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A. Demongeot, S. Mognier, S. Okada, C. Soulié-Ziakovic, F. Tournilhac. Coordination and catalysis of Zn $^{2+}$ in epoxy-based vitrimers. *Polymer Chemistry*, 2016, 7 (27), pp.4486-4493. 10.1039/C6PY00752J . hal-03229292

HAL Id: hal-03229292

<https://hal.science/hal-03229292>

Submitted on 25 Oct 2021

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Coordination and Catalysis of Zn^{2+} in Epoxy-Based Vitrimers†

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Accepted manuscript

DOI : 10.1039/C6PY00752J

To elucidate the catalytic action of $\text{Zn}(\text{II})$ in the exchange reactions taking place in epoxy-acid vitrimers, extended X-ray absorption fine structure (EXAFS) spectroscopy has been used as well as infrared (IR) spectroscopy on vitrimers. The presence of $\text{Zn}(\text{II})$ in vitrimers is revealed by characteristic alterations on both hydroxide groups and the ester carbonyl bond structures. EXAFS analysis shows that the zinc cation is coordinated by four oxygen atoms, two of them being negatively charged, i.e. alkoxide groups to satisfy the electroneutrality whereas IR analysis demonstrates the lengthening of the ester carbonyl bond. The catalytic effect of Zn^{2+} can be stated as follows: 1) shifting the alcohol/alkoxide equilibrium towards the more nucleophilic alkoxide form; 2) activating the carbonyl bonds by increasing the electrophilicity of the carbon atom; 3) bringing reacting centres close to each other. Zn^{2+} coordinated in the epoxy-based network therefore appears as a good option to catalyze exchange reactions by transesterification and bring vitrimer properties.

Introduction

Vitrimers¹⁻¹⁵ are networks in which the topology can be changed under thermal activation while keeping constant the number of bonds and crosslinks. Consequently, even though they are crosslinked and insoluble, vitrimers are able to flow.¹ Due to this combination of properties, vitrimers represent a third class of materials, aside from thermoplastics and thermosets. This unique feature lies in their chemical structure, characterized by the presence of exchangeable bonds, which make them also different from materials based on reversible bonds (either dynamic covalent or dynamic non-covalent systems). The concept is presented in Figure 1.

Two partners, A and B joined by an exchangeable bond do not thermally dissociate. However, if there is a third individual A', chemically similar to A, there is a possibility that A' takes the place of A, thereby changing the connection topology while keeping constant the number of A-B bonds. According to this picture, several types of chemical reactions may be recognized as exchange reactions,^{1-12,14-20} the thermodynamic feature that makes vitrimers different from systems with reversible covalent bonds²¹⁻²⁹ is that temperature does not influence the number of open and closed links but only the rate of exchanges. Indeed in a system like in figure 1, the initial state (before exchange) and the final state (after exchange) have the same energy. The difference between covalent adaptable networks, based on reversible bonds, and vitrimers, based on exchangeable bonds, has been well commented.³⁰ Existing chemical systems for vitrimers have also been the subject of a comprehensive review.³¹

Exchange reactions based on photoactivated radical processes were used by Bowman et al. to control stress relaxation in

Figure 1. Basic view of an exchange reaction.

crosslinked materials.³² In vitrimer, the principle is to rely on thermoactivation and catalysis to control the rate of exchange while keeping the connectivity of the network. In first vitrimer prototypes¹⁻³ the selected exchange reaction was transesterification, which follows an associative mechanism in basic conditions.³³ Thus, as shown in Scheme 1, the nucleophilic attack of an alkoxide on the carbonyl of an ester leads to a tetrahedral intermediate that in turn re-dissociates into a new alkoxide and a new ester.

In studies published to date on transesterification based vitrimers, epoxy chemistry has been widely put to contribution due to its technological relevance and the ability to easily produce β -hydroxy esters.¹⁻⁸ Noticeably, it has been shown in

Scheme 1. The mechanism of a non catalyzed transesterification reaction in basic conditions.

these systems that the exchange kinetics can be controlled by organic^{2,5-7} or metal^{1-4,9} catalysis. More particularly zinc catalysis was selected for its superior thermal toughness in a more technologically oriented study on continuous fiber

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See DOI : 10.1039/C6PY00752J

composites and commercial epoxy-based vitrimer formulations.⁵ For the further development of epoxy vitrimers, the rate of exchange has to be carefully controlled since, once properly catalyzed, vitrimer materials might be transformable in a tailor-made temperature range.

Even though metal catalysis fundamentally implies that the reactants get close together in an adequate orientation in the metal coordination shell, the mechanism of zinc-catalyzed transesterification in epoxy vitrimers is still to be determined. In order to elucidate the action mode of metal catalysts in the transesterification reaction and eventually, to correlate vitrimer performances with the nature of the catalytic system, we investigated vitrimers as well as model compounds. The curing of epoxy resins leads essentially to insoluble and amorphous systems. We selected spectroscopic methods adapted to insoluble materials. Infrared (IR) spectroscopy was used to control the achievement of curing reactions and to study the alteration of C=O and C-O stretching vibrations of ester groups upon complexation with Zn²⁺. Attenuated total reflection (ATR) was selected in order to produce quantitative measurements from thick elastomeric samples (typically 1mm here). Extended X-ray absorption fine structure (EXAFS) spectroscopy was used to measure the coordination number and the Zn-O distances in the first coordination shell. Model zinc compounds, which crystal structure is reported, were also recorded in the same conditions.

Experimental

Materials. 3-phenoxy-1,2-propanediol (95%), anhydrous pyridine (99%), acetyl chloride (99%), bisphenol A diglycidyl ether (DGEBA, DER 332), zinc acetate dihydrate (ACS Reagent, 98%) and chloroform (99%) were purchased from Sigma Aldrich and were used as received without further purification. Pripol® 1040 was kindly provided by Croda. It is a mixture of hydrogenated cycloaddition adducts of C₁₈ fatty acids derivatives, containing about 23 wt% of C₅₄ tricarboxylic acids and 77 wt% of C₃₆ dicarboxylic acids (see supplementary information).

Preparation of fatty acid-based zinc dicarboxylate (compound 1). The synthesis of fatty acid-based zinc dicarboxylate was performed according the synthetic procedure reported previously by Montarnal et al.¹: 20 g of the fatty acids (Pripol® 1040, 296 g/mol COOH) and the Zn²⁺ precursor (Zn(OAc)₂·2H₂O) at 10 mol% concentration with respect to the COOH groups were introduced in a 100mL round bottom flask. Temperature was gradually increased from 100 °C to 180 °C while maintaining the mixture under dynamic vacuum. Evolution of acetic acid indicated that the carboxylate of fatty acids replace the acetates as ligands of Zn²⁺. The mixture was left at 180 °C under dynamic vacuum until complete dissolution of Zn²⁺ precursor and acetic acid evolution.

Vitrimer synthesis: general procedure (compound 2). In a PTFE beaker compound 1 was mixed with the epoxy resin DGEBA (174 g/mol epoxy) at an acyl to epoxy ratio equal to 1:1. Typically, we used 15.7 g of the mixture of fatty acids (52.1 mmol) and catalyst (5.2 mmol) and 9.06 g (52.1 mmol) of

DGEBA. Initially biphasic, the mixture was heated to 130 °C until phase miscibility occurred, manually stirred, and then quickly poured into a 10cm × 10cm × 0.15cm brass mold sandwiched between two anti-adhesive silicone papers. The mold was placed in a heating press and left for at least 6 h at 130 °C.

3-phenoxy-1,2-propanediol (compound 3) was purchased from Sigma Aldrich with a purity of 95%. This compound is a liquid with a boiling point of 315 °C. Our analysis revealed the presence of about 7 mol% (NMR) or 6 wt% (GC-MS) of dimerized ether derivatives (mixture of isomers) with a molecular weight of $m/z = 318$ g/mol, together with the main product (racemic mixture) with $m/z = 168$ g/mol. In GC-monitored kinetic experiments, these oligomers reproduce at longer retention times the characteristic reaction pattern of the monomer. The data of figure 2 were determined from the monomer signals solely.

Synthesis of 3-phenoxypropylene diacetate (compound 4). In a 100 mL round-bottom flask bathed in ice water, 3-phenoxy-1,2-propanediol (5 g, 29.7 mmol) and anhydrous pyridine (5.17 mg, 65.4 mmol) were placed with 20mL of chloroform and stirred during 10 min. Then acetyl chloride (5.13 g, 65.4 mmol) was slowly added dropwise. Stirring was continued for 24 hours while the water bath was allowed returning to room temperature. Then distilled water (10mL) was added and the mixture was stirred during 30min. The organic layer was washed three times with NaCl-saturated water (30mL) and distilled water (30mL), dried over anhydrous magnesium sulfate, and filtered. Then the solvent was removed using a rotary evaporator. The final product is a viscous liquid and presents MS molecular peaks at $m/z = 252$ g/mol and $m/z = 402$ g/mol. NMR indicates 97% conversion of the oligomers identified in compound 3.

Methods. GC-MS were recorded using a Shimadzu GC-2010 gas chromatograph coupled to mass spectrometry (electron ionization at 70 eV). At any reaction time, the quantities of compound 3, compound 4 and their transesterification exchange products were identified by their characteristic MS molecular peak respectively at $m/z=168$ g/mol, $M/z=252$ g/mol and $m/z = 210$ g/mol. At higher retention times, the same scenario was observed for oligomers of compound 3 and compound 4 giving rise to characteristic m/z peaks at 318 g/mol, 360 g/mol and 402 g/mol.

FT-IR spectra were recorded at 4 cm⁻¹ resolution using a Bruker TENSOR 37 spectrometer fitted with a Specac Golden Gate ATR heating cell. Compound 2 samples were dried in vacuum oven at 120 °C during 2 h before analysis. In ATR measurements the depth of penetration of the IR evanescent wave in a non-absorbing medium is a nearly linear function of the wavelength.³⁴ On the OPUS software provided by Bruker, the transmitted signal, I/I_0 is converted into ATR units via the following expression: $ATR=-\log(I/I_0).x/1000$ where x is the wavenumber in cm⁻¹. For any given band, the ATR signal thus calculated increases linearly with the amount of the corresponding vibrator.

^1H NMR spectra were recorded from CDCl_3 solutions using a Bruker Avance 400 spectrometer and referenced internally to tetramethylsilane as a standard.

EXAFS measurements were made at the Dutch-Belgian beam line (DUBBLE), at BM26A of the European Synchrotron Radiation Facility (ESRF), Grenoble, France. Zn-catalyzed epoxy-acid vitrimer plates with a thickness of 1.5mm and reference powders, sandwiched between kapton tapes, were put into the X-ray beam. Spectra were acquired at room temperature in the transmission mode using a Si(111) double crystal monochromator above the Zn K-edge (9.659 KeV). The EXAFS data analysis was processed using the IFEFFIT library of numerical XAS algorithms written in Perl programming that utilizes the ab initio EXAFS code, FEFF 6.01.³⁵ In the EXAFS fitting procedure, we used the parameters determined from the first shell analysis of anhydrous zinc acetate, keeping solely r and N as adjustable parameters. We used crystal structure data of anhydrous zinc acetate and zinc acetate dihydrate proposed respectively by Clegg et al.³⁶ and Ishioka et al.³⁷

Results

Epoxy vitrimer synthesis

The method we use here is the same as the one previously reported.¹ The starting point is a mixture of dicarboxylic and tricarboxylic fatty acids (Pripol® 1040), added in large excess to a given quantity of zinc acetate. A ligand exchange is performed as shown in scheme 2-A.¹ Removing the acetic acid thereby formed shifts the equilibrium towards the formation of salts where Zn^{2+} is coordinated by two long chain carboxylate anions. Up to a concentration of 20 mol%, such salts are soluble in the fatty acid mixture. Thus the resulting material of this ligand exchange (compound 1) is a mixture of partially neutralized (typically 5 to 10 mol%) fatty polyacids wherein the counterion is zinc.

Compound 1 is then used to cure the epoxy resin that gives

rise to the vitrimer network (compound 2). In this second step, the major reaction is the addition of a carboxylic acid to the epoxy ring to form the already mentioned β -hydroxy-ester link, constitutive of the epoxy vitrimer network (Scheme 2-B). The number of epoxy functions is adjusted in order to consume all acylating species (carboxylic acid and carboxylate functions) and to leave on average, one OH group per each ester group formed. The time dependence of $\nu_{\text{C=O}}$ stretching vibrations associated to carboxylic acid (1710 cm^{-1}), carboxylate (1632 and 1550 cm^{-1}) and ester (1737 cm^{-1}) were followed by infrared spectroscopy (depicted in supplementary information) as well as the disappearance of δ_{COC} of the epoxide (914 cm^{-1}). It is evident on spectra that after 60 min the carboxylic acid, carboxylate and epoxide signals all disappear. Hence it logically follows that Zn^{2+} in epoxy vitrimer is no longer in the form of carboxylate at the end of the curing process. Consequently beside the neutral β -hydroxy-ester groups, β -oxyester anions are necessary to ensure the electroneutrality (Scheme 2-C).

Catalysis

In previous studies,^{1,2} the catalytic action of zinc was evidenced by stress relaxation experiments of networks and GC titrations on model molecules. As a result of epoxy-acid chemistry to produce model molecules, 2-methylimidazole was often present in the systems where it could play the role of co-catalyst. Here, we examine the efficiency of Zn^{2+} as a transesterification catalyst in model molecules not generated by epoxy-acid chemistry. The exchange reaction was studied by GC-MS. From 3-phenoxy-1,2-propanediol (compound 3), we synthesized 3-phenoxypropylene diacetate (compound 4). First a 1:1 mixture of compound 3 and compound 4 was prepared and the desired quantity of zinc acetate was added. Once heated to 130°C , zinc salt rapidly dissolved and gaseous acetic acid evolution was observed. Then the mixture was continuously stirred and transesterification products appear: two β -hydroxy-ester (isomers 5' and 5'') not distinguishable by GC. The total abundance of compound 5' and compound 5'', referred as compound 5, is plotted in Figure 2 as a function of Zn^{2+} concentration. Without the addition of any metal catalyst, the equilibrium took around 75 h at 150°C whereas the equilibrium is obtained in 1h or 5h when zinc acetate is added at a level of 5 mol% or 0.05 mol% respectively. Even a small amount of zinc catalyst accelerates the transesterification significantly. The rate of transesterification is a function of the metal concentration. In previous studies on epoxy vitrimers,¹⁻³ Zn^{2+} was used at molar concentration of 1% to 10%. Our data indicate that Zn^{2+} is efficient even at low concentration and in absence of any co-catalyst. This result confirms the use of zinc as an excellent option to catalyze exchange reactions in epoxy vitrimers.

Scheme 2. Preparation of Zn-catalyzed vitrimers. **A.** First step: ligand exchange reaction between fatty polyacids and zinc diacetate; acetic acid is eliminated by vacuum stripping. **B.** Second step, major reaction: generation of β -hydroxy-ester links by epoxy-acid curing. **C.** Second step, minor reaction: generation of β -oxy-ester anions. (X_1, X_2) = (H, R_2) or (R_2 , H)

Figure 2. Scheme of transesterification of model molecules (top). Transesterification kinetics results (bottom) at 150°C for the di-ester and di-hydroxy mix model molecules. Without metal catalyst added (triangle down, green), with added 0.05 mol% Zn(OAc)₂ (triangle up, blue), 0.5 mol% Zn(OAc)₂ (circle, red) and 5 mol% Zn(OAc)₂ (square, black).

EXAFS spectroscopy

We performed extended X-ray absorption fine structure spectroscopy (EXAFS) experiments at the Zn K-edge on vitrimer and model compounds. A special care was brought to the choice of references whose atomic positions are known with accuracy. We analyzed anhydrous zinc diacetate (compound 6) and zinc diacetate dihydrate (compound 7), two well-known zinc salts whose crystal structures are available in the literature (Figure 3). X-ray powder diffraction diagrams have been recorded to check the allomorphic purity of the samples (see supplementary information). In anhydrous zinc acetate, four oxygen atoms at an average Zn-O distance of 1.96 Å surround the zinc cation.³⁶ The bidentate acetate groups in this compound are bridging ligands in a syn-anti arrangement (Figure 3a). In contrast, in zinc acetate dihydrate, Zn²⁺ is hexacoordinated with an average Zn-O distance of 2.1 Å.³⁷ In this case the bidentate acetate groups are chelating ligands as depicted in Figure 3b. We took these compounds as references in the analysis of the EXAFS spectra obtained for our model vitrimer systems. The original X-Ray absorption spectra, $\mu(E)$ plots, and the detail of the analysis are reported in the supplementary information.

In Figure 4a, we show the normalized X-ray absorption spectra $\chi(k)$ of zinc acetate dihydrate, anhydrous zinc acetate and the 10 mol% Zn-catalyzed epoxy-acid vitrimer system (compound 2) at room temperature. In this plot, it is already apparent that compound 2 and the anhydrous zinc salt have very similar $\chi(k)$ patterns whereas spectrum of zinc acetate dihydrate show quite different oscillations. In Figure 4b, we present the result of EXAFS spectra analysis in the form of the radial structure function ($A(R)$ plot) of the 10 mol% Zn-catalyzed epoxy-acid vitrimer system. In this representation, the maxima correspond to the presence of neighboring atoms at path

Figure 3. Reference compounds used in EXAFS studies. a) In compound 6: (ZnAc₂) Zn²⁺ is tetraordinated with bridging acetate ligands. b) In compound 7 (ZnAc₂•2H₂O) Zn²⁺ is hexacoordinated with bidentate acetate ligands and water.

length R . By fitting the whole $\chi(k)$ curve we determine the radial distance r and the coordination numbers N of each type of atoms in the first shell. Note that the actual distance r is systematically larger than the position of the maximum in $A(R)$ plot due to the scattering phase shift. The general form of 10 mol% Zn-catalyzed epoxy-acid vitrimer $A(R)$ plot, and in particular the position and amplitude of the first shell maximum, are similar to those of anhydrous zinc acetate whereas the whole $A(R)$ plot is significantly different from that of zinc acetate dihydrate.

The result of the first shell simulation is plotted as a dotted line in Figure 4b. We determine by fitting a first shell coordination of $N=4.3 (\pm 0.6)$ oxygen atoms and an average Zn-O distance of $r=1.98 (\pm 0.01)$ Å at room temperature. At higher temperatures, i.e. 225 °C, this distance slightly decreases to a value of $r=1.95 (\pm 0.01)$ Å while the coordination number also slightly decreases to $N=3.8 (\pm 0.5)$. These distances and coordination numbers are close to those measured in anhydrous zinc acetate confirming that Zn²⁺ in vitrimers has a similar first shell environment i.e. same coordination number and same charges. In particular, it seems clear that the zinc cation in vitrimers is tetraordinated with two neutral and

Figure 4. a) EXAFS spectra of zinc acetate dihydrate (red), anhydrous zinc acetate (blue) and the 10 mol% Zn-catalyzed epoxy-acid vitrimer system (black). b) Radial structure functions of the 10 mol% Zn-catalyzed epoxy-acid vitrimer system (solid, black) and the first shell fit associated (dot, black) .

two negatively charged oxygen atoms.

Concerning the more distant environment of the metallic cation, it is interesting to note that in bridged complexes such as compound 6, the Zn-Zn distance is of about 4-5 Å. The existence of such bridges in vitrimers is also a question to consider. Unfortunately in all our data that were essentially collected at room temperature or above, we found impossible to confirm that small peaks arising in this range, even in model compounds, actually correspond to a signal and not to noise.

Infrared spectroscopy

Epoxy-acid networks, cured in the presence of a catalyst or in absence of any catalyst have different chemical structures.³⁸⁻³⁹ Thus, to investigate vibrational changes induced by the presence of Zn²⁺, we took as control a sample containing a lower concentration of Zn²⁺ (rather than a catalyst-free one) to ensure that the structure of the network is the same. Two epoxy vitrimer networks, with respectively 5 and 10 mol% of zinc, were analyzed by infrared (IR) spectroscopy in the attenuated total reflection (ATR) mode (Figure 5a). In both the ATR-IR spectra of compound 2 (vitrimer with 10 mol% Zn catalyst) and compound 2 bis (vitrimer with 5 mol% Zn catalyst) one can recognize several characteristic bands: $\nu_{C=O}$ of esters at 1737 cm⁻¹, ν_{C-C} of aromatic fragments at 1604, 1579 and 1508 cm⁻¹, δ_{C-H} of CH₂ at 1458 cm⁻¹, δ_{C-H} of C(CH₃)₂ at 1378 and 1363 cm⁻¹, γ of CH₂ at 1292 cm⁻¹, several stretching vibrations of the C-O single bonds ν_{C-O} between 1276 and 1018 cm⁻¹ and C-H deformation of aromatic at 827 cm⁻¹. To distinguish the characteristic vibrations associated with the presence of the metal, from those of the matrix, we also plotted in figure 5b the difference spectrum $ATR(10\%) - ATR(5\%)$. In this plot, a positive (resp. negative) ΔATR signal for a given wavenumber indicates an increase (resp. a decrease) of the corresponding vibrator concentration. On the difference spectrum, basically five characteristic features appear: a strong negative band at 1737 cm⁻¹, a large negative pattern between

Figure 5. a) ATR-IR spectra of 5 mol% Zn-catalyzed epoxy vitrimer (red) and 10 mol% Zn-catalyzed vitrimer (black). b) ATR-IR spectrum difference $A(10\%) - A(5\%)$.

1300 and 1100 cm⁻¹ with a minimum at 1167 cm⁻¹ and three positive signals centered at 1569, 1436 and 1026 cm⁻¹. These observations show that the presence of the metal modifies the environment of C-O and C=O vibrators. The negative bands correspond respectively to standard $\nu_{C=O}$ and ν_{C-O} stretching vibrations of organic oxygenated compounds. In contrast, the two positives bands centered at 1569 and 1436 cm⁻¹ are in the intermediate zone between $\nu_{C=O}$ and ν_{C-O} stretching vibration wavenumbers and the positive band centered at 1026 cm⁻¹ is at the edge of the classical ν_{C-O} stretching vibration region while all of them are clearly different from all signals of starting materials (see supporting information).

Discussion

Zinc being a transition metal, it comes naturally that neutral and negatively charged oxygen atoms are present in its first coordination sphere, i.e. attached directly to the metal, as clearly observed in EXAFS spectrum and IR spectrum. The structures of oxygenated Zn(II) complexes are reported in the literature.^{36,37,40-44} Zinc is usually tetraordinated, pentacoordinated or hexacoordinated, but the tetrahedral geometry is the most frequently encountered. From EXAFS measurements, we established that the coordination number of Zn²⁺ in the epoxy-acid vitrimer system is very close to 4. The positive band centered at 1026 cm⁻¹, observed in the IR difference spectrum, is characteristic of the vibration ν_{C-O-M} from a metallic alkoxide.⁴⁴⁻⁴⁶, represented as type III complex in figure 6. This confirms the presence of Zn-bound alkoxide groups, already inferred from the reaction sequence exposed scheme 2-C. The Zn-O mean distance in the epoxy-acid vitrimer system is the same as in anhydrous zinc acetate. It suggests a similar environment of oxygen atoms, i.e. two neutral oxygen atoms and two negatively charged oxygen atoms. The characteristic C-O and C=O vibrations of neutral oxygen atoms modified by zinc presence appear in the IR spectrum of figure 5b.

New bands observed in the IR subtraction spectrum differ completely from the bands of the carboxylate observed in compound 4. The two negative bands correspond to the stretching frequencies of the ester group at 1737 cm⁻¹ for C=O and in the 1100-1300 cm⁻¹ range for C-O. With the addition of zinc, these two bands diminish and two positive bands appear. The new stretching frequencies of the ester group in the presence of zinc fall outside the tabulated ranges of a free or

Figure 6. Possible complexation modes of a β -oxy ester group with a metal M. Experimental data suggest the presence of type I and type III complexation.

H-bonded ester ($1750\text{--}1735\text{ cm}^{-1}$ for C=O, $1300\text{--}1000\text{ cm}^{-1}$ for C-O). The important shifts of these bands suggest that the ester group is involved in a complexation. Metal complexes wherein the ligand is an ester have been widely reported in the literature.^{47–54} The two possible complexation modes of the ester group with a metal (type I and II) are presented figure 6. The C=O stretching frequency, as low as 1569 cm^{-1} , reveals a decrease of the bond order, consistent with a partial electron transfer onto an oxygen-metal bond, the metal cation having a withdrawing effect thus strongly support a complexation of type I. In the literature metal complexes wherein the ligand is an ester always induce a 50 to 150 cm^{-1} shift of the C=O band towards lower frequencies. Oppositely, the withdrawing effect of the metal cation shortens the C-O bond whose stretching mode is shifted to higher frequencies. This attribution, confirmed by computational simulations (see supporting information), reveals the presence of Zn^{2+} complexes of type I thus, interpretation as type I and type III complexes, supported by data seem the most appropriate. Type II was clearly not observed but we cannot exclude that other categories of sp^3 oxygen atoms (ether and hydroxyle) could be present in the first shell of Zn^{2+} . Hydroxyles in particular are certainly present. Indeed β -diol structures (like compound 3) are produced by transesterification and the conjugated base, a β -hydroxy-alkoxide anion is very likely to form a chelate with Zn^{2+} (see SI).

Thus, the surrounding of Zn^{2+} in the epoxy-acid vitrimer system as revealed by the above analysis can be represented, as tentatively done in Figure 7. Geometrically the best first shell approximation is the structure already shown in Figure 3a for compound 6: two negatively charged oxygen atoms from alkoxides, and two neutral oxygen atoms of sp^2 or sp^3 type are placed in the immediate neighborhood of the zinc atom. The sp^2 oxygen atoms from carbonyl get activated by electron withdrawing effect towards Zn^{2+} . If trying to estimate the distances from the data of compound 6, we find at equilibrium an interatomic distance of about $3.5\text{ \AA}$ between the carbonyl sp^2 carbon and the alkoxide oxygen atoms. Thus the exchange reaction is very likely to occur between these two sites whenever Zn^{2+} complexation bring them close to one another. Therefore, the action of Zn^{2+} , as transesterification catalyst, can be summarized as follows: 1) by shifting the acid-base alcohol/alkoxide balance toward the more nucleophilic species

alkoxide, 2) by activating the carbonyl via C=O link polarization, increasing the partial charge of the sp^3 carbon atom and 3) by bringing reacting centers close to each other.

Last, bringing reacting centers close to each other is the intrinsic role of transition metal catalysts ruled by their coordination number. Contrary to alkali or earth-alkaline metal catalysts, coordination bonds of transition metals are directional. Ligand positions in the coordination sphere are thus fixed and determine their proper orientation to react. The feature revealed here is that Zn^{2+} is chemically bonded to the vitrimer backbone, without intervention of another ligand. Thus, even though the coordination number does not exceed 4, there is probability to find exchangeable functions, alkoxide and ester, in the immediate environment of the zinc cation.

In the literature, binuclear zinc complexes are described,^{51,55,56} and frequently observed in natural^{55,57,58} and synthetic catalytic processes.^{51,54,56,59} Coates group showed that not only bimetallic complexes are more stable but monometallic complexes are unreactive during the copolymerization of epoxides and CO_2 .⁶⁰ According to their observations, zinc centers do not act independently but rather in a cooperative bimetallic mode. Last, Hammes et al. observed that zinc catalyzed transesterification did not occur when the complex was mononuclear.⁵¹ The existence of such structure in vitrimer is an open question but the present data cannot resolve this point.

Conclusion

Metal-catalyzed transesterification is a key reaction in polymer chemistry. Extensively used to make polyesters or seen as undesirable side reaction in ring-opening polymerizations,⁶¹ it was also recently recognized as an original route towards sequence-controlled copolymers.⁶² In this article, the catalytic role and the action modes of the zinc cation in epoxy-acid vitrimer networks has been investigated. Kinetic study revealed that zinc catalyzed transesterification reaction even at low concentration. EXAFS and IR studies revealed that zinc cations are tetracoordinated by oxygen atoms of the β -hydroxy-ester groups inherited from the epoxy-acid chemistry. Thus, Zn^{2+} is structurally bonded to the vitrimer network and this fact should be taken into account in the future for understanding the stress relaxation mechanism. Indeed, as the catalyst is not present everywhere, it has to diffuse to activate reactions throughout the sample. Chemical factors are analyzed here. The conclusions suggest that physical factors like the diffusivity of coordinated Zn^{2+} , not yet investigated could play a prominent role in the overall efficiency of the catalyst. As for chemical factors, Zn^{2+} acts as a catalyst by activating ester carbonyl, stabilizing alkoxide groups and moving them close to each other. It is known for organic catalysts that simultaneous activation of ester and alcohol strongly increases the rate of transesterification.⁶³ So apart the fact zinc is an environmentally friendly metallic atom, also present in many natural processes, it appears as a very attractive multifunctional option to catalyze exchange reactions in epoxy vitrimers.

Figure 7. The two main actions of zinc as a catalyst in epoxy-acid vitrimer network: mediation effect and ester activation designed from EXAFS and IR spectroscopy observations.

Acknowledgements

We thank L. Leibler for inspiration of this work and many advices. We gratefully acknowledge helpful discussions with J.G.P. Goossens, from Eindhoven University of Technology. We acknowledge ESRF (European Synchrotron Radiation Facility) for provision of X-Ray beam time and thank Dr. D. Banerjee for his help on Dutch-Belgian Beamline (DUBBLE). We acknowledge D. Montarnal and R. Fournier for preliminary work. S.-J. Mougner acknowledges the financial support from Arkema.

References

- 1 D. Montarnal, M. Capelot, F. Tournilhac, L. Leibler, *Science*, 2011, **334**, 965.
- 2 M. Capelot, M. Unterlass, F. Tournilhac, L. Leibler, *ACS Macro Lett.*, 2012, **1**, 789.
- 3 M. Capelot, D. Montarnal, F. Tournilhac, L. Leibler, *J. Am. Chem. Soc.*, 2012, **134**, 7664.
- 4 E. Chabert, J. Vial, J.P. Cauchois, M. Mihaluta, F. Tournilhac, *Soft Matter*, 2016, **12**, 4838.
- 5 Z. Y. Pei, Y. Yang, Q. Chen, E.M. Terentjev, Y. Wei, Y. Ji, *Nat. Mater.*, 2014, **13**, 36.
- 6 Y. Yang, Z. Pei, X. Zhang, L. Tao, Y. Wei, Y. Ji, *Chem. Sci.*, 2014, **5**, 3486.
- 7 Z. Pei, Y. Yang, Q. Chen, Y. Wei, Y. Ji, *Adv. Mater.*, 2015, **28**, 156.
- 8 F.I. Altuna, V. Pettarin, R.J. Williams, *Green Chem.*, 2013, **15**, 3360.
- 9 J.P. Brutman, P.A. Delgado, M.A. Hillmyer, *ACS Macro Lett.*, 2014, **3**, 607.
- 10 D.J. Fortman, J.P. Brutman, C.J. Cramer, M.A. Hillmyer, W.R. Dichtel, *J. Am. Chem. Soc.*, 2015, **137**, 14019.
- 11 Y.X. Lu, F. Tournilhac, L. Leibler, Z.J. Guan, *Am. Chem. Soc.*, 2012, **134**, 8424.
- 12 W. Denissen, G. Rivero, R. Nicolaÿ, L. Leibler, J.M. Winne, F.E. Du Prez, *Advanced Functional Materials*, 2015, **25**, 2451.
- 13 F. Romano, F. Sciortino, *Phys Rev Letters*, 2015, **114**, 78104.
- 14 F. Smalenburg, L. Leibler, F. Sciortino, *Phys. Rev. Letters*, 2013, **111**, 188002.
- 15 P. Zheng, T.J. McCarthy, *J. Am. Chem. Soc.*, 2012, **134**, 2024.
- 16 A. Rekondo, R. Martin, A. Ruiz de Luzuriaga, G. Cabanero, H.J. Grande, I. Odriozola, *Mater. Horiz.*, 2014, **1**, 237.
- 17 A.R. de Luzuriaga, R. Martin, N. Markaide, A. Rekondo, G. Cabañero, J. Rodríguez, I. Odriozola, *Mater. Horiz.*, 2016, DOI: 10.1039/C6MH00029K.
- 18 M. Pepels, I. Filot, B. Klumperman, H. Goossens, *Polym. Chem.*, 2013, **4**, 4955.
- 19 J. Canadell, H. Goossens, B. Klumperman, *Macromolecules*, 2011, **44**, 2536.
- 20 M.M. Obadia, B.P. Mudraboyina, A.S., D. Montarnal, E. Drockenmuller, *J. Am. Chem. Soc.*, 2015, **137**, 6078.
- 21 W.G. Skene, J.M. Lehn, *P. Proc. Natl. Acad. Sci. USA*, 2004, **101**, 8270.
- 22 X. Chen, M.A. Dam, K. Ono, A. Mal, H. Shen, S.R. Nutt, K. Sheran, F. Wudl, *Science*, 2002, **295**, 1698.
- 23 R. Nicolaÿ, J. Kamada, A. Van Wassen, K. Matyjaszewski, *Macromolecules*, 2010, **43**, 4355.
- 24 Y. Amamoto, H. Otsuka, A. Takahara, K. Matyjaszewski, *Adv. Mater.*, 2012, **24**, 3975.
- 25 H. Ying, Y. Zhang, J. Cheng, *Nat. Commun.*, 2014, **5**, 3218.
- 26 S. Telitel, Y. Amamoto, J. Poly, F. Morlet-Savary, O. Soppera, J. Lalevee, K. Matyjaszewski, *Polym. Chem.*, 2014, **5**, 921.
- 27 C.R. Fenoli, J.W. Wydra, C.N. Bowman, *Macromolecules*, 2014, **47**, 907.
- 28 C. Yuan, M.Q. Zhang, M.Z. Rong, *J. Mater. Chem. A*, 2014, **2**, 6558.
- 29 K. Imato, T. Ohishi, M. Nishihara, A. Takahara, H. Otsuka, *J. Am. Chem. Soc.*, 2014, **136**, 11839.
- 30 C.N. Bowman, C.J. Kloxin, *Angewandte Chemie International Edition*, 2012, **51**, 4272.
- 31 W. Denissen, J.M. Winne, F.E. Du Prez, *Chemical Science*, 2016, **7**, 30.
- 32 T.F. Scott, A.D., Schneider, W.D. Cook, C.N. Bowman, *Science* 2005, **308**, 1615.
- 33 R.V. Kudryavtsev, D.N. Kursanov, *Zhurnal Obshchei Khimii*, 1957, **27**, 1686.
- 34 N.J. Harrick, *Phys. Rev. Lett.*, 1960, **4**, 224.
- 35 B. Ravel, M. Newville, *J. Synchrotron Rad.*, 2005, **12**, 537.
- 36 W. Clegg, I. Little, B.P. Straughan, *Acta Crystallogr. Sect. C*, 1986, **42**, 1701.
- 37 T. Ishioka, Y. Shibata, M. Takahashi, I. Kanesaka, Y. Kitagawa T. Nakamura, *Spectroc. Acta A*, 1998, **54**, 1827.
- 38 Montarnal, D.; Tournilhac, F.; Hidalgo, M.; Leibler, L. *J. Polym. Sci. Part Polym. Chem.* 2010, **48**, 1133.
- 39 Sordo, F.; Mougner, S.-J.; Loureiro, N.; Tournilhac, F.; Michaud, V. *Macromolecules* 2015, **48**, 4394.
- 40 J.N. van Niekerk, F.R.L. Schoening, J.H. Talbot, *Acta Crystallogr.*, 1953, **6**, 720.
- 41 T. Ishioka, Y. Shibata, M. Takahashi, I. Kanesaka, *Spectroc. Acta A*, 1998, **54**, 1811.
- 42 Y. Marcus, *Chem. Rev.*, 1988, **88**, 1475.
- 43 I.L. Alberts, K. Nadassy, S.J. Wodak, *Protein Sci.*, 1998, **7**, 1700.
- 44 H.D. Kaesz, F.G.A. Stone, *Spectrochim. Acta*, 1959, **15**, 360.
- 45 F. K. Butcher, W. Gerrard, E.F. Mooney, R.G. Rees, H.A. Willis, *Spectrochim. Acta*, 1964, **20**, 51.
- 46 C.T. Lynch, K.S. Mazdiasni, J.S. Smith, W.J. Crawford, *Anal. Chem.*, 1964, **36**, 2332.
- 47 D.S. Bytsov, *J. Struct. Chem.*, 1964, **4**, 638.
- 48 M.F. Lappert, *J. Chem. Soc. (Resumed)*, 1962, **542**.
- 49 D.S. Bytsov, *J. Struct. Chem.* 1963, **4**, 501.
- 50 M.F. Lappert, *J. Chem. Soc. (Resumed)*, 1961, **817**.
- 51 B.S. Hammes, X. Luo, M.W. Carrano, C. Carrano, *J. Inorg. Chim. Acta*, 2002, **341**, 33.
- 52 I. Sóvágó, C. Kállay, K. Várnagy, *Coord. Chem. Rev.*, 2012, **256**, 2225.
- 53 B. Schreiner, W.Z. Beck, *Anorg. Allg. Chem.*, 2010, **636**, 499.
- 54 B.S. Hammes, X. Luo, B.S. Chohan, M.W. Carrano, C.J. Carrano, *J. Chem. Soc., Dalton Trans.*, 2002, **17**, 3374.
- 55 D.E. Wilcox, *Chem. Rev.*, 1996, **96**, 2435.
- 56 A.M. Baruah, A. Karmakar, J.B. Baruah, *Inorg. Chim. Acta*, 2008, **361**, 2777.
- 57 J.A. Cricco, E.G. Orellano, R.M. Rasia, E.A. Ceccarelli, A.J. Vila, A. J., *Coord. Chem. Rev.*, 1999, **190**, 519.
- 58 N. Sträter, W.N. Lipscomb, T. Klabunde, B. Krebs, *Angew. Chem. Int. Ed. Engl.*, 1996, **35**, 2024.
- 59 S.K. Papageorgiou, E.P. Kouvelos, E.P. Favvas, A.A. Sapolidis, G.E. Romanos, F.K. Katsaros, *Carbohydr. Res.*, 2010, **345**, 469.
- 60 D.R. Moore, M. Cheng, E.B. Lobkovsky, G.W. Coates, *J. Am. Chem. Soc.*, 2003, **125**, 11911.
- 61 M. Bouyahyi, R. Duchateau, *Macromolecules*, 2014, **47**, 517.
- 62 K. Nakatani, Y. Koda, T. Terashima, and M. Sawamoto, *J. Am. Chem. Soc.*, 2012, **134**, 4373.
- 63 J. Kadota, D. Pavlović, J.-P. Desvergne, B. Bibal, F. Peruch, A. Deffieux, *Macromolecules*, 2010, **43**, 8874.

Coordination and Catalysis of Zn^{2+} in Epoxy-Based Vitrimers

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Network-coordinated Zn^{2+} ions activate carbonyls, promote alkoxides and hold reactants close to each other for transesterification.