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Predictable and New Physics and Potential for Applications of Organic-based Magnets¹

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As discussed by Joel S. Miller in the previous talk, magnets utilizing organic groups with essential spin have been reported since the mid-1980's. Though initial organic-based magnets had magnetic ordering temperatures (T_c 's) below 5K, organic-based magnets now have T_c 's to above 400K. In addition to magnetic phenomena already known for conventional transition metal and rare earth magnets, organic-based magnets feature unique phenomena enabled by the shape and internal electronic structure of the organic molecules. Examples are illustrated with experimental results for magnets based on tetracyanethylene, [TCNE], which as an anion has spin $1/2$. For example, chains with spin containing molecules having relatively strong exchange within a chain and weak dipolar interaction with neighboring chains can have an unusual fractal ground state with unusual dynamics leading to 'coercive fields' approaching 3 tesla. In contrast to conventional magnets, the internal electronic structure of the molecules that make up a molecule-based magnet can be excited by light of the appropriate wavelength. This leads to changes of the spin state of the molecule and/or changes in the exchange interaction between molecules, opening up the concept of reversible light control of magnetism. Examples will be given from the $M^{++}[\text{TCNE}]_x^-$ ($x \sim 2$) ($M = \text{Mn}, \text{V}$) materials systems. Finally, we explore the new phenomena enabled by $V^{++}[\text{TCNE}]_x^-$ ($x \sim 2$), a material with T_c up to 400K and for which films may be prepared using low temperature CVD. It is a semiconductor (room temperature resistivity and activation energy similar to silicon) and magnetization $M(H,T)$ and coercive field are controlled by chemical composition. Magnetoresistance to 32 tesla supports that $V[\text{TCNE}]_2$ is a "half-semiconductor" with fully spin polarized valence and conduction bands of interest for spintronics applications.

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