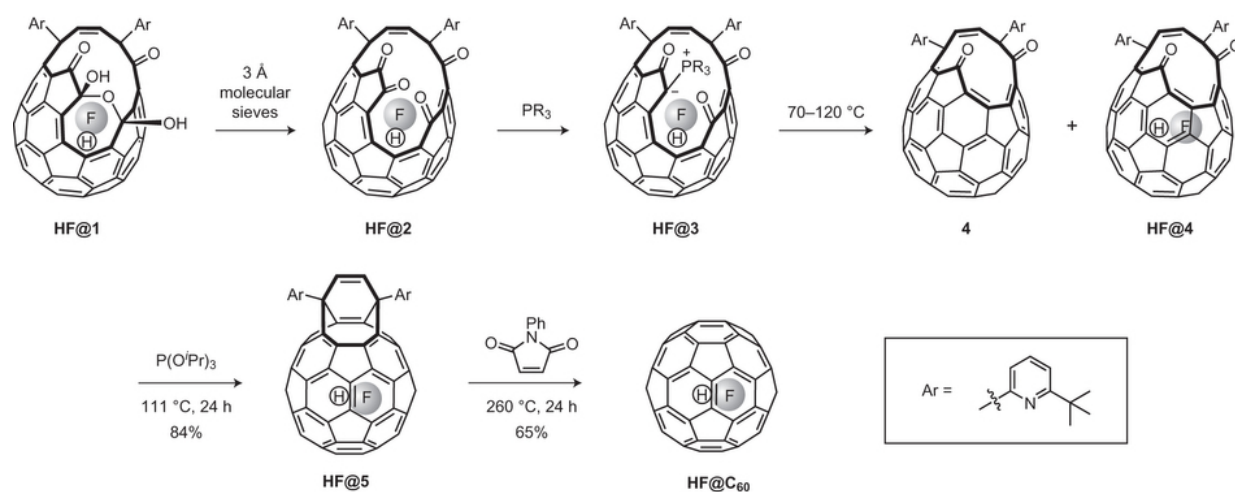


Synthesis and characterization of encapsulated single HF molecule

July 26 2016, by Heather Zeiger



Synthesis of HF@C₆₀ from HF@1. Credit: (c) Andrea Krachmalnicoff et al. The dipolar endofullerene HF@C₆₀, *Nature Chemistry* (2016). DOI: 10.1038/nchem.2563

(Phys.org)—Molecules are rarely found alone. In the real world, they are often networked to each other through hydrogen bonding or are bound to other molecules in the surrounding environment. One way to study an individual molecule is to trap it within a fullerene. A fullerene is an all-carbon, spherical molecule with carbons networked like the stitches of a soccer ball. The interior of the fullerene sphere is large enough to house small molecules, such as water or hydrogen gas.

In a recent study, Andrea Krachmalnicoff, et al. from the University of Southampton, the University of Nottingham, and Institutions in France and Estonia have trapped hydrogen fluoride in the cavity of a C_{60} fullerene ($HF@C_{60}$). Their characterization studies reveal that HF breaks the fullerene's icosahedral symmetry and that the fullerene cage serves to shield the strongly polar molecule. Their work appears in *Nature Chemistry*.

Researchers have been able to trap [molecules](#) within the fullerene cavity using a process called "molecular surgery." As the name implies, the fullerene molecule is chemically "cut open" and a small molecule is forced into the spherical cavity. The hole is then chemically "sutured."

Previous research by this group demonstrated the successful encapsulation of HF in an open-cage fullerene; however attempts to suture the open-cage resulted in the HF molecule escaping. The current study reports the successful encapsulation of HF by reacting their open-cage product with $PPh(Fu)_2$ ([2-furanyl] phenylphosphine) in the presence of molecular sieves at room temperature. They obtained the precursor product that can be sutured using methods employed in the synthesis $H_2O@C_{60}$ (i.e., reduction with triisopropylphosphite followed by reaction with N-phenyl-maleimide).

X-ray studies of $HF@C_{60}$ indicated some distortion of the [fullerene](#) cage but because the distortions were not outside of standard deviation values, the results were inconclusive. UV and electrochemical studies, however, did show that the HF molecule does not seem to show an electrostatic interaction with the C_{60} cage.

NMR studies using 1H , ^{13}C , and ^{19}F NMR as well as solid state 1H and ^{13}C NMR provided more compelling information on the behavior of HF. The J coupling for 1H - ^{19}F confirm that there is a single molecule of HF within the C_{60} cavity.

Krachmalnicoff, et al. then used inelastic neutron scattering and infrared absorption at low temperature to gain insight into the molecule's intrinsic properties. Their findings provide evidence that HF's rotational, vibrational, and translational motion is quantized with all transitions originating from the ground state. These studies showed that in the solid state, HF interact with the C₆₀ molecule both by breaking its icosahedral symmetry and through dipole interactions with the interior of the cage.

Hydrogen fluoride has a large dipole moment. Using capacitance studies to find the dielectric constant at low temperatures, Krachmalnicoff, et al. found that C₆₀ shields the dipole such that the dipole moment for encapsulated HF is about 25% of the calculated [dipole moment](#) for HF.

Overall, this research provides some interesting information on the properties of a single-molecule HF. Additional studies are needed to investigate why HF@C₆₀ breaks the endofullerene's icosahedral symmetry in the [solid state](#) as well as further investigation into HF@C₆₀'s electronic properties.

More information: Andrea Krachmalnicoff et al. The dipolar endofullerene HF@C₆₀, *Nature Chemistry* (2016). [DOI: 10.1038/nchem.2563](#)

Abstract

The cavity inside fullerenes provides a unique environment for the study of isolated atoms and molecules. We report the encapsulation of hydrogen fluoride inside C₆₀ using molecular surgery to give the endohedral fullerene HF@C₆₀. The key synthetic step is the closure of the open fullerene cage with the escape of HF minimized. The encapsulated HF molecule moves freely inside the cage and exhibits quantization of its translational and rotational degrees of freedom, as revealed by inelastic neutron scattering and infrared spectroscopy. The rotational and vibrational constants of the encapsulated HF molecules

were found to be redshifted relative to free HF. The NMR spectra display a large ^1H – ^{19}F J coupling typical of an isolated species. The dipole moment of HF@C₆₀ was estimated from the temperature dependence of the dielectric constant at cryogenic temperatures and showed that the cage shields around 75% of the HF dipole.

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