



Friction of Xenon monolayers on Copper

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Superlubricity 1

Dry friction assumes very small values when the (nano)contacts between two crystalline surfaces are incommensurate: The force coming from the mismatched atoms in the contact area point in all directions and sum up to zero. Hirano and K. Shinjo, PRB 90

Vanishing friction on a silicon surface was observed with ultrahigh vacuum (UHV) scanning tunneling microscopy Hirano et al., PRL 97





Superlubricity 2

First direct experimental confirmation of importance of lattice mismatch on sliding friction: rotating graphite flake above a graphite surface



Dienwiebel, PRL04

Small friction -> incommensurate state





Superlubricity in atomic systems

Xenon monolayers on Cu(111)

→ model system to study onset of frictional slip by domain nucleation

Reguzzoni, PNAS10





Commensurate phase Xe on Cu(111)

At temperatures between 50-80K Xe is found to form a $\int 3x \int 3R30^{\circ}$ registred phase on Cu(111)



Bulk Xe-Xe distance 4.38 Å √3x(Cu-Cu spacing) 4.41 Å





Nonlinear sliding dynamics

Xe deposition on Cu(111) at a constant temperature



Calculated static friction force per Xe atom at T~0 K F_s =6.5meV/Å for Cov=1/3 F_s =0.021meV/Å for Cov=0.36 Franchini, to appear in JPCM





Experimental results 1

Coverage scans of Ne on Pb(111) @ 6 K Pinning at low coverages followed by (structural?) depinning above ~ 0.3 ML



Lack of morphological data on Ne/Pb system to validate interpretation of observed depinning in terms of superlubricity





Experimental results 2

QCM study of Xe at 77K



FIG. 2. The slip times for xenon on Cu(111) (triangles), Ni(111) (squares), and graphene/Ni(111) (circles) vs coverage. The coverage (corrected for slip effects) is obtained by solving the right expression in Eq. (2) for $(\delta f_{\rm film})$ and then substituting the value into the left expression in Eq. (2). The frequency shift of a monolayer of xenon is calculated from the xenon mass and spacings (coverage) given in Table ¹. Compressed monolayer coverage is 6 atoms/nm², corresponding to $1.31 \times 10^{-12} \text{ ng/nm^2}$.

Coffey, PRL05



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Large slip times on Cu(111) are explained as due to the displacement of preexisting domain walls caused by vacancies in the Xe film or substrate defects

Reguzzoni, PNAS10



Quartz Crystal Microbalance (QCM)

Standard technique to measure mass (e.g. thickness) in evaporators



QCM as a nanofriction probe

In 1988 J. Krim suggested to use QCM to study sliding of adsorbed film



Xenon deposition on Cu(111)







Growth and characterization of Cu electrodes

Growth and characterization facilities available at SESAMO laboratory in Modena have been used to prepare crystalline copper in UHV conditions directly on a bare quartz crystal

in-situ STM scan



Average size crystallites ~ 50nm (111) texture

Room temperature deposition Cr: 30 nm - rate 5 Å/min Cu: 60 nm - rate 10 Å/min





Ne on Copper

Slip times of Ne films deposited on Cu @ 6K





Annealing of QCM at low temperatures

Slip times olf Ne films deposited on the Pb electrode of a QCM @ 6K







Deposition of Xe on Cu(111) at low T

No sliding of Xe up to 15 K and for coverages ≤ 2 layers



At low T (e.g. T≤15 K), only Ne is found to slide. All heavier adsorbates (N2, Ar, Kr) are always pinned to the electrodes

Pierno, PRB2010





CONCLUSIONS

•We have started a new experiment to study the sliding friction of rare gases on Cu(111) with a QCM technique

•Preliminary measurements on an oxidized Cu electrode confirm that only Ne films are found to reproducibly slide below 10K after thermal annealing of the QCM following initial cool down

OUTLOOK

•Repeat experiments with Cu electrodes of good crystalline quality transferred under vacuum at temperatures around 40-60K to search for superlubricity of Xe monolayers





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QCM as a nanofriction probe

From ΔV and Δf it is possible to determine slip time τ_s

Slip time is the time it takes the film to follow the electrode





