O- and H-induced surface core level shifts on Ru(0001): prevalence of the additivity rule

S Lizzit\textsuperscript{1}, Y Zhang\textsuperscript{2}, K L Kostov\textsuperscript{3}, L Petaccia\textsuperscript{1}, A Baraldi\textsuperscript{4,5,6}, D Menzel\textsuperscript{2,7} and K Reuter\textsuperscript{2}

\textsuperscript{1} Sincrotrone Trieste, Strada Statale 14 Km 163.5, I-34012 Trieste, Italy
\textsuperscript{2} Fritz-Haber-Institut der MPG, Faradayweg 4-6, D-14195 Berlin, Germany
\textsuperscript{3} Institute of General and Inorganic Chemistry, Bulgarian Academy of Science, 1113 Sofia, Bulgaria
\textsuperscript{4} Physics Department, University of Trieste, via A Valerio 2, I-34127 Trieste, Italy
\textsuperscript{5} Center of Excellence for Nanostructured Materials, University of Trieste, via A Valerio 2, I-34127 Trieste, Italy
\textsuperscript{6} Laboratorio Nazionale TASC INFM-CNR, in AREA Science Park, I-34012 Trieste, Italy
\textsuperscript{7} Physik-Department E20, Technische Universität München, D-85748 Garching, Germany

E-mail: dietrich.menzel@ph.tum.de and reuter@fhi-berlin.mpg.de

Received 14 October 2008, in final form 14 January 2009
Published 12 March 2009
Online at stacks.iop.org/JPhysCM/21/134009

Abstract
In previous work on adsorbate-induced surface core level shifts (SCLSs), the effects caused by O atom adsorption on Rh(111) and Ru(0001) were found to be additive: the measured shifts for first-layer Ru atoms depended linearly on the number of directly coordinated O atoms. Density-functional theory calculations quantitatively reproduced this effect, allowed separation of initial- and final-state contributions, and provided an explanation in terms of a roughly constant charge transfer per O atom. We have now conducted similar measurements and calculations for three well-defined adsorbate and coadsorbate layers containing O and H atoms: (1 \times 1)-H, (2 \times 2)-(O + H) and (2 \times 2)-(O + 3H) on Ru(0001). As H is stabilized in fcc sites in the prior two structures and in hcp sites in the latter, this enables us to not only study coverage and coadsorption effects on the adsorbate-induced SCLSs, but also the sensitivity to similar adsorption sites. Remarkably good agreement is obtained between experiment and calculations for the energies and geometries of the layers, as well as for all aspects of the SCLS values. The additivity of the next-neighbor adsorbate-induced SCLSs is found to prevail even for the coadsorbate structures. While this confirms the suggested use of SCLSs as fingerprints of the adsorbate configuration, their sensitivity is further demonstrated by the slightly different shifts unambiguously determined for H adsorption in either fcc or hcp hollow sites.

(Some figures in this article are in colour only in the electronic version)

1. Introduction
Core electron binding energies of solids are sensitive probes of the local atomic and geometric environment and, as such, markedly changed at surfaces. A quantitative understanding of the corresponding shifts, in particular of surface atoms coordinated to atomic or molecular adsorbates, provides a wealth of information, e.g. on the nature of the chemical bond [1], and may even be used for the interpretation and prediction of surface reactions [2]. Experimentally, the shifts can be accessed by high-resolution x-ray photoelectron spectroscopy (XPS) as part of the so-called surface core level shifts (SCLSs) [1]. Apart from the actual so-called initial-state shifts due to the changed electron density, and therewith electrostatic potential at the location of the core electrons, the measured SCLSs also contain so-called final-state contributions from variations in the screening of the core hole created in the XPS process, which is influenced by the surface or nearby adsorbates as well. At present,
a partitioning of the experimentally only accessible total shifts into initial- and final-state effects is only possible by first-principles electronic structure calculations. In particular for metal surfaces, approaches based on density-functional theory (DFT) have recently proven to be very successful. For instance, remarkable quantitative agreement could be achieved with SCLS measurements of the first-layer atoms of two close-packed transition metal surfaces, Rh(111) [3] and Ru(0001) [4], and of their modification by varying amounts of adsorbed O atoms in well-defined structures. Moreover, these calculations were then able to differentiate charge density and screening contributions and to analyze the respective physical basis in terms of the structures and the electronic properties of the systems studied.

Especially for adsorbate systems the high-resolution XPS spectra generally exhibit multi-peaked structures arising from different environments of the surface substrate atoms (first, second or deeper layers; neighbors to adsorbates of different type and number). Some assignments can often be reached by suitable spectroscopic means or by systematic variation and comparison of various sample conditions, e.g. in the form of different adsorbate coverages. However, a comprehensive interpretation requires detailed DFT calculations which in turn necessarily need to be based on a microscopic structural model. In view of a potential application to less well-characterized systems more general relations enabling a direct data interpretation are therefore of interest. In this respect, the mentioned SCLS work at Rh(111) [3] and Ru(0001) [4] identified an intriguing scaling in that the O-induced SCLSs of the first-layer metal atoms turned out to be linearly proportional to the number of directly coordinated O atoms to good accuracy. With the induced SCLS containing only a small and not much coverage-dependent final-state contribution, the source of this additivity could be traced back to a roughly constant amount of charge that is withdrawn from the surface atoms by each directly coordinated O neighbor [4]. With ensuing work on Rh(100) confirming this, then termed ‘additivity rule’, it was suggested that the latter may be a universal property of SCLSs which could be used for the fingerprinting of adsorbate populations in catalytic applications [2].

The cases mentioned all concern O atoms, a species with high electronegativity. SCLS results have also been obtained for H atoms on Rh(111), (100) and (110), including coverage dependences [5], but no results exist for mixed layers. It is therefore very interesting to examine whether such an additivity rule applies also to combinations of different adatoms. To this end we have selected H atom adsorption and O + H coadsorption on Ru(0001). This choice has the advantage that not only O [4, 6] and H [7] atomic overlayers, but also two well-defined (O + H) coadsorbate systems [8, 9], have already been characterized in detail in terms of adsorbate phases and geometries. On this basis we have measured the SCLS spectra for these systems and combined them with DFT calculations for ground state energies, geometries, electronic structure and SCLS values for the various distinguishable Ru surface atoms. We find excellent agreement between experiment and theory both for the geometries of the stable structures and for the SCLSs of all surface atoms. The central result we obtain is that we find the additivity rule to apply quite well also for H adsorbate atoms, and even for coadsorbate structures containing the strongly charge-transfer-inducing O atom and the roughly neutral H atom.

This paper is structured in the following way. An experimental section briefly summarizes the technique, the studied layers and their preparation, and displays the accumulated XPS data and the derived shift values. The fitting procedures to extract the latter from the measured spectra are briefly described in the appendix. Section 2.1 details the computational procedures and compares the obtained stable geometries in detail with existing experimental data. We then compare the total calculated shifts to the experimental ones and decompose these total SCLSs into initial- and final-state contributions. Concluding with a thorough discussion and analysis of the physical properties behind the observed ‘additivity rule’, we can trace the latter again back to an almost constant—albeit much smaller compared to O neighbors—amount of charge transferred to each directly coordinated H neighbor.

2. Experiment

2.1. Structures: geometry and preparation

The experimental procedures were similar to those of [4]. The SuperESCA beamline of the synchrotron radiation source ELETTRA in Trieste was used which works at base pressures in the low $10^{-10}$ mbar region and contains preparation facilities and an electron spectrometer with multichannel detection. The Ru(0001) surface was prepared by established procedures [4, 6, 7, 9]: cleanliness and order of the surface were checked by XPS and low-energy electron diffraction (LEED). The following adsorbate structures have been investigated: (1 × 1)-H, (2 × 2)-O, (2 × 2)-(O + H) and (2 × 2)-(O + 3H). For all structures, the adsorption sites and detailed geometries are known from previous quantitative IV-LEED or VLEED analysis [6, 9]. Figure 1 compiles the structures and
the sequence of preparation. In $(1 \times 1)$-H all hydrogen atoms sit in fcc sites. To prepare this saturated layer, the well-cleaned surface has to be exposed to high doses (several 1000 L) of molecular hydrogen, because the last few per cent to saturation coverage adsorb very slowly [10]. Low background pressure when hydrogen is introduced is important; otherwise the layer becomes contaminated with water [10]. For the two O + H coadsorbate structures, the starting point is a well-ordered $(2 \times 2)$-O structure with O adsorbed in hcp sites. The latter can be produced by slight overdosing with oxygen at 300 K to produce an O coverage of about 0.3 ML, heating to 1250 K, then heating to 400 K in 10⁻⁷ mbar hydrogen for about 10 min, and finally heating to 400 K for ordering [11]. Saturating this structure with hydrogen at 90 K, followed by a short anneal at 100–110 K for ordering, produces the $(2 \times 2)$-(O + 3H) structure. Upon heating the latter to 155 K with 2 K s⁻¹, or faster to 160 K, two-thirds of the hydrogen is desorbed and the $(2 \times 2)$-(O + H) structure remains, as was proven by thermal desorption spectra (TPD) [8] and quantitative LEED [9]. Quantitative LEED has also corroborated expectations from quantitative TPD [8] that, in the (O + 3H) structure, all H atoms sit in hcp sites, while in the (O + H) all H atoms sit in fcc sites [9]. This site selectivity in the two structures has been ascribed to repulsive O–H lateral interactions coupled with the tendency to maximize the coverage; we come back to this point below.

The interesting gist for the SCLS measurements is that the various structures contain Ru first-layer atoms in a variety of surroundings as also schematized in figure 1: with $3\text{H}_{\text{fcc}}$ neighbors in the $(1 \times 1)$-H layer; with 1 O or without adsorbate neighbor in the $(2 \times 2)$-O; with $1\text{O} + 2\text{H}_{\text{hcp}}$ or $3\text{H}_{\text{hcp}}$ in the $(2 \times 2)$-(O + 3H); and with 1 O + $1\text{H}_{\text{fcc}}$ or without adsorbate neighbor in the $(2 \times 2)$-(O + H). Together with the structures containing zero to threefold O-coordinated Ru surface atoms studied in [4], this variety offers a good basis to systematically check on the SCLS additivity with and without coadsorbates, as well as with H adsorbates in slightly different hollow sites.

2.2. SCLS measurements and analysis

The SCLS experiments were performed at three different photon energies (333, 345 and 400 eV) in order to exploit photoelectron diffraction effects to change the relative intensities of the surface components. The photon beam incidence angle was 70°. All photon energies (asymmetry $\alpha$, Lorentzian width $L$ and Gaussian width $G$) determined for the clean and $(2 \times 2)$-O surface are close to those published in [4]; small changes are explainable by the higher surface intensities and the improved energy resolution obtained in the present measurements. In any case, the central point is that all re-measured SCLSs for the clean and the $(2 \times 2)$-O surface are, within error bars, the same as in the previous work. The best-fit parameters determined for the H-containing structures are again similar, with further details given in the appendix. One notable difference is that the components corresponding to Ru atoms with H neighbors had to be given somewhat larger widths, which we interpret by for the other energies has been analyzed as well, without obtaining significantly different findings to those detailed below.

Figure 3 collects the SCLS measurements of all surfaces studied and shows the components derived from the fits; the numerical values are given in table 3 (vide infra). The fitting procedures used to distinguish and disentangle the different components in the spectra are identical to those discussed in [4], with further details given in the appendix. The best-fit parameters (asymmetry $\alpha$, Lorentzian width $L$ and Gaussian width $G$) determined for the clean and $(2 \times 2)$-O surface are close to those published in [4]; small changes are explainable by the higher surface intensities and the improved energy resolution obtained in the present measurements. In any case, the central point is that all re-measured SCLSs for the clean and the $(2 \times 2)$-O surface are, within error bars, the same as in the previous work. The best-fit parameters determined for the H-containing structures are again similar, with further details given in the appendix. One notable difference is that the components corresponding to Ru atoms with H neighbors had to be given somewhat larger widths, which we interpret by...
inversion symmetric slabs consisting of eight Ru(0001) layers in [4], in which the initial-state shifts are simply final-state contributions to the SCLSs is exactly the same as the xc functional [15]. The procedure to obtain initial- and generalized gradient approximation (GGA) using the PBE exchange–correlation (xc) energy is treated within the Slater–Janak transition state approach by an impurity calculation with half an electron screening is computed within the Slater–Janak transition state by an impurity calculation with half an electron.

The DFT calculations were performed using the full-potential augmented plane wave plus local orbital (L)APW + lo method [12, 13] as implemented in the WIEN2k code [14]. The exchange–correlation (xc) energy is treated with the generalized gradient approximation (GGA) using the PBE xc functional [15]. The procedure to obtain initial- and final-state contributions to the SCLSs is exactly the same as detailed in [4], in which the initial-state shifts are simply given by Kohn–Sham eigenvalue differences and the final-state screening is computed within the Slater–Janak transition state approach by an impurity calculation with half an electron removed from the core state considered.

All surfaces are modeled with supercell geometries, using inversion symmetric slabs consisting of eight Ru(0001) layers with O or H adsorption on both sides. The vacuum between consecutive slabs is at least 11 Å. Within the employed (2 × 2) surface unit cells the positions of all adsorbates and Ru atoms in the topmost three surface layers were fully relaxed for all structures considered. The gas-phase calculations for atomic H and O required for the binding energies are done in rectangular supercells with side lengths (13 × 14 × 15) bohr. The obtained total energies are then corrected by the gas-phase O2 [16] and H2 binding energies to obtain adsorbate binding energies with respect to the molecular references. The basis set parameters are briefly summarized as follows: the muffin-tin spheres for Ru, O and H are RMT = 2.0 bohr, RMT = 1.0 bohr and RMT = 0.8 bohr, respectively. Inside the muffin tins the wavefunctions are expanded up to lmax = 12 and the potential up to lmax = 6. The energy cutoff for the expansion of the wavefunction in the interstitial is Ewf = 20 Ryd and for the potential Ewf = 196 Ryd. For the (2 × 2) surface unit cells, a (6 × 6 × 1) Monkhorst–Pack grid was used for the Brillouin zone integrations, while for the gas-phase calculations in the rectangular supercells Γ-point sampling was employed.

The chosen computational approach is thus highly similar, if not superior, to the one used already in [4], with the notable exception that the now computationally feasible thicker slabs should enable a better description of the SCLSs of second-layer Ru atoms. Recomputing all SCLSs of all oxygen-containing structures reported in [4] with the new computational set-up, it is thus not surprising that we obtain, with one exception, values that are identical to the published ones to within 30 meV, thereby simply reconfirming the stringent convergence tests performed for the previous work. As expected, the sole disagreement concerns a second-layer shift in the (2 × 1)-O structure, denoted as S2(O) in [4], that was previously computed as −21 meV and is now obtained as +61 meV. This does not affect the published comparison to the experimental data though, as the latter peak was not resolved in the measurements.

3.2. Stable adsorption sites and adsorption geometries

In order to complement the preceding structure determination by quantitative LEED [6, 9] a first series of calculations addressed the stable O and H adsorption sites by comparing the binding energy at the high-symmetry sites offered by the Ru(0001) surface. Consistent with previous calculations [17] and in full agreement with the experimental analysis, the most favorable O adsorption site in purely oxygen-containing (2 × 2) overlayers with one up to four O atoms per surface unit cell is the hcp hollow site. In contrast, for H atoms in the same (2 × 2) overlayers the most stable site is the fcc hollow site, albeit overlayers with one up to four O atoms per surface unit cell is the hcp hollow site. Concerning the (2 × 2)-(O + H) and (2 × 2)-(O + 3H) coadsorption phases, we subjected all structural models considered in the recent quantitative LEED analysis of [9] to a full geometry optimization. The results are summarized in table I and are consistent with the preceding work in that the model with the lowest Pendry R factor is also the one with the highest binding energy and the detailed geometric parameters of these structures...
two phases summarized in Table 2 agree very well. This nicely confirms the assessment of [9] that LEED is indeed sensitive to the position of the H adsorbates. In these most stable structures the H atoms sit therefore, as already indicated in Figure 1, in the fcc sites in the (2×2)-(O + H) configuration and in the hcp sites in the (2×2)-(O + 3H) configuration.

4. SCLS results and discussion

4.1. Comparison of computed and measured SCLSs

From the geometric structures shown in Figure 1 one expects the following first-layer components in the measured SCLS spectra: a peak due to Ru atoms without adsorbate coordination (S1) in the clean surface, as well as peaks due to zero and onefold O-coordinated Ru atoms in the (2×2)-O structure, S1 and S2(O), respectively. In the (2×2)-(O + H) structure there are Ru atoms coordinated to 1O + 1Hfcc (S1(O + Hfcc)) and without adsorbate neighbor (S1); in the (2×2)-(O + 3H) structure there are Ru atoms coordinated to 1O + 2Hhcp (S1(O + 2Hhcp)) and Ru atoms coordinated to 3Hhcp (S1(3Hhcp)); and in the (1×1)-H layer there are Ru atoms coordinated to 3Hfcc neighbors (S1(3Hfcc)). In addition there can be non-negligible second-layer shifts (S2), where in the (2×2) layers one needs to furthermore distinguish between Ru atoms that are directly underneath an adsorbed O atom (S2(O)) and those that are not (S2). The SCLS data analysis and assignment for the clean surface and (2×2)-O layer has already been established in [4] and is the basis for the decomposition of the now measured spectra shown in Figure 3. Specifically, we followed the strategy explained in detail in [4] in which no attempt is made to resolve the small S2(O) second-layer component in the O-containing structure. Within this approach the assignment of the new peaks in the H overlayer and the two H-containing coadsorbate structures is straightforward. Also for the latter two only one non-negligible second-layer peak was considered and we come back to the consequences for the derived SCLS values below. However, we note already here that, although small, the shift of the first layer S1(O) component of the (2×2)-O overlayer when it becomes the S1(O + Hfcc) component in the (2×2)-(O + H) phase, and even further when it is turned into the S1(O + 2Hhcp) component in the (2×2)-(O + 3H) phase, is directly visible from the spectra and thus independent of the specific decomposition procedure employed. The same holds for the shift of the S1 peak of the clean surface when turned into the S1(3Hfcc) component of the (1×1)-H layer, cf Figure 3.

Table 3 compiles the SCLS values derived from the decomposition shown in Figure 3 and compares them to the calculated total shifts. The agreement is on the same remarkable level as that of [4], with maximum deviations of the order of 40 meV. As discussed in detail in this preceding work, one could presumably reach an even better agreement by explicitly considering the small S2(O) component in the experimental spectra, which is, however, not necessary for the scope of this work. Essentially, the experiment–theory comparison is thus flawless, which reconfirms not only that the employed GGA-PBE functional seems to allow for a quantitative determination of the quantity of interest in our study, but also fully supports the experimental peak assignment. On this basis it is intriguing to realize that the derived first-layer SCLS values are again quasi-linearly additive with respect to the coordination with H atoms. This is better visualized in Figure 4 which shows the H-induced changes of the SCLSs, i.e. the amount by which a first-layer

| 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 |
|---|---|---|---|---|---|---|---|
| \( E_{\text{tot}}^{(O-H)} \) | 3400 | 3146 | 3301 | model-2 | model-3 | model-2 | model-1 | 3270 |
| \( E_{\text{tot}}^{(O+3H)} \) | 3825 | 3049 | 3743 | 3752 | 2097 | model-1 | model-1 | model-1 |

Table 2. Computed geometric parameters of the stable (2×2)-(O + H) and (2×2)-(O + 3H) structures shown in Figure 1, compared to the results of the experimental LEED structure determination of [9]. d is the adatom–Ru bond length, \( d_i \) the mean vertical distance between layer i and j (with the adsorbate forming layer 0 and the topmost Ru layer forming layer 1, etc) and \( z \) is the vertical buckling of layer i. For the mean vertical layer distances, the values in brackets additionally indicate the relative changes with respect to the bulk interlayer distance.

|          | Theory | Experiment |
|----------|--------|------------|
| (2×2)-(O + H) |       |            |
| d (H–Ru) (Å) | 1.89   | 1.83 ± 0.13 |
| d (O–Ru) (Å) | 2.01   |            |
| \( \Delta d_{11} \) (H–Ru) (Å) | 1.15   | 1.03 ± 0.15/−0.08 |
| \( \Delta d_{10} \) (O–Ru) (Å) | 1.22   | 1.19 ± 0.02 |
| \( \Delta d_{12} \) (Å) | 2.13 (−1.2%) | 2.10 ± 0.02 (−1.9 ± 0.9 %) |
| \( \Delta d_{13} \) (Å) | 2.18 (+1.5%) | 2.14 ± 0.02 (0.0 ± 0.9 %) |
| \( \Delta d_{14} \) (Å) | 2.17 (+1.0%) |            |
| \( \Delta z_1 \) (Å) | 0.1    | 0.1        |
| \( \Delta z_2 \) (Å) | 0.05   | 0.01       |
| \( \Delta z_3 \) (Å) | 0.03   |            |

|          | Theory | Experiment |
|----------|--------|------------|
| (2×2)-(O + 3H) |       |            |
| d (H–Ru) (Å) | 1.86−1.88 | 1.89 ± 0.3 |
| d (O–Ru) (Å) | 2.0    |            |
| \( \Delta d_{11} \) (H–Ru) (Å) | 1.02   | 1.08 ± 0.15 |
| \( \Delta d_{10} \) (O–Ru) (Å) | 1.24   | 1.19 ± 0.025 |
| \( \Delta d_{12} \) (Å) | 2.18 (+1.2%) | 2.14 ± 0.02 (0.0 ± 0.9 %) |
| \( \Delta d_{13} \) (Å) | 2.16 (+0.3%) | 2.12 ± 0.02 (−0.9 ± 0.9 %) |
| \( \Delta d_{14} \) (Å) | 2.18 (+1.2%) |            |
| \( \Delta z_1 \) (Å) | 0.09   | 0.07       |
| \( \Delta z_2 \) (Å) | 0.02   | 0.03       |
| \( \Delta z_3 \) (Å) | 0.03   |            |
Table 3. Comparison of measured and calculated SCLS values for the five investigated structures. All shifts are given in meV. Error bars on the experimental values are between ±10 and ±30 meV (see the appendix). See text for an explanation of the employed nomenclature. No attempt was made to fit an S2(O) component in the experimental spectra (see the text).

| Exp./theory | Clean | (2 × 2)-O | (2 × 2)-(O + H) | (2 × 2)-(O + 3H) | (1 × 1)-H |
|-------------|-------|------------|----------------|-----------------|-----------|
| S13(3Hfcc)  | −371/−405 | −413/−423 | −414/−438 | −291/−331 | −210/−205 |
| S13(3Hhcp)  | −120/−120 | +24/+29 | +89/+91 | +120/+120 |          |
| S1(O)       | +133/+102 | +144/+155 | +135/+158 | +85/+54 | +78/+52 |
| S2(O)       | n.a./−20  | n.a./−22  |          |          |          |
| S1(O + Hfcc)|          |          |          |          |          |
| S1(O + 2Hhcp)|         |          |          |          |          |

Table 4. Calculated final-state contributions to the total SCLSs of first-layer Ru atoms directly coordinated to zero, one, two or three adsorbed H atoms. All values are given in meV.

|                | 0fold | 1fold | 2fold | 3fold |
|----------------|-------|-------|-------|-------|
| Clean          | −62   | −45   | −36   | −38   |
| (2 × 2)-O      |       |       |       |       |
| (2 × 2)-(O + Hfcc) | −75   | −38   | −38   | −44   |
| (2 × 2)-(O + 3Hhcp) |      |       |       |       |
| (2 × 2)-(4Hhcp) |       |       |       |       |

Figure 4. Experimental H-induced changes of the first-layer SCLSs as a function of the number of directly coordinated H atoms. Shown is the value by which the SCLS changes upon H neighbor addition, i.e. the reference is the S1 value of the clean Ru(0001) surface for the purely H-coordinated Ru atoms, the S1 value of the (2 × 2)-O structure for the Ru atoms coordinated solely to H in the coadsorbate phases and the S1(O) value of the (2 × 2)-O structure for the Ru atoms coordinated to both O and H in the coadsorbate phases.

Ru SCLS changes upon adding another H nearest neighbor. Strikingly, the observed additivity seems even independent of the presence of the O coadsorbate, and slightly different shifts are obtained for H neighbors in either fcc or hcp hollow sites. We furthermore mention that the O- and H-induced shifts are also closely the same as the ones observed in the (2 × 2)-(O + H) structure for a (2 × 2)-(O + H + CO) structure which additionally contains CO on top of those Ru surface atoms, which in the (2 × 2)-(O + H) have no direct adsorbate neighbors [18]. This also suggests a minor influence of next-nearest-neighbor occupation compared to the direct neighbor effects discussed here.

4.2. Analysis of the ‘additivity rule’

As a first step to analyze the physics governing the observed additivity rule it is useful to decompose the total calculated shifts into initial- and final-state contributions. Table 4 compiles the final-state contributions to all first-layer SCLSs of the five measured structures. Overall, the screening correction is found to be of the same order of magnitude as the actual total H-induced shifts presented in figure 4, as well as only marginally smaller than the screening contributions reported in [4] for the O-induced SCLSs at Ru(0001). This is distinctly different to the situation for the initial-state shifts, which for the latter O-containing structures were one order of magnitude larger than the now considered H-induced shifts.

The consistently obtained negative sign of the screening correction indicates that the screening capabilities of the core-ionized system are enhanced at the surface compared to the bulk. For the clean surface, the detailed analysis put forward in [4, 19] has already rationalized this by the narrowing of the local valence d-band induced by the lowered coordination of the surface atoms. This in turn leads in general to an enhancement of the density of states (DOS) at and above the Fermi level compared to the bulk situation, which—with the final-state effects at a transition metal surface like Ru(0001) largely due to the intra-atomic d-electron screening [19]—implies a more efficient screening at the surface. That this is not much affected by the presence of the adsorbates can be understood considering the Anderson–Grimley–Newns model of chemisorption [20]. Consistent with this model the computed adsorbate-induced DOS change concerns primarily the formation of a bound state at the bottom of the substrate d-band and a broad (antibonding) resonance at the upper edge of the d-band. With the Fermi level in Ru positioned at about 2/3 band-filling, this leaves the DOS at the Fermi level and thereby the screening capabilities largely unaffected compared to the situation at the clean surface. In this understanding one would also not expect a strong influence of the adsorbate-induced geometric relaxation on the magnitude of the SCLS screening.
correlation was observed before. 

As expected from the understanding developed in [4] we therefore arrive at the conclusion that the observed correlation of first-layer SCLSs with the number of H nearest neighbors is an initial-state effect. The formation of bonds between the adsorbate and the first-layer Ru atoms affects the local valence d-band, the center of which shifts to maintain local charge neutrality. This changes the local (near nucleus) electrostatic field, which in turn acts on the core electrons as well and induces the finite SCLSs. Effectively, the about linear dependence of the latter on the number of H neighbors thus simply reveals that the type of bonding, i.e. the bonds formed per H nearest neighbor, is roughly the same in all overlayers studied. With all the caveats given in [4] this may be interpreted in the conceptual language of charge transfer in that the amount of charge transferred to each H atom remains approximately constant in the different structures. The much smaller electronegativity of H then also rationalizes the much smaller observed H-induced initial-state shifts (≈50 meV/H atom) compared to the O-induced shifts reported before (≈400 meV/O atom [4]) and is fully consistent with the very different work function changes induced by the two adsorbates. In agreement with experiment [21] we compute the latter to be essentially zero even for the full monolayer H coverage, in stark contrast to measured [22] and computed [17] O-induced changes of up to ~1.5 eV for the full (1 × 1)-O monolayer. The thereby suggested near-neutrality of H on Ru(0001) may also explain why the additivity rule prevails even in the denser adsorbate overlayers. For an effectively charged adsorbate, some degree of depolarization, i.e. reduced charge transfer per adsorbate, would be favorable for higher coverages as it reduces the repulsive adsorbate–adsorbate interaction. This effect is indeed observed for dense O overlayers [3, 4], but is too small there to significantly affect the near-linear SCLS correlation with O coordination. In this respect, one would expect a noticeable violation of the additivity rule, particularly for adsorbates with only a modest bond strength (implying smaller total magnitudes of the adsorbate-induced initial-state SCLSs), but relatively strong induced dipole moment (suggesting more pronounced depolarization effects at decreased adsorbate–adsorbate distances). This situation could, for example, be given for molecular adsorbates like CO or NO, and we are currently analyzing coadsorbate overlayers containing O, H and these molecular species in this regard [18].

With the SCLSs sensitively reflecting the formed adsorbate–substrate bonds, it is clear that they can also be used as local probes of the adsorption site. This was already shown in the preceding work on the O-containing structures, where the computed Ru SCLSs for structures with O atoms in fcc sites were consistently about 100–200 meV smaller than the corresponding values in structures with O atoms in hcp sites and therewith irreconcilable with the experimental data [4]. For O adsorption into hcp sites the induced SCLS change per O neighbor is thus slightly larger, whereas, interestingly, for Rh(111) where the fcc site is the more stable adsorption site the situation is exactly reversed in that there O adsorption into fcc sites leads to slightly larger SCLS increases [3]. For both surfaces, a somewhat more ‘negatively charged’ oxygen atom is thus adsorbed in the more stable adsorption site, suggesting a preference for a stronger ionic bonding. In this respect, it is intriguing to notice from figure 4 that, also for the H overlayers, the slightly more stable fcc site is the one that leads to slightly larger induced SCLS changes. The less favorable hcp site is only populated in the O-containing dense (2 × 2)-(O + 3H) coadsorbate structure and it is tempting to attribute this stabilization in part to that a slightly less ionic hydrogen is preferred at the corresponding short inter-adsorbate distances. While in this structure the effect concerns only the position of the H atoms with the O atoms remaining in the hcp sites, coadsorption of CO is able to also shift the O atoms to fcc sites. Analyzing the correlation of the respective stabilities of different sites with the induced SCLS changes is thus another aspect of the mentioned on-going work on coadsorption structures containing O, H, CO and NO [18].

5. Conclusions

In conclusion we have presented a joint experimental and theoretical study of Ru(0001) surface core level shifts induced by O and H adsorbates. Remarkable quantitative agreement concerning the energetics, geometries and SCLS values of the investigated (1 × 1)-H, (2 × 2)-(O + H) and (2 × 2)-(O + 3H) (co)adsorbate overlayers on Ru(0001) is achieved. The quasilinear dependence of the adsorbate-induced SCLSs on the number of directly coordinated nearest neighbors reported before for O overlayers on Ru(0001) [4], Rh(111) [3] and Rh(100) [2], as well as for H adsorbate layers on all Rh low-index single-crystal surfaces [5], is found to prevail also in the H (co)adsorbate structures on Ru(0001). The computed decomposition of the total SCLSs into initial- and final-state contributions reveals an enhanced screening at the surface of the (co)adsorbate layers that is of the same order of magnitude as for the pure O overlayers studied before. Accounting for this final-state correction is thus necessary to obtain the reported quantitative agreement with the measured data, while its actual value exhibits only a small scatter for the differently coordinated surface atoms. In accordance with the detailed analysis put forward in [4], the observed additivity of the adsorbate-induced SCLSs is thus a pure initial-state effect. In
terms of the conceptual charge transfer language it merely reflects a roughly constant amount of charge transferred to each directly coordinated adsorbate neighbor. Not surprisingly, this amount of charge (and concomitantly the magnitude of the observed SCLS change) is significantly smaller for the almost electroneutral H adsorbates than for the electronegative O adsorbates. Intriguingly, the SCLS data reveals slightly larger induced changes for H atoms adsorbed in fcc sites than in hcp sites, consistent with the interpretation that a slightly more ionic H adsorbs in the prior sites. The established sensitivity even to such highly similar adsorption sites, as well as the prevalence of the additivity rule, confirms the suggested use of SCLSs as directly accessible and valuable fingerprints of the adsorbate populations.

Acknowledgments

The experimental work has been supported by the European Union through an I3 grant to Elettra, Trieste. DM thanks the German Fonds der Chemischen Industrie for support.

Appendix

Similar to the analysis performed in [4], the experimental data were fitted with a combination of Doniach–Šančič functions [23] convoluted with a Gaussian broadening and a linear background. The lineshape parameters are the Lorentzian \( L \), Gaussian \( G \) and asymmetry \( \alpha \). The following outlines the fitting procedure used to extract these parameters and the values of the Ru 3d\textsubscript{5/2} SCLSs.

The absolute binding energy of the peaks was calibrated to the Fermi level, measured for each spectrum. Moreover, the intensity of the spectra was normalized to the background at the low binding energy side (279.5 eV) in order to account for variations of the photon beam intensity. The best-fit parameters were extracted from the fit of the data taken at the three photon energies used, keeping for each structure the same number of peaks, lineshapes and peak positions, and letting only the relative peak intensities vary freely. First the spectrum of the clean surface was fitted with three components: bulk \( (L = 0.20 \text{ eV}, G = 0.06 \text{ eV}, \alpha = 0.07) \), \( S_1 \) \( (L = 0.24 \text{ eV}, G = 0.06 \text{ eV}, \alpha = 0.05) \) and \( S_2 \) \( (L = 0.20 \text{ eV}, G = 0.06 \text{ eV}, \alpha = 0.08) \). The absolute position of the bulk peak was determined as well and this position was kept fixed in all other fits. Similarly, the so-determined lineshape of bulk, first- and second-layer clean Ru atoms was maintained for all adsorbate structures. In this approach, we next fitted the second-layer clean Ru atoms was maintained for all adsorbate fits. Similarly, the so-determined lineshape of bulk, first- and second-layer clean Ru atoms was maintained for all adsorbate structures. In this approach, we next fitted the second-layer clean Ru atoms was maintained for all adsorbate fits. Similarly, the so-determined lineshape of bulk, first- and second-layer clean Ru atoms was maintained for all adsorbate structures.

We then continued with the \((1 \times 1)\)-H structure where again three components are expected: bulk, \( S_2 \) (same lineshape parameters as \( S_2 \) in the clean surface) and \( S_1 \) \( (3\text{H}_{\text{fcc}}) \) \( (L = 0.24 \text{ eV}, G = 0.10 \text{ eV}, \alpha = 0.05) \). It has to be noted here that the intensity of the different components is almost the same as for the clean surface, because the hydrogen atoms present on the surface have only small scattering effects on the electrons photoemitted from the Ru substrate. On the other hand, the presence of H at the surface seems to affect the peak shape in the spectra, and we obtain significantly improved fits when allowing for a larger Gaussian width. As mentioned in the text, we interpret this by inhomogeneous broadening due to a less than perfect ordering achieved for the H overlayer. The same situation (unchanged intensity with respect to the \((2 \times 2)\)-O, increased Gaussian width) is found for the coadsorbate structures, and the lineshape parameters determined are: \( S_1 \) \( (O+1\text{H}_{\text{fcc}}) \) \( (L = 0.24 \text{ eV}, G = 0.15 \text{ eV}, \alpha = 0.06) \) in the \((2 \times 2)-(O+H)\) structure; and \( S_1 \) \( (O+2\text{H}_{\text{hcp}}) \) \( (L = 0.26 \text{ eV}, G = 0.16 \text{ eV}, \alpha = 0.06) \) and \( S_1 \) \( (3\text{H}_{\text{hcp}}) \) \( (L = 0.24 \text{ eV}, G = 0.12 \text{ eV}, \alpha = 0.05) \) in the \((2 \times 2)-(O+3H)\) structure.

The high quality of the fits can be seen from the very small and structureless residuals as shown in figure 3. The remaining uncertainties lead to experimental error bars between \( \pm 10 \) and \( \pm 30 \) meV for the various peaks and depend mainly on the energetic distance from other components and their relative strengths. Specifically, they are \( \pm 10 \) meV for all \( S_1 \) components and for \( S_2 \) of the clean surface; \( \pm 20 \) meV for \( S_2 \) \( (O) \) and \( \pm 30 \) meV for \( S_1 \) \( (O+H) \) to \( S_1 \) \( (O+3H) \). Where overlap exists these errors constitute an improvement over those of [4].

References

[1] See, e.g., Mårtensson N and Nilsson A 1994 High Resolution Core-Level Photoelectron Spectroscopy of Surfaces and Adsorbates (Springer Series in Surface Science vol 35) (Berlin: Springer) p 65
[2] Baraldi A, Lizzit S, Comelli G, Kiskinova M, Rosei R, Honkala K and Nerskov J K 2004 Phys. Rev. Lett. 93 046101
[3] Ganduglia-Pirovano M V, Scheffler M, Baraldi A, Lizzit S, Comelli G, Paolucci G and Rosei R 2001 Phys. Rev. B 63 205415
[4] Lizzit S, Baraldi A, Grosso A, Reuter K, Pirovano-Ganduglia M V, Stampfl C, Scheffler M, Stüchler M, Keller C, Wurth W and Menzel D 2001 Phys. Rev. B 63 205419
[5] Vesselli E, Baraldi A, Bondino F, Comelli G, Peressi M and Rosei R 2004 Phys. Rev. B 70 115404
[6] Weestrate K J, Baraldi A, Runutz L, Lizzit S, Comelli G and Rosei R 2004 Surf. Sci. 566–566 486
[7] Baraldi A 2008 J. Phys.: Condens. Matter 20 93001
[8] Vesselli E, Campaniello M, Baraldi A, Bianchettin L, Africh C, Esch F, Lizzit S and Comelli G 2008 J. Phys. Chem. C 112 14475
[9] Lindroos M, Pfnür H, Held G and Menzel D 1989 Surf. Sci. 220 43
[10] Pfnür H, Held G, Lindroos M and Menzel D 1989 Surf. Sci. 222 451
[11] Lindroos M, Pfnür H, Feulner P and Menzel D 1987 Surf. Sci. 180 237
[12] Lindroos M, Pfnür H and Menzel D 1987 Surf. Sci. 192 421
[13] Held G and Menzel D 1994 Surf. Sci. 316 92
[14] Schiffer A, Jakob P and Menzel D 2000 Surf. Sci. 465 198
[15] Schiffer A 1998 PhD Thesis Technische Universität München
[16] Glöck M and Menzel D 2009 Surf. Sci. at press (doi:10.1016/j.susc.2008.07.045)
[10] Kostov K L, Widdra W and Menzel D 2004 Surf. Sci. 560 130 and references therein
[11] Koch M, Jakob P and Menzel D 1996 Surf. Sci. 367 293
[12] Sjöstedt E, Nordström L and Singh D J 2000 Solid State Commun. 114 15
[13] Madsen G K H, Blaha P, Schwarz K, Sjöstedt E and Nordström L 2001 Phys. Rev. B 64 195134
[14] Blaha P, Schwarz K, Madsen G K, Kvasnicka D and Luitz J 2001 WIEN2k Technische Universität Wien, Austria ISBN 3-9501031-1-2
[15] Perdew J P, Burke K and Ernzerhof M 1996 Phys. Rev. Lett. 77 3865
[16] Kiejna A, Kresse G, Rogal J, De Sarkar A, Reuter K and Scheffler M 2006 Phys. Rev. B 73 035404
[17] Reuter K, Ganduglia-Pirovano M V, Stampfl C and Scheffler M 2002 Phys. Rev. B 65 165403
[18] Lizzit S, Zhang Y, Kostov K L, Petaccia L, Baraldi A, Reuter K and Menzel D 2009 in preparation
[19] Andersen J N, Hennig D, Lundgren E, Methfessel M, Nyholm R and Scheffler M 1994 Phys. Rev. B 50 17525
Methfessel M, Hennig D and Scheffler M 1993 Surf. Sci. 287/288 785
Methfessel M, Hennig D and Scheffler M 1995 Surf. Rev. Lett. 2 197
[20] Grimley T B 1975 Prog. Surf. Membr. Sci. 9 71
Muscat J P and Newns D M 1978 Prog. Surf. Sci. 9 1
Muscat J P and Newns D M 1979 Phys. Rev. B 19 1270
[21] Feulner P and Menzel D 1985 Surf. Sci. 154 465
[22] Madey T E, Engelhardt H A and Menzel D 1975 Surf. Sci. 48 304
[23] Doniach S and Šunjic M 1970 J. Phys. C: Solid State Phys. 3 185