

Building quantum spintronic devices using the ferromagnetic metal /molecule interface

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Molecular spintronics is emerging as a vibrant field that utilizes the properties of magnetic molecules toward additional device functionalities that can exploit quantum physics. One spectacular property is the high spin polarization of the ferromagnetic metal/molecule interface, measured at 300K using spectroscopy [1] and magnetotransport [2] experiments.

The magnetism of this so-called spinterface [3] can be controlled using an underlying ferroelectric [4], and can in principle be combined with the spin crossover (SCO) molecular property by inserting a noble metal spacer layer [5,6]. This would enable single-molecule magnet [7] behavior on a much wider range of molecules, at higher temperatures. To improve our understanding of SCO device properties, device in operando experiments using synchrotron radiation shed light [8] into the transport path across the SCO thin film.

The spinterface formation entails changes to the magnetic properties of the interface constituents, from magnetic hardening of the metal [9] to the magnetic stabilization of otherwise paramagnetic molecular spin chains borne by phthalocyanine (Pc) molecules [10]. These effects enable the encoding of information into the quantum state of a molecular spin chain [11]. This also lays the groundwork for a molecular implementation [12] of our recent proposal [2] (www.spinengine.tech) to harvest the energy of thermal fluctuations on paramagnetic centers using spintronics.

- [1] F. Djeghloul et al, J. Phys. Chem. Lett. **7**, 2310 (2016).
- [2] K. Katcko et al, Commun. Phys. **2**, 116 (2019).
- [3] S. Delprat et al, J. Phys. Appl. Phys. **51**, 473001 (2018).
- [4] M. Studniarek et al, Adv. Funct. Mater. 1700259 (2017).
- [5] M. Gruber et al, Nano Lett. **15**, 7921 (2015).
- [6] E. Urbain et al, Adv. Funct. Mater. **28**, 1707123 (2018).
- [7] F.-S. Guo et al, Science eaav0652 (2018).
- [8] F. Schleicher et al, ACS Appl Mater Interfaces **10**, 31580 (2018).
- [9] K. V. Raman et al, Nature **493**, 509 (2013).
- [10] M. Gruber et al, Nat. Mater. **14**, 981 (2015).
- [11] K. Katcko and et al., Adv. Func. Mater. 2009467 (2021)
- [12] B. Chowrira, L. Kandpal et al, arXiv : 2009.10413