Stress engineering at the nanometer scale: Two-component adlayer stripes

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Abstract – Spontaneously formed equilibrium nanopatterns with long-range order are widely observed in a variety of systems, but their pronounced temperature dependence remains an impediment to maintain such patterns away from the temperature of formation. Here, we report on a highly ordered stress-induced stripe pattern in a two-component, Pd-O, adsorbate monolayer on W(110), produced at high temperature and identically preserved at lower temperatures. The pattern shows a tunable period (down to 16nm) and orientation, as predicted by a continuum model theory along with the surface stress and its anisotropy found in our DFT calculations. The control over thermal fluctuations in the stripe formation process is based on the breaking/restoring of ergodicity in a high-density lattice gas with long-range interactions upon turning off/on particle exchange with a heat bath.

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Spontaneous formation of periodic patterns is an example of nature’s tendency towards order. A class of such structures is induced by surface stress [1], and has been widely observed on single crystal surfaces [2–8]. It is well known that the formation of these equilibrium patterns is driven by a competition between interactions at different length scales [9]. The forces in action, due to short-range near-neighbour and long-range dipolar interactions, are of the most general type resulting in very similar phenomena occurring also in magnetic [10] and electrostatic systems [11]. This gives a strong motivation to include such equilibrium phases with mesoscopic order in the discussion of making self-assembly available to future applications [12].

To this date, the majority of the reports on stress-induced equilibrium patterns have focused on the characteristic feature shape and size. The temperature dependence of the pattern period appears as a common ingredient in the various systems studied, and the general explanation is based on a scaling of the interaction parameters due to thermal disorder [13]. The pronounced thermal fluctuations [5,7], derived from the high mobility of the adatoms, are essential in reaching the thermodynamic equilibrium phase. However, this mobility, which allows the pattern to almost instantaneously equilibrate, is a hindrance to preserving the pattern away from its formation temperature.

Here, we explore the possibility of controlling the thermal fluctuations by adding a second adspecies to an already stripe-forming system. The idea is based on the slow dynamics in a high-density binary lattice gas, which leads to a tendency towards glassy behaviour. In particular, we implement this by dosing small amounts of oxygen on submonolayer Pd/W(110). The oxygen-free surface is known to organize at high temperatures into alternating stripes of a dense Pd monolayer and a dilute Pd lattice-gas [13]. In the following, it will be shown that the mixed Pd-O layer necessitates special growth

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conditions in order to form a similar stripe phase. As we will describe shortly, this allows to identically preserve the equilibrium pattern, produced at high temperature, at lower temperatures. Moreover, the addition of oxygen leads to a progressive change in the surface stress, which is reflected by the changes in pattern period and anisotropy. The result is a highly ordered stripe pattern with a tunable period down to below 20 nm, and with an orientation that can be rotated by 90°. The density-functional theory (DFT) calculations, in conjunction with continuum model results, are used to predict the phase diagram of stripe shape and orientation of this system as a function of Pd and oxygen coverages. This allows us to identify the equilibrium phases and successfully account for the main observed changes in stripe pattern, including the 90° rotation in the stripe orientation.

Using low-energy electron microscopy (LEEM) [14–16] we studied Pd-O pattern formation on W(110) following different growth pathways and temperatures. The progressive changes to the Pd/W(110) stripes upon oxygen dosing and the kinetic factors in forming the new Pd-O stripe pattern are revealed in the following two approaches: i) dosing oxygen on a partially Pd-covered surface at 1200 K, and ii) depositing Pd on a partially oxygen-covered surface at 1240 K. The latter results in the Pd-O stripe phase as Pd is deposited at a temperature at which it partially desorbs. This new stripe phase, with perpendicular stripe orientation relative to the Pd stripes on the oxygen-free surface, is identified as an equilibrium configuration. The conditions under which the Pd-O stripes are obtained involve effective fluctuations via partial desorption of Pd, which enable to reach the minimum energy configuration.

Oxygen on W(110) forms a series of ordered phases as a function of coverage [17] and has been extensively studied as a model system for surface diffusion and island formation [18]. For coverages below 0.5 ML, relevant to this study, it is found in (1 × 2)-ordered islands. Importantly, oxygen has a strong tendency to laterally segregate from most transition metal adatoms coadsorbed on W(110) [19]. Furthermore, recent work has shown that oxygen induces a considerable stress change on tungsten [20]. These observations point towards the possibility of finding an ordered two-component (oxygen and metal) adsorbate phase.

Following this reasoning, we first dosed molecular oxygen on a partially Pd-covered W(110) surface at 1200 K, above the stripe disordering temperature. At this temperature, Pd shows only short-range order [13]. Upon adsorbing small amounts of oxygen the stripe phase appears along [1¯10] with a wavy morphology as shown in fig. 1a. With increasing O coverage the pattern period increases (fig. 1b), followed by segregation of Pd into large 2D islands surrounded by O (fig. 1c). Further increase in O coverage (above roughly 0.3 ML) results in the shrinking of the Pd islands (fig. 1d). This suggests that when the local density of O exceeds that of the (1 × 2)-ordered layer, Pd atoms are displaced into the second layer, from which they readily desorb at this high temperature. It is important to note that at this temperature the order of deposition has little influence on the final result. In other words, depositing Pd on a partially oxygen-precovered surface below 1200 K leads to similar structures as displayed in fig. 1d.

In the second approach, Pd was adsorbed at 1240 K (about 40 degrees higher compared to that in fig. 1) on a surface precovered with less than 0.5 ML of oxygen. This temperature corresponds to the tail of the thermal desorption peak of the first Pd adlayer [21], so that Pd adsorption is partially balanced by desorption. Under these conditions, a highly ordered stripe pattern develops along the [001] direction as illustrated in fig. 2. Crucially, partial desorption was established as the necessary condition for forming the mesoscopic pattern. The stripe pattern consists of alternating Pd and O phases. The stripe period strongly depends on the amount of Pd on the surface, and a minimum period of about 16 nm is reached at the highest Pd coverage (about 0.33 ML) allowed by the pre-adsorbed O. The new Pd-O stripe phase differs from the pure Pd stripes in the orientation of the stripes ([001] vs. [110]), i.e. by a 90° rotation in the stripe direction, as well
as in the improved directionality and reduced periodicity (16 vs. 60 nm). Moreover, the pattern can be frozen upon cooling and thus preserved at room temperature.

In order to elucidate whether this new phase represents the thermodynamic equilibrium, the energetics and anisotropy of the Pd-O adsorption system have been investigated by DFT calculations, providing the parameters to the continuum model arguments developed for stress-induced equilibrium patterns [22–24]. The DFT pseudopotential calculations were performed in a plane-wave basis by using the PWscf code [25]. The details of the surface stress calculations for W and O/W can be found in refs. [20] and [26]. The Pd-O boundary energy was calculated on a 6-layer slab consisting of a Pd-O surface layer and 5 W layers. The lateral size of the slab corresponds to 10 × 2 substrate unit cells, of which one half is covered by Pd and the other half by (1 × 2)O. In estimating the creation energy of a single boundary, only the first atoms adjacent to the Pd-O(1 × 2) boundary were allowed to relax (in the upper 4 layers) in order to minimize contribution from elastic interactions between neighbouring boundaries.

It should be noted that the atomic structures of both Pd and O stripes at high temperature are a challenge for modeling. On the one hand, the low-temperature ordered oxygen phases lose their long-range order several hundred degrees below the temperature at which stripes form (for example the (1 × 2) structure disorders at about 700 K [18]). However, the short-range order persists at high temperatures and influences the stripe phase profoundly. In particular, a local (1 × 2) density is the highest oxygen packing that can be attained in the presence of Pd islands, before Pd is displaced to the second and higher layers. Moreover, the oxygen stripe in the Pd-O stripe phase has the (1 × 2) structure when cooled down to room temperature. Therefore, at high temperature the oxygen-covered regions can be described as having short-range (1 × 2) order. The effect of disorder on the oxygen surface stress is studied in ref. [26], and the values obtained are reproduced in table 1.

On the other hand, at high temperature the submonolayer Pd is pseudomorphic on W(110) only along [001] and shows a mismatch structure along [110] (i.e. one-dimensional pseudomorphism) [21]. Using low-energy electron diffraction, we observe a very similar situation for Pd within the Pd-O stripes with a mismatch of approximately 5%. We have estimated the effect of this non-pseudomorphism on the surface stress by varying the mean lattice constant of the Pd overlayer along [110] for selected commensurate periodicities in DFT calculations using supercells with large [110] dimensions on asymmetric slabs [27]. The calculated values for 5% mismatch are displayed in the upper half of table 1. Note that small variations of the mismatch (less than 1%) due to changes in temperature or coverage are found to modify the calculated stresses only slightly and have no consequence in the following discussion.

The results of the calculations relevant to the stripe formation, i.e. the directional dependence of stress difference and boundary energy between the two phases (O and Pd), are summarized in the last two rows in table 1. For comparison, we include the stress anisotropy for the oxygen-free Pd stripes in the first column. An important finding is that the Pd-O boundary energy is nearly isotropic. The values in the table clearly identify

| Direction  | $\tau_{W(1\times1)}$ | $\tau_{O(1\times2)}^{\text{high}T}$ | $\tau_{\text{Pd}(5\%\text{mismatch})}$ |
|------------|---------------------|-------------------------------------|--------------------------------------|
| [110]      | 3.6                 | -0.2                                | 3.1                                  |
| [001]      | 5.2                 | 3.4                                 | 3.4                                  |

|          | $\Delta\tau_{\text{Pd-W}}$ | $\Delta\tau_{\text{Pd-O}}$ | $\beta_{\text{Pd-O}}$ |
|----------|-----------------------------|---------------------------|-----------------------|
| [110]    | -0.5                        | 3.3                       | 0.152                 |
| [001]    | -1.8                        | $\sim$ 0                 | 0.164                 |

Table 1: Calculated surface stresses ($\tau$) and the boundary energy ($\beta$) between Pd and O phases on W(110). The upper two rows list the stress values. The lower two rows display the directional dependence of stress differences and boundary energy. All stresses are given in N/m, and the boundary energy has units eV/Å. O(1 × 2) stress is taken from ref. [26].
the stress difference as the real source of anisotropy. The elastic energy gain is higher for a larger stress difference across the stripe boundary, i.e., perpendicular to the stripe direction. As a consequence, the expected stripe direction strikingly changes from $[1\bar{1}0]$ for oxygen-free Pd stripes to $[001]$ for Pd-O stripes on W(110).

The calculated stress anisotropy is found to be more than sufficient for driving the formation of the Pd-O stripes along $[001]$ according to the phase diagram (see fig. 3a) evaluated on the basis of the continuum model described by Suo and coworkers [22–24] and our computed stresses. The model predicts distinct stripe phases as the ratio of stress differences along the two high-symmetry crystallographic directions, $r = \Delta \tau_{[1\bar{1}0]} / \Delta \tau_{[001]}$, is varied. A critical value, $r_c = 1 + 2\nu$ (where $\nu$ is Poisson’s ratio), is found analytically by minimizing the free energy as a function of stripe orientation, which marks the onset of regular stripes [22]. In particular for an elastically isotropic substrate as W(110) [28], when $r$ is greater than $r_c = -0.44$ (or smaller than $-1/0.44$ for the perpendicular stripes) a linear stripe phase is expected. As can be seen from the results in table 1, this condition is satisfied both for Pd and Pd-O stripes. In order to check this quantitative criterion for all coverages displayed in fig. 2, the stress differences can be estimated by subtracting a weighted sum of oxygen and tungsten surface stresses from that of Pd. This is based on the observation that increasing Pd coverage squeezes the oxygen atoms towards a fully packed $(1 \times 2)$ order at the highest density. The corresponding stress ratios, shown as dark circles in fig. 3a, confirm the presence of $[001]$ stripes for 0.33 ML O coverage, regardless of the Pd coverage.

Based on the continuum model, a herringbone pattern (“θ-stripes” [22]) is expected when the stress differences are of similar magnitude but of opposite sign along orthogonal directions, with the $\Delta \tau$ ratio in the range $-1/0.44 < r < -0.44$. Inspecting table 1, this corresponds to an intermediate situation between O-free Pd stripes and fully packed Pd-O stripes. In agreement with this prediction, the herringbone pattern, already hinted by the wavy stripes shown in fig. 1b, is observed in perfect form when the experiment is repeated, always under simultaneous Pd adsorption-desorption, with much lower oxygen coverage. The resulting pattern is displayed in fig. 3b, together with the O-free Pd and Pd-O stripe phases. These three patterns correspond to the three different solid-state phases of the overlayer in the phase diagram of fig. 3a.

In addition to the pattern shape and orientation, the stripe period shows a strong coverage dependence. Using the expression from the continuum model [9], the period, $D$, is given by,

$$D(\theta_{Pd}, \theta_O) = \frac{A e^{2\frac{\Delta \tau_{Pd}}{\Delta \tau_{O}}} n}{\cos \left(\frac{\pi (1 - 2\theta O)}{2}\right)},$$

where $\theta_{Pd}$ and $\theta_O$ are the Pd and O coverages, $\Delta \tau$ is the coverage-dependent stress difference between the
fully packed Pd and the partially covered oxygen stripes, $A$ and $B$ are the fitting parameters corresponding to the effective boundary width and a combination of energetic factors, respectively. The denominator accounts for the effect of stripes with unequal widths [9]. The stress difference, $\Delta\tau$, is found by a weighted average of the calculated values displayed in table 1 as the Pd coverage is varied. The excellent fit to the experimental data is shown in fig. 4. The only fit parameters, $A$ and $B$, are found to be $0.38 \pm 0.04 \text{Å}$ and $55.6 \pm 1.0 \text{N}^2/\text{m}^2$, respectively.

The fact that the experimental stripe period, shape and orientation show very good agreement with the continuum model (along with the DFT-calculated energy and stress parameters) confirms that the observed stripes represent the minimum of the free energy, which is determined by the competition between short-range (atomic bonding) and long-range (elastic) interactions. In other words, the stripes are in thermodynamic equilibrium. However, as described in the context of the measurements, the necessary condition for obtaining this equilibrium phase is simultaneous Pd adsorption-desorption, i.e. an open system. On the other hand, in the closed Pd-O system the totally segregated state seen in fig. 1c is found, which is apparently a local minimum of the free energy. The kinetic barriers in this complex free-energy landscape are overcome only with increased fluctuations via exchange with a heat-bath (3D gas), allowing the system to reach the real equilibrium configuration, i.e. the stripe pattern.

It is to be expected that a system with long-range interactions should face heightened kinetic barriers in its search for the global minimum in the configuration space. The basis for this statement is apparent in the formation of the equilibrium state. Generally, the predominant mechanism in the decay towards the minimum energy phase is by nucleation of spatially confined droplets, which progressively increase in size [29]. Such a possibility does not exist for a phase stabilized by long-range forces, which necessitate its formation in a coherent manner within spatially extended regions. The difference is clear in studies of dense and viscous lattice gas models. In particular, systems with short-range interactions alone do not show glassy behaviour, in spite of their slow dynamics at high density [30]. On the other hand, long-range interactions in a thermodynamically closed system may lead to broken ergodicity and infinite-lifetime metastable states [31].

Importantly, it has recently been shown that the ergodicity, which is broken in a closed system due to long-range interactions, is restored by coupling it to a heat-bath [32]. This distinction between thermodynamically closed (microcanonical) and open (canonical) systems in their decay to the thermal equilibrium state provides a direct explanation to our observations [33]. In other words, the simultaneous Pd adsorption-desorption condition is identified as coupling the otherwise-closed two-dimensional adsorbate layer to a Pd reservoir (three-dimensional Pd gas), which acts as a heat-bath and allows the formation of the equilibrium stripe phase.

In conclusion, we have shown the presence of a stress-induced Pd-O stripe phase on W(110). The anisotropy and the period of the stripes depend sensitively on the surface stress, and can be tuned by changing the relative coverage of the two species. Due to the presence of long-range interactions, the high-density lattice gas is able to decay into the equilibrium stripe phase only via material exchange with the gas phase. As a consequence of suppressed diffusivity, the pattern can be preserved by cooling to lower temperatures. This opens a new path towards fabricating highly ordered periodic structures via self-organization at the nanometer scale.

REFERENCES

[1] IBACH H., Surf. Sci. Rep., 29 (1997) 195.
[2] KERN K., NIEHUS H., SCHATZ A., ZEPPENFELD P., GOERGE J. and COMSA G., Phys. Rev. Lett., 67 (1991) 855.
[3] JONES D. E., PELZ J. P., HONG Y., BAUER E. and TSONG I. S. T., Phys. Rev. Lett., 77 (1996) 330.
[4] HANSON J. B., BARTELT N. C., SWARZENTRUBER B. S., HAMILTON J. C. and KELLOGG G. L., Phys. Rev. Lett., 79 (1997) 4226.
[5] POHL K., BARTELT M. C., DE LA FIGUERA J., BARTELT N. C., HRBek J. and HWANG R. Q., Nature, 397 (1999) 238.
[6] PLASS R., LAST J., BARTELT N. C. and KELLOGG G. L., Nature, 412 (2001) 875.
[7] VAN GASTEL R., PLASS R., BARTELT N. C. and KELLOGG G. L., Phys. Rev. Lett., 91 (2003) 055503.
[8] DE LA FIGUERA J., LÉONARD F., BARTELT N. C., STUMPF R. and MCCARTY K. F., Phys. Rev. Lett., 100 (2008) 186102.
Alerhand O. L., Vanderbilt D., Meade R. D. and Joannopoulos J. D., Phys. Rev. Lett., 61 (1988) 1973.

De'Bell K., MacIsaac A. B. and Whitehead J. P., Rev. Mod. Phys., 72 (2000) 225.

Andelman D., Brochard F. and Joanny J.-F., J. Chem. Phys., 86 (1987) 3673.

Whitesides G. M. and Grzybowski B., Science, 295 (2002) 2418.

Mentes T. O., Locatelli A., Aballe L. and Bauer E., Phys. Rev. Lett., 101 (2008) 085701.

Bauer E., Rep. Prog. Phys., 57 (1994) 895.

Altman M., J. Phys.: Condens. Matter, 22 (2010) 084017.

Locatelli A., Aballe L., Mentes T. O., Kiskinova M. and Bauer E., Surf. Interface Anal., 38 (2006) 1554.

Johnson K. E., Wilson R. J. and Chiang S., Phys. Rev. Lett., 71 (1993) 1055.

Wu P. K., Tringides M. C. and Lagally M. G., Phys. Rev. B, 39 (1989) 7595.

Nahm T.-U. and Gomer R., Surf. Sci., 373 (1997) 237.

Mentes T. O., Stojic N., Binggeli N., Niño M. A., Locatelli A., Aballe L., Kiskinova M. and Bauer E., Phys. Rev. B, 77 (2008) 155414.

Schlenk W. and Bauer E., Surf. Sci., 93 (1980) 9.