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Elisa Meriggio, Rémi Lazzari, Christophe Méthivier, Pascal David, Stéphane Chenot, et al.. Chemical nature and thermal decomposition behavior of tartaric acid multilayers on rutile TiO₂ (110). Journal of Vacuum Science & Technology B Microelectronics and Nanometer Structures, 2019, 37 (5), pp.051803. 10.1116/1.5100957 . hal-02340226

HAL Id: hal-02340226

<https://hal.sorbonne-universite.fr/hal-02340226>

Submitted on 30 Oct 2019

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Citation: *Journal of Vacuum Science & Technology B* **37**, 051803 (2019); doi: 10.1116/1.5100957

View online: <https://doi.org/10.1116/1.5100957>

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Chemical nature and thermal decomposition behavior of tartaric acid multilayers on rutile TiO₂(110)

Elisa Meriggio,^{1,2} Rémi Lazzari,² Christophe Méthivier,¹ Pascal David,² Stéphane Chenot,² Xavier Carrier,¹ Gregory Cabailh,^{2,a)} and Vincent Humblot^{1,b)}

¹Laboratoire de Réactivité de Surface, Sorbonne Université, UMR CNRS 7197, 4 place Jussieu, 75252 Paris Cedex 05, France

²Institut des NanoSciences de Paris, Sorbonne Université, UMR CNRS 7588, 4 place Jussieu, 75252 Paris Cedex 05, France

(Received 22 April 2019; accepted 9 July 2019; published 12 August 2019)

R,R-tartaric acid (RR-TA) thermal stability and decomposition on the rutile TiO₂(110) surface was investigated by temperature programmed desorption. The authors show that a majority of RR-TA molecules are desorbed intact from multilayers at around 340 K, while they decompose from the first chemisorbed layer between 460 and 480 K. Complementary information on the chemical nature of RR-TA in the multilayer regime was gained by x-ray photoelectron spectroscopy, which shows that biacid molecules form the multilayer while they are monotartrate at the interface. *Published by the AVS.* <https://doi.org/10.1116/1.5100957>

I. INTRODUCTION

Chiral molecules, which are nonsuperimposable to their mirror image, are of prime importance for living organisms, since many biomolecules appear in only one handedness.¹ Biological activities of opposite enantiomers may, thus, lead to drastic effects, which explain why gaining control of enantioselectivity is a major challenge in many fields, including pharmaceutical and agrochemical industries.² Among several developed strategies for the production of enantiomerically pure compounds, asymmetric catalysis has been receiving increasing attention.^{3,4} Chiral molecules deposited in ultra-high vacuum (UHV) conditions either on plane metal surfaces or on oxide-supported metal catalysts are used in surface science as model systems to investigate at a microscopic level the catalyst structure and understand the key steps of the catalytic processes.^{2,3,5–7} While metal surfaces have been intensively explored, fundamental questions remain on the role of the oxide supports in oxide-supported metal catalysts.

In this context, we focus here on (R,R)-tartaric acid [HOOC–CH(OH)–CH(OH)–COOH, hereafter named RR-TA] and rutile TiO₂(110) single crystal as a model system of the interaction between a chiral molecule and an oxide substrate. The adsorption of RR-TA, one of the most efficient chiral modifier, has been investigated so far on several metal surfaces.^{5,8–19} Biacidic, monotartrate, or bitartrate molecules have been found upon adsorption, depending on how many COOH groups undergo deprotonation.⁵ The RR-TA thermal decomposition mechanism has been studied on Cu,^{10,17–19} Ni,^{11–15} and Pd (Ref. 16) surfaces and has been found to strongly depend on the surface structure. On some surfaces, such as Cu(110)^{19–22} and Ni(110),¹³ temperature programmed desorption (TPD) experiments have evidenced an explosive

decomposition behavior for RR-TA by tracing the molecular decomposition products H₂, H₂O, CO, and CO₂. However, the same desorption mechanism was not observed on other metal surfaces such as Pd(111)¹⁶ and Ni(111).¹¹ Different surface structures, surface roughness, contaminations, and experimental conditions (mainly the adsorption temperature) have been proposed as possible explanations for the different observed behaviors.¹⁷

The interaction of carboxylic acid molecules with the rutile TiO₂(110) surface has received considerable attention. Upon adsorption, they present a common predisposition to deprotonate one or multiple COOH groups, with the carboxylate O atoms bound to two five-fold coordinated surface titanium Ti_{5c} atoms.²³ In a previous work, we investigated RR-TA submonolayer and monolayer deposition on the rutile TiO₂(110) surface by combining several surface science techniques.²⁴ We have shown that RR-TA molecules adsorb on the oxide surface in a monotartrate form, i.e., only one of the two COOH groups deprotonates upon adsorption. The carboxylate oxygen atoms are bound to two adjacent Ti_{5c} atoms in the [001] direction and the remaining COOH group protrudes out of the surface. Moreover, at saturation of the first layer, low-energy electron diffraction (LEED) and STM evidenced the formation of an ordered (2 × 1) overlayer. Here, we extend the study to the RR-TA thermal stability and decomposition on TiO₂(110) as well as its adsorption behavior in the multilayer regime. It has already been observed experimentally that protruding carboxylic groups favor the formation of physisorbed multilayers.²⁵ Hence, the interface layer formed by monotartrate species on TiO₂(110) with a free carboxylic group available at the surface should lead to a multilayer formation via H-bond interactions.

II. EXPERIMENT

All experiments have been carried out in an UHV chamber at the Institut des Nanosciences de Paris laboratory, with a base pressure of 10^{–10} mbar or better. The rutile TiO₂(110) surface has been intensively investigated in surface science thanks to

Note: This paper is part of the Conference Collection from the AVS Pacific Rim Symposium on Surfaces, Coatings and Interfaces (PacSurf 2018) meeting.

^{a)}Electronic mail: gregory.cabailh@sorbonne-universite.fr

^{b)}Electronic mail: vincent.humblot@sorbonne-universite.fr

the possibility of increasing the crystal conductivity by reducing the samples under UHV conditions, thus allowing its characterization by electron spectroscopies and scanning tunneling microscopy.^{23,26} The TiO₂(110) single crystal (Mateck GmbH, Germany) was cleaned by several cycles of Ar⁺ sputtering at 1 keV and subsequent annealing at 1000 K. XPS and LEED were used to verify the absence of contamination and the surface crystallinity, respectively. After the last annealing, the TiO₂ sample was cooled down to room temperature, before RR-TA deposition.

Crystalline powder of (R,R)-tartaric acid was provided by Acros Organics (Fisher Scientific SAS France) and deposited from a glass crucible resistively heated by a metallic wire wrapped around it, after having been degassed for several hours (for details see Ref. 24). The typical cell temperature is between 370 and 395 K.^{8,16} During RR-TA evaporation, the pressure in the UHV chamber rose reproducibly between 2 and 3 × 10⁻⁹ mbar. Since the crucible is not equipped with a thermocouple, an accurate calibration is first performed by XPS. RR-TA coverage was estimated from the Ti2p/C1s XPS intensity ratio, assuming a homogeneous molecular film as detailed in Ref. 24. Knowing that in the submonolayer regime RR-TA binds to the TiO₂(110) surface in a monotartrate form (one RR-TA molecule every two surface Ti_{5c} atoms), the monolayer saturation coverage is defined as half of Ti_{5c} density of the exposed surface (5.2 × 10¹⁴ cm⁻²),²⁷ which corresponds to an equivalent film thickness of about 6–7 Å.

XPS was performed using a nonmonochromatic dual anode Al/Mg Kα x-ray source (hν = 1486.6 eV) and a hemispherical analyzer (Omicron EA 125). Multilayer TA molecules may desorb in UHV conditions during XPS analysis as observed in our previous study²⁴ leading to a monolayer regime. In the present case, rapid XPS scans (less than a minute) were recorded in order to avoid multilayer TA desorption and to analyze the chemical state of RR-TA molecules in the multilayer regime. Spectra were taken at normal emission using a pass energy of 50 eV. The spectra were normalized to the background level on the low binding energy (E_b) side of the peak. The XPS analysis was carried out with CASAXPS software, applying for the fit a Gaussian/Lorentzian ratio equal to 70/30.

TPD was carried out using a quadrupole mass spectrometer (QMG 220 from PFEIFFER) placed in a quartz tube differentially pumped by an ionic pump (for details see Ref. 28). The tube ends with a small cap pointing to the sample, the aperture of which is smaller than the sample surface. During TPD experiments, the tube is placed at about 1 mm from the TiO₂ surface to drastically limit the detection of species desorbing from the surroundings. A linear heating rate α = 0.5 K s⁻¹ was applied to the sample up to a temperature of 800 K, which was measured by a calibrated thermocouple spot-welded on the manipulator.

III. RESULTS AND DISCUSSION

A. Thermal stability of RR-TA on rutile TiO₂(110)

TPD experiments were carried out to gain information on RR-TA thermal stability and decomposition behavior on

TiO₂(110). Based on the mass spectrum of RR-TA from the NIST database,²⁹ the main ion fragments observed correspond to the mass to charge ratios (m/e) of 58 and 76 with relative intensities of 40 and 100, respectively. Hence, these fragments of the molecule have been tracked in the present TPD experiments. Moreover, m/e = 2 (H₂⁺), 18 (H₂O⁺), 28 (CO⁺), and 44 (CO₂⁺) were also monitored.

TPD spectra were recorded for the clean TiO₂(110) surface as a blank and following increasing doses of RR-TA at room temperature. The results obtained are summarized in Fig. 1 for m/e = 58 (HOC—COH⁺) and 76 (C₂O₃H₄⁺) corresponding to the main two fragments ascribed to the molecular form and in Fig. 2 for m/e = 2 (H₂⁺), 18 (H₂O⁺), 28 (CO⁺), and 44 (CO₂⁺). A particular attention has to be paid when considering H₂⁺, H₂O⁺, CO⁺, and CO₂⁺ fragments since they usually are the dominant residual gases in UHV chambers and their presence is inevitable when heating in UHV. To be sure that H₂⁺, H₂O⁺, CO⁺, and CO₂⁺ contributions truly arise from RR-TA molecular desorption, TPD

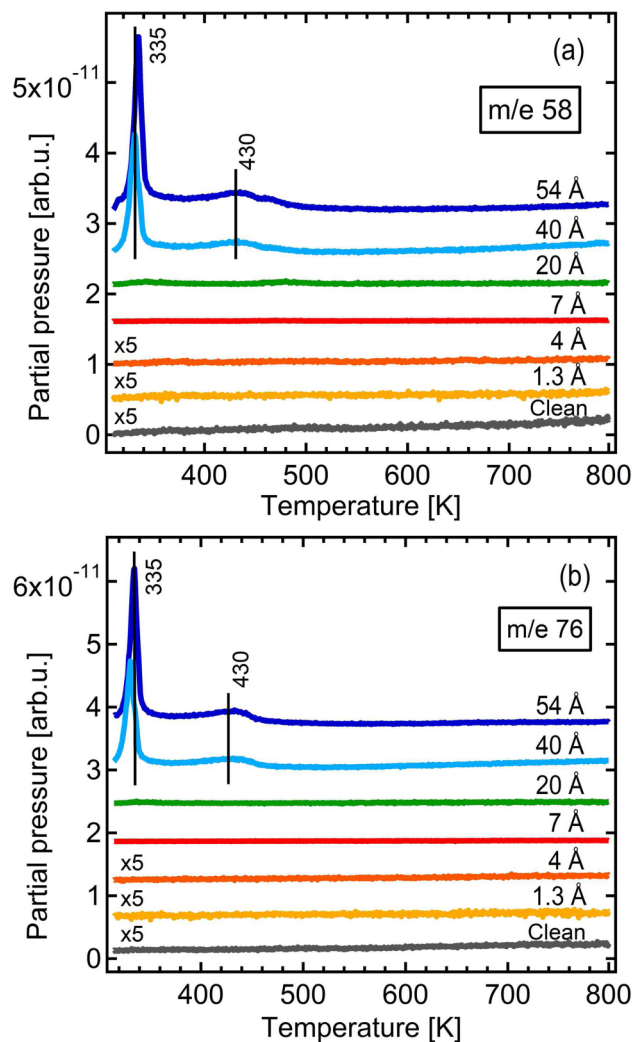


FIG. 1. TPD spectra of relative mass/charge (a) 58 and (b) 76 for increasing doses of RR-TA on TiO₂(110) at room temperature. The recorded spectra are vertically shifted for the sake of clarity. Scale factors are specified near the corresponding spectra.

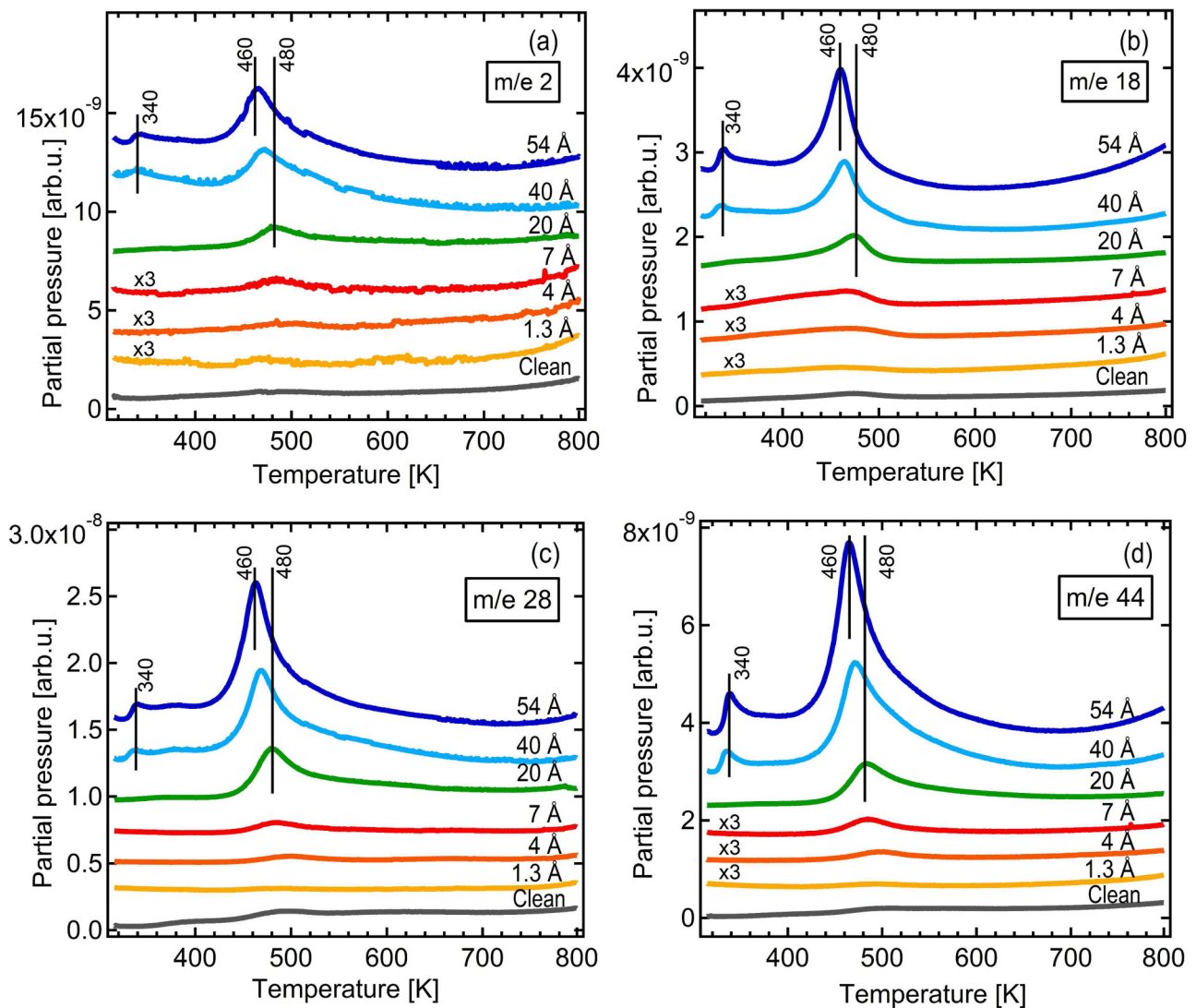


FIG. 2. TPD spectra of relative mass/charge (a) 2 (H_2^+), (b) 18 (H_2O^+), (c) 28 (CO^+), and (d) 44 (CO_2^+) for increasing doses of RR-TA on $\text{TiO}_2(110)$ at room temperature. The recorded spectra are vertically shifted for the sake of clarity. Scale factors are specified near the corresponding spectra.

spectra at increasing RR-TA coverage are compared to those recorded on the clean TiO_2 surface.

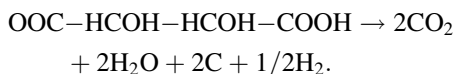
After RR-TA doses, two main features are identified at about 335–340 K and 430–480 K. The low temperature peak at 335–340 K appears very intense at high RR-TA doses for $m/e = 58$ and 76 (Fig. 1), which are specific to RR-TA molecules, and does not saturate as the molecular dose increases. In analogy to previous findings for RR-TA on $\text{Cu}(531)$ ¹⁷ and on $\text{Pd}(111)$,¹⁶ the peak at 335–340 K is, thus, attributed to multilayer desorption. Desorption of the second molecular layer was found to occur around 330 K on $\text{Pd}(111)$, in reasonable agreement with the present work. This peak is also far weaker for the molecular decomposition fragments H_2^+ , H_2O^+ , CO^+ , and CO_2^+ , again suggesting molecular desorption.

For higher temperatures, the desorption feature between 430 and 480 K is consequently expected to arise from monolayer desorption. This contribution appears much more intense in the H_2^+ , H_2O^+ , CO^+ , and CO_2^+ spectra and far less for the m/e specific to the desorption of the intact RR-TA molecule. The intensity of this feature does not completely saturate and

continues to slightly increase even for the higher doses; this fact may be explained by the formation of multilayers before the complete saturation of the first chemisorbed layer, as previously observed in the case of RR-TA on $\text{Cu}(531)$ ¹⁷ or from an incomplete multilayer desorption at 340 K. Moreover, a shift to lower temperature is observed at increasing molecular doses. The intensity of the desorption features observed in H_2^+ , H_2O^+ , CO^+ , and CO_2^+ spectra increases for increasing RR-TA doses, thus ruling out the attribution of their origin only to desorption of vacuum residuals. Moreover, in the corresponding $m/e = 58$ and 76 spectra, no desorption features are observed between 430 and 480 K for the lowest molecular coverages, confirming that the desorption peak comes from molecular desorption for the highest coverages.

Overall in the multilayer regime, large fragments of RR-TA molecules are detected suggesting desorption of intact molecular RR-TA, while RR-TA decomposes into smaller fragments when desorbed from a monolayer coverage. This observation is indicative of a strong interaction between the molecule and the oxide surface, stronger than

the intramolecular bonds. A rough estimate of the desorption temperature can be given by the Readhead formula. To break a C—C bond (90 kcal/mol), the expected temperature is enormous ~ 1300 K. The process of decomposition is probably a two-step process with an adsorbed intermediate. As previously reported, decomposition of RR-TA from the chemisorbed layer has been evidenced on other surfaces, such as Cu(110), Cu(531), Ni(110), and Pd(111), with the main decomposition products being hydrogen, carbon dioxide, and water as observed herein and where different decomposition pathways have been proposed.^{7,11,16,17} In the case of Cu(110), the following decomposition reaction was suggested:²²



However, the presence of CO from the TPD cannot be ascribed to CO₂ decomposition in the mass spectrometer. Indeed the relative ion currents for CO₂ are 100 and 13 for $m/e = 44$ and 28, respectively, at an impact energy of 90 eV. The $m/e = 28$ signal is stronger in our case and cannot come from CO₂ ionization only. This might be explained by another decomposition pathway, supported by the presence of C (285 eV) at the end of TPD experiments, as described later by XPS experiments (Fig. 4).

In the multilayer regime, RR-TA molecules are expected to form H bonds. The activation energy and order of desorption in the multilayer regime can be estimated using the leading edge analysis.^{28,30,31} The method is based on the ansatz of desorption rate of Polanyi and Wigner,^{28,31}

$$r_d = \frac{-d\theta}{dt} = \nu(\theta)\theta^{n(\theta)} e^{-\frac{E_a(\theta)}{k_B T}}, \quad (1)$$

where ν is the pre-exponential factor, θ is the adsorbate coverage, n is the desorption order, and E_a is the activation energy. At the desorption threshold, the adsorption coverage has not changed significantly yet so that ν , n , and E_a can be considered as constants. If the sample is linearly heated and with an efficient differential pumping, the ion current detected $i_d(T)$, proportional to the partial pressure, is considered to be proportional to $\frac{-d\theta}{dt}$. Equation (1) can be, thus, rearranged as

$$i_d(T) \propto \left(\int_T^\infty i_d(T') dT' \right)^n \times e^{-\frac{E_a}{k_B T}}. \quad (2)$$

Taking the logarithm of Eq. (2), defined as $I_d(n, T)$, the previous equation can be linearized as

$$I_d(n, T) = \ln[i_d(T)] - n \ln \left[\int_T^\infty i_d(T') dT' \right] = K - \frac{E_a}{k_B T}. \quad (3)$$

Based on these considerations, $I_d(n, T)$ should have a linear dependence on $1/T$ with a slope E_a . $I_d(n, T)$ vs $1/T$ is then drawn for different values of n [Fig. 3(a)] to establish the best value that linearizes the signal at the onset of desorption. This analysis is applied in this work for the highest doses (40 and 54 Å) and for the desorption peaks $m/e = 58$ and 76 at 335 K. An order $n = 0$ (Fig. 3) is found to linearize at best the signal leading to an activation energy in the range 37–39 kcal/mol. The zero order is typical for autocatalytic desorption as in the case of water ice and agrees with a peak shift to higher temperatures with increasing doses (Fig. 1). Assuming that layers of RR-TA are hydrogen bonded through carboxylic dimer rings, a value of 18.5–19.5 kcal/mol is

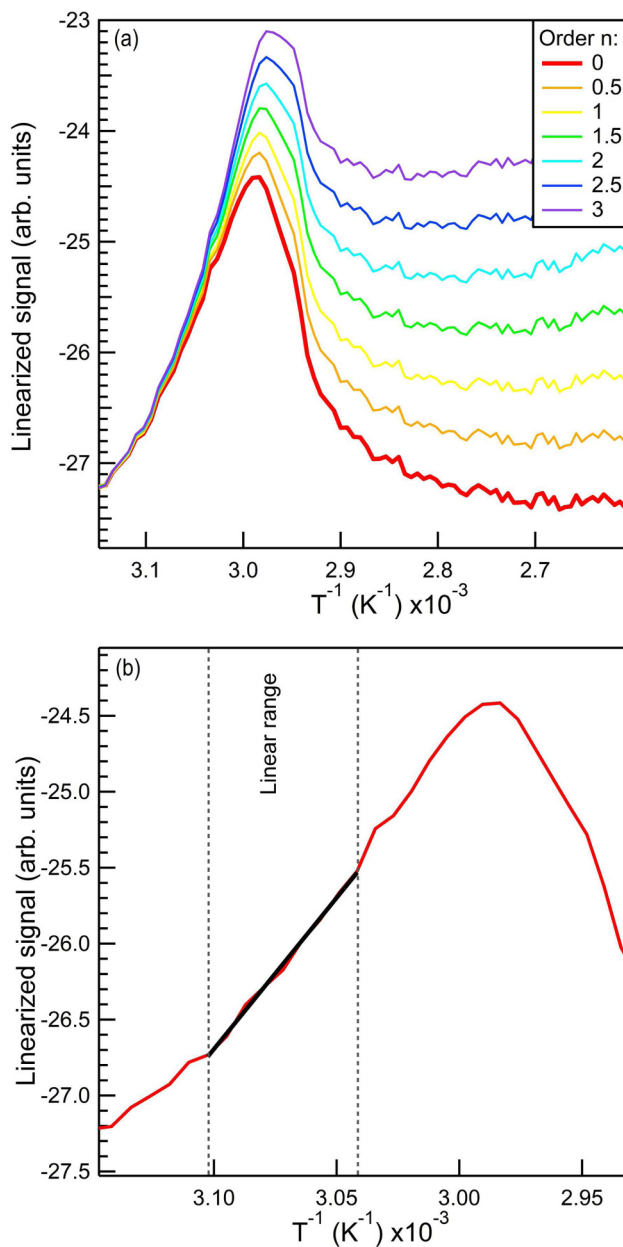


Fig. 3. (a) $I_d(n, T)$ linearized spectra plotted for different values of the desorption order n for $m/e = 58$ and 54 Å of TA. (b) Zoom on $I_d(n = 0, T)$. The linear fit of $I_d(n, T)$ at the edge of the desorption is represented by a black line.

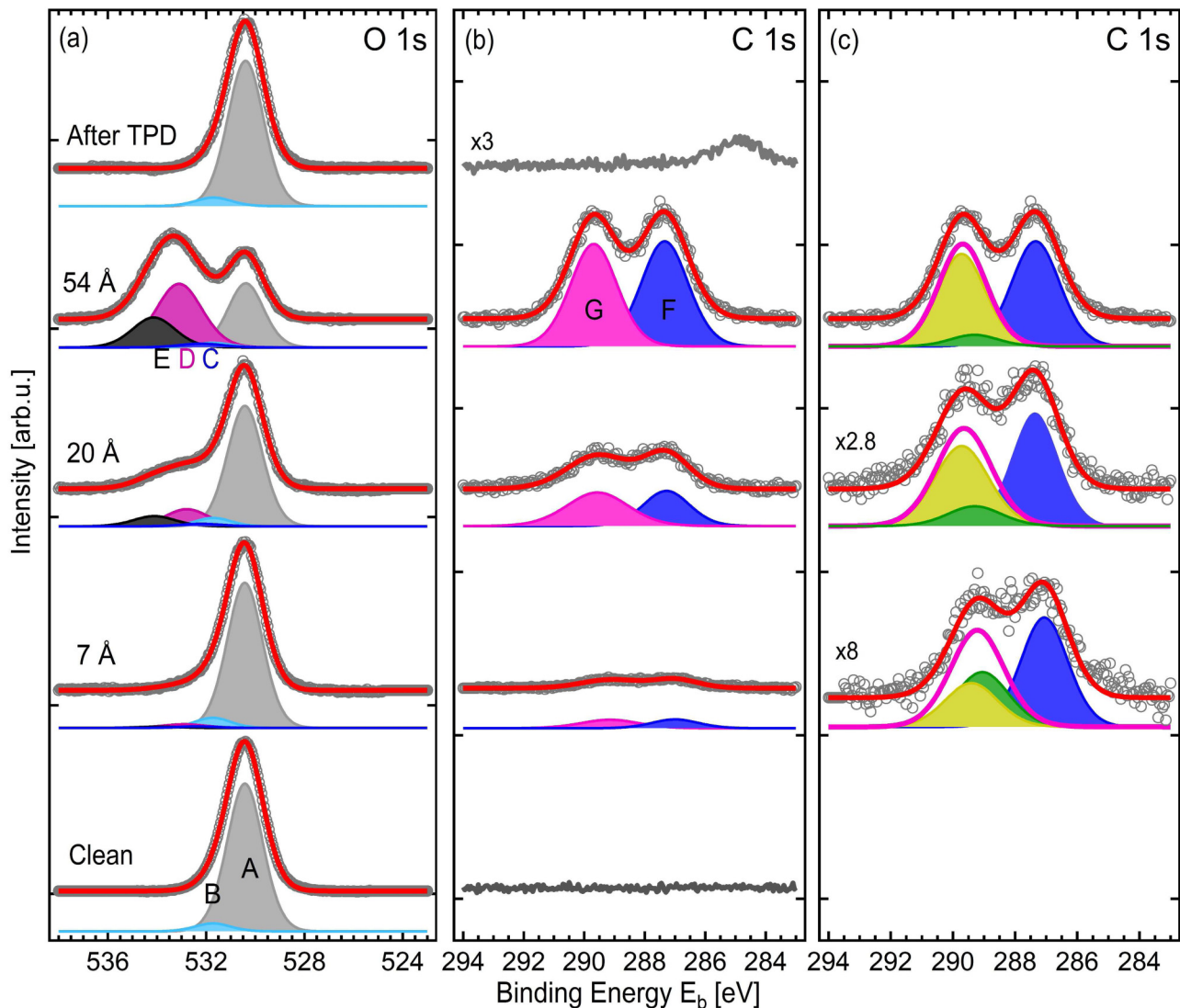


FIG. 4. Evolution of O1s (a) and C1s (b) peaks for RR-TA increasing doses on the rutile TiO₂(110) surface and after a TPD experiment; (c) same as (b) where the COO^- and COOH contributions are represented by the green (dark grey) the yellow (light grey) peaks, respectively. Spectra are vertically shifted for sake of clarity and normalized to the peak height. Raw data and fitted component sums are represented by grey dots and a red (solid) line, respectively. The peak fitting attribution is detailed in the text.

found per hydrogen bond, an order of magnitude in agreement with previous findings (14–15 kcal/mol).³²

B. Chemical nature of RR-TA adlayers on rutile TiO₂(110)

The chemical state of RR-TA molecules in the multilayer regime was investigated by XPS. The O1s and C1s spectra recorded for the clean TiO₂(110) surface and for increasing doses of RR-TA in the multilayer regime are presented in Fig. 4, and the corresponding fit parameters are summarized in Tables I and II. The fit performed is based on the previous considerations for RR-TA adsorption on rutile in the monolayer regime, where monotartrate molecules are observed. The O1s region of the clean rutile surface displays two contributions A and B at 530.4 and 531.7 eV, attributed to the TiO₂ O lattice atoms and the surface hydroxyl groups, respectively. Based on our previous work,²⁴ following

RR-TA deposition, three molecular contributions are identified, C, D, and E. Component C is related to the carboxylate functionality COO^- , D to a superposition of the C=O oxygen atoms and the OH alcohol groups (C—OH_{alc}), while E to the acidic OH groups (OH_{acid}). To substantiate this decomposition, the fit of the spectra at the highest coverage regime, the fit of the spectra is first performed on the highest coverage spectrum (54 Å), where the molecular contributions prevail over the TiO₂ substrate ones. The full-width at half-maximum (FWHM) and the peak binding energies determined are then kept fixed to fit the lower RR-TA dose spectra, where the presence of RR-TA results only in O1s peak shouldering at high binding energies. Some constraints on the peak areas are also used; for a 7 Å layer, close to the monolayer saturation, the analysis is consistent with a majority presence of monotartrate species. As the RR-TA dose increases, the intensities of peaks D and E, predominantly associated with the presence of protonated COOH groups, largely increases, suggesting the presence of fully

TABLE I. O1s peak parameters [binding energy (E_b) and relative area (RA)] used for the fit presented in Fig. 4. For components A and B, the FWHM is 1.7 eV, while for peaks C, D, and E, the FWHM is 2.0 eV. Error bars on E_b are of the order of 0.1 eV.

| Coverage | A—O _{lat} | | B—O _b H | | C—COO ⁻ | | D—C=O, C—OH _{alc} | | E—COOH | |
|-----------|--------------------|------------|--------------------|-----------|--------------------|-----------|----------------------------|------------|---------------|------------|
| | E_b (eV) | RA (%) | E_b (eV) | RA (%) | E_b (eV) | RA (%) | E_b (eV) | RA (%) | E_b (eV) | RA (%) |
| Clean | 530.4 | 94.9 ± 0.2 | 531.7 | 5.1 ± 0.2 | | | | | | |
| 7 Å | 530.4 | 89.0 ± 0.6 | 531.7 | 6.3 ± 0.1 | 532.3 | 1.1 ± 0.1 | 532.9 | 2.7 ± 0.4 | 533.9 | 0.9 ± 0.2 |
| 20 Å | 530.4 | 73.1 ± 3.5 | 531.7 | 5.1 ± 0.3 | 532.2 | 1.8 ± 0.3 | 532.9 | 12.5 ± 1.9 | 534.1 | 7.5 ± 2.5 |
| 54 Å | 530.4 | 33.6 ± 1.8 | 531.7 | 2.4 ± 1.9 | 532.3 | 3.4 ± 2.4 | 533.1 | 40.6 ± 3.4 | 533.9 | 20.0 ± 2.9 |
| After TPD | 530.4 | 94.4 ± 2.1 | 531.7 | 5.6 ± 2.1 | | | | | | |

protonated RR-TA molecules in the stacking of multilayers. Finally, following TPD, the O1s spectrum recovers its initial line shape.

For the C1s region, two contributions were identified at ~287 and ~289 eV.²⁴ The low binding energy peak (herein named F) was attributed to the central C—OH carbon atoms [C(2),C(3)], while the one at higher E_b (herein named G) to a superposition of COO⁻ and COOH groups. As a matter of fact, unlike what is observed for RR-TA molecules and other carboxylic acids adsorbed on Cu(110),^{24,33–36} components of carboxylate and carboxylic C atoms are not resolved on TiO₂(110), as previously observed for terephthalic acid adsorption on the same rutile surface.³⁷ Despite the superposition of COO⁻ and COOH contributions on TiO₂(110), information on the evolution of RR-TA chemical form may still be gained by considering the energy splitting between peaks F and G and the relative variation of their FWHM. Some constraints are used for the fit, namely, the intensity ratio F/G = 1, according to the molecular stoichiometry, and the FWHM of peak F, since no variation is expected for this peak which is related to the central C—OH carbon atoms. The fit is first performed for the highest coverage (54 Å), where the two peaks F and G are clearly discerned, to estimate the width of peak F and to use it for the fit of the lower coverage spectra. In the multilayer regime, upon increasing RR-TA doses, the energy splitting increases progressively from 2.1 to 2.4 eV. For the highest RR-TA dose, peak G is

found at 289.7 eV, in agreement with what was found for the carboxylic contribution of monotartrate RR-TA molecules on Cu(110).²⁴ At the same time, the width of the high binding energy peak reduces by 0.2 eV. These considerations overall suggest the prevalence of one species (COO⁻/COOH) over the other, and more precisely, the dominant contribution of carboxylic acid groups, as expected in the formation of multilayers of fully protonated RR-TA molecules. This conclusion matches what is observed for other carboxyl-containing molecules on the same rutile surface³⁷ and on other metallic surfaces.^{38,39} Moreover, for RR-TA adsorption on Cu(110), energy splitting values of 2.4 (Ref. 33) and 2.8 eV (Ref. 24) have been reported between the central C—OH and the COOH peaks. Splittings of 3.1 and 2.6 eV have also been measured on Cu(111)¹⁸ and Au(111).²⁴ The first value of 2.4 eV is in agreement with the present work for the highest RR-TA dose, again confirming the dominant contribution of COOH species in the multilayer regime. An energy splitting of 2.6 eV was also reported between the central COH and the COOH contribution for multilayer deposition of R malic acid on Cu(110).³⁶ To evidence the increasing contribution of the carboxylic component over the carboxylate one, two components are discerned in peak G, light and dark green (grey) peaks in Fig. 4(c). In the decomposition proposed the area ratio between COO⁻ and COOH peaks is fixed to the one obtained from the O1s analysis. It is clear how at coverages near the first layer saturation, the COO⁻ and COOH components are almost equivalent, while at increasing doses, the COOH prevails, as expected in nondeprotonated RR-TA multilayers. At last, a graphitic like C1s component at ~285 eV remains at the surface after the thermal decomposition of the molecule.

TABLE II. C1s peak parameters [binding energy (E_b), FWHM, and relative area (RA)] used for the spectra presented in Fig. 4. The binding energy difference between peaks B and A, $\Delta(B-A)$ (eV), is also reported. Error bars on E_b are of the order of 0.1 eV.

| Coverage | F—C(2),C(3) | | | G—COO ⁻ , COOH | | | $\Delta(G-F)$ [COO ⁻ , COOH —C(2), C(3)] |
|----------|---------------|--------------|-----------|---------------------------|--------------|-----------|---|
| | E_b (eV) | FWHM (eV) | RA (%) | E_b (eV) | FWHM (eV) | RA (%) | |
| Clean | 0 | | | 0 | | | 0 |
| 7 Å | 287.1 | 1.8 | 50 ± 9 | 289.2 | 2.1 | 50 ± 9 | 2.1 |
| 20 Å | 287.4 | 1.8 | 50 ± 7 | 289.6 | 2.1 | 50 ± 7 | 2.2 |
| 54 Å | 287.3 | 1.8 | 50 ± 7 | 289.7 | 1.8 | 50 ± 7 | 2.4 |

IV. SUMMARY AND CONCLUSIONS

The desorption and decomposition behavior of RR-TA molecules on TiO₂(110) was investigated by TPD. While molecules desorb more or less intact from the multilayer, they decompose in the monolayer regime, which indicates a strong interaction with the substrate. Information on the evolution of the RR-TA chemical state in the multilayer regime is gained by XPS and overall from the analysis of the O1s and C1s core levels; a majority presence of intact biacid molecules is observed in the multilayer regime.

ACKNOWLEDGMENTS

This work was supported by the French State Fund managed by the ANR within the Investissements d'Avenir program under Reference No. ANR-11-IDEX-0004-02, and more specifically within the framework of the Cluster of Excellence MATISSE.

- ¹C. J. Baddeley and N. V. Richardson, "Scanning tunneling microscopy in surface science," in *Nanoscience and Catalysis*, edited by M. Bowker and P. R. Davies (Wiley-VCH, Weinheim, 2010), Vol. 1, pp. 1–27.
- ²A. J. Gellman, W. T. Tysse, and F. Zaera, *Catal. Lett.* **145**, 220 (2015).
- ³D. Y. Murzin, P. Mäki-Arvela, E. Toukonniitty, and T. Salmi, *Catal. Rev.* **47**, 175 (2005).
- ⁴T. Mallat, E. Orglmeister, and A. Baiker, *Chem. Rev.* **107**, 4863 (2007).
- ⁵S. M. Barlow and R. Raval, *Surf. Sci. Rep.* **50**, 201 (2003).
- ⁶R. Raval, *J. Phys. Condens. Matter* **14**, 4119 (2002).
- ⁷K.-H. Ernst, *Phys. Status Solidi B* **249**, 2057 (2012).
- ⁸M. Ortega Lorenzo, S. Haq, T. Bertrams, P. Murray, R. Raval, and C. J. Baddeley, *J. Phys. Chem. B* **103**, 10661 (1999).
- ⁹M. Ortega Lorenzo, C. J. Baddeley, C. Muryn, and R. Raval, *Nature* **404**, 376 (2000).
- ¹⁰M. Ortega Lorenzo, V. Humblot, P. Murray, C. J. Baddeley, S. Haq, and R. Raval, *J. Catal.* **205**, 123 (2002).
- ¹¹T. E. Jones and C. J. Baddeley, *Surf. Sci.* **513**, 453 (2002).
- ¹²T. E. Jones and C. J. Baddeley, *J. Mol. Catal. A Chem.* **216**, 223 (2004).
- ¹³V. Humblot, S. Haq, C. Muryn, W. A. Hofer, and R. Raval, *J. Am. Chem. Soc.* **124**, 503 (2002).
- ¹⁴W. A. Hofer, V. Humblot, and R. Raval, *Surf. Sci.* **554**, 141 (2004).
- ¹⁵V. Humblot, S. Haq, C. Muryn, and R. Raval, *J. Catal.* **228**, 130 (2004).
- ¹⁶M. Mahapatra and W. T. Tysse, *Surf. Sci.* **629**, 132 (2014).
- ¹⁷S. Baldanza, J. Ardini, A. Giglia, and G. Held, *Surf. Sci.* **643**, 108 (2016).
- ¹⁸A. Rieger, C. Sax, T. Bauert, C. Wäckerlin, and K.-H. Ernst, *Chirality* **30**, 369 (2018).
- ¹⁹A. J. Gellman, Y. Huang, X. Feng, V. V. Pushkarev, B. Holsclaw, and B. S. Mhatre, *J. Am. Chem. Soc.* **135**, 19208 (2013).
- ²⁰V. Humblot, M. Ortega Lorenzo, C. J. Baddeley, S. Haq, and R. Raval, *J. Am. Chem. Soc.* **126**, 6460 (2004).
- ²¹B. S. Mhatre, V. Pushkarev, B. Holsclaw, T. J. Lawton, E. C. H. Sykes, and A. J. Gellman, *J. Phys. Chem. C* **117**, 7577 (2013).
- ²²B. Behzadi, S. Romer, R. Fasel, and K.-H. Ernst, *J. Am. Chem. Soc.* **126**, 9176 (2004).
- ²³C. L. Pang, R. Lindsay, and G. Thornton, *Chem. Rev.* **113**, 3887 (2013).
- ²⁴E. Meriggio, R. Lazzari, S. Chenot, P. David, C. Méthivier, X. Carrier, G. Cabailh, and V. Humblot, *Appl. Surf. Sci.* **493**, 1134 (2019).
- ²⁵C. Méthivier, V. Humblot, and C.-M. Pradier, *J. Phys. Chem. C* **120**, 27364 (2016).
- ²⁶U. Diebold, *Surf. Sci. Rep.* **48**, 53 (2003).
- ²⁷H. Onishi, T. Aruga, and Y. Iwasawa, *J. Catal.* **146**, 557 (1994).
- ²⁸H.-L. Thi Le, R. Lazzari, J. Goniakowski, R. Cavallotti, S. Chenot, C. Noguera, J. Jupille, A. Koltsov, and J.-M. Maigne, *J. Phys. Chem. C* **121**, 11464 (2017).
- ²⁹See: <https://webbook.nist.gov/cgi/cbook.cgi?ID=C87694&Mask=200#Mass-Spec>.
- ³⁰P. A. Redhead, *Vacuum* **12**, 203 (1962).
- ³¹D. A. King, *Surf. Sci.* **47**, 384 (1975).
- ³²S. Scheiner, *Hydrogen Bonding: A Theoretical Perspective* (Oxford University, New York, 1997).
- ³³M. Parschau, B. Behzadi, S. Romer, and K.-H. Ernst, *Surf. Interface Anal.* **38**, 1607 (2006).
- ³⁴C. Roth and K.-H. Ernst, *Top. Catal.* **54**, 1378 (2011).
- ³⁵C. Roth, M. Parschau, and K.-H. Ernst, *Chem. Phys. Chem.* **12**, 1572 (2011).
- ³⁶C. Roth, D. Passerone, L. Merz, M. Parschau, and K.-H. Ernst, *J. Phys. Chem. C* **115**, 1240 (2011).
- ³⁷W. Zhang, L. Cao, L. Wan, L. Liu, and F. Xu, *J. Phys. Chem. C* **117**, 21351 (2013).
- ³⁸M. Wühh, J. Weckesser, and C. Wöll, *Langmuir* **17**, 7605 (2001).
- ³⁹W. Zhang, A. Nefedov, M. Naboka, L. Cao, and C. Wöll, *Phys. Chem. Chem. Phys.* **14**, 10125 (2012).