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# Environmental Polymer Degradation: Using the Distonic Radical Ion Approach to Study the Gas-Phase Reactions of Model Polyester Radicals

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# Abstract

A novel precursor to the distonic *O*- and *C*-centred radical cations  $Oxo^+O\bullet$  and  $Oxo^+C\bullet$  was designed and synthesized, which represent model systems for radicals produced during polyester degradation. The precursor is equipped with a nitrate functional group, which serves as a masked site for these alkoxyl and carbon radicals that are unleashed through collision-induced dissociation (CID).  $Oxo^+O\bullet$  and  $Oxo^+C\bullet$  feature a cyclic carboxonium ion as permanent charge tag to enable monitoring their ion-molecule reactions on the millisecond to second timescale in the ion trap of the mass spectrometer. The reactions of  $Oxo^+O\bullet$  and  $Oxo^+C\bullet$  with cyclohexene,

cyclohexadiene, ethyl acetate, 1,1-dimethoxyethane and 1,2-dimethoxyethane, which exhibit structural features present in both intact and defective polyesters, were explored through product and kinetic studies to identify 'hot spots' for radical-induced damage in polyesters. All reactions with  $Oxo^+O$ • were extremely fast and proceeded predominantly through HAT.  $Oxo^+C$ • was about two orders of magnitude less reactive and did not noticeably damage aliphatic ester moieties through hydrogen abstraction on the timescale of our experiments. Radical addition to alkene  $\pi$  systems was identified as an important pathway for *C*-radicals, which needs to be included in polymer degradation mechanisms.

# Introduction

Synthetic polymers are amongst the most widely used materials due to their extremely diverse properties, which are controlled by their composition, the synthetic and processing techniques as well as their chain length, in conjunction with any secondary bonding interactions between polymer molecules. Environmental exposure of polymers leads to dramatic changes of their properties resulting in decreased service life and limited usage.

Environmental polymer degradation can occur hydrolytically, thermally, and by UV-light.<sup>1</sup> Both thermal and photochemical degradation can proceed through radical pathways. Key intermediates are polymer-derived peroxyl radicals, ROO•, resulting from reaction of a carboncentred polymer radical, R• (which can be formed, for example, through mechanical stress),<sup>2</sup> with oxygen, O<sub>2</sub>. ROO• are believed to act as carrier in a radical chain process that propagates damage through abstraction of a hydrogen atom from a neighbouring polymer strand to form a hydroperoxide, ROOH, and a new polymer radical R•. Although this 'autoxidation' mechanism

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was originally only proposed for the degradation of rubber,<sup>3-6</sup> it has been generally adapted to rationalize radical degradation in any polymer, irrespective of their structure. However, a recent high-level computational study clearly showed that this radical propagation is energetically unfavourable in polymers without activated C–H bonds,<sup>7</sup> which suggests that important pathways in environmental polymer degradation remain unrecognized. In fact, early degradation products, which could shed light onto the mechanism(s), have been rarely identified.<sup>8-10</sup>

Since reactions between O<sub>2</sub> and polymer radicals R• occur at the interface between gas-phase and polymer surface, gas-phase physical organic chemistry studies should provide valuable insights into reactive intermediates associated with the formation of early degradation products. A general way of investigating gas-phase radical reactions is by taking advantage of the multistage mass spectrometry capability of ion traps to generate distonic radical ions and then study their reactivity.<sup>11</sup> The chemistry of any charged species in the reaction system can be directly monitored in a time resolved fashion to obtain kinetic data for individual reaction steps. This approach has been used to investigate radical reactions of relevance to numerous fields, for example organic synthesis,<sup>12-13</sup> atmospheric chemistry,<sup>14-18</sup> and biological chemistry.<sup>19,20</sup> We have recently employed this methodology to study the reaction of distonic *N*-methylpyridinium peroxyl radical cations (Pyr<sup>+</sup>OO•),<sup>12,13</sup> which were used as a simplified model for ROO•, with various substrates that exhibit typical structural motifs in intact or defective polyesters.<sup>21</sup> Scheme 1 outlines the reaction of Pyr<sup>+</sup>OO• with cyclohexene, which shows that allylic hydrogen atom transfer (HAT; pathway (a)) is only a minor pathway.

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The major reaction channel involves addition of Pyr<sup>+</sup>OO• to the alkene  $\pi$  system, followed by rapid homolytic O–O bond fragmentation to yield an epoxide and Pyr<sup>+</sup>O• (pathway (b)), which is a reactive oxygen-centred radical species (RO•) that reacts rapidly through HAT and radical addition.<sup>21</sup> Formation of highly reactive RO• through reaction of ROO• with  $\pi$  systems in alkenes has not been considered as pathway in radical polymer degradation so far. This not only highlights the deficiencies of the current mechanistic rationale, but also demonstrates the power of the distonic radical ion approach, which allows the discovery of new, previously unidentified degradation pathways.

In this work, we have designed and synthesised the novel distonic radical cation precursor (1) to perform a mass spectrometric study of the reaction kinetics and products of two aliphatic alkoxyl and alkyl radicals, which mimic important radical intermediates formed during polyester degradation. This precursor contains glycerol monobenzoate as simplified model for the phthalate structures typically seen in polyesters, and an *N*-protonated glycine moiety, which acts as a temporary charge tag to enable its isolation in the ion trap of the mass spectrometer (Scheme 2).

**Scheme 2.** Generation of the distonic radical cations  $Oxo^+O\bullet$  and  $Oxo^+C\bullet$  in the mass spectrometer and the ion-molecule reactions studied in this work.



The nitrate group in **1** serves as a masked radical site that is unleashed through collisioninduced dissociation (CID) of the labile O–NO<sub>2</sub> bond.<sup>22,23</sup> Upon CID, first loss of glycine occurs through a neighbouring group reaction<sup>24,25</sup> to produce nitrate **2** at m/z 224, which features a cyclic tertiary carboxonium ion<sup>26-28</sup> as the fixed charge tag. A second stage of CID generates the desired alkoxyl radical cation Oxo<sup>+</sup>O• at m/z 178. Subsequent β-fragmentation with elimination of formaldehyde produces the *C*-centred radical cation Oxo<sup>+</sup>C• at m/z 148.

The chemistry of  $Oxo^+O\bullet$  and  $Oxo^+C\bullet$  was explored through their ion-molecule reactions with cyclohexene (CH), cyclohexadiene (CHD), ethyl acetate (EA), 1,1-dimethoxyethane (1,1-DME) and 1,2-dimethoxyethane (1,2-DME), which exhibit important structural features present in both intact and defective polyesters and are sufficiently volatile model systems to be studied in the gas phase. In particular, CH and CHD provide detailed insight into the reactivity of alkene  $\pi$  systems

that could be formed through thermal decomposition of polyesters via ester pyrolysis during the manufacturing process.<sup>29</sup> EA enables exploration of the reactivity of aliphatic ester linkages towards radical damage, whereas 1,1- and 1,2-DME represent different glycol motifs. For selected reactions, the experiments were augmented by density functional theory (DFT) calculations of the potential energy surface to identify the regioselectivity.

#### **Experimental Methods**

**Synthesis of starting materials.** Reactions were monitored using thin layer chromatography (tlc) on commercial silica gel 60 aluminium-backed plates coated with fluorescent indicator F254 (Merck). Plates were visualised using UV light (254 nm) or in conjunction with ninhydrin-, potassium permanganate- or phosphomolybdic acid-based stains. Purification by silica gel chromatography was performed using Davisil Chromatographic Silica Media LC60A 40-63 micron. NMR spectra were recorded on a Varian INOVA 400 instrument, with operating frequencies of 400 MHz for <sup>1</sup>H NMR and 101 MHz for <sup>13</sup>C NMR. Chemical shifts (δ) are in ppm, with residual undeuterated solvent peaks used as an internal reference. High resolution mass spectra (HRMS) were collected via electrospray ionisation (ESI) mass spectrometry, using either a Thermo Scientific Exactive Plus Orbitrap mass spectrometer or a Thermo hybrid LTQ-FTICR mass spectrometer (Thermo Scientific, Bremen, Germany).

The precursor **1** was synthesized as trifluoroacetate (TFA) salt according to Scheme 3. The NMR spectra are shown in the Supporting Information (SI).





*1-Benzoyl-3-bromo-rac-glycerol (6).* The procedure reported by Dawe *et al.*<sup>30</sup> was followed to synthesise Li<sub>2</sub>NiBr<sub>4</sub> as a 0.4 M solution in tetrahydrofuran. Epoxide  $5^{31}$  (24 mmol) was dissolved in tetrahydrofuran (71 mL) under nitrogen and cooled to 0°C, followed by dropwise addition of the Li<sub>2</sub>NiBr<sub>4</sub> solution (76 mL, ~30 mmol). Consumption of the epoxide was monitored by tlc and complete after 35 mins. The reaction mixture was then treated with phosphate buffer (110 mL, pH 7) and extracted with dichloromethane, dried and concentrated *in vacuo*. The crude product was purified by column chromatography (petroleum spirits:ethyl acetate 3:1,  $R_f = 0.34$ ) to provide **6** as a white powder (82%). <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>):  $\delta = 8.08 - 8.02$  (m, 2H, aromatic), 7.62 – 7.56 (m, 1H), 7.49 – 7.42 (m, 2H), 4.53 – 4.43 (m, 2H), 4.21 (h, <sup>3</sup>J = 5.2 Hz,

1H), 3.61 (dd,  ${}^{3}J = 10.5$ , 4.9 Hz, 1H), 3.55 (dd,  ${}^{3}J = 10.6$ , 5.9 Hz, 1H), 2.66 (s, 1H);  ${}^{13}C$  NMR (101 MHz, CDCl<sub>3</sub>):  $\delta = 166.58$ , 133.40, 129.72, 129.43, 128.49, 69.41, 66.32, 35.08; HRMS (ESI, *m/z*): [M+H]<sup>+</sup> calcd for C<sub>10</sub>H<sub>12</sub>O<sub>3</sub>Br, 258.99643 and 260.99448; found 258.99639 and 260.99436.

2-Hydroxy-3-nitrooxypropyl benzoate (7). **6** (6.5 mmol) was suspended in dry acetonitrile (12 mL) under nitrogen along with silver nitrate (12 mmol). The mixture was kept at 50°C in the dark for 7 d, with regular filtration to remove the silver bromide precipitate. Once complete, the mixture was cooled to room temperature and filtered into saturated brine (40 mL). The resulting silver chloride precipitate was separated and the filtrate extracted with ethyl acetate, dried (magnesium sulfate), and concentrated *in vacuo*. The crude material was purified by column chromatography (petroleum spirits:ethyl acetate 1:1,  $R_f = 0.65$ ) to provide **7** as a waxy solid (29%). <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>):  $\delta = 8.07 - 7.96$  (m, 2H), 7.63 - 7.53 (m, 1H), 7.49 - 7.40 (m, 2H), 5.42 - 5.36 (m, 1H), 4.87 (ddd, <sup>3</sup>J = 12.3, 3.7, 0.6 Hz, 1H), 4.78 (ddd, <sup>3</sup>J = 12.4, 6.4, 0.6 Hz, 1H), 3.99 - 3.93 (m, 1H), 3.93 - 3.87 (m, 1H), 2.26 (s, 1H); <sup>13</sup>C NMR (101 MHz, CDCl<sub>3</sub>):  $\delta = 165.87$ , 133.57, 129.79, 129.15, 128.50, 71.27, 70.66, 61.20. HRMS (ESI, *m/z*): [M+H]<sup>+</sup> calcd for C<sub>10</sub>H<sub>12</sub>NO<sub>6</sub>, 242.06591; found 242.06586; [M+Na]<sup>+</sup> calcd for C<sub>10</sub>H<sub>11</sub>NO<sub>6</sub>Na, 264.04786; found 264.04802.

2-(2-(tert-Butoxycarbonylamino)acetoxy)-3-nitrooxypropyl benzoate (8). 8 was synthesised from 7 (1 equiv.), *N*-Boc-protected glycine (1.5 equiv.), dimethylaminopyridine (DMAP; 0.1 equiv.) and *N*,*N*-dicyclohexylcarbodiimide (DCC; 1.5 equiv.), following the "*General Esterification Procedure*" (see Supporting Information). 8 was obtained as a white solid (95%), which was used without further purification. <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>):  $\delta = 8.05 - 7.98$  (m, 2H), 7.63 - 7.56 (m, 1H), 7.51 - 7.42 (m, 2H), 5.52 (p, <sup>3</sup>J = 4.9 Hz, 1H), 4.98 (s, 1H), 4.82 (dd,

 ${}^{3}J = 12.5, 3.8$  Hz, 1H), 4.69 – 4.62 (m, 1H), 4.62 – 4.56 (m, 1H), 4.46 (dd,  ${}^{3}J = 12.2, 5.3$  Hz, 1H), 3.97 (d,  ${}^{3}J = 5.8$  Hz, 2H), 1.44 (s, 9H);  ${}^{13}$ C NMR (101 MHz, CDCl<sub>3</sub>):  $\delta = 165.86, 157.38$  155.55, 133.55, 129.73, 129.04, 128.57, 79.66, 70.12, 68.74, 62.08, 42.33, 28.25. HRMS (ESI, *m/z*): [M+H]<sup>+</sup> calcd for C<sub>17</sub>H<sub>23</sub>N<sub>2</sub>O<sub>9</sub>, 399.13981; found 399.13973; [M+Na]<sup>+</sup> calcd for C<sub>17</sub>H<sub>22</sub>N<sub>2</sub>O<sub>9</sub>Na, 421.12175; found 421.12177.

2-(1-Benzoyloxy-3-nitrooxy)propan-2-yloxy-2-oxoethanaminium (1). **8** was dissolved in dichloromethane at room temperature, and cooled to 0°C. With stirring, trifluoroacetic acid (TFA; 6 equiv.) was added dropwise, and the mixture was stirred at room temperature and monitored by tlc. When consumption of **8** was complete, the solvent was evaporated under vacuum, using toluene as an azeotrope to remove any remaining acid. The resulting TFA salt was directly used in the mass spectrometric experiments without further purification. <sup>1</sup>H NMR (400 MHz, DMSO-*d*<sub>6</sub>):  $\delta = 8.23$  (s, 3H), 8.01 - 7.94 (m, 2H), 7.73 - 7.64 (m, 1H), 7.58 - 7.49 (m, 2H), 5.59 (ddd, J = 9.7, 6.5, 3.2 Hz, 1H), 4.95 (dd,  $^{3}J = 12.4$ , 3.1 Hz, 1H), 4.83 (dd,  $^{3}J = 12.4$ , 7.0 Hz, 1H), 4.57 (dd,  $^{3}J = 12.2$ , 3.8 Hz, 1H), 4.46 (dd, J = 12.3, 5.8 Hz, 1H), 3.90 (s, 2H); HRMS (ESI, m/z):  $[M+H]^+$  calcd for  $C_{12}H_{15}N_2O_7$ , 299.08738; found 299.08727.

**Mass Spectrometric Experiments.** Gas-phase studies were conducted on a Thermo Scientific (Bremen, Germany) LTQ ESI mass spectrometer, modified to enable the introduction of volatile neutral reagents into the linear ion trap to perform ion-molecule reactions (IMRs). Briefly, a suitably volatile liquid reagent was injected at a rate of  $5 - 50 \mu L \min^{-1}$  directly into the helium line supplying helium bath gas to the linear ion trap. The helium line was heated and maintained at a temperature above the boiling point of the reagent, volatilising the liquid before it reached the ion trap. The pressure regulator controlling the flow of helium into the ion trap under normal operating conditions was bypassed, with the helium pressure controlled manually to maintain an

ion gauge pressure the same as under normal operating conditions ( $0.69 \times 10^{-5}$  Torr). Further details are given in refs.<sup>12,13,32,33</sup> Ions undergoing ion-molecule reactions in the ion trap of the mass spectrometer are quasi thermalized to the temperature of the helium bath gas (298 K).<sup>32</sup> When performing IMRs, an ion of interest was isolated inside the ion trap, where it could react with the neutral. Each neutral compound used in the IMRs was purified prior to use via distillation, with purity checked by NMR.

General Mass Spectrometer Settings. Compound samples were prepared to 75-100 µM in methanol, and were injected into the ESI source at a flow rate of 5  $\mu$ L min<sup>-1</sup>. The instrument was operated in positive ion mode, with the settings tuned to optimise the signal of the initial parent ion peak at the beginning of each data collection session. ESI source conditions included needle potentials between 2.5-4.7 kV, a capillary temperature of 250°C, capillary voltages between 0.0-34.0, and tube lens voltages between 0.0-55.0 V. The Helium bath gas was adjusted to maintain an ion gauge pressure of  $0.69 \times 10^{-5}$  Torr. Ions of interest for MS<sup>n</sup> experiments were isolated with a 2-5 m/z window. The CID parameter was selected so the parent ion being fragmented was reduced to ~15% abundance. Data was collected using three micro-scans and taking between 10-100 duplicate spectra.

**Kinetic Studies.** Absolute rate coefficients for the consumption of  $Oxo^+O_{\bullet}/C_{\bullet}$  through reaction with the neutrals CH, CHD, EA, 1,1-DME and 1,2-DME according to:  $Oxo^+O_{\bullet}/C_{\bullet} + C_{\bullet}$ neutral  $\rightarrow$  products, were obtained at 298 K by monitoring the decay of the Oxo<sup>+</sup>O<sub>•</sub>/C<sub>•</sub> signal at m/z 178 and m/z 148, respectively ([M]<sup>+</sup>) as a function of reaction time under pseudo-first order conditions. Four to seven different excess concentrations of the neutral were used on time scales that allowed the peak intensity of Oxo<sup>+</sup>O<sub>•</sub>/C<sub>•</sub> to decrease to ca. 20% and to build up timeresolved decay profiles (8-17 reaction times). The pseudo-first order rate coefficient  $k_{obs}$  was

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obtained from the slope of the semilogarithmic plot  $\ln[Oxo^+O\cdot/C\cdot]$  vs. reaction time. The second-order rate coefficient *k* was determined by plotting  $k_{obs}$  vs. [neutral], see Figure S1 in the SI. Since the reactions of  $Oxo^+O\cdot$  with both CHD and 1,2-DME were extremely fast,  $k_{obs}$  could only be obtained for a narrow substrate concentration range. The second order rate coefficients were therefore determined for each concentration-time profile according to  $k_{obs}/[CHD \text{ or } 1,2-DME]$ , followed by averaging the resulting data.

The rate coefficients for product formation, i.e., Oxo<sup>+</sup>OH/CH and the radical adducts **11a**,**b** and **21a**, **b** were obtained from kinetic modelling of the experimental concentration-time profiles using the program Dynafit 4.<sup>34</sup> The concentration of the various ionic species was directly reflected by their peak areas, since mass discrimination could be excluded. The pseudo-first order rate coefficients for product formation,  $k'_{obs}$ , were fitted for the HAT pathway:  $Oxo^+O_{\bullet}/C_{\bullet}$ + neutral  $\rightarrow$  Oxo<sup>+</sup>OH/CH + [neutral-H], at m/z 179 and m/z 149, respectively, and for the  $Oxo^+O_{\bullet}/C_{\bullet}$  addition pathway:  $Oxo^+O_{\bullet}/C_{\bullet}$  + neutral  $\rightarrow$  [Oxo<sup>+</sup>O-neutral adduct], at m/z 260 (11a) and m/z 258 (11b); and  $[Oxo^+C-neutral adduct]$  at m/z 230 (21a) and m/z 228 (21b). The second-order rate coefficient k' for formation of each individual product was obtained from plotting  $k'_{obs}$  vs. [neutral], except for the low-yielding adducts **11b** and **21b** in the reaction of CHD with  $Oxo^+O \bullet$  and  $Oxo^+C \bullet$ , respectively, where k' was determined from the average of the individual concentration-time profiles, i.e., k'obs/[CHD]. It was necessary to extend the kinetic scheme for simulating the rate coefficients for formation of the products Oxo<sup>+</sup>CH and [Oxo<sup>+</sup>C– neutral adduct]• by the reaction:  $Oxo^+C \bullet \to PhC \equiv O^+$ , in order to account for the competing unimolecular decomposition of  $Oxo^+C\bullet$ . The data are shown in Figures S2 – S5 in the SI.

**Computational Studies.** DFT calculations were carried out utilizing the Gaussian 09 program<sup>35</sup> at the M062X/6-31+G\* level of theory. All ground and transition states were verified

by vibrational frequency analysis at the same level of theory, and all identified transition states showed only one imaginary frequency. The spin expectation value,  $\langle s^2 \rangle$ , was very close to 0.75 after spin annihilation. All energies are given as enthalpies, which have been shown to best describe chemical reactivity under low-pressure bimolecular conditions.<sup>36</sup> We also explored the BHandHLYP/6-31+G\* method, but found that it calculates higher energies for the stationary points in these gas-phase reactions than M062X/6-31+G\*. In some cases, the BHandHLYP computed transition states even indicated slow reactions, in contradiction with the experimental findings. The M062X/6-31+G\* data were considered sufficiently precise to obtain qualitative information about radical reactivity and selectivity for the different pathways. The Gaussian 09 archive entries for all M062X/6-31+G\* optimized ground and transition states are given in the SI.

#### **Results and Discussion**

### **1. Mass spectrometry experiments**

Figure 1 shows the CID mass spectrum of the distonic radical cation precursor 1 (see Experimental Methods for details). As outlined in Scheme 2, the initial CID process produces nitrate 2 at m/z 224 as the major product, which results from loss of glycine (Figure 1a). DFT calculations revealed that the positive charge in 2 is stabilized through formation of a cyclic carboxonium ion,<sup>26-28</sup> suggesting that the cleavage of the glycine moiety is facilitated by neighbouring group effects.<sup>24,25</sup> Attempts to locate a stable ground state structure with an uncyclised carbocation by DFT were not successful.



Figure 1. Mass spectra resulting from (a) CID of precursor 1 (m/z 299) to produce cyclic nitrate 2 at m/z 224; (b) CID of 2 leads to the distonic radical cations Oxo<sup>+</sup>O• (m/z 178) and Oxo<sup>+</sup>C• (m/z 148). The mass-selected precursor ion is indicated by an asterisk (\*).

Isolation of **2** in the ion trap and CID leads to cleavage of the O–NO<sub>2</sub> bond to form the alkoxyl radical Oxo<sup>+</sup>O• at m/z 178 (Figure 1b). Subsequent spontaneous or CID induced fragmentation of Oxo<sup>+</sup>O• produces the carbon-centred radical Oxo<sup>+</sup>C• (m/z 148). In addition to this, fragmentation to the benzoyl cation PhC=O<sup>+</sup> at m/z 105 also occurred, which likely results from a sequential ring-opening/dissociation process in Oxo<sup>+</sup>C• *via* an intermediate radical cation **9** (Scheme 4).

Scheme 4. Proposed mechanism of formation of  $PhC=O^+$  at m/z 105 through ringopening/fragmentation of  $Oxo^+C\bullet$ .



The minor product at m/z 123 in Figure 1b could result from a secondary reaction of PhC=O<sup>+</sup> with residual water present in the ion trap, in particular, since CID of the isolated m/z 123 species produced only PhC=O<sup>+</sup>. However, isolation of PhC=O<sup>+</sup> in the ion trap revealed no reaction even after an exposure time of ten seconds, which suggests that m/z 123 is a direct fragmentation product.

The site of the unpaired electron in  $Oxo^+O_{\bullet}$  and  $Oxo^+C_{\bullet}$  was confirmed by mass spectrometry through their reaction with  $O_2$ , using Helium bath gas that contained 0.001%  $O_2$ . Reaction with  $Oxo^+O_{\bullet}$  led to formation of a product at m/z 177 in accordance with aldehyde **10**, which results from abstraction of a  $\alpha$ -hydrogen atom (Scheme 5).<sup>37</sup>

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Scheme 5. Experimental and computational analysis of the location of the unpaired spin and charge in  $Oxo^+O\bullet$  and  $Oxo^+C\bullet$ . Spin densities and charges (in square brackets) calculated with  $M062X/6-31+G^*$ .



According to DFT calculations (see below), the unpaired electron in  $Oxo^+O_{\bullet}$  is located practically exclusively (94%) on the oxygen atom, confirming its nature as *O*-centred radical. The spin density in  $Oxo^+C_{\bullet}$  is also predominantly (89%) located on the expected carbon atom with some delocalisation onto the benzylic carbon atom and the phenyl ring. Analysis of the Mulliken charges further revealed that charge and spin are not fully separated and that the radical centre in  $Oxo^+C_{\bullet}$  and the carbon atom adjacent to the radical centre in  $Oxo^+O_{\bullet}$  exhibit a partial positive charge. It should be noted that we have also designed and synthesized a series of other precursor molecules structurally related to **1**, but in the mass spectrometer none of these underwent CID to produce distonic *O*- and *C*-centred radical cations (see SI for details). This shows that seemingly small structural differences have a profound and difficult-to-predict impact on the molecule's behaviour upon CID, making the design of suitable radical precursors a challenge.

# **1.1 Ion-molecule reactions with Oxo<sup>+</sup>O**•

The reaction of  $Oxo^+O$ • with CH was found to be very fast with the peak intensity dropping to ~20% after a maximum of 400 ms. The major pathway occurs through HAT, likely from the allylic CH<sub>2</sub> moiety, to give  $Oxo^+OH$  at *m*/*z* 179 (Figure 2a). The corresponding  $\alpha$ -cyclohexenyl radical (not shown) is not charged and cannot be detected.



**Figure 2.** Mass spectrum of the reaction of  $Oxo^+O\bullet$  with (a) CH after 280 ms ([CH] ca.  $7.0 \times 10^9$  molecules cm<sup>-3</sup>) and with (b) CHD after 180 ms ([CHD] ca.  $2.5 \times 10^9$  molecules cm<sup>-3</sup>). The mass-selected precursor ion is indicated by an asterisk (\*).

Minor formation of the addition product between  $Oxo^+O\bullet$  and CH at m/z 260 (i.e., **11a**) also occurred. The finding that aliphatic alkoxyl radicals react with alkenes with high preference through allylic HAT, rather than by addition to the  $\pi$  system, is consistent with literature data<sup>38,39</sup> and verifies our experimental set-up.

Due to the presence of highly activated bis-allylic C–H bonds, the reaction of Oxo<sup>+</sup>O• with CHD was extremely fast and only low excess [CHD] was required for the reaction to be

completed on a millisecond timescale.  $Oxo^+OH$  was formed as the major product (Figure 2b), whereas only trace amounts of the adduct **11b** (at *m*/*z* 258) resulting from addition of  $Oxo^+O\bullet$  to one of the  $\pi$  systems in CHD were observed.

In the  $Oxo^+O\bullet$  reactions with both CH and CHD formation of small amounts of by-products at m/z 148 and m/z 105 also occurred, which result from the competing unimolecular decay of  $Oxo^+O\bullet$  (see Figure 1b).

The minor by-products at m/z 155 and m/z 138 in the reaction involving CH and their lighter analogues in the reaction with CHD (m/z 153 and m/z 136) are likely formed through fragmentation of the radical adducts **11a** and **11b**, respectively, which confirms that Oxo<sup>+</sup>O• addition to the  $\pi$  system in CHD indeed occurs, but only as a very minor pathway. Scheme 6 proposes a possible mechanism initiated by 1,6- or 1,5-HAT, which transfers the radical site onto the carboxonium moiety to give **12** and **17**, respectively. Homolytic fragmentation leads to the respective isomeric radical cations **13'** and **18'**. **13'** can undergo a homolytic dissociation (pathway (a)) to release the vinyloxy cation **15** at m/z 153/155 and acyl radical **14**. Heterolytic cleavage (pathway (b)) leads to radical **16** and PhC=O<sup>+</sup>. Radical cation **18'** could rearrange through 1,5-HAT from the cyclohex(en)yl ring to give **19**, followed by heterolytic fragmentation into benzoic acid and the cationic radical fragment **20** at m/z 138/136. Thus, although formation of the *C*-centred radical adducts **11a/b** are only minor pathways in the reaction of Oxo<sup>+</sup>O• with CH and CHD, our findings clearly show that such intermediates rapidly propagate radical damage through rearrangement and fragmentation processes.

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Scheme 6. Proposed mechanism for formation of the by-products at m/z 155/153 and m/z 138/136 in the reactions of Oxo<sup>+</sup>O• with CH and CHD.



The reaction of  $Oxo^+O$ • with EA was considerably slower and required monitoring on the seconds timescale for the  $Oxo^+O$ • signal to decrease to ~20%. Formation of  $Oxo^+OH$  was the major pathway (Figure 3a), but with increasing reaction time the competing unimolecular decay  $Oxo^+O$ •  $\rightarrow Oxo^+C$ • became significant, as shown in the concentration/time profiles in Figure 3b. As will be outlined below, the product ion at *m*/*z* 177 results from the secondary reaction of  $Oxo^+C$ • with EA.



**Figure 3.** Reaction of  $Oxo^+O\bullet$  with EA ([EA] ca.  $7.6 \times 10^9$  molecules cm<sup>-3</sup>). (a) Mass spectrum after 960 ms; the mass-selected precursor ion is indicated by an asterisk (\*). (b) Concentration-time profile for  $Oxo^+O\bullet$  ( $\bullet$ , m/z = 178),  $Oxo^+OH$  ( $\circ$ , m/z = 179),  $Oxo^+C\bullet$  ( $\times$ , m/z = 148) and PhC=O<sup>+</sup> ( $\triangle$ , m/z = 105).

In contrast to this, the reaction of  $Oxo^+O$ • with 1,1-DME was very fast and required only about 450 ms for the  $Oxo^+O$ • signal to drop to ~20%. Formation of  $Oxo^+OH$  was identified as the major pathway (see Figure S6). The reaction of  $Oxo^+O$ • with 1,2-DME was even faster, and the  $Oxo^+O$ • signal dropped to < 20% in less than 150 ms at the lowest experimentally possible excess [1,2-DME].  $Oxo^+OH$  was the only product (see Figure S7). These findings clearly demonstrate the high susceptibility of aliphatic ethers to damage by alkoxyl radicals. The

regioselectivity in the reactions with EA, 1,1-DME and 1,2-DME was identified using computational studies, which will be discussed in section 3.

We also determined the absolute rate coefficients for various the ion-molecules reactions of  $Oxo^+O_{\bullet}$ , which are compiled in Table 1 (see Experimental Methods for details).

**Table 1.** Absolute second-order rate coefficients k for the reaction of  $Oxo^+O \bullet$  with the neutral model systems at 298 K.<sup>*a,b*</sup>

Neutral	$k (\text{cm}^3 \text{ molecule}^{-1} \text{ s}^{-1})$			
	Oxo <sup>+</sup> O• consumption	Oxo <sup>+</sup> OH formation	Oxo <sup>+</sup> O• addition	
СН	$(3.0\pm0.8)\times10^{-10}$	$(1.7 \pm 0.4) \times 10^{-10}$	$(1.1 \pm 0.3) \times 10^{-11} c$	
$\operatorname{CHD}^{d}$	$(1.5 \pm 0.4) \times 10^{-9}$	$(1.2 \pm 0.3) \times 10^{-9}$	$(7.0 \pm 2.1) \times 10^{-12} e$	
EA	$(9.2 \pm 2.3) \times 10^{-11}$	$(7.6 \pm 1.9) \times 10^{-11}$		
1,1-DME	$(4.6 \pm 1.2) \times 10^{-10}$	$(3.4 \pm 0.9) \times 10^{-10}$		
1,2 <b>-</b> DME <sup><i>d</i></sup>	$(1.6 \pm 0.4) \times 10^{-9}$	$(1.5 \pm 0.4) \times 10^{-9}$		

<sup>a</sup>Rate data for Oxo<sup>+</sup>O• consumption from pseudo-first order measurements; rate data for Oxo<sup>+</sup>OH formation and Oxo<sup>+</sup>O• addition were fitted using the Dynafit 4 program (see Experimental Methods). <sup>b</sup>Experimental error ( $\geq 25\%$ ) included. <sup>c</sup>Formation of product **11a** at m/z260. <sup>d</sup>Determined by averaging k obtained from individual concentration-time profiles (see Experimental Methods). <sup>*e*</sup>Formation of product **11b** at m/z 258.

The rate data determined for the consumption of Oxo<sup>+</sup>O• confirm its high reactivity, in particular with CHD and 1,2-DME, which are about 30 times higher than for EA, the least reactive substrate explored in this study. The rate coefficient for the reaction with CH is one order of magnitude lower than with CHD. This difference in reactivity can be rationalised by the presence of four bis-allylic hydrogen atoms in CHD, which are highly susceptible for HAT. Comparison of the kinetic data for product formation in the reaction of CHD revealed similar rate coefficients for Oxo<sup>+</sup>OH formation and Oxo<sup>+</sup>O• consumption, clearly confirming that HAT

is the dominant pathway ( $k_{\text{HAT}}/k_{add} \approx 170$ ). In contrast to this, radical addition is considerably more favourable in the reaction of  $\text{Oxo}^+\text{O} \cdot$  with CH, as revealed by the ratio of  $k_{\text{HAT}}/k_{add} \approx 15$ . The rate coefficients for  $\text{Oxo}^+\text{O} \cdot$  consumption and  $\text{Oxo}^+\text{OH}$  formation in the reactions involving 1,1-DME and 1,2-DME are, within error, the same. This shows that HAT is the exclusive reaction, in agreement with the findings from the product studies. The slightly lower rate coefficient for  $\text{Oxo}^+\text{OH}$  formation compared to  $\text{Oxo}^+\text{O} \cdot$  decay in the comparably slow reaction with EA can be rationalised by the fact that unimolecular fragmentation in  $\text{Oxo}^+\text{O} \cdot$  became a competitive pathway. Overall, the excellent agreement between the rate data for  $\text{Oxo}^+\text{O} \cdot$ consumption and product formation in all reactions provide evidence that there are no unidentified major sinks for  $\text{Oxo}^+\text{O} \cdot$ , for example through fragmentation with loss of the permanent charge tag, which would render these products 'invisible' by mass spectrometry.

# **1.2 Ion-molecule reactions with Oxo<sup>+</sup>C**•

 $Oxo^+C\bullet$  was considerably less reactive in its bimolecular reactions than  $Oxo^+O\bullet$ , and monitoring on the time scale of several seconds was required for the  $Oxo^+C\bullet$  signal to drop to ~20%. Under these conditions the fragmentation to PhC=O<sup>+</sup> (see Scheme 4) became an important side reaction.

In the reaction of  $Oxo^+C \cdot$  with CH the HAT product  $Oxo^+CH$  (at m/z 149) and the radical adduct **21a** (at m/z 230) were both obtained as major products (Figure 4a). However, the most abundant species at m/z 187 could be assigned to the adduct **22a**, which was formed through electrophilic addition of the fragmentation product  $PhC \equiv O^+$  to CH. This was confirmed through independent experiments where  $PhC \equiv O^+$  was isolated in the ion trap and reacted with CH.



**Figure 4.** Mass spectrum of the reaction of  $Oxo^+C \cdot$  with (a) CH after 5000 ms ([CH] ca. 8.7 × 10<sup>9</sup> molecules cm<sup>-3</sup>) and with (b) CHD after 5000 ms ([CHD] ca.  $3.3 \times 10^9$  molecules cm<sup>-3</sup>). The mass-selected precursor ion is indicated by an asterisk (\*).

 $Oxo^+CH$  was also obtained as major product in the reaction of CHD with  $Oxo^+C_{\bullet}$ , due to the higher susceptibility of the bisallylic methylene groups for HAT, whereas radical addition to give the adduct **21b** at *m*/*z* 228 was less favourable (Figure 4b). Formation of small amounts of the CHD – PhC=O<sup>+</sup> adduct **22b** at *m*/*z* 185 was also observed. No attempts were made to identify the

product at m/z 200, since it did not seem to have a 'counterpart' in the reaction of  $Oxo^+C\bullet$  with CH.

Interestingly,  $Oxo^+C\bullet$  did not react with EA through HAT to a noticeable extent on the timescale of our experiment. The only significant products were PhC=O<sup>+</sup>,  $Oxo^+COO\bullet$  (*m/z* 180; resulting from reaction of  $Oxo^+C\bullet$  with residual O<sub>2</sub> in the ion trap) and a product at *m/z* 177 (Figure 5a).



**Figure 5.** Mass spectrum of the reaction of  $Oxo^+C_{\bullet}$  with (a) EA after 5000 ms ([EA] ca. 7.6 × 10<sup>9</sup> molecules cm<sup>-3</sup>), (b) 1,1-DME after 5000 ms ([1,1-DME] ca. 7.1 × 10<sup>9</sup> molecules cm<sup>-3</sup>) and with (c) 1,2-DME after 5000 ms ([1,2-DME] ca. 5.6 × 10<sup>9</sup> molecules cm<sup>-3</sup>). The mass-selected precursor ion is indicated by an asterisk (\*).

 Trapping experiments of both  $PhC=O^+$  and  $Oxo^+COO_\bullet$  revealed that the *m/z* 177 product was not formed through reaction of EA with either of these species. It is therefore suggested that  $Oxo^+C_\bullet$  reacts with EA through a homolytic substitution, as shown in Scheme 7, leading to the closed-shell cation 23 and carboxyl radical 24 (which could subsequently decarboxylate). Future studies with isotopically labelled EA will reveal further insight into this unusual radical alkyl transfer reaction.

Scheme 7. Proposed mechanism for formation of the product at m/z 177 in the reaction of  $Oxo^+C\bullet$  with EA.



In contrast to this,  $Oxo^+C\bullet$  reacted readily with 1,1-DME through HAT and formation of  $Oxo^+CH$  (Figure 5b). In addition, small amounts of  $Oxo^+COO\bullet$  were also produced. The notable absence of PhC=O<sup>+</sup> could indicate a fast reaction with 1,2-DME that might lead to formation of the products at *m*/*z* 85 and *m*/*z* 117, which were not further identified.

The reaction of  $Oxo^+C \cdot$  with 1,2-DME was considerably faster than with the isomeric 1,1-DME and led to  $Oxo^+CH$  as major product (Figure 5c). Additional products resulted from the competing reactions of  $Oxo^+C \cdot$ , e.g. the decay to  $PhC \equiv O^+$  and formation of  $Oxo^+COO \cdot$ . The latter likely underwent a secondary HAT reaction with 1,2-DME to produce the hydroperoxide  $Oxo^+COOH$  at m/z 181, which clearly highlights the high susceptibility of ethylene glycol moieties to radical-induced damage. The decay rate data determined for the  $Oxo^+C_{\bullet}$  reactions with the various neutral model systems are about 1 - 2 orders of magnitude lower than for  $Oxo^+O_{\bullet}$  (Table 2). Similar to  $Oxo^+O_{\bullet}$ , the fastest rate coefficients were found for CHD and 1,2-DME, whereas the other neutrals are about ten times less reactive.

**Table 2.** Absolute second-order rate coefficients for the reaction of  $Oxo^+C\bullet$  with the neutral model systems at 298 K.<sup>*a,b*</sup>

	k (cm <sup>3</sup> molecule <sup>-1</sup> s <sup>-1</sup> )			
Neutral	Oxo <sup>+</sup> C• consumption	Oxo <sup>+</sup> CH formation	Oxo <sup>+</sup> C• addition	
СН	$(7.6 \pm 1.9) \times 10^{-12}$	$(3.3 \pm 0.8) \times 10^{-12}$	$(3.3 \pm 0.8) \times 10^{-12}  c$	
CHD	$(3.1 \pm 0.8) \times 10^{-11}$	$(1.8 \pm 0.4) \times 10^{-11}$	$(1.9\pm1.9) \times 10^{-12  d,e}$	
EA	$(5.7 \pm 1.4) \times 10^{-12}$	$< 9 \times 10^{-14}$	$(2.1 \pm 0.5) \times 10^{-12f}$	
1,1 <b>-DM</b> E	$(3.8 \pm 1.0) \times 10^{-12}$	$(3.5 \pm 0.9) \times 10^{-12}$		
1, <b>2-DM</b> E	$(1.6 \pm 0.4) \times 10^{-11}$	$(1.4 \pm 0.4) \times 10^{-11}$		

<sup>*a*</sup>Rate data for  $Oxo^+C_{\bullet}$  consumption from pseudo-first order measurements; rate data for  $Oxo^+CH$  formation and  $Oxo^+C_{\bullet}$  addition were fitted using the Dynafit 4 program (see Experimental Methods). <sup>*b*</sup>Experimental error ( $\geq 25\%$ ) included. <sup>*c*</sup>Formation of product **21a** at m/z 230. <sup>*d*</sup>Determined by averaging *k* obtained from individual concentration–time profiles (see Experimental Methods). <sup>*e*</sup>Formation of product **21b** at m/z 228. <sup>*f*</sup>Formation of product **23** at m/z 177 (see text).

The data show a ratio of  $k_{\text{HAT}}/k_{\text{add}} \approx 1$  for the reaction of  $\text{Oxo}^+\text{C}\bullet$  with CH, whereas for the reaction with CHD the HAT pathway is considerable more favourable ( $k_{\text{HAT}}/k_{\text{add}} \approx 10$ ). Similar to  $\text{Oxo}^+\text{O}\bullet$ , the rate coefficients for  $\text{Oxo}^+\text{C}\bullet$  consumption and  $\text{Oxo}^+\text{CH}$  formation in the reactions involving 1,1-DME and 1,2-DME, respectively, are the same, confirming that HAT is the exclusive bimolecular reaction pathway for  $\text{Oxo}^+\text{C}\bullet$ .

While the rate coefficient determined from the consumption of  $Oxo^+C\bullet$  in the reaction with EA suggests a fast reaction, this is not reflected by the rate coefficient for  $Oxo^+CH$  formation, which is lower by more than two orders of magnitude (it should be noted that the rate of  $Oxo^+CH$  formation in this reaction,  $k_{HAT}$ , is at the low end of our experimental set-up and only an upper limit can be given). As outlined in the product studies, the dominant loss channels for  $Oxo^+C\bullet$  are unimolecular decay to  $PhC=O^+$  and formation of the closed-shell compound 23 through an unusual homolytic ethyl transfer from EA, which is much faster than HAT ( $k_{HAT}/k_{transfer} < 0.04$ ; see Table 2).

# 2. Computational studies

DFT calculations were carried out to determine the most reactive site for the Oxo<sup>+</sup>O•/C• induced HAT in EA, 1,1-DME and 1,2-DME. All reactions proceed through formation of an initial ion-molecule association complex (reactant complex), which is at least 60 kJ mol<sup>-1</sup> lower in energy than the free reactants. The energy of the free reactants, or 'entrance channel', will be defined as reference point, which is set to 0 kJ mol<sup>-1</sup>. Due to the low pressure in the ion trap of the mass spectrometer, energy exchange through collisions with the surroundings can be excluded. Thus, the reactant complex has excess energy resulting from the internal and kinetic energy of the free reactants and the electrostatic energy that is released upon complex formation. Any subsequent reaction of the reactant complex is therefore only possible, if the activation barrier for this process is smaller or equal to the energy difference between the entrance channel and the association complex.<sup>40,41</sup> Reaction pathways with energies above the entrance channel cannot be accessed.

Generally, the HAT reactions involving  $Oxo^+O_{\bullet}$  are more exothermic than with  $Oxo^+C_{\bullet}$ . Figure 6 shows the calculated potential energy diagram for the reaction of  $Oxo^+O_{\bullet}$  and  $Oxo^+C_{\bullet}$  with EA. HAT from the  $\alpha$ -,  $\beta$ -, and  $\gamma$ -position and formation of the isomeric EA radicals (as product complex with  $Oxo^+OH$  or  $Oxo^+CH$ , respectively) is exothermic for both  $Oxo^+O_{\bullet}$  and  $Oxo^+C_{\bullet}$ .



**Figure 6.** Potential energy diagram for the reaction of  $Oxo^+O\bullet$  and  $Oxo^+C\bullet$  with EA. M062X/6-31+G\* enthalpies are in kJ mol<sup>-1</sup> for the reaction involving  $Oxo^+O\bullet$  (normal) and  $Oxo^+C\bullet$  (in italics).

The  $\alpha$ - and  $\beta$ -radicals are the thermodynamically preferred products, whereas formation of the  $\gamma$ -radical is least favourable. HAT from the  $\beta$ -position is the kinetically preferred pathway for  $Oxo^+O\bullet$ , which is associated with an activation barrier of 31 kJ mol<sup>-1</sup>. In contrast to this, any of the possible  $TS_{HAT}$  for the reaction with  $Oxo^+C\bullet$  are located either close to or above the entrance channel, indicating a slow reaction through HAT, in agreement with the experimental data.

The potential energy diagram for the reaction of  $Oxo^+O\bullet$  and  $Oxo^+C\bullet$  with 1,1-DME is shown in Figure 7.



**Figure 7.** Potential energy diagram for the reaction of  $Oxo^+O\bullet$  and  $Oxo^+C\bullet$  with 1,1-DME. M062X/6-31+G\* enthalpies are in kJ mol<sup>-1</sup> for the reaction involving  $Oxo^+O\bullet$  (normal) and  $Oxo^+C\bullet$  (in italics).

Formation of  $\gamma$ -1,1-DME is thermodynamically least favourable, whereas both  $\alpha$ - and  $\beta$ -1,1-DME are more stable by about 20 kJ mol<sup>-1</sup> (as product complexes with Oxo<sup>+</sup>OH/CH). Although the energies of the isomeric transition states are low for the reaction with Oxo<sup>+</sup>O•, overall the  $\beta$ -HAT is kinetically the most favourable pathway with a barrier of ca. 22 kJ mol<sup>-1</sup>. The lower reactivity of Oxo<sup>+</sup>C• leads to a higher selectivity, and a reaction occurs only through TS<sub>HAT( $\alpha$ ) and TS<sub>HAT( $\beta$ )</sub>, whereas a pathway via TS<sub>HAT( $\gamma$ )</sub> is energetically not possible.</sub>

Figure 8 presents the computed potential energy diagram for the reaction of  $Oxo^+O\bullet$  and  $Oxo^+C\bullet$  with 1,2-DME.



**Figure 8.** Potential energy diagram for the reaction of  $Oxo^+O\bullet$  and  $Oxo^+C\bullet$  with 1,2-DME. M062X/6-31+G\* enthalpies are in kJ mol<sup>-1</sup> for the reaction involving  $Oxo^+O\bullet$  (normal) and  $Oxo^+C\bullet$  (in italics).

Formation of both  $\alpha$ -1,2-DME and  $\beta$ -1,2-DME (as product complexes with Oxo<sup>+</sup>OH and Oxo<sup>+</sup>CH, respectively) is thermodynamically favourable, with the secondary radical  $\beta$ -1,2-DME being more stable by about 24 ± 2 kJ mol<sup>-1</sup>. Oxo<sup>+</sup>O• could abstract hydrogen from both  $\alpha$ - and  $\beta$ -position, which are associated with barriers of ca. 44 kJ mol<sup>-1</sup> ( $\alpha$  abstraction) and 35 kJ mol<sup>-1</sup> ( $\beta$  abstraction), respectively. In contrast to this, only  $\beta$ -1,2-DME is formed in the reaction involving

 $Oxo^+C\bullet$ , since  $TS_{HAT(\alpha)}$ , which leads to  $\alpha$ -1,2-DME, is located some 23 kJ mol<sup>-1</sup> above the entrance channel and cannot be accessed.

It should be noted that the calculations predict for  $Oxo^+O\bullet$  a faster reaction with 1,1-DME than with 1,2-DME (*via* the lowest-energy pathway along  $TS_{HAT(\beta)}$  in both cases), which seems to contradict the experimental data. However, since 1,1-DME possesses only one reactive  $\beta$ hydrogen, whereas 1,2-DME has four, the higher rate coefficient determined experimentally for the reaction of  $Oxo^+O\bullet$  with 1,2-DME likely results from a higher probability of successful encounters.

Computational studies were also employed to explore the role of the positive charge tag in  $Oxo^+O_*/C_*$  on the radical reactivity by comparing them with their neutral counterparts  $OxoO_*/C_*$ , which have a neutral dioxolane instead of the dioxonium ring. Using the reaction with CH as example, the activation barriers for both allylic HAT and radical addition to the  $\pi$  system are about 26 – 37 kJ mol<sup>-1</sup> higher for  $OxoO_*/C_*$  compared to  $Oxo^+O_*/C_*$ . The calculations also revealed that the charge tag does not significantly affect the preference for one pathway over the other (data not shown). This confirms that the distonic radical cations studied in this work are excellent model systems for *O*- and *C*-centred polymer-derived radicals that enable to study their reactions on the short timescales of ion trap mass spectrometry experiments.

# Conclusions

A novel precursor (1) to the distonic *O*- and *C*-centred radical cations  $Oxo^+O_{\bullet}$  and  $Oxo^+C_{\bullet}$  was designed and synthesized to provide model systems for radicals produced during polyester degradation. The mass spectrometric studies revealed that reactions involving  $Oxo^+O_{\bullet}$  were extremely fast. Allylic HAT was the predominant reaction pathway with CH and CHD, whereas

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radical addition to the  $\pi$  system occurred only to a very minor extent. Aliphatic esters (EA) also reacted rapidly with Oxo<sup>+</sup>O• through HAT, preferentially from the ester O-CH<sub>2</sub>- moiety, followed by CH<sub>3</sub>-C=O. The central -O-C<sub>2</sub>H<sub>4</sub>-O- structure in 1,2-DME is particularly susceptible to radical-induced damage, which is significant, since ethylene glycol moieties are constituents of many large-scale industrial polyesters, for example polyethylene terephthalate (PET).

The reactivity of  $Oxo^+C\bullet$  was about two orders of magnitude lower than that of  $Oxo^+O\bullet$ . An unprecedented alkyl transfer in the reaction of  $Oxo^+C\bullet$  with EA was discovered, which could propagate radical damage in polyesters through a novel homolytic substitution pathway.

In contrast to  $Oxo^+O$ ,  $Oxo^+C$  more readily undergoes addition to alkenes. Comparison of the rate coefficients for the competing pathways clearly shows that both allylic HAT and radical addition to alkene  $\pi$  systems need to be considered as reaction pathways of *C*-centred radicals during polymer degradation. We will in the future employ this strategy to explore the chemistry of secondary radicals, in particular the role of oxygen on the degradation mechanism in step-growth and chain-growth polymers under environmental conditions.

Supporting Information. The following files are available free of charge.

Kinetic data (Figures S1 – S5); mass spectra (Figures S6 and S7); synthetic procedures to alternative radical precursors; general synthetic procedures, <sup>1</sup>H and <sup>13</sup> NMR spectra; Gaussian 09 archive entries for Figures 6-8 (PDF).

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The manuscript was written through contributions of all authors. All authors have given approval to the final version of the manuscript.

#### Notes

The authors declare no financial interest.

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