

## PAPER

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[View Journal](#) | [View Issue](#)Cite this: *Nanoscale Adv.*, 2022, 4, 4517MoS<sub>2</sub> oxidative etching caught in the act: formation of single (MoO<sub>3</sub>)<sub>n</sub> molecules†Saeed Sovizi, <sup>a</sup> Sergio Tosoni <sup>\*b</sup> and Robert Szoszkiewicz <sup>\*a</sup>

We report the presence of sub-nm MoO<sub>x</sub> clusters formed on basal planes of the 2H MoS<sub>2</sub> crystals during thermal oxidative etching in air at a temperature of 370 °C. Using high resolution non-contact atomic force microscopy (AFM) we provide a histogram of their preferred heights. The AFM results combined with density functional theory (DFT) simulations show remarkably well that the MoO<sub>x</sub> clusters are predominantly single MoO<sub>3</sub> molecules and their dimers at the sulfur vacancies. Additional Raman spectroscopy, and energy and wavelength dispersive X-ray spectroscopies as well as Kelvin probe AFM investigations confirmed the presence of the MoO<sub>3</sub>/MoO<sub>x</sub> species covering the MoS<sub>2</sub> surface only sparsely. The X-ray absorption near edge spectroscopy data confirm the MoO<sub>3</sub> stoichiometry. Taken together, our results show that oxidative etching and removal of Mo atoms at the atomic level follow predominantly *via* formation of single MoO<sub>3</sub> molecules. Such findings confirm the previously only proposed oxidative etching stoichiometry.

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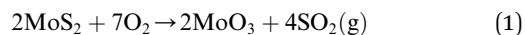
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## Introduction

Thin MoS<sub>2</sub> crystals have become the subject of intense research in the last 10 years due to many applications in sensing, optoelectronics and energy harvesting.<sup>1,2</sup> In addition, various methods have been devised for obtaining wafer-scale poly- and single-crystalline MoS<sub>2</sub> monolayers<sup>3,4</sup> as well as their heterogeneous integration.<sup>5,6</sup> The temperature within MoS<sub>2</sub> based devices has been measured to locally surpass 250 °C,<sup>7</sup> which besides affecting performance, makes them prone to oxidation. While most abundant 2H MoS<sub>2</sub> crystals do not oxidase easily,<sup>2</sup> several seminal papers about their local oxidative etching above 300 °C were reported in 2013.<sup>8–10</sup> Till now, many types of surface reactions of 2H MoS<sub>2</sub> crystals with oxygen, much beyond oxidative etching, have been discovered *via* high-resolution tools, such as atomic force microscopy (AFM), scanning tunneling microscopy (STM) and scanning transmission electron microscopy (S-TEM). These reactions involve chemical oxygen adsorption, oxygen dissociative reactions, and other oxidation mechanisms, as well as oxygen penetration between stacked MoS<sub>2</sub> layers.<sup>2</sup> Selected products of such reactions have been identified *via* X-ray photoelectron spectroscopy (XPS), X-

ray diffraction (XRD), X-ray absorption spectroscopy (XAS) and Raman studies.<sup>2,11–13</sup> However, despite substantial experimental and computational efforts, oxidative etching is still not well understood due to various competing reaction mechanisms.<sup>2</sup>

It has been proposed but not confirmed on the atomic level that MoS<sub>2</sub> oxidative etching at elevated temperatures obeys the following stoichiometry:



The products of such a reaction leave the MoS<sub>2</sub> surface, which results in formation of triangular etch pits due to hexagonal surface symmetry.<sup>8,14,15</sup> Most of the computational studies have been focused on the initial steps of defective or perfect MoS<sub>2</sub> surface oxidation, namely, dissociation of O<sub>2</sub> molecules, and adsorption of oxygen atoms and their complexes with S and/or Mo atoms, as well as formation of sulfur vacancies by the action of an attacking oxygen molecule.<sup>11,16,17</sup> However, according to our knowledge, it has never been confirmed that at the atomic level, oxidative etching proceeds through eqn (1).

Herein, we report the surface presence of the sub-nm MoO<sub>x</sub> clusters in the oxidative etching of the MoS<sub>2</sub> samples at 370 °C, where the Mo oxides start to form on the MoS<sub>2</sub> surface. Through accurate AFM investigations together with density functional theory (DFT) simulations as well as local Raman spectroscopy, we discuss the origin, chemical composition and most probable conformations of the sub-nm MoO<sub>x</sub> clusters. Additional energy dispersive X-ray spectroscopy (EDS), wavelength dispersive X-ray spectroscopy (WDS), X-ray absorption near edge spectroscopy (XANES) and Kelvin probe AFM investigations agreed with the AFM and DFT findings.

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† Electronic supplementary information (ESI) available: Topography images of the samples, separate height histograms of the samples, discussion of fitting parameters, additional DFT simulations, more details on Raman maps, and additional SEM-EDS and SEM-WDS investigations. See <https://doi.org/10.1039/d2na00374k>

## Experimental

Laboratory grade 2H molybdenite crystals were bought from SPI Supplies, West Chester, PA, USA, catalogue number #429MM-AB. The MoS<sub>2</sub> samples have been mechanically exfoliated using a standard blue tape and transferred on fine polished and basically undoped <111> Si crystals with a resistivity more than 10 000 Ω cm, bought from ITME, Warsaw, Poland. Prior to a sample transfer, the Si substrates were ultrasonically cleaned in sequential steps with deionized water, acetone, ethanol and isopropanol for 10 min each. After each ultrasonication step, the substrate was dried with pure N<sub>2</sub>. At the end of ultrasonic cleaning, oxygen plasma (PE-25, Plasma Etch, 150 mTorr O<sub>2</sub>, 100 W RF power, 180 s) was used to get rid of any remaining residues/pollutants. Due to subsequent heating studies at temperatures of more than 350 °C, no annealing or any other cleaning methods have been used to remove any remaining traces of the blue tape from the Si substrates. After exfoliation, the samples were heated in air at 370 °C for 15 minutes on a standard hot plate covered with a quartz Petri dish to ensure a controlled atmosphere and proper distribution of temperatures within a heating zone. The hot plate was calibrated with a standard thermocouple and a Pt thermometer to yield a temperature error of ±2 °C on the sample surface. For cluster analysis, two separate samples were investigated. 1100+ clusters were selected from sample 1, and 350+ clusters were selected from sample 2.

### AFM measurements

A Cypher S AFM (Asylum Research, Goleta, CA, USA) in a contact mode in air and with an MLCT-F cantilever (Bruker, Santa Barbara, CA, USA) was utilized to perform scratching/removal of the MoO<sub>3</sub> from the thermally oxidized MoS<sub>2</sub> flakes. Each time, a specific area was scratched several times with a scanning rate between 2.60 and 4.88 Hz at loads between 0.6 ± 0.1 and 12 ± 1 nN. Loads were calibrated from a spring constant and sensitivity factors, which were obtained from thermal resonance of the used cantilever using routines available within the data acquisition software. The same AFM but in a light tapping mode in air was utilized for topography imaging of the sample before and after scratching. An AC160TS-R3 cantilever (Asylum Research) was used to collect high-resolution AFM images with at least 1536 points per line. All raw AFM data were treated with the Gwyddion software (v.2.51, Czech Metrology Institute, Brno, Czech Republic).<sup>18</sup> Standard line-by-line first or second-order horizontal flattening together with removal of occasional glitches was applied. No other image treatment and/or conditioning were performed.

### DFT calculations

DFT calculations were performed on periodic slab models with the code VASP 6.2.<sup>19,20</sup> Mo(4p, 4d, and 5s), S(3s and 3p) and O(2s and 2p) electrons were treated explicitly, and the interaction between the inner electrons and the nuclei was treated with the projector augmented wave method.<sup>21,22</sup> The exchange-correlation functional proposed by Perdew, Burke and

Ernzerhof (PBE)<sup>23</sup> was adopted. The long-range dispersion was added to the potential according to the DFT+D3 scheme with a Becke–Johnson damping function.<sup>24,25</sup> Thresholds of 10<sup>−5</sup> eV and 0.01 eV Å<sup>−1</sup> were adopted to truncate the electronic and ionic loops. A smearing of 0.05 eV was used to ensure convergence of the self-consistent field process. The lattice relaxation of the MoS<sub>2</sub> monolayer was done with a mesh of 6 × 6 × 6 special *k*-points and a kinetic cutoff of 600 eV, yielding lattice parameters in good agreement with the experiment (*a* and *b* = 0.315 nm, and *c* = 1.207 nm). Static relaxations of (Mo<sub>x</sub>O<sub>y</sub>)<sub>*n*</sub>/MoS<sub>2</sub> structures were then performed at the *Γ* point with a cutoff of 400 eV. Spin-polarization effects were included in all calculations. Dipole and quadrupole corrections to the total energy were applied along the non-periodic direction for all calculations on slab models.<sup>26</sup> In order to avoid spurious interactions between the replica of the slab models, a vacuum region of at least 1.5 nm was included along the non-periodic direction in the supercell. All calculations were carried out adopting a (5 × 2√3) MoS<sub>2</sub> supercell. This box is large enough (1.58 nm × 1.09 nm) to simulate the formation of isolated mono- and dimer species. The (MoO<sub>3</sub>)<sub>ML</sub>/MoS<sub>2</sub> interface is built superimposing a (4 × 3) supercell of MoO<sub>3</sub> onto the aforementioned (5 × 2√3) MoS<sub>2</sub> supercell. The residual strain arising from the lattice mismatch (0.2% and −1.3% along *a* and *b*) is released adapting MoO<sub>3</sub> to the MoS<sub>2</sub> lattice. The formation energy, *E<sub>f</sub>*, has been calculated through the following equation:

$$E_f = \{E(\text{Mo}_x\text{O}_y/\text{MoS}_2) - E(\text{MoS}_2)\} - x\mu_{\text{Mo}} - y\mu_{\text{O}} \quad (2)$$

*E*(Mo<sub>x</sub>O<sub>y</sub>/MoS<sub>2</sub>) is the total energy of an oxidized MoS<sub>2</sub> supercell, which has one MoO<sub>x</sub> monomer or dimer. *E*(MoS<sub>2</sub>) is the total energy of a pure MoS<sub>2</sub> supercell with or without a sulphur vacancy. *μ<sub>Mo</sub>* and *μ<sub>O</sub>* are chemical potentials of Mo and O atoms, respectively.

The structures reported in the paper were obtained by static relaxations, starting from initial guess structures designed by chemical intuition. The choices to consider among all possible Mo<sub>x</sub>O<sub>y</sub> species were the MoO<sub>1</sub>O<sub>2</sub> and MoO<sub>1</sub>O<sub>3</sub> monomers, as well as Mo<sub>2</sub>O<sub>4</sub> and Mo<sub>2</sub>O<sub>6</sub> dimers, as is discussed in the subsequent sections.

The height of the adsorbed Mo<sub>x</sub>O<sub>y</sub> clusters was defined as the difference between the *Z* coordinate of the apical O atom of the cluster with respect to the average of the S atoms from the MoS<sub>2</sub> basal plane.

### Raman measurements

The Raman map was collected using a Horiba Jobin Yvon LabRAM HR800 Raman spectrometer. Measurements were made using a 532.0 nm laser excitation line with a 100× objective lens and a 1800 g mm<sup>−1</sup> grating. The power was set below 5 mW to minimize any oxidative effects during Raman mapping. An area of 15 × 15 μm consisting of 30 × 30 points with a 0.5 μm step size was chosen for mapping. At each point, data in each spectral window were collected for 30 s. Two spectral windows at each point were taken.



### Kelvin-probe force microscopy (KPFM) measurements

A Cypher S AFM double-pass amplitude-modulated KPFM with ASYELEC-01 cantilevers (Asylum Research) was utilized. During the KPFM measurements, the bias voltage was applied to the tip, while the sample was grounded. The raw, *i.e.*, untreated KPFM tip-sample CPD data in volts ("Nap potential") were analyzed with Gwyddion.<sup>18</sup>

### XANES measurements

The chemical state and local structure of MoS<sub>2</sub> flakes on Si substrates were examined using X-ray absorption fine structure (XAFS) measurements. XAFS spectra were collected at the PEEM/XAS end station at the Solaris National Synchrotron Radiation Centre in Cracow, Poland. XAS beam parameters involved a photon energy range of 300–1600 eV with a relative energy resolution ( $\Delta E/E$ ) not worse than 0.00 025, a beam size at the sample of 0.200 mm horizontally and 0.050 mm vertically and a photon flux at the sample of 109–1010 [ph/s/100 mA]. The sample was mounted on a standard nonmagnetic and conductive Elmitec PEEM sample holder for scanning electron microscopy (SEM)-EDS measurements.

The studies were performed with a scanning electron-ion microscope Crossbeam 540× (Zeiss) equipped with an EDS X-MAX detector (Oxford Instruments) with an active area of 80 mm<sup>2</sup>. An EDS spectrometer was calibrated for the Cu K $\alpha$  line and controlled with a Smart SEM (ZEISS) and the Aztec 4.2 (Oxford Instruments) software. The surface morphology was imaged using an SESI and In-Lens detectors (mix signal) at an accelerating voltage of 5 kV, a working distance from the column WD = 5 mm and a stage tilt of 0°.

### SEM-WDS measurements

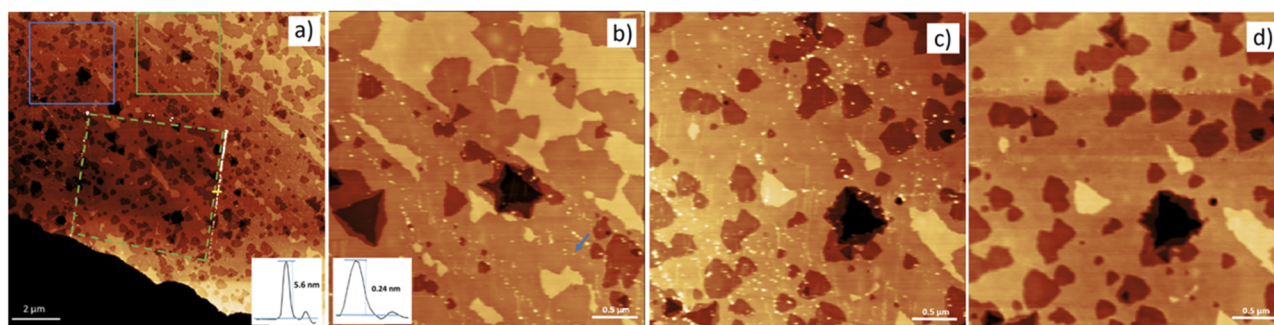
A Cameca SX100 Electron Probe Microanalyzer (EPMA, manufactured at Courbevoie Cedex, France, 2000) equipped with 4 WDS detectors was used. Electrons were accelerated to 7.5 keV and regulated to a 10 nA beam current. The four WDS detectors were set to PC0-TAP-PET-LPET diffracting crystal configuration for point analysis to find O–Si–S–Mo concentrations on

modified flakes. The position and reference intensities of O K $\alpha$  were calibrated on a LiNbO<sub>3</sub> crystal (Li: 4.6947 wt%, Nb: 62.8395 wt%, and O: 32.4658%). A molybdenite crystal (S: 40.06 wt%; Mo: 59.86 wt%) was used for referencing the positions and intensities of the S K $\alpha$  and Mo L $\alpha$  peaks. The Si K $\alpha$  peak position and intensity were calibrated by using a silicon substrate as a reference sample. Counting times were set to 20 s on the peak and 10 s on single background position with a precise slope. Optimal single-background positions and precise slopes were graphically estimated using the HussariX software (<https://github.com/sem-geologist/HussariX>). X-ray intensities were calculated in wt% using the Cameca Peak-sight 5.1 software with a built-in X-PHI matrix correction method in conjunction with the built-in database of Mass Attenuation Coefficients. To eliminate the sample charging, conductive contacts were made with the sample through a standard conductive carbon tape (SPM supplies).

## Results and discussion

Fig. 1 shows a high-resolution non-contact AFM topography image of a thick MoS<sub>2</sub> flake which was heated at 370 °C for 15 minutes. Under such conditions, the basal MoS<sub>2</sub> layer has been etched at a large portion of the flake. Simultaneously, underlying MoS<sub>2</sub> layers have been incompletely etched. Etching resulted in formation of triangular etch pits of various depths, between one to seven MoS<sub>2</sub> layers. In addition, some small and isolated spots are barely visible in Fig. 1(a). To investigate whether the spots are not AFM imaging artefacts, part of the image in Fig. 1(a) – marked by a dashed green box – was square rastered several times with an AFM tip.

As a result of rastering, a collection of white "debris" with a typical height of *ca.* 6 nm appeared at the border of the rastered zone. Thus, the etched surface was indeed covered with a considerable number of "particles". To investigate these "particles" in greater detail, magnified AFM topographs of non-rastered zones are shown in Fig. 1(b) and (c). Therein, many clearly distinguishable and sub-nm thick clusters can be identified. An inset in Fig. 1(b) shows the typical height of such



**Fig. 1** Non-contact AFM topography of a thermally etched MoS<sub>2</sub> flake. (a) Image taken after rastering the area depicted by a dashed-green box. Inset: the height profile of piled-up debris along the yellow line. Image Z-scale 6.5 nm. (b) Magnified area from a solid-green box in (a). Inset: a typical cross-section along a single MoO<sub>x</sub> cluster marked by a blue arrow in (b). Image Z-scale 4.2 nm. (c and d) Magnified area depicted by a blue box in (a) before (c) and after (d) rastering the whole flake. Image Z-scale is 3.8 nm in (c) and 3.3 nm in (d). Brighter colors represent higher Z-values.





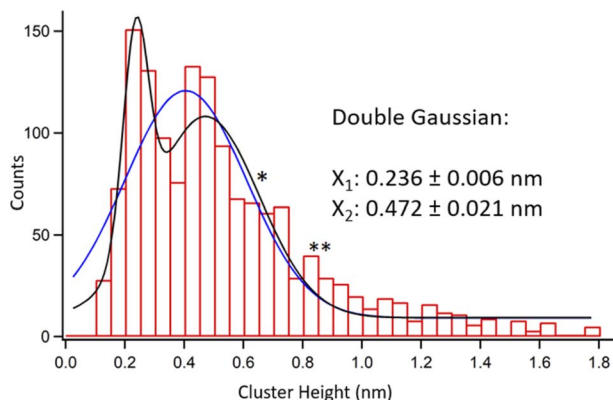


Fig. 2 Height histogram of randomly chosen 1469 oxide clusters on two thermally etched MoS<sub>2</sub> flakes.

clusters to be only *ca.* 0.2 nm. These observations suggest that the “debris” discussed earlier originated from the piling up many of such sub-nm clusters.

Next, the sample was rastered in a contact mode to detect only loosely surface-bound clusters. Then, a high resolution non-contact AFM image of the same area was collected for further studies, see Fig. S1, ESI†. By comparing the topography before and after rastering, see Fig. 1(c) and (d), one concludes that AFM scratching displaced a great majority of sub-nm clusters without inducing any damage to the basal plane of the MoS<sub>2</sub> flake. For further analyses, 1469 movable clusters from two separately prepared samples were randomly chosen, and their height profiles were obtained.

Fig. 2 shows a collective height histogram of the chosen clusters. The maximum height reading error in AFM experiments was only 0.03 nm. Therefore, a 0.05 nm bin size was chosen for the histogram. See Fig. S2–S4 in the ESI† regarding reproducibility of the results. A considerable portion of the clusters in Fig. 2 is shorter than the typical thickness of a single MoS<sub>2</sub> layer, which is *ca.* 0.7 nm. This proves that movable clusters are not remainders of the non-etched MoS<sub>2</sub>. Next, Gaussian fits to the histogram were produced. Single Gaussian yielded one peak at  $0.405 \pm 0.020$  nm. Double Gaussian produced two peaks at  $0.236 \pm 0.006$  nm and  $0.472 \pm 0.021$  nm. Both visually and *via* a  $\chi^2$  test, the double Gaussian fit is much better than the single Gaussian, see Table S1, ESI†. Noteworthy, due to the large number of the data points within the histogram, its maxima are obtained with smaller errors than the AFM height reading errors for individual data points.

The clusters in Fig. 2 were either reabsorbed or originated from single etching events on the MoS<sub>2</sub> samples. The latter means that they were produced and trapped at the same spots on the surface. More than 99% of the clusters were found exclusively within the etched regions, *i.e.*, away from any intact MoS<sub>2</sub> basal planes, see Fig. 1(a) and (b) as well as S2 in the ESI†. If such clusters detached and reabsorbed elsewhere, then they would appear also on the intact basal surface. Moreover, most of the reabsorbed species would locate at edges of the etched zones, which is also not the case here. Due to sub-nm heights of

the obtained clusters and their exclusive presence only within the etched zones, we hypothesize that they are oxidation products caught in the act just after single etching events.

In order to elucidate the chemical composition of the clusters, DFT simulations were performed to investigate conformations taken by sub-nanometric Mo<sub>x</sub>O<sub>y</sub> fragments onto the MoS<sub>2</sub> monolayer. We did not aim at modelling the whole oxidation pathway, which is still unclear and has been the subject of several studies.<sup>2,11,12,17</sup> Thus, we concentrated on the final outcomes of the oxidative events and classified them by topographic heights as well as thermodynamic stability of the resulting Mo<sub>x</sub>O<sub>y</sub> species, see eqn (2). Four types of molybdenum oxide species were considered: Mo<sub>1</sub>O<sub>2</sub> and Mo<sub>1</sub>O<sub>3</sub> monomers and their dimers, *i.e.*, Mo<sub>2</sub>O<sub>4</sub> and Mo<sub>2</sub>O<sub>6</sub>. They agree with XPS and XAS experimental observations which show that MoO<sub>3</sub> is the most expected oxide, and MoO<sub>2</sub> could also be marginally

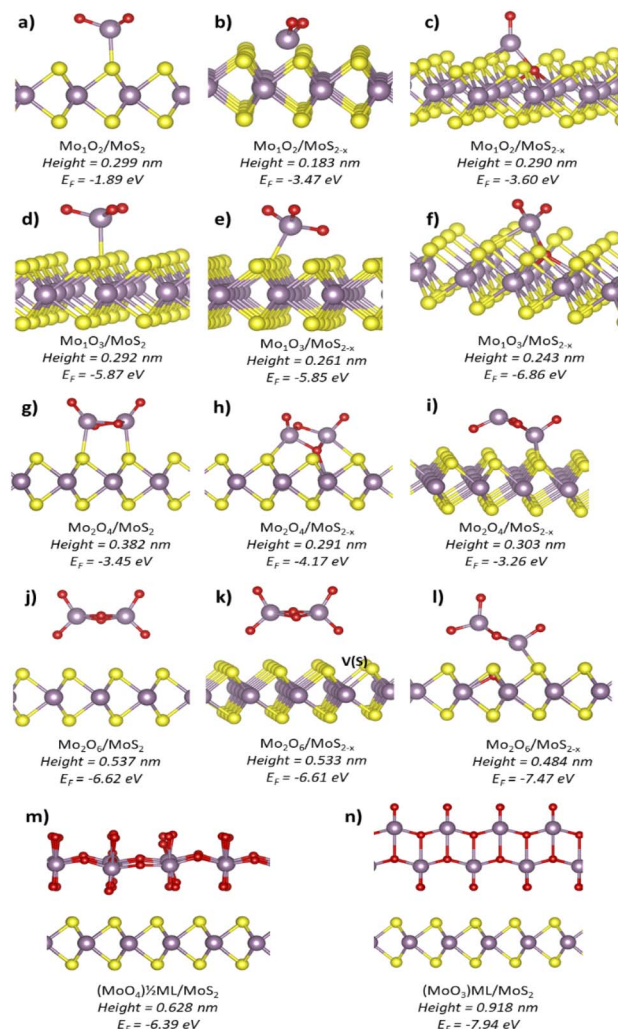


Fig. 3 Side views of DFT structures along with their expected heights and formation energies (EF) per considered Mo<sub>x</sub>O<sub>y</sub> unit. (a, d, g and j) Mo<sub>x</sub>O<sub>y</sub> species on stoichiometric MoS<sub>2</sub>. (b, e, h and k) Mo<sub>x</sub>O<sub>y</sub> species adsorbed near the SSV. (c, f, i and l) Mo<sub>x</sub>O<sub>y</sub> species adsorbed at the SSV. (m and n) Continuous MoO<sub>3</sub> structures: the half-monolayer and full monolayer, respectively.



present.<sup>2,27,28</sup> Furthermore, out of all the Mo oxides, the  $\text{MoO}_2$  and  $\text{MoO}_3$  are thermodynamically preferred, and the  $\text{MoO}_3$  crystals are the most stable.<sup>29</sup>

In order to verify the role of point defects as active centers for oxidation, each  $\text{Mo}_x\text{O}_y$  type was simulated onto three  $\text{MoS}_2$  surface spots: (i) stoichiometric  $\text{MoS}_2$ , (ii) near single sulfur vacancy (SSV) and (iii) at the SSV. The cluster height above the  $\text{MoS}_2$  monolayer was calculated referring to its topmost atom and the average  $Z$  coordinate of the atoms from the top S layer in the dichalcogenide slab.

Fig. 3 presents our DFT simulation results. For the  $\text{Mo}_1\text{O}_2$  molecule, in Fig. 1(a)–(c), the simulated heights changed from 0.299 nm on pristine  $\text{MoS}_2$  to 0.183 nm near the SSV and to 0.290 nm at the SSV. The structures near and at the SSV had comparable thermodynamic stability, while the structure on pristine  $\text{MoS}_2$  was the least stable. In the case of  $\text{Mo}_1\text{O}_3$  monomers, Fig. 1(d)–(f), more stable species were obtained compared to  $\text{Mo}_1\text{O}_2$ . Their simulated heights changed from 0.292 nm on pristine  $\text{MoS}_2$  to 0.261 nm near the SSV and 0.243 nm at the SSV. The  $\text{Mo}_1\text{O}_3$  structure at the sulfur vacancy was the most stable and matched very well with our experimental results.

A remarkable increase in height was reported when the dimers were considered. For  $\text{Mo}_2\text{O}_4$ , the height varied from 0.382 nm on stoichiometric  $\text{MoS}_2$  to 0.291–0.303 nm on defective  $\text{MoS}_2$ . The  $\text{Mo}_2\text{O}_6$  species positioned themselves at larger distances from the  $\text{MoS}_2$  surface than  $\text{Mo}_2\text{O}_4$  dimers. Such

distances decreased from 0.537 nm on a pristine sample to 0.533 nm near the SSV and 0.484 nm at the SSV. The last structure, shown in Fig. 3(l), resulted from spontaneous reorganization of the  $\text{Mo}_2\text{O}_6$  cluster at the SSV. During such a process, the cluster lost an O atom, which filled the S vacancy. The resulting structure was the most stable out of all considered dimers. The curious feature of complete detachment of an O atom from the cluster has been double-checked. Despite several attempts, it was not possible to identify a stable structure with the O atom entering the S vacancy and the  $\text{Mo}_2\text{O}_6$  molecule remaining intact. Interestingly, similarly to the case of monomers, the  $(\text{MoO}_3)_n$  species being more stable than the corresponding  $(\text{MoO}_2)_n$  species.

Formation of continuous  $\text{MoO}_x$  layers was also simulated. Fig. 3(m) shows a half monolayer formed by single  $\text{MoO}_6$  octahedra with a resulting stoichiometry of  $\text{MoO}_4$  and a height of 0.628 nm above  $\text{MoS}_2$ . Fig. 3(n) shows a full  $\alpha\text{-MoO}_3$  monolayer with two stacked planes of  $\text{MoO}_6$  octahedra and a height of 0.918 nm above  $\text{MoS}_2$ . For comparison, the height of a single  $\text{MoS}_2$  layer was 0.60 nm at a coherent level of calculations.

Comparing the heights obtained by simulations with the results of the AFM measurements, the most prominent type of the clusters in Fig. 2 corresponds to the  $\text{MoO}_3$  monomer at the SSV. Its simulated height of  $0.243 \pm 0.001$  nm matches extremely well with  $0.236 \pm 0.006$  nm observed in the experiments. The simulated structure of the  $\text{MoO}_3$  species where a Mo atom is extracted from the  $\text{MoS}_2$  surface is also provided, see

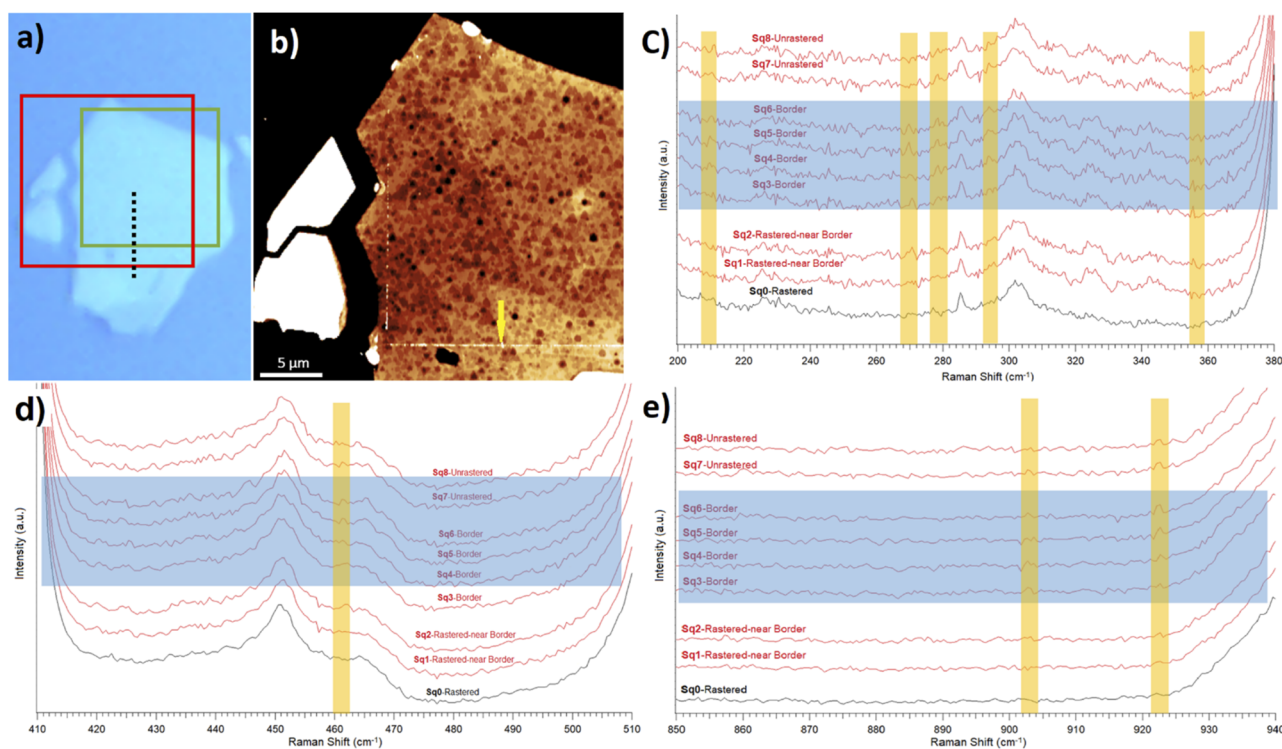


Fig. 4 Raman and AFM studies on a selected  $\text{MoS}_2$  flake. (a) Optical image of the investigated flake. A green box shows the rastered region. (b) NC-AFM topography image (6.2 nm Z-scale) of an area depicted by a red box in (a). Brighter colors represent a higher Z-scale. A yellow arrow shows a border between rastered and unrastered zones, where Mo oxide clusters are piled up. The scale bar is 5  $\mu\text{m}$ . (c–e) Raman spectra obtained from the flake along the dashed black line in (a). Blue rectangles show the border area in the overlaid spectra, and yellow windows mark Raman shifts with noticeable changes between rastered and border zones.



Fig. S5 in the ESI.† Such a structure was coordinated by three oxygen atoms and showed the height of 0.222 nm, which is still close to the first peak in Fig. 2. Other types of MoO<sub>3</sub> adsorption modes yielded substantially different heights. Such a spectacular agreement between the experimental AFM height data and simulated heights of the expected clusters seems truly significant. It strongly points towards single MoO<sub>3</sub> molecules produced on the MoS<sub>2</sub> surface predominantly at sulfur vacancies during its oxidative etching.

It has been already shown that during oxidative etching, atmospheric oxygen reacts with a MoS<sub>2</sub> basal plane at defect sites to produce volatile oxide compounds. Two kinds of molecular mechanisms were proposed, *i.e.*, either etching along the zig-zag (ZZ)-Mo or ZZ-S edges.<sup>8,15</sup> Exclusive presence of the MoO<sub>3</sub> molecules within the single sulfur vacancies is strong proof for such mechanistic assumptions since it confirms the stoichiometry of eqn (1) and shows the MoO<sub>3</sub> caught on the surface just before leaving it.

Next, we discuss the second peak in the height histogram. Its best match from simulations is the MoO<sub>3</sub> dimer at the SSV, which also is the most stable out of all the considered dimers. To ponder on a role of sulfur vacancies in more detail, we calculated the formation energies of dimers aggregating from two MoO<sub>3</sub> monomers adsorbed separately, see Table S2 in the ESI.† We obtained an energy gain of 0.69 eV if the dimers aggregated at the SSV compared to aggregation on a pristine surface. This suggests that SSVs act as nucleation centers both for monomers and dimers.

The second histogram peak in Fig. 2 is lower in amplitude and much wider than the first peak. Its lower amplitude relates to lower likelihood of obtaining a MoO<sub>3</sub> dimer *vs.* monomer at the reaction spot. The width of the second peak might as well relate to other chemical forms of the MoO<sub>3</sub> expected on the surface. For example, a single star (\*) marks the height associated with half of the MoO<sub>3</sub> layer. This structure is presented in Fig. 3(m) and is also known as a metastable monoclinic β-MoO<sub>3</sub> structure.<sup>29</sup> A similar height is also expected from thermodynamically favored chain-like structures of the MoO<sub>6</sub> octahedra, *a.k.a.* molybdenum polyoxides.<sup>30</sup> Two stars (\*\*) in Fig. 2 mark the expected height of fully formed fragments of the most stable α-MoO<sub>3</sub> monolayer, see Fig. 3(n).

To further report the MoO<sub>3</sub> stoichiometry of the clusters, we performed a detailed Raman analysis. We concentrated on

comparisons between the high quality Raman spectra collected around the border between rastered (without oxide) and unrastered (with oxide clusters) MoS<sub>2</sub> surfaces. We extracted such data from the Raman map, see the ESI.† Fig. 4(a) and (b) show the optical and topography images of the selected area. Fig. 4(c)–(e) display the Raman spectra taken along a dashed line from Fig. 4(a). Such a line originates within a rastered zone, passes through a border region and ends up in an unrastered zone. Similarly, the Raman spectra displayed in Fig. 4(d) and (e) are arranged from the ones collected on unrastered zone (on top) towards the ones collected on the rastered zone (on the bottom).

While it is difficult to claim any presence of typical Raman peaks associated with the MoO<sub>3</sub> oxide, the data in Fig. 4 show differences between rastered and unrastered zones in the case of several particular Raman shifts. For example, there is a clear difference between border and rastered regions at around the 278 cm<sup>−1</sup> Raman shift in Fig. 4(c). Namely, the peaks from within the MoO<sub>3</sub> containing zone in the first six spectra (from top) collected in unrastered and border regions are absent in the last two Raman spectra collected in the MoO<sub>3</sub> depleted zone. Similar observations can be extended to other Raman shifts associated in the literature, see Table 1, with the presence of MoO<sub>3</sub> and MoO<sub>3−x</sub>. Often, the differences are not striking, which confirms difficulties associated with collecting Raman spectra of (sub) monolayers. Till now – at least according to our knowledge – obtaining the Raman signature of thin MoO<sub>3</sub> layers onto MoS<sub>2</sub> flakes has not been reported. At best, some authors were able to report Raman shifts only for fully established MoO<sub>3</sub> crystals on Si substrates.<sup>2</sup>

As mentioned earlier, we did not observe any noticeable differences in the Raman spectra between rastered and unrastered regions around 666, 821, and 996 cm<sup>−1</sup>, which are associated with pure MoO<sub>3</sub>.<sup>10,31,32</sup> The 666 cm<sup>−1</sup> mode is assigned to the triply coordinated oxygen stretching mode, ν(OMo<sub>3</sub>),<sup>31,33</sup> which is specific to a full α-MoO<sub>3</sub> monolayer. This is not the case here, since even our border region consists of randomly oriented but yet separated MoO<sub>3</sub>/MoO<sub>3−x</sub> clusters. Raman peaks at 821 and 996 cm<sup>−1</sup> come from a double-coordinated oxygen stretching mode, ν(OMo<sub>2</sub>), and a terminal oxygen stretching mode, ν(O=Mo), respectively. However, a weak MoS<sub>2</sub> peak at around 820 cm<sup>−1</sup> (second order Raman mode of A<sub>1g</sub> in

**Table 1** Comparison of the MoO<sub>3−x</sub>/MoO<sub>3</sub> Raman peaks obtained in this work with other publications

Raman shift (cm <sup>−1</sup> )	Peak attribution	Raman shifts observed in the literature (cm <sup>−1</sup> )
210	A <sub>g</sub> -δ(OMo <sub>2</sub> ) in MoO <sub>3</sub> (ref. 31 and 38)	211, <sup>38</sup> 218, <sup>31</sup> and 217 <sup>39</sup>
270	B <sub>2u</sub> δ(O=Mo) in MoO <sub>3</sub> (ref. 31 and 40)	270 and <sup>31</sup> 267 (calculation) <sup>40</sup>
278	B <sub>1u</sub> -δ(O=Mo) in MoO <sub>3</sub> (ref. 31 and 40) and MoO <sub>3−x</sub> (ref. 38)	280, 282, <sup>38</sup> 270, <sup>31</sup> and 276 (calculation) <sup>40</sup>
294	B <sub>3g</sub> δ(O=Mo) in MoO <sub>3</sub> (ref. 31 and 40)	293, <sup>38,41</sup> 291, <sup>31,39</sup> and 290 (calculation) <sup>40</sup>
356–359	B <sub>3u</sub> δ(O=Mo) in MoO <sub>3</sub> (ref. 31 and 40) and MoO <sub>3−x</sub> (ref. 38)	356, <sup>38</sup> 358, <sup>31</sup> and 358 (calculation) <sup>40</sup>
461	B <sub>1g</sub> -ν(OMoO <sub>3</sub> ) in MoO <sub>3−x</sub> (ref. 38)	462, <sup>38</sup> and 461 <sup>42,43</sup>
903	β-MoO <sub>3</sub> (ref. 44 and 45) m-MoO <sub>3</sub> (ref. 31) α-MoO <sub>3</sub> within MoO <sub>3−x</sub> (ref. 46)	902 <sup>31,44–46</sup>
922–923	V(O=Mo) in α-MoO <sub>3</sub> hydrides <sup>31,47</sup> and hydrates <sup>48</sup> Mo–O <sub>2</sub> mode in MoO <sub>3−x</sub> quantum dots <sup>49</sup>	922, <sup>31,49</sup> 923, <sup>47</sup> and 920 <sup>48</sup>





MoS<sub>2</sub>)<sup>34–36</sup> and a broad silicon peak between 930 and 1000 cm<sup>−1</sup> (second order Raman scattering of Si)<sup>37</sup> effectively mask them.

Overall, Raman spectroscopy produced evidence for the surface presence of the MoO<sub>3</sub> stoichiometries in the unrastered and border regions, *i.e.*, the ones containing the AFM-detected clusters.

To shed more light in the presence of MoO<sub>3</sub> clusters within the unrastered and border zones, we performed the KPFM study. Fig. 5 shows the topography and corresponding contact potential difference (CPD) for an area around the border between rastered and unrastered regions of the MoS<sub>2</sub> flake from Fig. 4(b). Sub-nm white spots in the unrastered region are well visible – as single white dots – at the right side of the topography image in Fig. 5(a). The 5 mV (on average) difference in the CPD values between unrastered *vs.* rastered zones, see Fig. 5(c), yields the same difference in the work function (WF). The border region, where Mo oxide clusters were piled up show even higher values of the WF. Qualitatively, the same trends were observed in other studies. In particular, Zhang *et al.*<sup>50</sup> found out that a homogeneous 6 nm MoO<sub>3</sub> layer atop of a MoS<sub>2</sub> flake showed a 260 mV higher WF than pure MoS<sub>2</sub>. Rogala *et al.*<sup>27</sup> showed that a continuous 2 nm MoO<sub>3</sub>/MoO<sub>x</sub> layer atop of a MoS<sub>2</sub> flake resulted in a 70 to 100 mV higher WF than pure MoS<sub>2</sub>. Following this trend, very small WF differences are expected between the MoS<sub>2</sub> surface and isolated MoO<sub>3</sub>/MoO<sub>x</sub> clusters covering it only sparsely. However, while the thickness of the piled-up Mo oxides at the border is between 4 and 15 nm, it showed only about a 25 mV higher WF than on rastered MoS<sub>2</sub>. We expect that such a low WF difference is due to the inhomogeneity and small width of the border area. The latter produces only a small contribution to the overall KPFM signal, which is collected mostly from the MoS<sub>2</sub> surface.

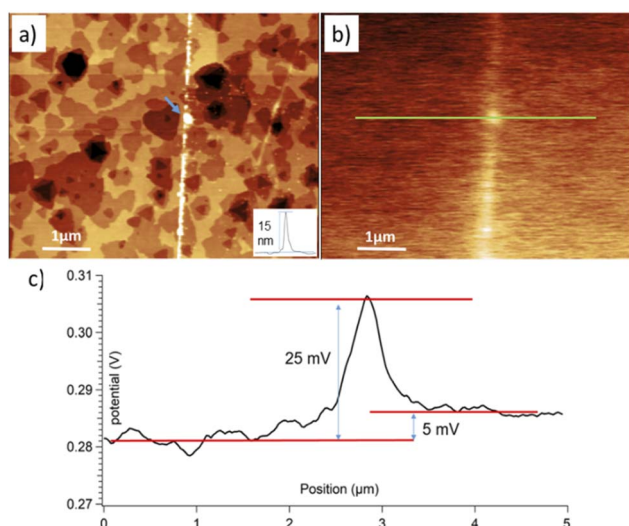


Fig. 5 (a) Topography (4.9 nm Z-scale) and (b) contact potential difference, CPD, (0.047 V Z-scale) for an area around the border between rastered (left) and unrastered (right) regions of the flake depicted in Fig. S8.† The inset in (a) shows the height profile along the border marked by a blue arrow. (c) Cross-section taken along a green line in (b). Notably different CPD values are noticed in each of the regions, *i.e.*, rastered, border, and unrastered.

In conclusion, while the KPFM did not produce a direct chemical identification of the scanned species, we observed differences between MoO<sub>3</sub> rich and lacking zones, which agreed with some previously published results.

To elucidate coordination of the oxidation products in more detail, an untreated (as-exfoliated) and thermally etched/oxidized flakes were chosen for XANES measurements. Fig. 6 shows the comparative XANES spectra. Mo M<sub>2</sub> and M<sub>3</sub>-edges in an unmodified MoS<sub>2</sub> flake produced two peaks at 413.5 eV and 396.4 eV, respectively. In comparison to the untreated sample, the XANES spectra of an etched/oxidized flake shows a 0.5 eV blue shift of the Mo M<sub>2</sub> edge and a new M<sub>3</sub> post peak at 399.8 eV. These two important features of the Mo M<sub>2</sub> and M<sub>3</sub> edges have been attributed to the presence of MoO<sub>3</sub> by Rogala *et al.*<sup>27</sup> Moreover, there is a strong peak at 531.5 eV in the XANES spectra of a modified flake which is absent in the case of the untreated sample. Lajaunie *et al.*<sup>51</sup> studied the XANES spectra of the oxygen K-edges in MoO<sub>2</sub> and MoO<sub>3</sub>. They found out that oxygen in MoO<sub>3</sub> produced a strong peak at 531.2 eV, while the strongest peak of the O K-edge in MoO<sub>2</sub> was at 534.5 eV. Based on these data, the peak at 531.5 eV in our XANES spectra of the modified flake shows that the oxidation products are MoO<sub>3</sub> rather than MoO<sub>2</sub>.

Finally, additional SEM-EDS and SEM-WDS studies presented in the ESI† show that the presence of isolated and

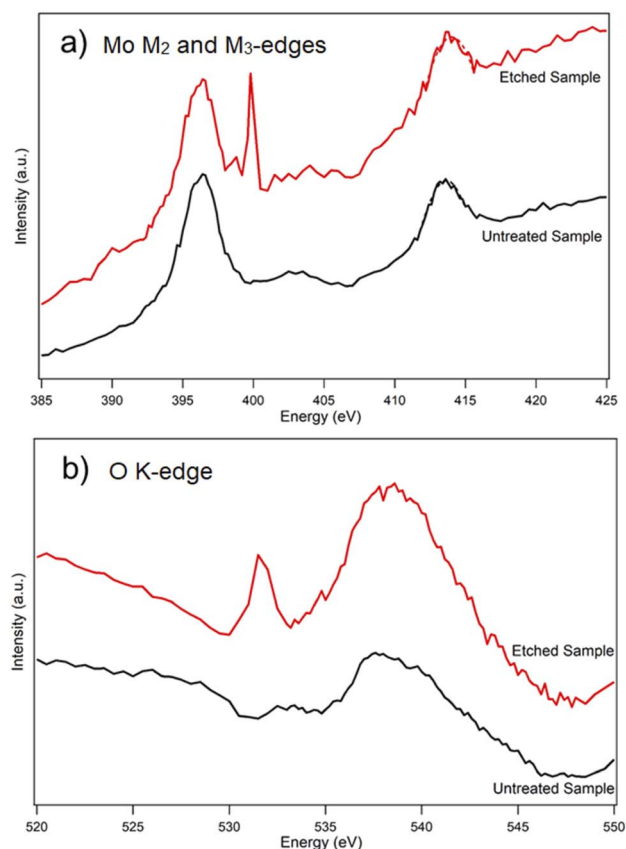


Fig. 6 XANES spectra of (a) an untreated MoS<sub>2</sub> flake and (b) a modified MoS<sub>2</sub> flake heated at 370 °C in air.



sparsely present sub-nm MoO<sub>x</sub> clusters produced a detectable increase in the O/Mo ratio on investigated MoS<sub>2</sub> flakes.

## Conclusions

Thermal etching of a single MoS<sub>2</sub> flake at 370 °C produced not only etch pits, but also some sub-nm oxide clusters. The high resolution AFM data together with DFT simulations show that majority of the clusters were MoO<sub>3</sub> monomers and dimers caught in the act of being produced at the single sulfur vacancies. Raman spectroscopy confirmed the surface presence of the MoO<sub>3</sub> stoichiometry. The KPFM and SEM-EDS/WDS data strengthened this hypothesis, although were not able to directly differentiate the MoO<sub>3</sub> clusters from an underlying MoS<sub>2</sub> surface. The XANES results contributed to confirming the MoO<sub>3</sub> stoichiometries present on the MoS<sub>2</sub> surface. Our findings confirm that during MoS<sub>2</sub> thermal oxidative etching, Mo atoms leave the surface mainly as MoO<sub>3</sub> molecules rather than any other forms of Mo oxides. This confirms the stoichiometry of the thermal etching process exemplified in eqn (1). Moreover, sulfur vacancies were suggested by our DFT results to act as nucleation sites for the thermally induced oxidative etching.

Beyond fundamental research, these findings are expected to find applications in nanoscale catalysis, since defected MoS<sub>2</sub> as well as MoO<sub>3</sub> clusters are potent catalysts for numerous important reactions including the hydrogen evolution reaction and oxygen reduction reaction.

## Conflicts of interest

There are no conflicts to declare.

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