

OPTICAL (OR LASER ENHANCED) NMR, INTERPRETATION USING  
THE PHOTOMAGNETON OF LIGHT

by

M. W. Evans  
Department of Physics  
University of North Carolina  
Charlotte, NC 28223

In a recent technical comment [1] on the remarkable optical NMR experiments reported by Warren *et al.* [2], Harris and Tinoco give a conventional calculation of optically induced NMR shifts. In their reply [3], Warren *et al.* report careful and repeated experimental checks which confirm beyond doubt that the experimentally observed shifts are *many orders of magnitude* larger than those given by conventional theory [1]. The observed shifts [2, 3] are of the order 0.1 to 1.0 Hz for laser intensities of 1 to 3  $W\ cm^{-2}$ , while Harris and Tinoco [1] assert that the shift for 10  $W\ cm^{-2}$  is theoretically no larger than about  $10^{-15}$  Hz. There is therefore an enormous discrepancy between experiment and conventional theory, clearly signalling the presence of a novel property of light.

I would like to suggest that the novel [4-8] photomagneton of light,  $\hat{B}^{(3)}$ , is the source of the shifts seen by Warren *et al.* [2, 3] with circularly polarized light.

Extensive theoretical work [4-8] has shown that the spin angular momentum of the photon, the operator  $\hat{J}$ , generates the phase free photomagneton,

$$\hat{B}^{(3)} = B^{(0)} \frac{\hat{J}}{\hbar}, \quad (1)$$

per photon in vacuo. Here  $B^{(0)}$  is the scalar magnitude of magnetic flux density in vacuo (tesla), calculable from the intensity  $I_0$  of the light beam through the relation  $B^{(0)} \sim 10^{-7} I_0^{\frac{1}{2}}$ . For a beam intensity of 1,0  $W\ cm^{-2}$ ,  $B^{(0)}$  is about  $10^{-5}$  tesla, which for a 270 MHz ( $\sim 10$  tesla) spectrometer is expected to shift resonances by about  $\sim \pm 27$  Hz, depending on the sense of circular polarity of the light beam. This theoretical figure is about an order of magnitude too large, but the conventional theory [1] produces results some fifteen to sixteen orders too small. The operator  $\hat{B}^{(3)}$  is zero in linearly polarized

or incoherent light [4-8] and switches sign with the sense of circular polarity of light. This switch of sign is consistent with experimental data [2, 3]. The fact that  $\hat{B}^{(3)}$  appears to produce shifts which are about an order too large could easily be due to the fact that only a fraction of available beam intensity reaches the sample, or may be due to some novel, incompletely understood, mechanism of shielding by the electrons, a novel type of useful chemical shift.

The photomagneton  $\hat{B}^{(3)}$  is a novel and fundamental [4-8] photon property, which is responsible for magnetization by light, as in the inverse Faraday effect [9-13], which can be expressed very simply as

$$\mathbf{M} = \langle \hat{M} \rangle = AB^{(0)} \langle \hat{B}^{(3)} \rangle. \quad (2)$$

Here A is an ensemble averaged molecular property tensor, a hyperpolarizability [14].

## References

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