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# Model studies on heterogeneous catalysts at the atomic scale: From supported metal particles to two-dimensional zeolites



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# ABSTRACT

After a general introduction to model studies in catalysis, results from different research areas are presented: (a) adsorption and reaction on nanoparticles supported on thin oxide films. It is shown that carbonaceous deposits on specific nanoparticle sites influence hydrogenation reactions through control of hydrogen surface–subsurface diffusion. (b) 2D-3D-morphology, geometric, and electronic structure of supported metal nanoparticles partially in relation to doping of the support. We learn how dopants, as electron sources, even if they are well beneath the active surface influence nanoparticle morphology on surfaces. (c) Strong metal–support interaction (SMSI). Ultrathin oxide films are proven to become chemically active in CO oxidation as they form on deposited nanoparticles dependent on the oxygen chemical potential in the gas phase. (d) Adsorption and reaction on two-dimensional silicates and alumino silicates in ordered and vitreous phases. Bilayer silica films on Ru(0001) are transformed by Al doping into 2D-aluminosilicates and shown to exhibit binding hydroxyl with an acidity close to 3D-zeolites, such as chabazite.

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# 1. Introduction

Heterogeneous catalysts are complex materials consisting of combinations of different components, such as an oxide support and an active disperse metal. The classical model system for these materials, in the past, has been the metal single crystal [1–3]. The reason for this choice can be made transparent by inspecting Fig. 1 [4]. It shows STM images of Pd nanoparticles, which were grown by physical vapor deposition of Pd onto a thin alumina film. The left image showing a larger area reveals the presence of nanoscopic Pd single crystals exposing well-defined facets. The image at atomic resolution supports the idea that the topmost facet, indeed, exhibits (111) morphology and structure. Exactly, those observations, made in the 1960s of the last century, at the time with transmission electron microscopy [1], were the basis for the development of single-crystal model systems of defined orientation within surface science, in order to understand the elementary steps of chemical reactions at surfaces [5,6]. Our understanding of catalysis, and heterogeneous catalysis, in particular, is, to a large extent, based on these investigations of model systems. The enormous impact of metal single-crystal model surface chemistry, pioneered by physical chemists that led to the 2007 Nobel Prize in chemistry for Gerhard Ertl, has been the basis for an outstanding

\* Corresponding author. *E-mail address:* freund@fhi-berlin.mpg.de (H.-J. Freund). success story [6]. The simplicity of this model approach is particularly appealing, and it is this aspect that has led to some of the groundbreaking ideas influencing our thinking about catalytic reactions. However, it is also very obvious that a full understanding of all factors influencing catalytic reactions requires a considerable increase in complexity in the structure and morphology of the model systems, actually, taking the finite size of the disperse metal particles and the support-metal interface into account [7,8]. The important question is how much complexity is necessary to catch the relevant aspects [9]. It is obvious, however, that the only strategy that will lead to an understanding at the atomic level is to proceed from the simple to the complex and not the other way around!

We have developed concepts to prepare more complex model systems based on single-crystalline oxide films, which are used as supports for metal and oxide nanoparticles [10–13]. Together with the development of novel instrumentation [14], this approach allows us to study the factors influencing catalysis at the atomic level using the tools that have been developed, and so successfully applied, during the last thirty years in surface science, including those based on scanning probe techniques [15,16].

Here, we report on different areas, where such an approach has allowed us to directly establish structure/morphology-reactivity relations previously impossible and thus explain some surprising observations made in real catalysis. Using a model system similar to the one shown in Fig. 1, we investigate the question, why one

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Fig. 1. Scanning tunneling microscopic images of Pd deposited on a thin alumina film at low (left) and high (right) resolution [4].

needs the presence of deposited carbon on nanoparticles to sustain hydrogenation catalysis in the conversion of alkenes to alkanes [17,18], and how does this relate to the higher degree of flexibility of nanoparticles as compared to metal single crystals influencing the adsorption enthalpy of molecules [19,20]. We address in a further case study the morphology and structure of nanoparticulate supported gold occurring as small, flat raft-like aggregates, and the role of the oxide metal interface in relation to its unmatched reactivity in CO oxidation [21-23]. Then, we pursue the study of charge transfer through oxide films as a function of their morphology and thickness to reveal novel mechanisms of molecular activation on oxides in the case of encapsulation of nanoparticles and strong metal-support interaction (SMSI) [24]. In a final case study, we reveal our initial experiments to prepare two-dimensional well-ordered silica and alumino-silica surfaces in an attempt to study the surface science of zeolites [25,26].

#### 2. Experimental

The experiments were carried out in a number of custom made ultrahigh vacuum (uhv) systems, including equipment to bring samples to ambient conditions with direct transfer. A broad variety of experimental techniques have been applied [14]. These include scanning tunneling microscopy in uhv and ambient conditions, non-contact atomic force microscopy, elastic and inelastic electron scattering, photon STM, infrared Fourier transform spectroscopy, non-linear optical techniques, electron spin resonance spectroscopy, resonant nuclear reaction analysis, molecular beam techniques, microcalorimetry, and combinations thereof. The materials were prepared by epitaxial growth of oxides by reactive physical vapor deposition in uhv. Surface modifications and exposure to gases were carried out via specifically designed gas-handling systems.

# 3. Results and discussion

#### 3.1. Carbon deposits on supported Pd nanoparticles and its flexibility

Fig. 2 shows a comparison of hydrogenation rates of trans-2-butene to  $d_2$ -butane obtained from a series of pulsed molecular beam experiments, in which  $Fe_3O_4(111)$  supported Pd nanoparticles of 4 nm average diameter (similar to the Pd particles on alumina imaged in Fig. 1) were exposed to a continuous  $D_2$  molecular beam and a modulated cis-2-butene (which was chosen as the model di-methyl substituted ethene) molecular beam at 260 K [17]. The figure shows that the product response curves obtained for hydrogenation under steady-state conditions in comparison with similar

results on Pd(111). The rates were calculated by averaging the reaction rates from the last 30 long pulses of the experiment and were normalized to the number of the surface Pd atoms for quantitative comparison. Shown are the hydrogenation rates plotted as a function of time on Pd(111) (upper row) and Pd particles (lower row) on the C-free (black curves) and C-containing (gray curves) surfaces. It is noteworthy that on both C-free surfaces, no sustained hydrogenation is observed [17,18,27,28]. Fig. 3 shows the production rate of butane-d<sub>2</sub> resulting from the reaction of cis-2-butene with D<sub>2</sub> over a C-free (Fig. 3a) and C-precovered (Fig. 3b) Pd-supported model catalysts at 260 K, which were pre-saturated with D<sub>2</sub> prior to the olefin exposure. A sequence of cis-2-butene pulses, consisting of 50 short (4 s on, 4 s off) and 30 long (20 s on, 10 s off) ones, was applied using an independent beam source concomitant with the continuous D<sub>2</sub> exposure. The reaction was carried out in a large excess of D<sub>2</sub> with the ratio of D<sub>2</sub>:cis-2-butene molecules amounting to 570. On the pristine Pd particles (Fig. 3a), the initial period of high hydrogenation activity on D-saturated catalyst is followed by a decrease in the reaction rate to zero under steadystate conditions. Remarkably, carbon deposition prevents the suppression of the hydrogenation pathway in the steady state and results in persisting hydrogenation activity at the initially high level (Fig. 3b). It should be noted that the reaction rate of the



**Fig. 2.** Average hydrogenation rates on Pd/Fe<sub>3</sub>O<sub>4</sub>(111)/Pt(111) model catalyst, in comparison to a Pd(111) single crystal. The panel shows the situation for the model catalysts without pre-deposition of carbon on the left and after carbon deposition on the right [17].

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**Fig. 3.** The hydrogenation rates of cis-2-butene at 260 K over initially  $D_2$ -saturated (a) clean and (b) carbon-precovered Pd/Fe<sub>3</sub>O<sub>4</sub>(111) model catalysts [17]. The cartoons on the right indicate how we envision carbon decoration on the edges of the Pd particle.

cis-trans isomerization/H-D exchange, i.e., the reaction competing with hydrogenation, according to the Horiuti–Polanyi mechanism [29], in the steady state, remains very similar on both C-free and C-precovered Pd particles, indicating that the surface concentrations of D and butyl-d1 are not substantially affected by C deposition under the investigated conditions. The microscopic reasons for the vanishing hydrogenation activity on the C-free catalyst could be deduced recently from the combination of H-depth profiling nuclear reaction analysis (NRA) measurements, and transient molecular beam experiments [26,30] carried out at different H<sub>2</sub>/D<sub>2</sub> pressures [18], which provided experimental evidence that the presence of subsurface H species [30] is necessary for hydrogenation. In view of this finding, the initially high hydrogenation rate (probed by the initial short pulses) on both C-free and C-precovered Pd particles can be ascribed to the high availability of subsurface D atoms directly after pre-saturation. However, the abundance of these species on the C-free particles cannot be maintained at the initially high level after prolonged olefin exposure, resulting in vanishing hydrogenation rates. This behavior was suggested [18] to be a consequence of the slow D subsurface diffusion hindered by co-adsorbed hydrocarbons. In contrast, sustained hydrogenation activity of the C-precovered particles indicates effective replenishment of the subsurface D species even under steady-state reaction conditions. It is possible that hydrogen subsurface diffusion through the C-modified edge sites can be the reason for this observation, as was recently supported by theoretical calculations [31]. Summarizing, we find, that due to the considerably higher flexibility of the more or less strained lattice of the supported small particle as compared with a single crystal, and due to the size of the object, the interior of the particle actively participates in hydrogenation reactions, by hydrogen accommodation at conditions different from the bulk material. Although the important role of carbonaceous deposits in the activity and selectivity of transition metals has been recognized for a long time in the catalytic community [32-34], it only via such model studies that one reaches an understanding at the atomic level. On single crystals, the deposition of several well-defined carbonaceous species, among them ethylidyne, has been identified [35,36], but they have been merely considered to be spectators blocking sites: Now, we know that such dehydrogenated hydrocarbon derivatives may fulfill different functions on appropriate systems. While hydrogenation is an example for the influence of flexibility under reaction conditions, there are more subtle effects for nanoparticles, which

already show up for relatively large supported nanoparticles of 4-8 nm in adsorption experiments, i.e., the variation in the adsorption enthalpy with particle size in comparison with single crystals [19]. In order to study such subtle effects, one cannot use temperature-programmed desorption techniques - otherwise typically applied techniques in surface science to determine adsorption energies - because, during heating, the supported particles undergo considerable changes. The method of choice in this situation is single-crystal adsorption calorimetry (SCAC). This technique, described in detail elsewhere [37,38], allows one to determine adsorption enthalpies of molecular adsorption as well as dissociative adsorption in a thermodynamically profound fashion on a small area single-crystal-based sample. The experiments summarized in the following have been conducted on the same kind of model systems used in the reactivity studies discussed above [20,38]. To address the dependence of the initial O adsorption energies on particle size in more detail [20], for example, four different supported model systems with different Pd particle sizes ranging from  $\sim$ 2 nm to  $\sim$ 8 nm were investigated by SCAC. The initial heats of adsorption on all supported systems are displayed in Fig. 4d as a function of the nominal Pd deposition thickness together with data measured on Pd(111). Each point in this curve represents an average of 4-6 independent measurements on freshly prepared model systems. The dependence of the O adsorption energy on the particle size shows a clear trend: It strongly rises from  $\sim 205 \text{ kJ} \text{ mol}^{-1}$  to 250–275 kJ mol<sup>-1</sup> when changing from Pd(111), where oxygen adsorbs at threefold-hollow sites, to large Pd nanoclusters, where oxygen first occupies particles edges, as evidenced by IRAS data, which have been shown elsewhere [20]. However, if the local adsorption site (the edge/corner site) was preserved, and only the particles size was reduced, the initial oxygen adsorption energy was observed to strongly decrease, reaching the value of 205 kJ mol<sup>-1</sup> on the smallest Pd nanoparticles. Thus, the reduction in the Pd cluster size results in an opposite trend - decrease in the adsorption energy – with the magnitude that is comparable with the effect of the change in the local adsorption environment. Note that the total number of oxygen atoms, contributing to the initial adsorption energies (on the pristine Pd nanoparticles) reported in Fig. 4a, amounts typically to only a few O atoms per particle or less than 0.04 per surface Pd atom. Since the number of surface Pd atoms on each particle is much larger (~140 on the smallest and ~1100 on the largest Pd clusters) than the number of adsorbed O atoms, we can safely assume that the repulsive interaction between adsorbed O atoms does not play any significant role for these low oxygen surface coverages. The latter trend - decreasing adsorption energy with decreasing particle size- coincides with the trend previously observed in our studies for CO adsorption in direct calorimetric experiments on the same model system [19]. The initial CO adsorption energy on the same Pd nanoparticles is plotted in Fig. 4b for comparison, showing a pronounced decrease in the adsorption energy for small clusters. In contrast, the change in the adsorption site from a threefold hollow on Pd(111) to the strongest binding site on Pd nanoparticles obviously does not result in an increase in the adsorption energy as observed for oxygen. This observation agrees well with the literature data, suggesting that the degree of coordination of the surface Pd atoms both on low Miller index surfaces and on stepped Pd surfaces does not significantly affect the CO binding energy (for the full discussion see Ref. [39]).

Two alternative microscopic effects can be put forward to rationalize the observed decrease in the initial adsorption energy on Pd nanoparticles for both oxygen and CO adsorption: weakening of the chemisorptive interaction [40–46] and reduction in the van der Waals attraction [47,48]. The decrease in the chemisorption energy has been previously predicted in a theoretical study for Pd clusters [40–42] in the size range corresponding to a scalable





**Fig. 4.** Initial adsorption energies (i.e. at lowest coverage) for  $O_2$  (a) and CO (b) molecules plotted as a function of the nominal thickness of the deposited Pd. The sizes are given as determined from STM measurements on Pd/Fe<sub>3</sub>O<sub>4</sub>(111)/Pt(111) model catalysts in comparison to adsorption energies on Pd(111). Error bars show the standard of the mean of several measurements.

regime. These computational results showed that interatomic Pd-Pd bond length in small metal particles decreases [49] with decreasing particle size, and this effect results in systematically lower adsorption energies than on the clusters with a bulk terminated geometry, i.e., with all interatomic distances d (Pd-Pd) = 275 pm. This finding agrees well with the principle of bond order conservation [45,46]: In the bond-contracted particles, one expects weaker adsorption bonds due to stronger binding within the particle. A second reason for the decrease in adsorption energy of a gas-phase molecule on the small metal clusters is a weakening of the van der Waals interaction strength [47,48]. This type of interaction is induced by the dynamic response of bulk electrons of the metal to charge density fluctuations in an adsorbed molecule, and since smaller clusters contain fewer electrons available for the dynamic response, the dispersive interaction strength weakens. This effect is expected to play a more important role for the interaction of CO with Pd clusters and be of minor importance in case of oxygen.

Via such studies, one is forced to acknowledge that the flexibility of supported metal nanoparticles as a consequence of its limited size and the lattice strain induced through bonding to the support, results in complications for the description of adsorption and reaction at surfaces that need to be taken into account, also by theory, if one wants to be able to properly catch the details of a heterogeneous catalytic reaction.

#### 3.2. Reactivity of supported flat Au nanoparticles

In the 1980s of the last century, Masatake Haruta [50] discovered the high catalytic CO oxidation activity of Au nanoparticles anchored to a reducible support. The report initiated a very active search for the reasons of this unexpected discovery, as bulk Au is not exactly known for its chemical activity. The research group of

the late Wayne Goodman [51] reported the first model studies on this system by preparing Au nanoparticles on a bulk TiO<sub>2</sub>(110) surface and on thin TiO<sub>2</sub> films and studying their relative CO oxidation activity. He compared flat raft-like (two-dimensional) Au particles with three-dimensional or spherical particles and found that the latter were more than three times less active than the former. This turned out to be in line with observations using aberration-corrected TEM studies on TiO2 and Fe2O3 supported Au particles [52,53], indicating that the most active particles exhibit a raft-like morphology and very limited size. On the one hand, those were very interesting observations, and on the other hand, they also led to questions, such as why would a metal aggregate that usually exhibits three-dimensional morphology when grown on an oxide surface due to surface energetics (as in Fig. 1 for Pd on alumina) in the case of Au grow two-dimensional on an oxide substrate? A publication from the groups of Gianfranco Pacchioni and Uzi Landmann pointed toward a solution on the basis of a theoretical prediction [54,55]: If one adsorbs a Au<sub>20</sub> cluster (which Lei-Shang Wang showed to have the structure of an almost ideal tetrahedron having been cut out of a Au lattice [56]) onto a MgO(100) surface, it keeps its three-dimensional structure as expected based on total energy density functional calculations. If, however, one adsorbs the same moiety on a two layer thick MgO(100) film, supported on a Mo(100) metal surface, it prefers, based on the same calculations, to adopt a two-dimensional, raftlike structure or in other words a monolayer island, while the three-dimensional structure, stable on the bulk oxide, is higher in energy. As reason, the authors argue [54,55] that electron transfer from the oxide support-metal interface to the electronegative Au particle leads to this morphology, and it is the electron transfer that then induces the local wetting of the oxide. If this prediction was true, we would have to look for sources of electrons within the support to induce the raft-like structure. We will come back



Fig. 5. STM images and height profiles (right panel) of Au islands grown on MgO(100) films of two thicknesses (as indicated in monolayers, ML). The oxide films have been grown on a Ag(001) surface.

to this after we have experimentally verified the above theoretical prediction.

In Fig. 5, two sets of panels are depicted, each showing two STM images and a STM contour along the white lines indicated in the images [57-59]. The upper panel has been recorded for Au deposited from an evaporation source onto three layer thick MgO(100) film supported by an Ag(100) substrate at low temperature and then annealed to 210 K and to room temperature. An equivalent set is reproduced below for an eight layer thick MgO(100) film on Ag(100). The comparison of the line profiles on the right reveals the interesting result: The Au islands on the 3-layer film are all two-dimensional monolayer islands, and the Au islands on the 8layer film are three-dimensional, half-spherical in shape! This appears to corroborate the theoretical prediction [54,55]. However, it also implies that an 8-layer MgO(100) film, indeed, represents the situation encountered on a bulk oxide surface. This appears a bit surprising and needs further attention. As shown in Fig. 5, both systems can be imaged using STM, i.e., tunneling through films of both thicknesses is possible. Why, then, does the 8-layer film represent the bulk, so that there is no stable electron transfer leading to two-dimensional growth not possible? The answer may be provided, if we analyze the different contributions that contribute to the process of electron transfer, which are schematically summarized in Fig. 6 [54,60]. The cartoon on the left-hand side shows a

single Au atom located on top of a double-layer MgO(100) on either Ag or Mo. The energetic ingredients in an electron transfer from the oxide support-metal interface are the ionization potential for an electron from this interface, the electron affinity of the adsorbed Au atom, and then any further energetic contribution that could lead to a stabilization of a transferred electron on the Au atom. One such contribution is the polarization of the substrate, which establishes an image charge. Another one is a possible polaronic distortion of the oxide film that would additionally stabilize the negative charge on the surface. It is this contribution that is decisive between the 3-layer and the 8-layer film. A polaronic distortion involves the phonons of the oxide [60]. Oxides, in general, and MgO, in particular, exhibit a spectrum of rather high energetic phonons. A film of 8 layers of MgO exhibits a phonon spectrum similar to the bulk, while the phonons of a 3-layer film are considerably softer, i.e., they may be excited with much lower energy as shown by Mario Rocca's group [61]. Therefore, the 3-layer film is able to respond to the charge and distort, while the 8-layer film is not [62]. It is this subtle difference that stabilizes the negative charge on the thin film. It is possible to provide experimental evidence for this polaronic distortion for a slightly different adsorbate system via electron spin resonance spectroscopy [60]. In this case,  $O_2$  forms an  $O_2^-$  on a thin MgO(100) film, which is in effect the proposed precursor for thin oxide film growth as discussed by Cabrera



Fig. 6. Schematic representation of the charge transfer process from the metal substrate through the thin oxide film to an adsorbed Au atom [54,60].

and Mott [63]. From the measured g-tensor components, the polaronic distortion can be seen in the g<sub>zz</sub>-tensor-component anomaly [60]. Coming back to Au, one may convince oneself that it is really the high electron affinity of Au that enables electron transfer by comparing individual Au atoms on a 3-layer MgO(100) film with Pd atoms on the same substrate. This is shown in Fig. 7. In the left panel, the Au atoms reveal an unusual next neighbor distribution which is not diffusion limited, while the Pd atoms in the right-hand panel show a typical diffusion limited distribution [58]. The reason for this difference is obvious: Au is electronegative (2.54 on the Pauling scale) and thus negatively charged which leads to a spatial distribution where neighboring Au atoms avoid contact, while Pd atoms are much less electronegative (2.2 on the Pauling scale) and remain neutral, and thus show the classical pattern of a diffusing adsorbate. The collected information supports the idea of electron transfer through thin films that can lead to wetting of oxides by a metal as suggested by Pacchioni and Landmann [55]. However, how does this relate to the situation on a bulk reducible oxide, addressed in the beginning of this section, where raft-like structures have been suggested to be the active state of Au particles in CO oxidation? In other words, where is the source of electrons in the bulk material that induces the wetting phenomenon? The answer is dopants! Imagine an oxide lattice with rock salt structure. A metal ion of this host lattice is replaced by another metal ion, which may also adopt oxidation states different from the one sufficient to replace the host ion. If the system provides electron traps with proper energy balance, then an electron transfer may occur involving the oxidation of the dopant. If the electron trap is located on the surface, i.e., Au atoms and the energy balance is favorable, an electron may be transferred to the Au inducing wetting of the oxide surface by the metal. Fig. 8 shows STM images of Au nanoparticles which have been grown on a CaO(100) surface. In the left image, the substrate represents a thick (60 layers) pristine CaO(100) substrate, in the image on the right, the same substrate has been doped with a very small percentage of Mo ions [64-66]. Mo may adopt the oxidation state 2+ of the Ca ions, but may also be oxidized to Mo<sup>3+</sup> or higher oxidation states. The insets in the large-scale images reveal the morphology of the individual nanoparticles. While the particles on the pristine substrate are three-dimensional, the particles on the doped substrate are two-dimensional raft-like, as indicated by the Moiré pattern on the particles. It is, therefore, clear that dopants in support materials may represent electron sources to induce two-dimensional growths of metal particles on bulk oxide surfaces and explain the surprising experimental finding for the high activity of the raft-like structures reported by a number of laboratories. A particularly interesting observation is the fact that the influence of the dopants is not local, i.e., dopants in the deeper subsurface region influence adsorption [64].

This finding also has a number of implications for controlling the transfer of charge. These implications may be deduced from the rules that have been developed in the mature field of semiconductor physics, where dopants are regularly used to control electronic properties [67]. For example, if the lattice contains other electron traps such as color centers, those will compete for the charge. If, for example, Li, as an under-valent dopant (opposed to Mo which would be called over-valent) is introduced into the CaO lattice, the electron transfer to the surface is suppressed and the Au nanoparticles grow again in three-dimensional morphology [66]. After those experimental results had been published, Hannu Häkkinen's group has published a density functional based analysis of the various energy contributions that may lead to charge transfer [68]. Their concept is similar to the acid-base concept put forward by Horia Metiu [69]. They basically show via an appropriately constructed Born-Haber cycle, that the local redox properties of the Au/surface and the Dopant/oxide pairs are responsible for the electron transfer. With this in mind, one may predict dopants that are effective in electron transfer to the surface, given that no other electron traps interfere. Another consequence of the laid out scenario is that the presence of gas-phase species could influence particle growth, in particular, if the gas-phase species had a large electron affinity, as, for example, oxygen molecules [23]. In fact, if one attempts to grow Au nanoparticles on doped CaO(100) surfaces in the presence of molecular oxygen, a competition between Au and O<sub>2</sub> for the dopant charge is observed, which leads to a switch over in growth morphology from two-dimensional to three-dimensional depending on oxygen coverage (and pressure). This is shown in Fig. 9 where STM topographic images of 0.5 ML gold deposited onto (a) 50 ML thick CaO without dopants, (b) self-doped CaO films of 25 ML thickness in UHV and (c-f) in increasing  $O_2$  partial pressures that are (c)  $5 \times 10-8$  mbar, (d)  $5 \times 10-7$  mbar, (e)  $5 \times 10-6$  mbar and (f)  $2 \times 10-5$  mbar are depicted [23]. Note that the two-dimensional Au deposits appear as depressions on the thick CaO films at the used positive bias. To support the above statement, in Fig. 10, histograms [23] of Au particle shapes for growth conditions identical to the ones shown in Fig. 9 are collected: (a) non-doped CaO in UHV, Mo-doped CaO (b) in UHV, (c) in  $5 \times 10-8$  mbar, (d) in  $5 \times 10-7$  mbar and (e) in  $5 \times 10-6$  mbar of oxygen. The histograms in (c) and (d) display bimodal shape distributions due to the coexistence of two-dimensional and three-dimensional deposits. Fig. 10f indicates the evolution of the number density of two-dimensional and threedimensional deposits as a function of the O<sub>2</sub> background pressure during deposition. Even though this does not provide concrete



Fig. 7. STM images of Au atoms adsorbed on a thin (3 ML) MgO film on Ag(100) in comparison to Pd atoms adsorbed on the same supports [57].



Fig. 8. STM images of Au islands on thick pristine (left) and Mo-doped CaO(100) films (right). The small insets show the higher resolution [64].



Fig. 9. STM typographic images of 0.5 ML deposited onto (a) 50 ML thick Cao without dopants, (b) Mo-doped CaO films of 25 ML in UHV and (e) and (f) in increasing  $O_2$  partial pressures from  $5 \times 10^{-8}$ – $2 \times 10^{-5}$  mbar [23].



**Fig. 10.** Histograms of Au particle shapes for growth conditions identical to the ones in Fig. 9a–f shows the number densities of 2D and 3D deposits as a function of the O<sub>2</sub> background pressure during deposition [23].

proof, these results imply that if we would anneal two-dimensional Au nanoparticles in the presence of an appropriate oxygen pressure, they would switch to three-dimensional growth [70]. The latter experiments imply that  $O_2$  and Au compete for the

electronic charge, which, in turn indicates that  $O_2$  may be transformed to  $O_2^-$  which, if this could be proven, might be an active component in interesting oxidative hydrocarbon reaction.

#### 3.3. Strong metal-support interaction

Strong metal-support interactions (SMSI) observed with particular catalyst systems, in which metal particles (such as Pd and Pt) strongly interact with a reducible support (such as titania), and are covered by a thin oxide film upon heating to elevated temperature, usually lead to reduced catalytic activity. As a consequence of SMSI, the oxide film exhibits a strong attenuation of adsorption capacity and a deactivation of the system [71-73]. There have been many attempts to elucidate, even on model systems, the nature of the migrating oxide film [74,75]. The best studied system is Pt/  $TiO_2(110)$ , but in this case, the attempts have been rather unsuccessful. Very recently, we succeeded in preparing such a SMSI model system for which we are able to identify the atomic structure of the encapsulating oxide film [76–78]. The system consists of Pt supported on a Fe<sub>3</sub>O<sub>4</sub>(111) film grown on a Pt(111) single crystal. Fig. 11 shows an STM image of this system after heating to 850 K. After this treatment, the CO adsorption capacity is drastically reduced, which is typical for a SMSI effect. A close look at the STM images reveals well structured and facetted nanoparticles. Moreover, atomically resolved images reveal corrugation that does not stem from Pt, but rather from a well-ordered bilayer FeO(111) film, well described and characterized in the literature [79–82]. As the oxide film that covers Pt particles has been identified, one may reduce the complexity of the model system by studying the properties of the bilayer FeO(111) film on a Pt(111) single crystal. Its structure has been studied in detail and characterized at the atomic level. The 10% misfit between the FeO(111) lattice constant and that of Pt(111) gives rise to a typical Moiré-pattern in the STM image. This film is unreactive under ultrahigh vacuum conditions [83]. The situation changes, however, dramatically if one tests the system with respect to CO oxidation at ambient conditions (1 atm) in a reactor with careful control of the relative amounts of oxygen (one part, 20 mbar), carbon monoxide (two parts, 40 mbar), and helium as buffer gas [83,84]. CO<sub>2</sub> production was measured by gas chromatography. Only initial reaction rates (toward zero conversion) were used to compare data on different systems. If one ramps the temperature linear with 1 K min<sup>-1</sup> from 300



Fig. 11. STM image of Pt nanoparticles on  $Fe_3O_4(111)$ . The inset shows an image with atomic resolution taken on the top facet of one of the Pt particles.

to 455 K, CO oxidation ignites at 430 K. The interesting observation is that this FeO/Pt(111) system at this temperature is more than an order of magnitude more reactive than clean Pt(111). Usually, SMSI leads to an attenuated activity, while here we observe a strong enhancement! Further studies as a function of different gas compositions, as well as thermal desorption studies, STM investigations, and detailed DFT model calculations reveal an interesting scenario that allows us to understand this phenomenon [78]. The scenario is depicted in Fig. 12 [78,85,86]. The gas phase sets the chemical potential of the system. The shown steps are based on density functional calculations. Oxygen interacts with the bilayer FeO film on Pt(111) by pulling an iron atom up above the oxygen layer. This lowers the work function at the interface locally to allow for an electron transfer toward oxygen accompanied by the formation of a transient  $O_2^-$  molecule, which dissociates and results, at higher oxygen coverage, in the formation of a local O-Fe–O trilayer. There is a range of experimental evidence for the existence of such a trilayer [78,85,86]. The middle panel in Fig. 12 shows an STM image of such a trilayer formed in situ at elevated O<sub>2</sub> pressure in a microscope. The trilayers appearance is mostly determined by the Moiré structure of the FeO bilayer and fills 80-90% of the surface as thermal desorption spectra indicate. The STM images are completely consistent with the structure suggested by the calculation, although the latter does not reproduce the patched morphology due to the enormous size of the unit cell, which was impossible to implement, but necessary to fully reproduce the details. Nevertheless, if the trilayer is exposed to carbon monoxide it oxidizes the incoming CO into CO<sub>2</sub> by means of an Eley-Rideal mechanism, leaving behind an oxygen vacancy in the film. At sufficiently high oxygen pressure the oxygen vacancy is filled again and the trilayer is sustained. If, however, the gas phase is oxygen-poor, the reaction finally stops because the trilayer is destroyed. Experimentally, we have confirmed [84] that the iron oxide film de-wets the Pt(111) surface under CO-rich reaction conditions by forming small iron oxide particles and exposing the Pt(111) surface, which then determines the reactivity of the systems, in contrast to a recent study in the literature [87]. Heating the de-wetted surface in vacuum again leads to the formation of the FeO double layer, which then, at higher oxygen pressure, may be transformed into the trilayer again.

As suggested in this section thin oxide films exhibit very interesting catalytic activity which will depend on the materials combination between the metal and the covering oxide film. It may be possible to design, by choosing appropriate combinations of materials, systems with specific catalytic reactivities. Temperature-programmed desorption results indeed reveal that a trilayer film contains much more weakly bound oxygen than in a pristine bilayer FeO film [78]. Therefore, it appears that the binding energy of the oxygen in the top-most layer governs the activity of the system. To verify this hypothesis, we have investigated other transition metal oxide films on the same Pt(111) support. For comparison, we also studied reactivity of oxide films prepared on Ru(0001) [88].

Fig. 13 collects 32 amu (O<sub>2</sub>) traces in TPD spectra resulting from re-combinative atomic oxygen desorption of a series of ultrathin film systems, all exposed to 20 mbar of O<sub>2</sub> at 450 K for 10 min. Except ZnO, these films showed desorption signals at temperatures, which are much below the "decomposition" temperature (1000–1200 K) and follow a series Ru  $\ll$  Fe < Mn. In turn, the CO oxidation rate measured under identical conditions (10 mbar CO; 50 mbar O<sub>2</sub>; He balance to 1 bar at 450 K) over these closed films follows the opposite row, i.e. Ru  $\gg$  Fe > Mn > Zn [89]. Clearly, there is a correlation between the activity and the presence of weakly bound oxygen species on the film surfaces at high chemical potential of oxygen which is basically set by oxygen partial pressure.



**Fig. 12.** STM image of the active trilayer iron oxide phase. The inset shows an image at higher resolution. The panels surrounding the image represent the individual steps in forming the active trilayer phase starting from FeO(111)/Pt(111) and its reaction with CO to form CO<sub>2</sub>, as revealed by DFT calculations [75,86].



**Fig. 13.** A 32 amu ( $O_2$ ) signal in thermal desorption spectra of ultrathin oxide films grown on Pt(111) and exposed to 20 mbar  $O_2$  at 450 K for 10 min. Data for a "native" Ru oxide films (RuO<sup>\*</sup>) grown on Ru(0001) are shown for comparison.

# 3.4. Two-dimensional silica and aluminosilicates toward a surface science of zeolites

Silica belongs to the most important supports in catalysis [1]. For more than a decade has the group been involved in the preparation of well-ordered silica films [25,90–98]. The first successful attempt led to a monolayer SiO<sub>2.5</sub> film on Mo(112) that has been used frequently by a number of groups as a model support [90,99]. This film consists of a hexagonal network of corner

shared SiO<sub>4</sub> tetrahedra, with the one Si-O bond, not associated with the intra silica network establishing a strong bond to the Mo(112) substrate. Since the stoichiometry of this film did not properly represent silica with a 1:2 Si:O ratio, the search continued and led to the preparation of a double-layer SiO<sub>2</sub> silica film, which consists of two single layer corner shared SiO<sub>4</sub> networks as in the case of the film on Mo(112) where, however, the Si-O bond formed to bind to Mo is now used to bind to the other identical monolayer silica network. This double-layer film is only weakly bound to a Ru(0001) substrate [91]. It comes as a crystalline film, and, depending on preparation conditions also as a vitreous film [25,95–97]. An equivalent to the originally prepared monolayer film on Mo(112) may also be prepared on Ru(0001) and the properties may directly be compared with the double-layer film [25]. Fig. 14 shows the structures of the two thin film systems, addressed above together with their vibrational spectra. The spectra are shown in order to provide evidence that the two systems, although similar in symmetry, may be clearly differentiated via their vibrational properties [100]. The experimental observations are very well supported by density functional calculations from the group of Joachim Sauer [90,91,100]. In fact, the structural parameters are, to a large extent, based on the theoretical studies [90,91,100]. While we are not going through a detailed analysis of the spectra, which has been presented elsewhere [90,91,100], it may suffice to mention that the strongest phonons at  $\sim 1100$ and 1300 cm<sup>-1</sup> refers to the Si-O-Si and Si-O-metal bonds, respectively. Fig. 15 collects STM images of the crystalline (a) and vitreous (f) structures together with schematic representations of the structures [25]. While the crystalline film consists of six membered rings only (b and c), the vitreous film exhibits a variation in ring sizes varying between 4 and 9 membered rings (g and h) still with a maximum at the six membered ring size, as predicted by Zachariasen in 1932 [95-97,101]. The sizes follow a



**Fig. 14.** Schematic structures (a) of the  $SiO_{2.5}$  monolayer and (b) the  $SiO_2$  double layer on Ru(0001) as derived from DFT calculations. Both, the top and the side views are presented. Below the experimental IRAS spectra of the two systems are shown for comparison [100].

log-normal distribution. Both films are being explored for their use as model supports for metals. In fact, metal atoms have already been adsorbed onto and into them, and their electronic structure is explored at present (d, e, g, j).

A direct comparison between Au (Fig. 15d and i) and Pd (Fig. 15e and j) adsorbed on freshly prepared silica films with cut-outs of crystalline (Fig. 15d and e) and vitreous (Fig. 15i and j) patches is provided. Au adsorbs only on vitreous while Pd can adsorb on both phases. Au seems to favor larger pores as adsorption sites [102].

Another interesting research area for our group with respect to silica films is to replace silicon atoms by other metal atoms. One particular direction is the exploration of aluminosilicates as models for two-dimensional zeolites [103-105]. As it turns out, both, the monolayer as well as the double-layer silica films allow for the preparation of aluminosilicates following the Löwenstein rule for nearest Al-Al distances. The expectation is that due to the substitution of the quatrivalent Si by the trivalent Al the oxygen atoms in bridging positions try to compensate the valence imbalance by picking up hydrogen [106]. However, due to the presence of the metal substrate, the valence imbalance is screened by the metal for the monolayer film, so that hydrogen take up cannot be detected [103]. On the other hand, the double-layer film in the top layer does, indeed, take up hydrogen as may be unambiguously proven by isotopic hydrogen/deuterium exchange [104]. This renders the aluminosilicate double-layer film to be a flat or two-dimensional model for zeo-



**Fig. 15.** (a and f) Zachariasen's schemes for the atomic arrangements of crystal (a) and a glass (f) form [101]b, c, g, and h. Atomically resolved crystalline and vitreous regions of the thin silica film (the scan area of these images is  $3.5 \text{ nm} \times 2.4 \text{ nm}$ ). Green balls represent the assigned position of Si atoms, red balls for O atoms. (b) STM image of a crystalline area showing the position of Si atoms ( $V_S = 3.0 \text{ mV}$ ,  $I_T = 100 \text{ pA}$ ). (c) STM image of a crystal line patch showing the arrangement of O atoms ( $V_S = 100 \text{ mV}$ ,  $I_T = 100 \text{ pA}$ ). (g) STM image of a vitreous area revealing the positions of Si atoms ( $V_S = 2.0 \text{ V}$ ,  $I_T = 50 \text{ pA}$ ). (h) STM image of a vitreous area showing the arrangement of O atoms ( $V_S = 1000 \text{ mV}$ ,  $I_T = 100 \text{ pA}$ ). (g) STM image of a vitreous area showing the arrangement of O atoms ( $V_S = 1000 \text{ mV}$ ,  $I_T = 100 \text{ pA}$ ). (h) STM image of a vitreous area showing the arrangement of O atoms ( $V_S = 1000 \text{ mV}$ ,  $I_T = 100 \text{ pA}$ ). (g) STM image of a vitreous area showing the arrangement of O atoms ( $V_S = 1000 \text{ mV}$ ,  $I_T = 100 \text{ pA}$ ). (g) atomet of  $V_S = 1000 \text{ mV}$ ,  $I_T = 100 \text{ pA}$ ) [97]. In (d, e, i, and j) STM images of sample preparations with adsorbed metal particles are shown (scan size of  $5.3 \text{ mx} \times 3.3 \text{ mm}$ ,  $V_S = 2 \text{ V}$ ,  $I_T = 100 \text{ pA}$ ). A direct comparison between Au (d and i) and Pd (e and j) adsorbed on freshly prepared silica films with cut-outs of crystalline (d and e) and vitreous (i and j) patches is provided [102].

lites, given that it would be possible to prove the high acidity of the adsorbed hydrogen [104].

Fig. 16 collects infrared spectra taken on the aluminosilicate film. The topmost trace refers to the hydrogen/deuterium exchange. "Positive" peaks are indicative of species that have been replaced on the surface. "Negative" peaks are indicative of newly adsorbed species, and clearly reveal the result of the H-D exchange process. In order to evaluate the acidity of the hydroxyl groups we resort to the adsorption of probe molecules, a procedure well known in zeolite characterization. The second and third traces from the top are infrared spectra recorded after exposure of the hydrogenated alumino silicate surface to CO.



**Fig. 16.** IRAS difference spectra taken on a hydrogenated aluminosilicate double layer as schematically shown in the inset. Upper trace: after H-D exchange, second trace: CO adsorption on the hydrogenated surface, lowest trace: after adsorption of ammonia on the hydrogenated surface [104,107].

The CO interacts preferentially with the hydroxyl groups and their frequency is shifted in a characteristic way. In addition, the stretching frequency of the adsorbed CO molecules is visible at 2183 cm<sup>-1</sup>, which is shifted to higher frequencies from the gas-phase value. The absolute value of the shift is a measure for the acidity of the hydroxyls at the surface. The observed shifts of 249 and 391 cm<sup>-1</sup> for v(OD) and v(OH), respectively, are relatively large and are indicative of an acidity of the hydroxyls on the two-dimensional aluminosilicate similar to chabasite. Another weak base, i.e. ethene, has been tried [107]. The results are similar. Two strong bases have also been tested, i.e. ammonia and pyridine, which lead to the formation of ammonium and pyridinium species [107]. The spectrum after adsorption of ammonia is shown as the forth trace in Fig. 16.

The hydroxyl groups are being consumed, and the two new bands occurring at 3398 and  $1502 \text{ cm}^{-1}$  are consistent, according to density functional calculations from Joachim Sauer's group [107], with the formation of ammonium species adsorbed atop the hexagonal rings.

Fig. 17 shows an STM image containing a variety of distinct features [104]. Similar to the case of an ordered pure silica bilayer system the majority of the framework is composed of double 6-membered rings (d6r). In the case of the ordered silica film, as well as the vitreous one, bright protrusions arranged as triangles were assigned to the position of the top-most oxygen atoms in the film, where these three O atoms are bound to the same tetrahedral atom [95-97]. Although only the immediate surface can be seen in the image, evidence of the same structure, symmetrically repeating below, is given by IRAS, as in the case of the pure silica films [105]. Fig. 17 shows the presence of structures lacking long-range order. Close-ups of  $1.5 \times 1.5$  nm sections of the large-scale image showing double N-membered rings are shown in insets a (d6r), b (d4r and d8r) and c (d5r and d7r). In these insets, the position of the tetrahedral T atoms is shown in green while the O atoms are shown in red. Insets a', b' and c' show connected-dots representations where the different ring sizes are emphasized by different colors. Summarizing a large body of statistical information we come to conclude that polygonal prisms of tetrahedral atoms, which constitute the building blocks of significant number of zeolites are identified with atomic resolution in a flat aluminosilicate. A substantial increase is observed in the population of prisms with even-numbered rings, when compared to analogous pure silica systems. These units had been previously proposed as precursors for the zeolite formation. In particular, the number of double four-membered rings (cubes) shows a significant increase. In addition, we note a close topological relation between this planar aluminosilicate and zeolite A [108].

It is our feeling that the study of two-dimensional zeolite film opens up an avenue for surface science to provide some evidence for details of chemical reactions on zeolites using a combination of scanning probe and spectroscopic techniques that are difficult to obtain otherwise.



**Fig. 17.** STM image showing the morphology of the surface. Insets (a)–(c) correspond to regions containing d6r, d4r + d8r and d5r + d7r, respectively. Red dots show the location of O atoms while the green atoms represent the tetrahedrally coordinated atoms. Insets a', b', c' show a different representation where the dots correspond to the location of the tetrahedrally coordinated atoms [104]. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

# 4. Conclusion and outlook

The four case studies presented were supposed to demonstrate and underline how model studies may be employed to investigate complex problems in catalysis leading to unexpected points of view that may influence our thinking for future research. The first case study could stimulate our thinking with respect to enantiospecific hydrogenation reactions at nanoparticles. The second one could initiate further studies toward a more systematic analysis of the influence of dopants and the effect of electron traps in more complex powder materials. Case study number three may motivate an investigation of materials combinations that could profitably be used as SMSI systems. In the last example we presented a case study that might open up possibilities to study reactions on zeolite like surfaces using local probe techniques.

We have only chosen a small number of case studies; however, model catalysis holds the potential to contribute in many areas of catalysis, in particular when techniques are employed to work under ambient conditions. Our group has started recently to look into those processes that are involved in the preparation of metal nanoparticles from a wet chemical environment using a combination of spectroscopic (non-linear optical) techniques and (electrochemical) scanning probe techniques. We believe there is much more to come to close the gap between models and real catalysts.

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