This is an Accepted Manuscript, which has been through the Royal Society of Chemistry peer review process and has been accepted for publication.

Accepted Manuscripts are published online shortly after acceptance, before technical editing, formatting and proof reading. Using this free service, authors can make their results available to the community, in citable form, before we publish the edited article. We will replace this Accepted Manuscript with the edited and formatted Advance Article as soon as it is available.

You can find more information about Accepted Manuscripts in the Information for Authors.

Please note that technical editing may introduce minor changes to the text and/or graphics, which may alter content. The journal’s standard Terms & Conditions and the Ethical guidelines still apply. In no event shall the Royal Society of Chemistry be held responsible for any errors or omissions in this Accepted Manuscript or any consequences arising from the use of any information it contains.
1 Introduction

Fullerenes and their compounds remain among the most actively studied materials in the world of carbon nanostructures for their potential application in energy storage or conversion. In the former case, they have been successfully applied as reversible H2 solid-state storage media, in the latter, they are ubiquitously considered suitable acceptors in photoinduced electron processes because of their high electron affinity and ability to transport charge effectively. Polymeric networks of fullerenes, in particular, are gaining increasing attention after the remarkable report of superhard C60-based amorphous phase obtained under pressure, with bulk modulus exceeding that of diamond. C60 polymerization can either be photoactivated, caused by external pressure or induced by the intercalation of alkali metals that promote Coulomb interaction, thus contracting the fullerene lattice. This leads to the formation of either 1D, 2D or 3D networks of fullerenes (some examples are, respectively) whose bonding schemes are mainly driven by the electronic structure of C60 molecules. Such versatile structures are due to the different possible addition reactions: the most common network is produced in a [2+2] cycloaddition reaction driving the formation of four-membered carbon rings between adjacent fullerenes. This is normally observed in pressure- and photo-polymerized structures, while single C-C bonds are favoured when C60 LUMO is populated. Mixed architectures are also observed. This is the case of magnesium intercalated fullerene salt Mg2C60. With a relatively high magnesium stoichiometry x=4, a 2D polymer is created, where C60 molecules connect with a purely [2+2] cycloaddition bonding network. For the lower stoichiometry, x=2, a monoclinic lattice is stabilized and C60 units are interconnected by four member carbon rings and single C-C bonds along orthogonal directions. The observation of ionic conductivity in Mg2C60 makes the study of such system timely because of its great attractiveness for rechargeable Mg-ion batteries, as already highlighted by the high rate capability of fullerene-magnesium clusters as cathodes. The decomposition steps, their chemistry, and electronic Supplementary Information (ESI) available: [Optical Raman spectra of the Mg2C60 decomposed compound]. See DOI: 10.1039/b000000x/

Probing the thermal stability and decomposition mechanism of a magnesium-fullerene polymer via X-ray Raman spectroscopy, X-ray diffraction and molecular dynamics simulations

Matteo Aramini, a Johannes Niskanen, a Chiara Cavallari, b,c Daniele Pontiroli, c Abdurrahman Musazay, a Michael Krisch, d Mikko Hakala a Simo Huotari a

We report the microscopic view of the thermal structural stability of the magnesium intercalated fullerene polymer Mg2C60. With the application of X-ray Raman spectroscopy and X-ray diffraction, we study in detail the decomposition pathways of the polymer system upon annealing with temperatures between 300-700°C. We show that there are at least two energy scales involved in the decomposition reaction. Intermolecular carbon bonds, which are responsible for the formation of a 2D fullerene polymer, are broken with a relatively modest thermal energy, while the long-range order of the original polymer remains intact. With an increased thermal energy, the crystal structure in turn is found to undergo a transition to a novel intercalated cubic phase that is stable up to the highest temperature studied here. The local structure surrounding magnesium ions gets severely modified close to, possibly at, the phase transition. We used density-functional-theory-based calculations to study the thermodynamic and kinetic aspects of the collapse of the fullerene network, and to explain the intermediate steps as well as the reaction pathways in the break-up of this peculiar C60 intermolecular bonding architecture.
and activation energies are crucial information when considering the material's safe use in Mg-ion electrochemistry. Furthermore, a thorough understanding of the polymer C_{60} formation and decomposition has a paramount importance for the successful exploitation of the superhard materials that C_{60} can create. In this work we explore the structural stability of Mg_{2}C_{60} with a combined X-ray Raman and X-ray diffraction study, and address the thermal decomposition of the polymer and its products. These different probes are sensitive to the long range order (XRD) and to the local structure (XRS) yielding a contrast to size-scales in the decomposition process. We complementarily apply DFT and shed light on the thermodynamics and dynamics of the decomposition pathway after the exposure of the polymer to high temperatures.

2 Methods

2.1 X-ray Raman Spectroscopy

X-ray Raman spectroscopy (XRS) is a technique based on the measurement of the energy of X-rays and the evaluation of their energy loss yields the energy transfer in the inelastic scattering process. The scattering event can be modelled with the first order perturbation theory describing the transition of an electron from its ground state |i⟩ with energy E_{i} to the final state |f⟩ whose energy is E_{f}. During the excitation, the incoming photon with polarization e_{i} and energy o_{i} is transformed to the outgoing one with polarization e_{f} and energy o_{f}. The transition is fully described by the double differential cross section given by:

\[
\frac{d^2\sigma}{d\Omega d\omega} = n_0 \left( \frac{o_f}{o_i} \right) |e_i \cdot e_f| S(\vec{q}, \omega),
\]

where n_0 refers to the classical electron radius, whereas the term S(\vec{q}, \omega) is the dynamic structure factor that is expressed by:

\[
S(\vec{q}, \omega) = \sum_{f} \sum_{i} e^{-i\vec{q}\cdot\vec{r}_i} \delta (E_f - E_i - h\omega).
\]

The summation accounts for all possible final states f and electrons j of the system and h\vec{q}, h\omega are, respectively, the momentum and energy transferred to the system in the scattering process. XRS yields an information which is fully equivalent to X-ray absorption spectroscopy but includes the advantages of 1- sensitivity to low energy exchange (exploring the region from fractions of meV to keV) and 2- probing depth of hard X-rays which can bring unique bulk chemical information from low-energy absorption edges.

The XRS experiment was performed using the 72-element X-ray Raman spectrometer on the beamline ID20/UPBL6 of the European Synchrotron Radiation Facility (ESRF). The beam was monochromated with a Si(111) double crystal and a Si(311) channel-cut monochromators, and focused to a spot size of 20×20 μm. The spectrometer employs Si (660) analyzers that reflect X-rays with a fixed photon energy of 9.69 keV with an energy resolution of 0.6 eV. Spectra were collected at room temperature on various thermally treated samples. A suitable background was subtracted from the spectra and they were normalized over the whole edge intensity to a fixed area in the range 280-310 eV for C K edge and 45-68 eV for Mg L_{2,3} edge.

2.2 Materials

The air sensitivity of the materials required their handling under strict oxygen and moisture free condition (less than 1 ppm O_{2} and H_{2}O). Mg_{2}C_{60} fullerene polymer was prepared using the solid state reaction between magnesium (Alfa Aesar, purity 99.8%) and fullerene powders (MER corp., purity 99.9%) described in a previous paper by some of the authors. A few milligrams of this material were carefully sealed in quartz capillaries under argon atmosphere, prior to their exposure to high temperature for the decomposition treatment. Temperature was increased with a constant rate of 100°C/hour from normal conditions up to the temperature target, where dwelling time of 30 minutes was applied in all the treatments unless differently specified. Decompositions were performed between 300°C and 700°C; at the end all samples were quenched to room temperature. Powder diffraction was performed on different diffractometers both employing Cu Kα radiation.

2.3 DFT calculations

First-principles quantum mechanical simulations were performed with the Castep module of the Materials Studio software, using density functional theory and local density approximation with ultrasoft pseudopotentials. In addition, high temperature dynamics was explored up to 0.5 picosecond with femtosecond steps. Polymer fragments consisting of four fullerene units were enclosed in triclinic (space group P1) unit cell and their dangling bonds were saturated with hydrogen atoms. In the static calculations, and as the initial structure for the MD calculations, experimental atomic positions and lattice parameters were used for the polymer and, likewise, experimental values were set for the C-H distances. The system was simulated as an NVT ensemble using a Nose thermostat and 1000°C temperature.

3 Results and Discussion

3.1 Carbon K edge XRS

X-ray Raman Spectroscopy results of the carbon K edge are shown in Figure 1. In pristine Mg_{2}C_{60} the π∗-type peak is clearly visible at the energy of 285 eV composed of two partially overlapping features located at 284.5 and 286 eV. The edge profile rises at 290 eV and is composed of a broad structure at 292 eV and a more dispersed contribution with its centre of mass around 300 eV. The spectral signatures strongly resemble the ones of the 2D polymer form of C_{60} created under high pressure and measured with XRS. We can therefore assign the merged shape of peaks labelled I and II to the polymer network. When the sample is treated at increasing temperatures peak I is slightly shifted to lower energies and the peaks I and II become clearly separated. A further peak (III) emerges at 286.5 eV. Furthermore, peak IV slightly changes from a broad single feature to a double-peak whose separation increases with increasing annealing temperature. The pre-edge peaks labelled I, II and III correspond to the 1s → π∗ transitions and IV, V correspond to the 1s → σ∗ transitions, respectively. The spectral features of the pre-edge were originally discussed by Rueff et al. and attributed to excited states with t_{1u}, t_{2u}, and a_{g} symmetries. The a_{g}
state, in particular, appears in fullerene monomer only and can be therefore considered as the fingerprint of the breakage of the interlinking C₆₀ molecular bonds, which leads to the formation of C₆₀ monomers at relatively modest temperatures: in this case between 300-550°C.

3.2 X-ray Diffraction

Selected XRD patterns are reported in Figure 2 and the conservation of the polymer can be clearly noticed by the strong (011) reflection at 2Θ = 13.7° after treatments up to 600°C. Interestingly, this clearly shows that while the C₆₀ molecular bonds are already compromised at those temperatures, the long-range order takes more energy to be destroyed. Only treatments at T higher than 600°C result in the decomposition of the polymer shown by the disappearance of the polymer-related diffraction peak. In those cases, we observe an XRD pattern that indicates a cubic structure compatible with the symmetry of C₆₀ (s.g. Fm3m) and a lattice parameter a=14.11Å. In order to exclude a possible uncertainty in the calibration of the instrument, and thus excluding those peaks belonging to pure C₆₀, whose s.g. Fm3m and a=14.16Å, the measurements have been repeated on two different diffractometers yielding identical results. This cubic phase emerges from the polymer collapse for all temperatures above 450°C and it appears to be an intermediate decomposition product with a limited thermodynamic stability: decomposition at temperature higher than 600°C strongly affects its crystallinity and drives the formation of a mainly amorphous structure where only very weak and broad diffraction peaks can be seen.

3.3 Time evolution of the decomposition reaction

In order to ascertain the character of the final state of the decomposition reaction, we measured samples treated under an extended thermal treatment of 60 minutes. The resulting XRD and carbon K edge spectra are reported in figure 3. The picture presented by XRD shows that the same monomer structure previously achieved can be obtained via a longer treatment at more moderate temperatures. The cubic phase is stabilized at 450°C and treatments at higher temperature accordingly result in the same monomer structure. Its crystallinity, however, is strongly affected and diffraction peaks appear much weaker than in the case of shorter treatments for temperatures higher than 600°C. XRS in agreement with XRD and shows that longer treatments result in breaking the polymer network and creating C₆₀ monomers. The spectra of the annealed system indeed resemble that of the C₆₀ reference. When comparing the intensity of peak III and the width of I and II after 60-min long exposure to 450°C and 30-min exposure to either 550 or 600°C, we notice that they are more intense and sharper, respectively. This shows that the decomposition reaction is further developed towards the release of C₆₀ monomer. As a conclusion, we can state that within the decomposition process of the network of the C₆₀ molecules, there are two different energy scales involved: the individual C₆₀ short-range bonds can be lost with less thermal energy than is needed.
the long-range order does not necessarily imply structural stability on the inter-fullerene scale. This may set new standards in the understanding of the physics and chemistry of fullerene polymers with traditional characterization tools.

3.4 Magnesium $L_{2,3}$ edge XRS
Even further insight into the decomposition reaction is offered by magnesium $L_{2,3}$ edge XRS spectra in the polymer and in the decomposed phase that we report in figure 4. The spectrum of the pristine sample reveals the onset of absorption around 51 eV and a rather featureless EXAFS region with a weak broad feature around 4 eV from the onset. The spectrum remains unchanged after 550 °C treatment, where the polymer is still present, according to XRD, while interlinking C$_{60}$ bonds are already broken, according to XRS. The spectrum changes considerably for the sample treated at 700 °C. From XRD we know that, at that point, the polymer is fully decomposed and Mg $L_{2,3}$ absorption probes the magnesium environment in the cubic phase. The near edge feature at 51 eV seems strongly suppressed while the peak is shifted to higher energies around 52 eV. The most intriguing aspect of the behaviour of magnesium across the decomposition comes from the parallel observation of carbon K edges at the same temperatures. For the pristine sample both C and Mg spectra appear descriptive of the inter-linked C$_{60}$ and intercalated Mg present in the polymer structure, but their results surprisingly differ at 550 °C. The Mg $L_{2,3}$ edge is comparable to the one recorded in the pristine sample, which describes magnesium in the polymer phase, while carbon already presents the features (see peak III in Figure 1) that characterize the monomer C$_{60}$. This defines the mechanism of Mg$_{2}$C$_{60}$ depolymerization: Mg migration and the rearrangement of its local environment takes a higher activation energy than the breaking of the individual C$_{60}$ bonds. Cautiously estimating, taking into account the limited number of data points, the change of the local structure surrounding Mg ions may coincide with the change in the crystal structure, or at least lies in its vicinity. Depolymerization of the fullerenes conserves the symmetry of the system detected by XRD and does not affect Mg ions. For higher temperatures, fullerenes organize in a different cubic lattice for the monomer phase and Mg ions are stabilized in a new local structure, which determines the blue-shift of the spectrum of the sample decomposed at 700 °C. This feature is not to be attributed to a segregation of magnesium. Recorded spectra present a large shift towards higher energies with respect to the edge position in magnesium metal: such a reference can exclude the existence of segregated metallic Mg. At the same time we know from XRD that the polymer is still present in Mg$_{2}$C$_{60}$ treated at 550 °C; thus the spectra of the pristine compound and of the sample treated at 550 °C describe Mg in oxidation state +2 within the polymer. Therefore, the Mg $L_{2,3}$ edges provide convincing indication of the intercalated nature of Mg ions (in agreement with the picture of intercalated magnesium confirmed by optical Raman measurements which are reported in the electronic supplementary information).

3.5 DFT calculations
In order to get a complementary insight, we addressed the decomposition of Mg$_{2}$C$_{60}$ with a static calculation of energetics and with molecular dynamics simulations. A fragment of the polymer consisting of four fullerene molecules, terminated with hydrogen atoms, was built according to the polymer binding network and enclosed in a primitive triclinic cell (P1 symmetry). Since the XRS results showed a little or no influence of magnesium on the decomposition mechanism of the fullerene network, it is justified to omit the intercalated ion from the simulation and hence we explored the dynamics of the bare C$_{60}$ layer. Double and single inter-fullerene bridging bonds were alternatively cut and their energies were calculated. Figure 5 reports the energy levels for the different structures, relative to the pristine polymer network. The configuration with residual double bonds lies at an energy 0.28 eV higher than the pristine structure while the other with residual single bonds has an energy higher by 7.44 eV and is close to the fully decomposed polymer (7.89 eV). Since the minimum energy required for the transition to the structure with double-bond is much lower than the one with single-bond, the total energy calculations of the system suggest the cleavage of the single bond as the first step of the depolymerization. However it should be noted that this static picture does not take into account the activation energy required for the transition. For this reason we performed molecular dynamics calculations to get further insights in the decomposition pathways. We calculated the dynamics of the pristine polymer fragment at 1000 °C and we present the results in Figure 6. The simulated temperature was intentionally set higher than the experimental ones in order to speed up the structural decomposition within the time window explorable by DFT.

The main panel and the inset report the distance between carbon atoms involved in inter-fullerene bonding: single inter-C$_{60}$ bonds are shown with circles while double inter-C$_{60}$ bonds with
dashed lines. Vertical thin lines mark the time necessary for the stabilization process (estimated around 17 fs) while the other inset graphically helps attributing the results. The most intriguing aspect of the dynamical calculation is the agreement with the thermodynamics of the system. During the run both single bonds (whose initial length is 1.584 Å) start stretching and after a quick exposure of 40 fs the distance between C atoms exceeds 2 Å, clearly suggesting the path towards the collapse of the network. The distance of the single bonds keeps increasing during the whole simulation; double bonds, instead, keep oscillating around their average values (1.606, 1.594, 1.578 and 1.588 Å), which are similar to the bond length at room temperature (1.595 Å), thus suggesting that the decomposition has not yet addressed the double bond network. We may therefore conclude that the decomposition of the fullerene polymer Mg$_2$C$_{60}$ begins via the collapse of the weaker single-bond network, as highlighted by both energetic and kinetic perspectives.

4 Conclusions

By the complementary application of X-ray Raman spectroscopy, X-ray diffraction, as well as DFT calculations for both energetics and kinetics, we provided a thorough description of the decomposition for 2D intercalated fullerene polymer Mg$_2$C$_{60}$. The interfullerene bonding is broken with the lowest activation energy but the system preserves its crystal structure. This takes place at temperatures even below 450°C, and its detection requires a local probe that is sensitive to the chemical bonds, such as XRS, since the crystal structure is retained. The polymer crystal structure takes more energy to be modified, and the final decomposition product can be identified to consist of fullerene monomers in a cubic phase, which is stable up to the highest temperatures reached here (700°C). In this compound Mg still conserves its intercalated nature in the fullerene structure, but its local environment is however modified from that of the original compound. Depolymerization, investigated with molecular dynamics simulations, proceeds with a kinetic mechanism starting from the dissociation of the single-bond network. The system evolves into a completely monomeric structure through a successive detachment of double bonds. These results and the complementarity of spectroscopies, diffraction and dynamical methods, sensitive to both short range interactions and long-range order, offer a new scenario for the study of the physics and chemistry of carbon nanostructures.

The authors would like to acknowledge financial support from the Academy of Finland (grants No. 1254065, 1283136, 1259526, 1259599 and 1260204), European Synchrotron Radiation Facility laboratory for beamtime allocation, Dr. Christoph Sahle for support during the X-Ray Raman scattering experiment and Dr. Tiziano Rimoldi for providing optical Raman measurements.

References

See Supplemental Material at [URL to be inserted by publisher] for Raman measurements of decomposed material.