

HIGH MAGNETIC FIELD STUDIES OF THE SPIN-PEIERLS STATE IN THE ORGANIC CONDUCTOR $\text{Per}_2[\text{Pt}(\text{MNT})_2]$

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$\text{Per}_2[\text{Pt}(\text{mnt})_2]$ is a quasi-one dimensional organic conductor that consists of metallic perylene chains and insulating $\text{Pt}(\text{mnt})_2$ chains with localized Pt spins ($S = 1/2$). Under ambient conditions, the perylene chains undergo a metal-insulator transition at $T_c \sim 8$ K due to the formation of a charge density wave (CDW), while the $\text{Pt}(\text{mnt})_2$ chains undergo a spin-Peierls transition at the sample temperature. To investigate the magnetic properties of the spin chains in this material a tunnel diode oscillator (TDO) was used. A TDO is a self-resonating LC circuit drive by a tunnel diode. For this work, aligned single crystals of $\text{Per}_2[\text{Pt}(\text{mnt})_2]$ were placed inside of the inductor coil of the TDO, which resulted in a change in the resonant frequency of the circuit due to changes in the magnetic susceptibility of the sample. [1]

The experimental results obtained by using the TDO technique, in conjunction with previous resistivity studies [2], suggest a coupling of the spin-Peierls and charge density wave states below 8 K and up to 20 T, above which both states are suppressed. At fields above 20 T a transition into a second spin state occurs, which coincides with a field induced insulating state in the perylene chains. These results not only suggest a strong coupling between the segregated charge and spin chains but also provide support to the idea that the charge density wave is needed to support the spin-Peierls state even in the low-temperature, low-magnetic field state. [3, 4]

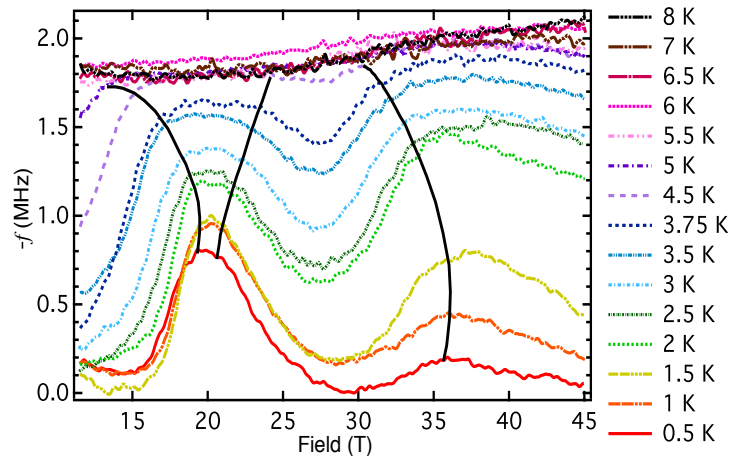


Figure 1. The change in TDO frequency as a function of magnetic field for a range of temperatures. The black lines are guides to the eye to indicate transitions into and out of the spin-Peierls state.

[1] L.E. Winter *et al.*, EPL **103**, 37008 (2013).

[2] D. Graf, *et al.*, Phys. Rev. Lett. **93**, 076406 (1-4) (2004).

[3] E. L. Green, *et al.*, Phys. Rev. B **84** 121101(R) (1-4) (2011).

[4] E. L. Green, *et al.*, Crystals, **2**, 1116-1135 (2012).

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