

Abstract Submitted
for the MAR15 Meeting of
The American Physical Society

Low Temperature Scanning Tunneling Spectroscopy of isolated Mn₁₂-Ph Single Molecule Magnets¹ K. REAVES, Depts Materials Science and Engineering and Physics and Astronomy, Texas A&M University; WPI-AIMR, Tohoku University, Sendai, Japan, P. HAN, K. IWAYA, T. HITOSUGI, D. PACKWOOD, WPI-AIMR, Tohoku University, Sendai, Japan, H.G. KATZGRABER, Depts Materials Science and Engineering and Physics and Astronomy, Texas A&M University; Santa Fe Institute, H. ZHAO, K.R. DUNBAR, Dept of Chemistry, Texas A&M University, K. KIM, W. TEIZER, Depts Materials Science and Engineering and Physics and Astronomy, Texas A&M University; WPI-AIMR, Tohoku University, Sendai, Japan — We study Mn₁₂O₁₂(C₆H₅COO)₁₆(H₂O)₄ (Mn₁₂-Ph) single-molecule magnets on a Cu(111) surface using scanning tunneling microscopy and scanning tunneling spectroscopy at cryogenic temperatures ($T < 6\text{K}$). We report the observation of Mn₁₂-Ph in isolation and in thin films, deposited through *in situ* vacuum spray deposition onto clean Cu(111). The tunneling current of isolated Mn₁₂-Ph, normalized with respect to the Cu background, shows a strong bias voltage dependence within the molecular interior. The qualitative features of these I vs.V curves differ by spatial location in several intriguing ways (e.g. fixed junction impedance with increasing bias voltages). We explore these normalized I vs. V curves and present a phenomenological explanation for the observed behaviors, corresponding to the physical and electronic structure within the molecule.

¹Funding from WPI-AIMR

Kelley Reaves
Texas A&M University Materials Science and Engineering

Date submitted: 13 Nov 2014

Electronic form version 1.4