

## Giant Fullerenes

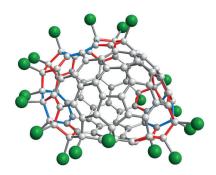
S.-F. Yang,\* S. Wang, E. Kemnitz,\*
S. I. Troyanov\* \_\_\_\_\_\_ 2460 – 2463



Chlorination of IPR  $C_{100}$  Fullerene Affords Unconventional  $C_{96}Cl_{20}$  with a Nonclassical Cage Containing Three Heptagons

## Three heptagons in the fullerene cage

were found for the first time in  $C_{96}Cl_{20}$  with a nonclassical cage, which was obtained by chlorination of isolated-pentagon-rule  $C_{100}$  fullerene. Most of 15 cage pentagons are fused resulting fused pentagon pairs and triples (see structure: gray C, green Cl; pentagons red, heptagons blue).

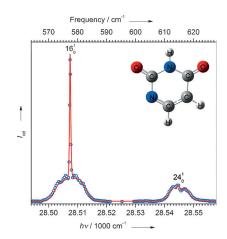


## Vibrational Spectroscopy

H. T. Liu, C. G. Ning, D. L. Huang, L. S. Wang\* \_\_\_\_\_\_ 2464 – 2468



Vibrational Spectroscopy of the Dehydrogenated Uracil Radical by Autodetachment of Dipole-Bound Excited States of Cold Anions The vibrational spectrum of the dehydrogenated uracil radical has been measured by autodetachment from dipole-bound states of cold deprotonated uracil anions. The spectrum shows observed rotational profiles (blue circles) for mode  $\nu_{16}$  at 577 cm<sup>-1</sup> (c-type) and mode  $\nu_{24}$  at 615 cm<sup>-1</sup> (b-type), in comparison with the simulated rotational profiles (red line).



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## Flashback: 50 Years Ago ...

Vitamin B<sub>12</sub> had at that time been only partially chemically synthesized (the total synthesis was first published in 1973), and a Review by K. Bernhauer et al. summarized the latest developments in the field, including partial syntheses, coenzyme forms, and enzymatic functions. A Review by Albert Eschenmoser on the origin of the molecular structure of vitamin B<sub>12</sub> was published in our Angewandte's Centenary Issue (Angew. Chem. Int. Ed. Engl. 1988, 27, 5–39).

Emanuel Vogel published a Communication on the properties of the cyclodeca-

pentaene system. The system contains  $10\,\pi$  electrons and can thus be expected to have an aromatic character. The proton NMR spectrum of 1,6-methanocyclodecapentane showed it to have a strong ring current, however, it shows olefinic reactivity and cannot be classified as a classical aromatic compound.

Georg Wittig reported a new synthesis of tetraarylphosphonium salts. Reactions of triarylphosphines and *o*-fluorophenyllithium with either fluorene or methyl iodide at -40°C resulted in the corresponding tetraarylphosphonium salt or its *o*-methyl derivative.

Max Schmidt reported the synthesis of 1,1,1,3,3,3-hexamethyldigermazane ((CH<sub>3</sub>)<sub>3</sub>Ge-NH-Ge(CH<sub>3</sub>)<sub>3</sub>). The reaction of trimethylgermanium chloride with ammonia produced the target compound as a colorless liquid. Reaction of dimethylgermanium dichloride under the same conditions resulted in the formation of [(CH<sub>3</sub>)<sub>2</sub>GeCl]<sub>3</sub>N.

Read more in Issue 3/1964.