Unexpected benzene oxidation in collisions with superoxide anions

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Superoxide anions colliding with benzene molecules at impact energies from 200 to 900 eV are reported for the first time to form massive complexes. With the aid of quantum chemistry calculations, we propose a mechanism in which a sudden double ionization of benzene and the subsequent electrostatic attraction between the dication and the anion form a stable covalently bonded $\text{C}_6\text{H}_6\text{O}_2^+$ molecule, that evolves towards the formation of benzene-diol conformers. These findings lend support to a model presenting a new high energy anion-driven chemistry as an alternative way to form complex molecules.

Benzene is one of the simplest and more stable aromatic ring molecules. It has been considered as a prototype for the study of chemical reactions involving biomolecules (pyridine1, carbamates2), a building block for the synthesis of carbon nano-cages3 and a precursor of hydrocarbon structures (phenol, toluene, aniline, anthracene) with important technological applications4. Oxygen superoxide anion ($\text{O}_2^-$) is one of the reactive oxygen species (ROS) which is responsible for several biochemical processes leading to oxidative damage in living organisms5 and materials6. In particular, 8-oxoguanine is frequently formed by the interaction of ROS with the guanine base in DNA under conditions of oxidative stress yielding an efficient way of damaging DNA7.

Oxidative processes in benzene have been extensively studied from different points of view of scientific and technical relevance, including e.g. atmospheric reactions8,9, surface catalyzed reactions10 or direct mechanisms induced by metallic oxide cations11. Most of these procedures fall into the domain of room temperature chemical reactions, i.e. no significant kinetic energy of the reactants is required to trigger such processes. Yet, if the kinetic energy involved in these collisions increases, new channels are open yielding excitation, ionization or even molecular dissociation, compatible with the transferred energy. Ascenzi et al.12 showed that oxygen-benzene reactions in atmospheric pressure plasmas (corona and dielectric barrier discharges) may induce energetically disfavored chemical processes leading to formation of phenol cations ($\text{C}_6\text{H}_5\text{OH}^+$) and neutral phenol ($\text{C}_6\text{H}_5\text{OH}$). Additionally, relevant ion chemistry processes in the interstellar medium (ISM) have been reported from low-energy anion induced reactions with different types of hydrocarbons yielding increasingly complex molecules13.

Here we show for the first time that gas-phase energetic interactions of a superoxide anion with a benzene molecule lead to an unexpectedly and quite efficient oxidation process of the neutral molecule. Under the current experimental collision energy range probed (200–900 eV), significant neutral and cationic fragmentation of benzene is reported, whilst a comprehensive description of the underlying collision dynamics mechanisms is investigated. In addition, the experimental conditions provide binary collisions between the incoming projectile ($\text{O}_2^-$) and the neutral target molecule ($\text{C}_6\text{H}_6$), where no positive ions with higher mass-to-charge ratio ($m/z$) than that of the parent ion ($m/z = 78$ u) are expected to be formed. However, as we will discuss below, the mass/charge analysis of the collision products reveals the presence of a prominent feature at $m/z \sim 110$ u for specific impact energies between 200 and 900 eV.

The remainder of this paper is organized as follows. In the next section we briefly present our experimental and theoretical methods, then the corresponding results are presented and discussed and finally some conclusions from this study are drawn in the last section.

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Methods

The experimental arrangement has been reported elsewhere\(^1\). Briefly, the primary \(\text{O}_2^-\) beam is generated in the afterglow of a pulsed hollow cathode discharge. A schematic diagram of the experimental set-up is shown in Fig. 1.

The precursor gas projectile (\(\text{O}_2\)) is admitted into vacuum through a commercial Parker pulsed valve (VAC1250) operated at 350-μs pulse-width in an 80-ms duty cycle and at a gas pressure of \(4.0 \times 10^{-5}\) mbar. It produces a supersonic expansion neutral beam that triggers the discharge when it passes along the 20 mm long hollow cathode which is negatively biased (-500 V). Positive ions initially formed in the discharge are neutralized during the plasma extinction and negative ions are finally formed by electron attachment processes. Negatively charged particles are extracted from the cathode and focused into the entrance aperture of the collision chamber (electrons are removed from the primary beam with the help of two permanent magnets placed along the beam path). The collision chamber (CC) is a 30 mm length metallic cube that contains the target \(\text{C}_6\text{H}_6\) gas at a well-known pressure (as measured with a MKS Baratron 627DX). A parallel plate system along the axis, inside the collision chamber, allows the extraction of the positive ions formed during the \(\text{O}_2^-\cdot\text{C}_6\text{H}_6\) interactions by applying a 50–900 V pulse which is synchronized with the control unit of the supersonic valve. Positive ions enter a 1.5 m drift tube and their time-of-flight (TOF) is analyzed with a digital oscilloscope (Tektronix MSO 3034, 2.5 GS/s). Ion signals are formed with a microchannel plate detector (MCP-1) in single pulse mode operation with typical counting rates below \(10^2\) s\(^{-1}\). Primary particles emerging from the CC are detected by another microchannel plate detector (MCP-2). The maximum counting rate of the primary beam was typically \(10^3\) s\(^{-1}\). The \(\text{C}_6\text{H}_6\) pressure was maintained within 0.1–1.5 mTorr. These conditions ensure a binary collision regime and multiple scattering processes are negligible. Beneath the gas cell, an electron gun provides an energy-controlled electron beam (0–1000 eV) entering the CC normal to the anion beam and opposite to the cation TOF mass analyzer. This electron beam is used to analyze the molecular composition of the background and the gas target as well as to provide a reference fragmentation pattern of the molecular target induced by electron impact\(^1\). The uncertainty on the point along the 2 cm cathode where anions are produced is spreading the arrival time distribution. This time uncertainty introduces an inherent mass resolution limitation which has been estimated to be \(\Delta m/m = 0.05\). Typical time of flight (TOF) mass spectra, of the primary anion beam (see Ref.\(^1\) for details), are shown in Fig. 2 for 700 eV. This figure reveals that the anion beam is mainly composed by two species assigned to \(\text{O}_2^-\) and \(\text{O}_3^-\). Once the extraction pulse is applied, the area of the primary beam removed from the beam (see Fig. 2a) corresponds to the colliding charged species which generated the extracted ion fragments. By proper tuning the extraction pulse delay, this area can be selected all along the primary beam profile, thus allowing the choice of the projectile to collide with the target. The time scale is energy calibrated for different anion incident energies and the energy distribution corresponding to two different target conditions (0 and 1.3 mTorr of benzene in the scattering chamber, respectively) is plotted in Fig. 2b.

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**Figure 1.** Schematics of the experimental setup. V, pulsed supersonic valve; C, hollow cathode discharge; A1 and A2, anodes (grounded); L1, L2, and L3, Einzel lenses; D1, D2, D3, and D4, deflecting plates; M1 and M2, magnets; E1, E2, E3, and E4, extraction plates; V1 and V2 voltage chamber; G2, retarding field analyzer grids; EG, electron gun; GC, gas cell; MCP 1 and MCP 2, multichannel plate detectors; QMS, quadrupole mass spectrometer; SEM, secondary electron multiplier detector; P1, P2, P3, and P4, turbomolecular pumps.
Calculations of the potential energy surfaces were performed in the frame of the density functional theory, in particular using the hybrid long-range corrected CAM-B3LYP functional in combination with the cc-pVDZ basis set. This functional shows a variable amount of HF and exchange interactions at short- and long-range and has been shown to predict exited states with better accuracy than other widely used functionals, in particular performing well for charge transfer excitations thanks to the addition of the long-range correction. We have checked the quality of the basis set by comparing absorption spectra computed with a bigger basis, aug-cc-pVTZ, obtaining energy differences for the states of interest of less than 0.1 eV. In the exploration of the potential energy surface the critical points, i.e. minima and transition states, have been accurately located. Harmonic frequencies were computed to confirm the nature of the minima and the transition states (TSs). Furthermore, intrinsic reaction coordinate calculations were carried out to verify connectivity between TSs and adjacent minima. Harmonic frequencies were also used to correct relative energies with the Zero Point Energy (ZPE). Those simulations involving electronic excited states were performed within the time dependent DFT (TD-DFT), by using the same functional and basis set. We have also carried out a Natural Transition Orbital (NTO) analysis in the relevant charge transfer excited state. Details on molecular dynamics simulations are given in the Supplementary Information. All simulations have been carried out with the Gaussian16 program.

Results and discussion
A typical TOF spectrum of the cations extracted after the interaction of $\text{O}_2^-$ is shown in Fig. 3a for 700 eV incident energy and 1.2 mTorr of benzene in the scattering chamber. The electron impact mass spectrum, generated in such conditions but with a 700 eV electron beam normal to the target sample, is also shown in this figure for comparison. Both spectra are normalized to the parent ion (C$_6$H$_6^+$, 78u) intensity. A close inspection of this figure shows that the cation fragmentation induced by the anion beam is qualitatively the same as with the electron beam for m/z smaller than 78 u. Notwithstanding the impact energy being the same in both cases, the collision time, projectile velocity and the energy transferred are quite different in both beams, thus lending support to the changes in the yields from both spectra. However, the main differences appear for generated molecular cation species larger than 78 u, where in Fig. 3b we depict the difference between the anion and electron induced fragmentation spectra normalized to the parent ion intensity. This subtraction procedure ensures that features shown in Fig. 3b are only induced by the anion beam without background contributions. In this figure, the most intense feature is found between 110 and 120 u together with other minor contributions at 94 and 99 u. In order to assign the 110–120 u structure’s composition, we have first proved that for low pressure (<1.5 mTorr) conditions, only single binary collisions prevail. In this pressure range, no dependence of the measured attenuation cross-section (total electron detachment cross-section) with the gas pressure has been observed so ensuring that multiple scattering processes are negligible. In addition, we have verified that the intensity of the 110–120 u feature remains proportional to that of the parent ion for benzene pressures ranging from 0.1 to 1.5 mTorr. Under these conditions, the only possible assignment for the ion formed after collision of $\text{O}_2^-$ and C$_6$H$_6$ with 110 u, is to consider that the projectile forms with the target a positively charged complex: C$_6$H$_6\text{O}_2^+$. Yet, the quite broad nature of the 110–120 u feature shows that it can accommodate an extra contribution (126 u in
In order to describe the underlying molecular mechanism, several electronic excited states were computed in a frozen scan on the quasi-molecule \([\text{C}_6\text{H}_6\ldots\text{O}_2]^+\). Figure 4b shows the potential energy curves of the 250 computed electronic states as a function of the benzene-oxygen distance. States below \(-24\) eV (referred to the collision \(\text{O}_2^-+\text{C}_6\text{H}_6\)) lie under the second ionization threshold of benzene.

Interestingly, we observe attractive...
curves, that correspond to a Coulomb-type attraction between the doubly positively charged benzene and the negatively charged oxygen in the entrance channel. These curves then collapse at a minimum close to ~ 2 Å. In order to further our knowledge about the collision dynamics, a more refined examination of the potential energy surfaces has been implemented. To this, we have performed relaxed scans in the ground state, where the charge is redistributed over the whole $\text{[C}_6\text{H}_6\ldots\text{O}_2]^+$ system, and in the excited state of the lowest energy showing a charge transfer character, i.e. where it still presents the nature of the entrance channel with the superoxide anion approaching the doubly ionized benzene $[\text{O}_2^– \rightarrow \text{C}_6\text{H}_6^{2+}]$. Figure 4c shows the difference in electron density distribution between these two states. A major change is appreciated with an increase of electron population of the $\text{O}_2^–$ antibonding π* molecular orbital, which is filled by an electron from the $\text{C}_6\text{H}_6$ occupied π molecular orbital. The Natural Transition Orbital (NTO) analysis shown in Fig. 5 confirms the charge transfer character of this state. The relaxed scans in both electronic states are shown in Fig. 4d, from which while in the excited state oxygen-carbon bonding appears as a barrier less process, a barrier of ~ 0.7 eV is shown in the ground state. Both calculations allow us to locate the corresponding minima in the potential energy surfaces, with strong covalent oxygen-carbon bonds stabilizing such structures, and the excited state showing charge transfer character. We can assume that non-radiative decay processes may be operative in such stabilization, since both states present similar structures and the corresponding potential energy curves are not far apart in energy, thus leading the system in the electronic ground state. It is precisely from this point where we continue our examination of the potential energy surface. Figure 4e shows the critical points, minima and transition states, in several reaction paths that lead to the production of stable diol structures (labeled as Min6 and Min8 in the figure). Energy barriers of ~ 2–3 eV in the forward direction, yielding diol structures, can be easily reached if we assume that the entrance channel is at higher energy. However, much higher energy barriers of ~ 7–8 eV prevent the diol structures to come back and to dissociate into $\text{O}_2/\text{C}_6\text{H}_6$. Thus, these are kept trapped in these potential wells. Interestingly, these two diol structures appear only at ~ 3.7 eV above the initial point before the double ionization, i.e. $\text{C}_6\text{H}_6^+ + \text{O}_2^–$. Such great stability seems to indicate that the 110 u feature assigned to $\text{C}_6\text{H}_6\text{O}_2^+$ most likely corresponds to any of these two diol compounds. Nevertheless, these structures may carry enough internal energy to continue evolving through low-energy barrier processes, e.g. with the loss of OH; this is supported by the experiment where fragments 93 u assigned to $\text{C}_6\text{H}_6\text{O}^+$ and/or 94 u to $\text{C}_6\text{H}_6\text{O}^+$ are detected as combined features (see Fig. 3b) due
to the lack of mass resolution. Complementary molecular dynamics studies on the C₆H₆O₂⁺ compound (Min1), confirm the stability of this structure, thus verifying the validity of the proposed mechanism (see details in the Supplementary Information). The formation of the complex should be a fast process taking place in a fs time scale. Considering strong coulomb attraction between doubly ionized benzene and O₂⁻, and assuming excitation energies of about 15-20 eV, a few tens of femtoseconds is enough to cleavage and form covalent bonds.

It is worth mentioning in this point that benzene dication undergoes rearrangement to a more stable pyramidal isomer with a C₅H₅ base and CH at the apex. It has been proposed that the structure is composed of a donor–acceptor bond from a C₅H₅⁻ to a CH⁺ moiety. The isomerization barrier between the dicationic canonical form of benzene and the pyramidal one is of 100 kJ·mol⁻¹, but we consider that in our case this possibility can be ruled out, since the Coulomb attraction between C₆H₆²⁺ and O₂⁻ leads to a very rapid formation of C₆H₆-O₂⁻ complex.

A careful analysis of the experimental mass spectra can provide additional information on the observed peak composition. Note that the time (mass) resolution of the present experiment (Δm/m ≈ 0.05) is limited by the gas cell configuration and the inherent characteristic of the hollow cathode discharge. Nonetheless, the decomposition of the observed peaks into single Gaussian functions, representing the different cationic fragment masses, may enlighten which ion species are contributing to the observed peaks. As mentioned above, Fig. 3b shows the difference between the anion and electron induced fragmentation spectra once normalized to the parent ion. The decompositions into Gaussian functions reveals that the m/z = 110–120 u is formed by the contribution of C₆H₆-O₂⁻ (n = 0–5) ion fragments. However, there is a clear enhancement of the peaks around 60 u and 38 u, corresponding to C₅H₅⁺ (n = 1–6) and C₃H₃⁺ (n = 0–5), respectively, as well as new weaker structures appearing below 30 u. This suggests that anion collisions are favoring further parent ion fragmentation either by losing CH₃⁺ (n = 0–4) fragments or breaking it into two similar size fragments.

For ions corresponding to m/z > 78 u, in which this study is mainly focused, we found the aforementioned m/z = 110–120 u structure together with some smaller peaks around 94 and 99 u. The analysis of these structures using the decomposition into Gaussian functions reveals that m/z = 110–120 u is formed by the contribution of two features centered at 110 and 126 u, respectively. The 110 u is assigned to C₆H₆O₂⁺ with its formation mechanism supported by theoretical calculations described above. The 126 u ion can be explained under the same formation rationale as the 110u feature but now being formed by the O₃⁻ anion, also present in the projectile beam. For ions corresponding to m/z > 78 u, double ionization of benzene has proven to be particularly likely, single ionizing processes contributing to the 78 m/z peak should be even more pronounced. Nonetheless, this can be explained within our proposed model. Note that the Coulomb attraction between the formed cation and the anion projectile in the case of single ionization will give a neutral compound (C₆H₆O₂) which will not be detected by our TOF spectrometer but will contribute to reduce the observed 78 m/z peak intensity. Thus, an alternative way for the formation of the 110–120 m/z structure can be considered if the neutral C₆H₆O₂ compound is formed in an excited autoionizing
state that finally decays to \( \text{C}_6\text{H}_6\text{O}_2^+ \), whose evolution would follow the potential energy surface shown in Fig. 4e. Note that the lifetime of this excited state will naturally contribute to form a broad structure as that shown here. In good agreement with the proposed model, the two pathways to form neutral and cation benzene diols via \( \text{O}_2^- \) collisions with \( \text{C}_6\text{H}_6 \) are summarized in Fig. 6.

**Conclusions**

In conclusion, we have experimentally shown the formation of large complex molecules (larger than the target molecule) in relatively high energy (200–900 eV) collisions of oxygen anions with neutral benzene molecules. Quantum chemical calculations have been performed within the framework of the density functional theory, to further our knowledge on the possible paths involved to generate these molecules. We have then proposed the formation of “quasi-molecular” compounds resulting from the electrostatic attraction between a doubly ionized target molecule and the negatively charged projectile. These electrostatically bonded complexes finally decay via non-radiative processes resulting in stable diol/triol structures in collisions with \( \text{O}_2^-/\text{O}_3^- \). We have also shown that an alternative way to generate these complexes is to single ionize the benzene molecule, forming the “quasi-molecular” compound in an autoionizing state of its neutral configuration which finally decays to the corresponding diol cation. We can then confirm that the present experimental evidences are supported by the theoretical calculations. This model presents a new high energy anion-driven chemistry as an alternative way to form complex molecules.

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Author contributions
PLV and GG conceived and conducted the experiment, CG, MM and AIL did the measurements, SDT and FAG did the calculations, FAG, SDT, PLV and GG analyzed the data. All the authors reviewed the manuscript.

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Competing interests
The authors declare no competing interests.
