

Atomic-scale friction sensed by a single organic molecule

R. Pawlak¹, E. Gnecco², S. Kawai¹, T. Glatzel¹, S. Fremy¹, H. Fang³, L.-A. Fendt³, F. Diederich³, A. Baratoff¹ and E. Meyer¹

1: Departement of Physics, University of Basel, Klingelbergstrasse 82, 4056 Basel, Switzerland, 2, Instituto Madrilenio de Estudios Avanzados en Nanociencia (IMDEA Nanociencia), Cantoblanco, 28049 Madrid, Spain, 3, Laboratory of Organic Chemistry, Department of Chemistry and Applied Biosciences, ETH Zürich, Honggerberg, HCI, 8093 Zürich, Switzerland.

The frictional behavior of nano-objects such as organic molecules has also a fundamental interest for understanding their diffusive processes on surfaces or designing molecular nano-machines [1-2]. In previous works, we use a tuning fork based AFM operated at low temperature to investigate the mechanical properties of single molecules and developed force-induced manipulations [3-5]. Here, we intentionally functionalized the tip apex with a single porphyrin molecule and performed friction experiment (Fig.a). In this configuration, the molecule acts as a spring attached to the apex and dynamically interacts with the surface lattice potential. Scanning the Cu(111) surface at constant height while recording the frequency shift systematically reveals atomic-scale friction patterns (Fig.b). The molecule stiffness as well as the chemical reactivity of specific end-groups give rise to the typical stick-slip behavior (Fig.c). Numerical calculations based on a gas-atom solid interaction potential [6] further support the experimental data.

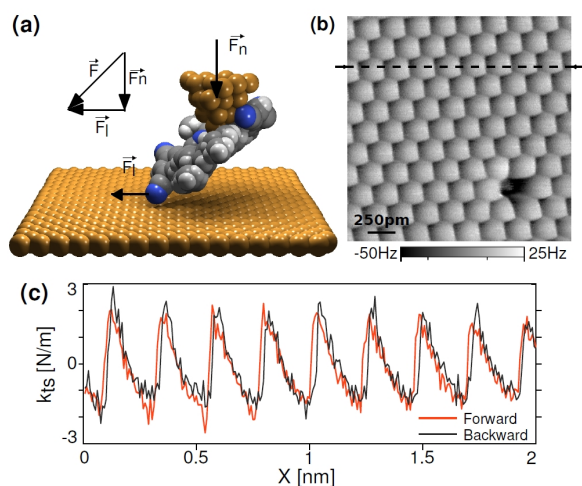


Figure 1: (a) Illustration of the friction experiment : the porphyrin-terminated tip is brought into contact to the Cu(111) while oscillating at its resonance frequency. (b). Constant-height $Df(x,y)$ maps of the surface showing the atomic lattice. (c) Tip-sample stiffness profile k_{TS} along the dashed line of (b) revealing a typical stick-slip pattern. (parameters : $f_0 = 25956$ kHz, $A_0 = 50$ pm, scan speed = 1 nm/s).

References

- [1] Urbakh, M.; Meyer, E. *Nature Mat*, **9**, 8-9 (2010).
- [2] M. Kisiel *et al.*, *Nature Mat.*, **10**, 119122 (2011).
- [3] R. Pawlak *et al.*, *ACS Nano*, **6**, 6318-6324 (2012).
- [4] R. Pawlak *et al.*, *ACS Nano* **5**, 6349-6354 (2011).
- [5] R. Pawlak *et al.*, *J. Phys.: Condens. Matter* **24**, 084005 (2012).
- [6] W.A. Steele, *Surf. Sci.* **36**, 317 (1973)