponding cross-linking reaction of 4EP-pPDA precursors, they also exhibit different catalytic activities. $\text{Ce}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$ was found to be the most efficient catalyst to reduce the curing temperature of 4EP-pPDA precursors. This result was mainly related to its better dispersibility within the benzoxazine matrix. It seems that the most important parameter is to obtain a dispersion of the initiator at a molecular scale to achieve optimal catalytic efficiency.

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References

1. Ning, X.; Ishida, H. Phenolic materials via ring-opening polymerization: Synthesis and characterization of bisphenol-A based benzoxazines and their polymers. *J. Polym. Sci. Part A: Polym. Chem.* **1994**, *32*, 1121–1129. [[CrossRef](#)]
2. Yagci, Y.; Kiskan, B.; Ghosh, N.N. Recent advancement on polybenzoxazine-A newly developed high performance thermoset. *J. Polym. Sci. Part A: Polym. Chem.* **2009**, *47*, 5565–5576. [[CrossRef](#)]
3. Ishida, H. Overview and Historical Background of Polybenzoxazine Research. In *Handbook of Benzoxazine Resins*; Elsevier BV: Amsterdam, the Netherlands, 2011; pp. 3–81.
4. Ishida, H. Process for Preparation of Benzoxazine Compounds in Solventless Systems. United State Patent 5543516, 6 August 1996.
5. Ghosh, N.; Kiskan, B.; Yagci, Y. Polybenzoxazines—New high performance thermosetting resins: Synthesis and properties. *Prog. Polym. Sci.* **2007**, *32*, 1344–1391. [[CrossRef](#)]
6. Ishida, H.; Liu, J.-P. Benzoxazine Chemistry in Solution and Melt. In *Handbook of Benzoxazine Resins*; Elsevier BV: Amsterdam, the Netherlands, 2011; Volume Ch. 2, pp. 85–102.
7. Arslan, M.; Kiskan, B.; Yagci, Y. Recycling and Self-Healing of Polybenzoxazines with Dynamic Sulfide Linkages. *Sci. Rep.* **2017**, *7*, 5207. [[CrossRef](#)]
8. Caldon, E.B.; De Leon, A.C.C.; Thomas, P.G.; Naylor, D.F.; Pajarito, B.B.; Advincula, R.C. Superhydrophobic Rubber-Modified Polybenzoxazine/SiO₂ Nanocomposite Coating with Anticorrosion, Anti-Ice, and Superoleophilicity Properties. *Ind. Eng. Chem. Res.* **2017**, *56*, 1485–1497. [[CrossRef](#)]
9. Kiskan, B.; Yagci, Y. Thermally curable benzoxazine monomer with a photodimerizable coumarin group. *J. Polym. Sci. Part A: Polym. Chem.* **2007**, *45*, 1670–1676. [[CrossRef](#)]
10. Nair, C. Advances in addition-cure phenolic resins. *Prog. Polym. Sci.* **2004**, *29*, 401–498. [[CrossRef](#)]
11. Takeichi, T.; Kawauchi, T.; Agag, T. High Performance Polybenzoxazines as a Novel Type of Phenolic Resin. *Polym. J.* **2008**, *40*, 1121–1131. [[CrossRef](#)]
12. Dumas, L.; Bonnaud, L.; Olivier, M.; Poorteman, M.; Dubois, P. Facile preparation of a novel high performance benzoxazine–CNT based nano-hybrid network exhibiting outstanding thermo-mechanical properties. *Chem. Commun.* **2013**, *49*, 9543–9545. [[CrossRef](#)]
13. Dumas, L.; Bonnaud, L.; Olivier, M.; Poorteman, M.; Dubois, P. Multiscale benzoxazine composites: The role of pristine CNTs as efficient reinforcing agents for high-performance applications. *Compos. Part B Eng.* **2017**, *112*, 57–65. [[CrossRef](#)]

14. Renaud, A.; Poorteman, M.; Escobar, J.; Dumas, L.; Paint, Y.; Bonnaud, L.; Dubois, P.; Olivier, M. A new corrosion protection approach for aeronautical applications combining a phenol-paraphenylenediamine benzoxazine resin applied on sulfo-tartaric anodized aluminum. *Prog. Org. Coat.* **2017**, *112*, 278–287. [[CrossRef](#)]
15. Poorteman, M.; Renaud, A.; Escobar, J.; Dumas, L.; Bonnaud, L.; Dubois, P.; Olivier, M.-G. Thermal curing of para -phenylenediamine benzoxazine for barrier coating applications on 1050 aluminum alloys. *Prog. Org. Coat.* **2016**, *97*, 99–109. [[CrossRef](#)]
16. Renaud, A.; Paint, Y.; Lanzutti, A.; Bonnaud, L.; Fedrizzi, L.; Dubois, P.; Poorteman, M.; Olivier, M. Sealing porous anodic layers on AA2024-T3 with a low viscosity benzoxazine resin for corrosion protection in aeronautical applications. *RSC Adv.* **2019**, *9*, 16819–16830. [[CrossRef](#)]
17. Liu, C.; Chen, Q.-Y. Catalytic Accelerated Polymerization of Benzoxazines and Their Mechanistic Considerations. In *Advanced and Emerging Polybenzoxazine Science and Technology*; Elsevier BV: Amsterdam, the Netherlands, 2017; Volume Ch. 2, pp. 9–21.
18. Rishwana, S.S.; Pitchaimari, G.; Vijayakumar, C.T. Studies on structurally different diamines and bisphenol benzoxazines: synthesis and curing kinetics. *High Perform. Poly.* **2016**, *28*, 466–478. [[CrossRef](#)]
19. Wang, X.; Chen, F.; Gu, Y. Influence of electronic effects from bridging groups on synthetic reaction and thermally activated polymerization of bisphenol-based benzoxazines. *J. Polym. Sci. Part A: Polym. Chem.* **2011**, *49*, 1443–1452. [[CrossRef](#)]
20. Allen, D.J.; Ishida, H. Polymerization of linear aliphatic diamine-based benzoxazine resins under inert and oxidative environments. *Polymer* **2007**, *48*, 6763–6772. [[CrossRef](#)]
21. Soto, M.; Hiller, M.; Oschkinat, H.; Koschek, K. Multifunctional benzoxazines feature low polymerization temperature and diverse polymer structures. *Polymers* **2016**, *8*, 278. [[CrossRef](#)]
22. Sini, N.K.; Endo, T. Toward Elucidating the Role of Number of Oxazine Rings and Intermediates in the Benzoxazine Backbone on Their Thermal Characteristics. *Macromolecules* **2016**, *49*, 8466–8478. [[CrossRef](#)]
23. Baquar, M.; Agag, T.; Huang, R.; Maia, J.; Qutubuddin, S.; Ishida, H. Mechanistic pathways for the polymerization of methylol-functional benzoxazine monomers. *Macromolecules* **2012**, *45*, 8119–8125. [[CrossRef](#)]
24. Xu, H.; Zhang, W.; Lu, Z.; Zhang, G. Hybrid polybenzoxazine with tunable properties. *RSC Adv.* **2013**, *3*, 3677. [[CrossRef](#)]
25. Yang, P.; Gu, Y. A novel benzimidazole moiety-containing benzoxazine: Synthesis, polymerization, and thermal properties. *J. Polym. Sci. Part A: Polym. Chem.* **2012**, *50*, 1261–1271. [[CrossRef](#)]
26. Agag, T.; Arza, C.R.; Maurer, F.H.J.; Ishida, H. Primary Amine-Functional Benzoxazine Monomers and Their Use for Amide-Containing Monomeric Benzoxazines. *Macromolecules* **2010**, *43*, 2748–2758. [[CrossRef](#)]
27. Baqar, M.; Agag, T.; Qutubuddin, S.; Ishida, H. Effect of Neighboring Groups on Enhancing Benzoxazine Autocatalytic Polymerization. In *Handbook of Benzoxazine Resins*; Elsevier BV: Amsterdam, the Netherlands, 2011; pp. 193–210.
28. Andreu, R.; Reina, J.A.; Ronda, J.C. Carboxylic acid-containing benzoxazines as efficient catalysts in the thermal polymerization of benzoxazines. *J. Polym. Sci. Part A: Polym. Chem.* **2008**, *46*, 6091–6101. [[CrossRef](#)]
29. Kudoh, R.; Sudo, A.; Endo, T. A Highly Reactive Benzoxazine Monomer, 1-(2-Hydroxyethyl)-1,3-Benzoxazine: Activation of Benzoxazine by Neighboring Group Participation of Hydroxyl Group. *Macromolecules* **2010**, *43*, 1185–1187. [[CrossRef](#)]
30. Zhang, K.; Ishida, H. Smart synthesis of high-performance thermosets based on ortho-amide-imide functional benzoxazines. *Front. Mat.* **2015**, *2*, 5. [[CrossRef](#)]
31. Zhang, K.; Froimowicz, P.; Han, L.; Ishida, H. Hydrogen-bonding characteristics and unique ring-opening polymerization behavior of Ortho-methylol functional benzoxazine. *J. Polym. Sci. Part A: Polym. Chem.* **2016**, *54*, 3635–3642. [[CrossRef](#)]
32. Sun, J.; Wei, W.; Xu, Y.; Qu, J.; Liu, X.; Endo, T. A curing system of benzoxazine with amine: reactivity, reaction mechanism and material properties. *RSC Adv.* **2015**, *5*, 19048–19057. [[CrossRef](#)]
33. Zhang, L.; Mao, J.; Wang, S.; Yang, Y.; Chen, Y.; Lu, H. Meta-phenylenediamine formaldehyde oligomer: A new accelerator for benzoxazine resin. *React. Funct. Polym.* **2017**, *121*, 51–57. [[CrossRef](#)]
34. Ran, Q.-C.; Zhang, D.-X.; Zhu, R.-Q.; Gu, Y. The structural transformation during polymerization of benzoxazine/FeCl₃ and the effect on the thermal stability. *Polymer* **2012**, *53*, 4119–4127. [[CrossRef](#)]

35. Kocaarslan, A.; Kiskan, B.; Yagci, Y. Ammonium salt catalyzed ring-opening polymerization of 1,3-benzoxazines. *Polymer* **2017**, *122*, 340–346. [[CrossRef](#)]
36. Liu, C.; Shen, D.; Sebastián, R.M.; Marquet, J.; Schönfeld, R. Catalyst effects on the ring-opening polymerization of 1,3-benzoxazine and on the polymer structure. *Polymer* **2013**, *54*, 2873–2878. [[CrossRef](#)]
37. Liu, C.; Shen, D.; Sebastián, R.M.; Marquet, J.; Schönfeld, R. Mechanistic Studies on Ring-Opening Polymerization of Benzoxazines: A Mechanistically Based Catalyst Design. *Macromolecules* **2011**, *44*, 4616–4622. [[CrossRef](#)]
38. Akkus, B.; Kiskan, B.; Yagci, Y. Counterion Effect of Amine Salts on Ring-Opening Polymerization of 1,3-Benzoxazines. *Macromol. Chem. Phys.* **2018**, *220*. [[CrossRef](#)]
39. Sudo, A.; Hirayama, S.; Endo, T. Highly efficient catalysts-acetylacetonato complexes of transition metals in the 4th period for ring-opening polymerization of 1,3-benzoxazine. *J. Polym. Sci. Part A: Polym. Chem.* **2010**, *48*, 479–484. [[CrossRef](#)]
40. Sudo, A.; Mori, A.; Endo, T. Promoting effects of urethane derivatives of phenols on the ring-opening polymerization of 1,3-benzoxazines. *J. Polym. Sci. Part A: Polym. Chem.* **2011**, *49*, 2183–2190. [[CrossRef](#)]
41. Zhang, D.; Yue, J.; Li, H.; Li, Y.; Zhao, C. Curing kinetics study of benzoxazine using diaryliodonium salts as thermal initiators. *Thermochim. Acta* **2016**, *643*, 13–22. [[CrossRef](#)]
42. Chatonnet, P.; Dubourdie, D.; Boidron, J.-N.; Pons, M. The origin of ethylphenols in wines. *J. Sci. Food Agric.* **1992**, *60*, 165–178. [[CrossRef](#)]
43. Gao, S.; Feng, S.; Lu, Z.; Liu, Y. Synthesis of borosiloxane/polybenzoxazine hybrids as highly efficient and environmentally friendly flame retardant materials. *J. Polym. Sci. Part A: Polym. Chem.* **2017**, *55*, 2390–2396. [[CrossRef](#)]
44. Bonnaud, L.; Chollet, B.; Dumas, L.; Peru, A.A.M.; Flourat, A.L.; Allais, F.; Dubois, P. High-Performance Bio-Based Benzoxazines from Enzymatic Synthesis of Diphenols. *Macromol. Chem. Phys.* **2018**, *220*, 1800312. [[CrossRef](#)]
45. Dumas, L.; Bonnaud, L.; Olivier, M.; Poorteman, M.; Dubois, P. Chavicol benzoxazine: Ultrahigh Tg biobased thermoset with tunable extended network. *Eur. Polym. J.* **2016**, *81*, 337–346. [[CrossRef](#)]
46. Froimowicz, P.; Zhang, K.; Ishida, H. Intramolecular Hydrogen Bonding in Benzoxazines: When Structural Design Becomes Functional. *Chem. A Eur. J.* **2016**, *22*, 2691–2707. [[CrossRef](#)]
47. Dumas, L.; Bonnaud, L.; Olivier, M.; Poorteman, M.; Dubois, P. High performance benzoxazine/CNT nanohybrid network – An easy and scalable way to combine attractive properties. *Eur. Polym. J.* **2014**, *58*, 218–225. [[CrossRef](#)]
48. Wattanathana, W.; Veranitisagul, C.; Koonsaeng, N.; Laobuthee, A. 3, 4-Dihydro-1, 3-2H-Benzoxazines: Uses Other Than Making Polybenzoxazines. In *Advanced and Emerging Polybenzoxazine Science and Technology*; Elsevier: Amsterdam, the Netherlands, 2017; pp. 75–88.
49. Veranitisagul, C.; Kaewvilai, A.; Sangngern, S.; Wattanathana, W.; Suramitr, S.; Koonsaeng, N.; Laobuthee, A. Novel Recovery of Nano-Structured Ceria (CeO₂) from Ce(III)-Benzoxazine Dimer Complexes via Thermal Decomposition. *Int. J. Mol. Sci.* **2011**, *12*, 4365–4377. [[CrossRef](#)]

