Seminář odd. 26 Tenkých vrstev a nanostruktur

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TÉMA

Charge distribution of adsorbed single molecules: the effect of theunderlying surface

Tobias Meier, Rémy Pawlak

Department of Physics, University of Basel, Klingelbergstr. 82, 4056 Basel, Switzerland

Electron donor-acceptor molecules are promising candidates for molecular electronic devices since they may generate separated electron-hole pairs upon absorption of photons. To preserve the intrinsic electronic character of such complexes upon adsorption, a particular attention must be focused on their interaction with the underlying surface. In this contribution, we will first show the adsorption of TTF-dppz [1], a fused piconjugated donor-acceptor molecule, deposited on insulating NaCl films on Cu(111). To investigate the interplay between the atomic structures of TTF-dppz and NaCl, we performed high-resolution scanning tunneling (STM) and atomic force microscopy (AFM) measurements at 5 K with CO-terminated tips. The charge distribution above individual molecules has been studied by local contact potential mapping [2,3] which reveals the coexistence of two charge redistribution depending on the adsorption site. In agreement with numerical calculations [4], such variation of the TTF-dppz properties is found to arise from the formation or not of a complex with a single Na⁺ ion of the surface. In a second example, we will describe the adsorption of porphyrin derivatives on hydroxylated rutile-TiO₂(110). The interaction of the single molecules with the hydroxyl groups (OH) of the rutile surface has been particularly regarded by a combined STM/AFM study. We found that a relevant change of the charge distribution of those molecules is provoked by the presence or not of OH groups underneath [5]. Finally, we will discuss our first results on the adsorption of phtalocyanin molecules on a topological insulator (Bi₂Te₃).

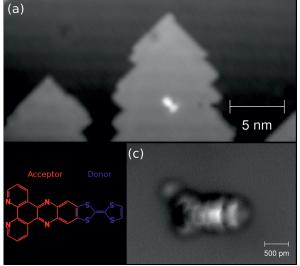


Figure 1: (a) STM topographic image of a single TTF-dppz molecule on NaCl thin film deposited on Cu(111).(b) chemical structure of TTF-dppz. (c) constant height AFM image ofTTF-dppz.

References:

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