

Isolation and characterisation of the  
oxahomofluorofullerenol C<sub>60</sub>F<sub>17</sub>O.OH

Adam D. Darwish,<sup>a</sup> Ala'a K. Abdul-Sada,<sup>a</sup> Anthony G. Avent,<sup>a</sup> Joan M. Street<sup>b</sup> and Roger Taylor<sup>a</sup>

<sup>a</sup> Chemistry Department, The University, Southampton  
SO17 1BJ, UK

<sup>b</sup> The Chemistry Laboratory, CPES School, Sussex  
University, Brighton BN1 9QJ, UK

We report the isolation and characterisation of the first oxahomofluorofullerenol C<sub>60</sub>F<sub>17</sub>O.OH (a hydroxyether), from UV irradiation of a toluene solution of C<sub>60</sub>F<sub>18</sub> in air.

This is formed *via* the S<sub>N</sub>2' substitution [1] of a peripheral fluorine by OH, followed by insertion of oxygen αβ to the hydroxy group. No product is obtained if the reaction is performed under argon. Evidence will be presented that the structure is a particularly stable one and can be produced under quite different conditions.

The structure was determined from both EI mass spectrum fragmentation pattern and from 1 D and 2 D <sup>19</sup>F NMR spectroscopy. Analysis of these latter enabled the specific location of all of the fluorines to be assigned.

[1] A. G. Avent, A. K. Abdul-Sada, B. W. Clare, D. L. Kepert J. M. Street and R. Taylor, submitted.