



KCl acts as a flux to assist the growth of sub-millimeter-scale metallic 2D non-layered molybdenum dioxide

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Abstract Two-dimensional (2D) metal oxides (2DMOs), such as MoO₂, have made impressive strides in recent years, and their applicability in a number of fields such as electronic devices, optoelectronic devices and lasers has been demonstrated. However, 2DMOs present challenges in their synthesis using conventional methods due to their non-van der Waals nature. We report that KCl acts as a flux to prepare large-area 2DMOs with sub-millimeter scale. We systematically investigate the effects of temperature, homogeneous time and cooling rate on the products in the

flux method, demonstrating that in this reaction a saturated homogenous solution is obtained upon the melting of the salt and precursor. Afterward, the cooling rate was adjusted to regulate the thickness of the target crystals, leading to the precipitation of 2D non-layered material from the supersaturated solution; by applying this method, the highly crystalline non-layered 2D MoO₂ flakes with so far the largest lateral size of up to sub-millimeter scale (~ 464 μm) were yielded. Electrical studies have revealed that the 2D MoO₂ features metallic properties, with an excellent sheet resistance as low as 99 Ω·square⁻¹ at room temperature, and exhibits a property of charge density wave in the measurement of resistivity as a function of temperature.

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1 Introduction

Two-dimensional (2D) materials of atomic scale exhibit tremendous potential for future semiconductor device applications due to their captivating physical attributes and outstanding optoelectronic capabilities. Based on their microstructure, 2D materials can roughly be divided into two categories: 2D layered and 2D non-layered materials [1–3]. 2D non-layered materials are formed by chemical bonds in three-dimensional (3D) directions, leading to unsaturated dangling bonds on the surface and inducing highly active and energetic surfaces, making them have broad prospects as components of efficient and practical devices [4, 5]. The major categories of 2D non-layered materials include metal dichalcogenides, metals, metal oxides, III-V semiconductors, organic–inorganic perovskites and others [6]. 2D metal oxides (2DMOs) have attracted significant interest in the realms of electronics, electret technology, optoelectronics, lasers, and related fields, primarily due to their unique electrical architectures and diverse surface chemistry [7–13]. MoO₂, exemplifying the characteristics of 2DMOs, has showcased its utility in magnetism, piezoelectricity, and sensor technologies. For example, Zhang et al. [14] reported that the 2D MoO₂ flakes exhibited a large linear magnetoresistance of up to 455% at 3 K and –9 T and a nonlinear hall effect. Moreover, 2D MoO₂ reveals an unexpected piezoelectric-like response. As reported by Ajayan et al., the 2D MoO₂ exhibited an out-of-plane piezoresponse of 0.56 pm·V⁻¹, which is comparable to that of standard 2D piezoelectric materials [15]. In addition to its superior properties resulting from its intrinsic structures, 2D MoO₂ is also

promising to be integrated into heterojunctions to perform some advanced functions. For example, Zou et al. revealed that 2D MoO₂ which is vertically stacked with MoSe₂ as 2D heterostructures exhibited a photoresponsivity of 100.86 mA·W⁻¹ and a detectivity of 23.4 × 10⁹ Jones [16]. However, owing to the inherent non-van der Waals structure of 2D MoO₂, there has been a lack of effective strategies for the precise cultivation of extensive single crystals, posing challenges to conventional top-down synthesis methods [12]. The lateral size of 2D MoO₂ synthesized via bottom-up methodologies like chemical vapor deposition (CVD) is typically constrained to 10 μm, thereby hindering its subsequent exploration and practical utilization [9, 16–22]. It is essential to develop a versatile growth method for large-area 2D MoO₂ single crystals.

The flux method is considered to be a promising and economical method for the formation of single crystals [23]. The flux method has the benefit of producing crystals with excellent crystallinity and form at a reduced crystallization temperature. Compared with crystals grown from their melt, these crystals typically have fewer defects and are under less strain [24, 25]. In flux growth, the flux is generally a molten salt or oxide that serves as a solvent to dissolve the components of the desired substance [26]. During the reaction, the flux and the precursor are heated to the molten state and the temperature is maintained above the saturation temperature of the solution, after which the target single crystal will precipitate from the supersaturation solution during the cooling down process [27]. The flux method was recorded against various single crystals, from rare earth aluminum borate crystals [28] and insulators [29], to oxide materials [30], etc., but the resultant growth products are typically bulk single crystals, typically millimeters in size on all three dimensions.

While historically, the flux method has been confined to fabricating bulk single crystals utilizing specialized equipment, and there is a lack of explicit reports on its utilization in CVD for synthesizing 2D materials, we do believe it is a feasible approach. The current growth mechanism for 2D materials can be summed up in one of two ways [31–34]: either through a surface-catalyzed process, in which the substrate is essentially not involved in the reaction; or through a precipitation process, in which the target product's constituents are dissolved in the substrate and precipitate out upon cooling. The main distinctions between the two are: (a) The solubility of the target component in the substrate or precursor at the reaction temperature. If the solubility is high, the growth is predominantly precipitation-based. (b) The effect of cooling rate on the thickness of the target product. The cooling rate has a significant impact on thickness in the precipitation process but has little effect on the catalysis process. According to the dissolution-based reaction mechanism of

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the flux method and its product size can be controlled by the cooling rates [35], we believe that flux growth belongs to the precipitation method.

Considering the ability to lower the melting point of the reactants and the use of salt in the precursor, similarities can be found between the flux method and the “molten-salt-assisted method” (MSAM) frequently reported in the literature for the preparation of large-area 2D materials. Here, we endeavor to illustrate that the MSAM fundamentally aligns with the flux approach rather than representing a catalytic methodology, as the salt typically dissolves the precursor and engages in a reactive process with it. In fact, there are some MSAM reports demonstrating that their target low-dimensional single crystals are precipitated from supersaturated droplets [36–38], in agreement with the flux method we claim. Although the MSAM has been widely employed for the synthesis of various 2D materials, particularly transition-metal chalcogenides [39], there is a notable scarcity of reports pertaining to non-layered 2D materials. This may be ascribed, at least in part, to the lack of understanding of the detailed mechanism, such as how the MSAM regulates the thickness of 2D materials. Indeed, the capability of the flux method to modulate thickness through manipulation of the temperature span and cooling rate underscores its promise in synthesizing non-layered 2D materials such as 2DMOs.

Here, we apply the flux growth strategy in CVD to realize the large-scale preparation of 2D non-layered MoO₂, elucidating that the essence of reducing the thickness and enlarging the area of the products lies in a supersaturated homogeneous solution and an appropriate cooling rate. In our flux method, the 2D non-layered MoO₂ single crystals precipitated out of the supersaturated solution formed by the precursors and the salt. Through meticulous control of the reaction temperature, homogeneous time and cooling rate, we successfully achieved the largest known non-layered 2DMOs crystals to date [7, 9, 16–21, 40], with thickness effectively regulated by the temperature range and cooling rate. Electrical studies revealed that the obtained 2D MoO₂ crystals featured a metallic nature, with a sheet resistance as low as 99 Ω-square⁻¹ at room temperature. Moreover, we observed the occurrence of the charge density wave phenomenon at low temperatures. We thus clarify that the prevailing MSAM is in essence a flux method. Armed with this pivotal comprehension, the experimental conditions can be deliberately optimized toward the flux method. This approach offers an expeditious avenue for the fabrication of large-scale 2D MoO₂ and harbors potential for expansion to additional 2DMOs or layered materials, thereby enhancing their attributes and accelerating their utilization across various domains.

2 Experimental

2.1 Synthesis of MoO₂ flakes

The flux-grown MoO₂ flakes were grown via CVD, 20 mg MoO₃ (Shanghai Aladdin Bio-Chem Technology Co., LTD, 99.9%), and 5 mg KCl powder was positioned in an alumina crucible placed in the middle of a 1-inch-diameter quartz tube furnace. The SiO₂/Si substrate (cleaned with acetone, alcohol and deionized water) was set on the same crucible and kept directly above the mixture powder. During the first ten min, the quartz tube was cleaned with 300 ml·min⁻¹ of highly pure Ar carrier gas. To eliminate pollutants (moisture or organic) that may have been present in the system, the furnace was kept at 150 °C for 10 min. Following this, the temperature is raised to 900 °C in fifty minutes with 30 ml·min⁻¹ of Ar flow. The temperature is maintained for 30 min and then cooled to room temperature. The flux-free MoO₂ flakes were grown via CVD, too. The experimental conditions were the same as for the flux method, except that no flux (KCl) was added.

2.2 Characterization of MoO₂ flakes

Atomic force microscope (AFM, NT-MDTNEXT), optical microscope (OM, OLYMPUS BX51M), Raman spectroscopy (Invia Reflex with an excitation wavelength of 532 nm laser), scanning electron microscope (SEM, Helios G4 CX), X-ray diffraction (XRD, Rigaku Ultima IV), thermogravimetric study (TG, STA449C/6/G), X-ray photoelectron spectroscopy (XPS, Thermo Scientific K-Alpha), transmission electron microscopy (TEM, JEM2800 and Talos F200S), and aberration-corrected scanning transmission electron microscopy (AC-STEM, Themis Z) were used. For TEM sample, the flakes were transferred to the Cu grid via the polymethyl methacrylate (PMMA)-assisted transfer method. Spin-coating PMMA (950 A4) onto MoO₂ of SiO₂/Si (1500 r·min⁻¹, 1 min, twice) was followed by annealing at 150 °C for 5 min. A solid piece of polydimethylsiloxane (PDMS) then adhered to MoO₂ covered by the PMMA and cured at 80 °C for 1 min. After a day of immersion in deionized water (DI water) which caused the spontaneous penetration of the water into the interface between PDMS/PMMA/MoO₂ and SiO₂/Si, the PDMS/PMMA/MoO₂ was then carefully delaminated from SiO₂/Si with the assistance of tweezers. The acquired film was then fished by a Cu grid. Lastly, the PDMS was separated from the PMMA after heating at 150 °C for 5 min, and then the PMMA was removed after being immersed in acetone for 20 min. Before further processing, the transferred MoO₂ was washed two times with DI water and ethanol and, finally, dried in N₂ gas.

Schematic illustrations of the crystal structures were drawn using the program VESTA [41].

2.3 Device fabrication and measurements

The 150 nm-thick gold electrodes were transferred onto the MoO₂ flakes by probes to construct the two-terminal devices. Using a probe station accompanied by a semiconductor property analyzer, the electrical performance of the MoO₂ devices was measured in ambient conditions with a standard setting (Model: 4200, Keithley, USA). For the temperature-dependent measurement of resistivity, the MoO₂ devices were firstly bonded in open air followed by assembling on a cryostat with a 1-Pa low-pressure chamber. The temperature of the cryostat can be adjusted between 77 and 267 K.

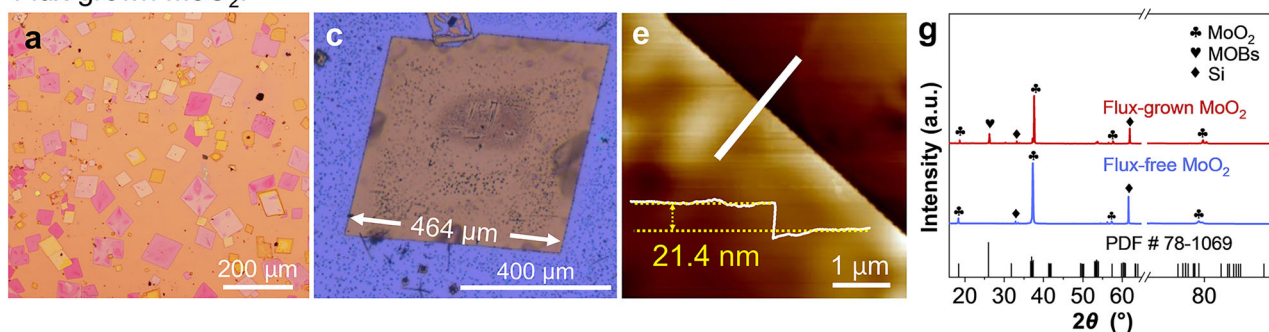
3 Results and discussion

3.1 Characterizations of MoO₂

MoO₂ prepared with and without the flux method is denoted as flux-grown and flux-free MoO₂. The diagram of two CVD installations is shown in Fig. S1a, b. Figure 1 illustrates the differences between the two kinds of MoO₂. In sharp contrast to the dense nanometer-level cubes of the

flux-free MoO₂ (OM image of Figs. 1b, S1c, d), the lateral length of the flux-grown MoO₂ is tens or even hundreds of microns (Fig. 1a). As shown in OM image in Fig. 1c, the lateral size of the flux-grown sample is as large as 464 μm , which to our knowledge is largest in size of non-layered 2DMO single crystals ever prepared by CVD, and much larger than the numerous studied WO₂ and K-MnO₂, and so on (Table S1) [7]. The typical lateral size of flux-free MoO₂ is only 257 nm as shown in SEM image in Fig. 1d. It demonstrates that the flux method has the powerful ability to obtain large-area materials in comparison to the conventional CVD method. In addition, it also shows the ability to decrease the dimension of the products from 3 to 2D. As shown in Fig. 1e, f, AFM images show single-crystal flakes with thicknesses of 21.4 nm for flux-grown MoO₂ and 192.5 nm for flux-free MoO₂ (Fig. S2). To illustrate the differences between our flux-free MoO₂ samples and the flux-grown samples more intuitively, we tallied the lateral size and thickness of the products as shown in Fig. S3. The size of the flux-grown MoO₂ is in the range of 20–460 μm with an average lateral size of around 95.7 μm (Fig. S3a), and the thickness distribution reveals the average thickness of flux-grown MoO₂ is 15.8 nm with a range of 5–30 nm (Fig. S3b), and the thinnest 2D MoO₂ we obtained is 6.5 nm, as shown in Fig. S3c. However, for the flux-free MoO₂, the average lateral size is only about 0.32 μm in the range of 0–1.2 μm (Fig. S3d), while for the

Flux-grown MoO₂:



Flux-free MoO₂:

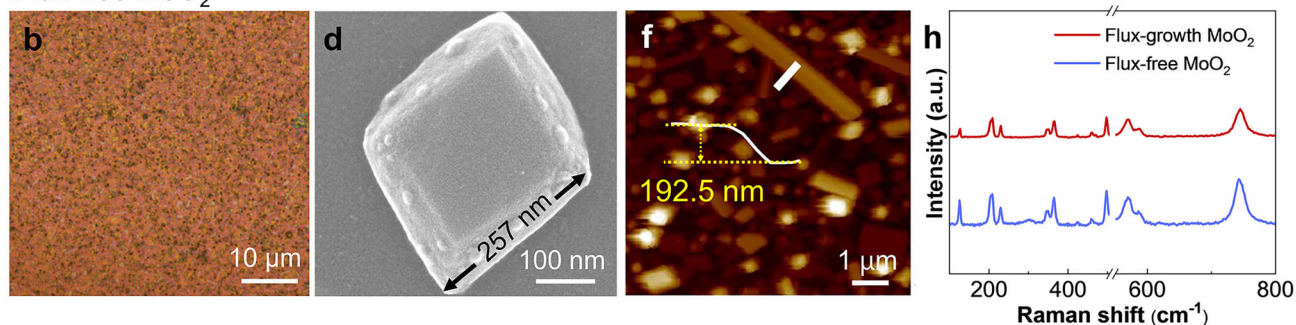


Fig. 1 Characterizations of MoO₂ flakes with and without flux method: typical OM images of **a** flux-grown MoO₂ flakes and **b** flux-free MoO₂ flakes; typical SEM images of **c** flux-grown MoO₂ flake and **d** flux-free MoO₂ flake; AFM images of **e** single flux-grown MoO₂ flake and **f** flux-free MoO₂ flake; **g** XRD patterns and **h** Raman spectra of flux-grown and flux-free MoO₂ flakes

flux-free MoO₂ an average thickness of ~ 110.5 nm is in the range of 0–300 nm (Fig. S3). XRD result in Fig. 1g demonstrates that the flux-grown MoO₂ and the flux-free MoO₂ flakes have identical crystal structures, which match well with the MoO₂ monoclinic symmetry crystal structure. Thus, the flux method is promising to produce high-quality 2D non-layered single crystals. In Raman spectra (Fig. 1h), we observed the same MoO₂ peaks at 125, 206, 229, 345, 364, 460, 497, 569, 587 and 744 cm⁻¹ in both two samples [42]. The bands at 587 and 744 cm⁻¹ are characteristics of MoO₂ and can be attributed to the Mo–O bond stretching

vibrations [43]. From the comparison, it is found that the use of the flux method facilitates the successful fabrication of thinner and larger MoO₂ flakes which is more suitable for device applications.

Following the initial assessment of the quality of the samples, we proceeded with the characterization of its atomic structure to gain a comprehensive understanding of its internal arrangement, aiming to provide insights into the underlying growth mechanisms. The crystalline and surface structure of the flux-grown MoO₂ flakes was studied using AC-STEM. Figure 2a shows a typical rhombic MoO₂

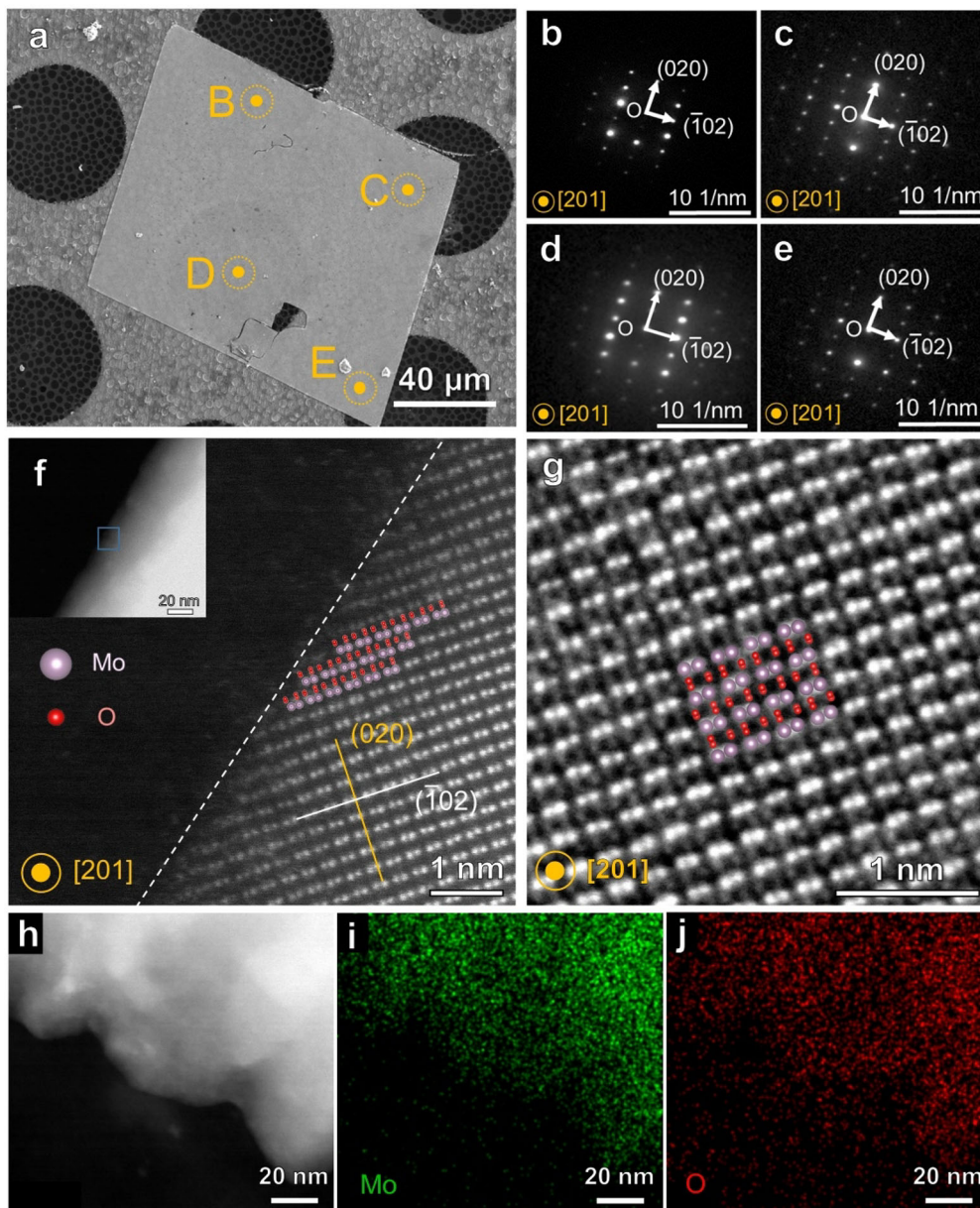


Fig. 2 STEM images of transferred flux-grown MoO₂ flakes: **a** SEM image of transferred MoO₂ domain on a Cu grid; **b–e** corresponding SAED patterns extracted from different regions as outlined by yellow dots B, C, D and E in **a**; **f, g** AC-IDPC and HAADF images of MoO₂ flake viewed along [201] zone axis and (inset) low-magnification HAADF image; **h** low-magnified HAADF image on edge of flux-grown MoO₂ flake and **i, j** corresponding EDS maps of Mo and O

domain with an edge length of 114 μm . Selected area electron diffraction (SAED) patterns (Fig. 2b–e) extracted from its different areas (yellow dots in Fig. 2a) exhibit well-resolved 2D lattices, where the spacings of 0.28 and 0.24 nm correspond to the (102) and (020) planes of MoO₃, respectively. The integrated differential phase contrast (iDPC) technique, which offers the advantages of simultaneous imaging of heavy and light atoms [44], was employed to image Mo and O columns. As seen in Fig. 2f, g, the iDPC combined with the high-angle annular dark-field STEM (HAADF-STEM) clearly resolves the periodic arrangement of Mo and O columns, without any discernible vacancy (Fig. S4). Finally, EDS maps confirm that Mo and O elements are homogeneously distributed within the MoO₂ flake, as shown in Fig. 2h–j.

The dominantly exposed surface of these highly crystalline flakes, also defined as their basal habit planes, is assigned to the (100) plane, after considering the crystallography of MoO₂ (see the shaded planes from two orthogonal directions in Fig. 3a, b). The surface energies of 15 possible reconstructed (001), (010) and (100) surfaces of MoO₂ crystal as a function of oxygen chemical potential are tested by density functional theory (DFT) calculations

in Fig. 3c. It is evident that the MoO₂ (100)-1 surface terminated by excess O atoms (Fig. 3c) is the most stable one over a large oxygen chemical potential range, from O-poor limit to O-rich environment (see Calculation Methods in Supplementary Information). This agrees well with the fact that MoO₂ platelets prefer to expose (100) basal facets; therefore, we depicted the schematic illustration of MoO₂ crystal growth in Fig. 3d, vividly depicting the dynamic formation process of MoO₂ crystal with (100) facets as preferential surface.

3.2 Mechanism of flux-grown 2D MoO₂

What makes the flux method successfully obtain the 2D MoO₂? Here, we propose the reaction mechanism. First, the reaction temperature needs to be high enough to completely dissolve all the constituent parts and form a saturated solution, as only then can the crystals precipitate out from the solution as the supersaturation point is reached when the temperature is lowered [24]. Second, it is essential to keep the reaction temperature a sufficient time until a homogeneous solution is obtained [26]. The appropriate cooling rate is the third consideration. The flux

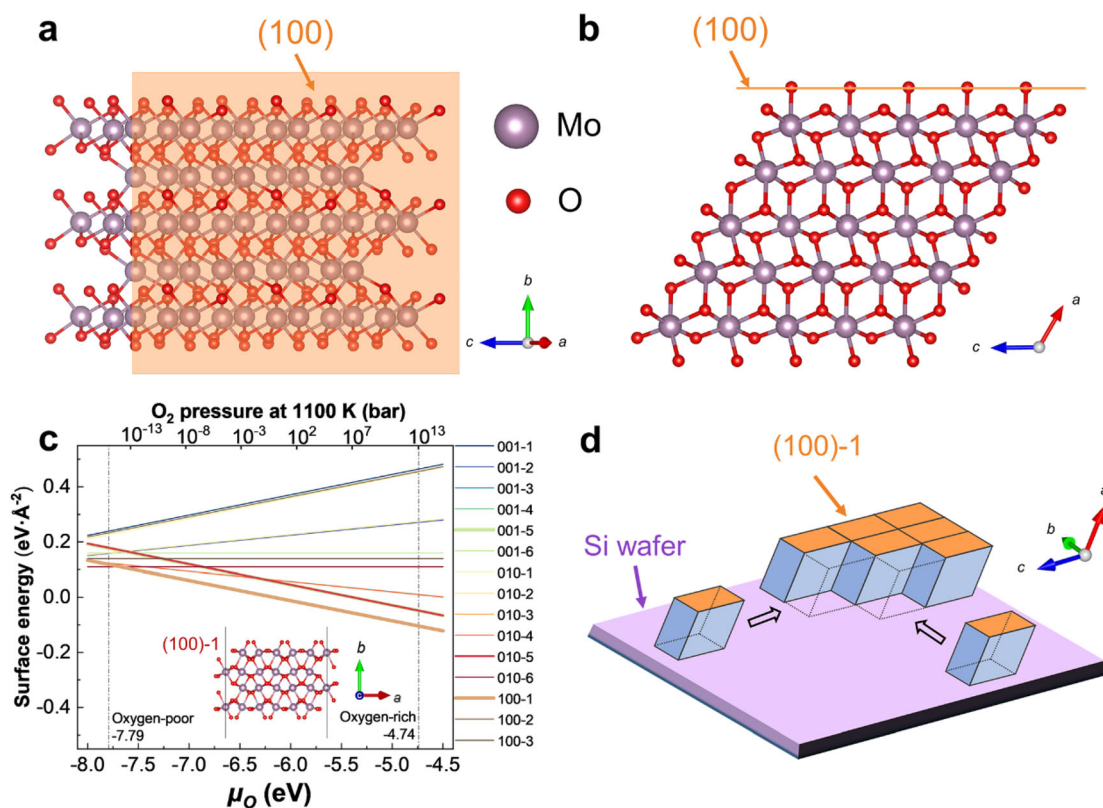


Fig. 3 Atomic structure of flux-grown MoO₂ flakes: **a** crystal structure of MoO₂ projected along [201] direction, with the electron beam oriented perpendicular to surfaces of platelet, as illustrated by the shaded plane; **b** shade plane rotated 90° around beam direction and (100) plane projected as a line in right panel; **c** surface energies of 15 possible reconstructed (001), (010) and (100) surfaces of MoO₂ crystal as a function of oxygen chemical potential and (inset) illustration of MoO₂ (100)-1 surface; **d** schematic illustration of MoO₂ crystal growth

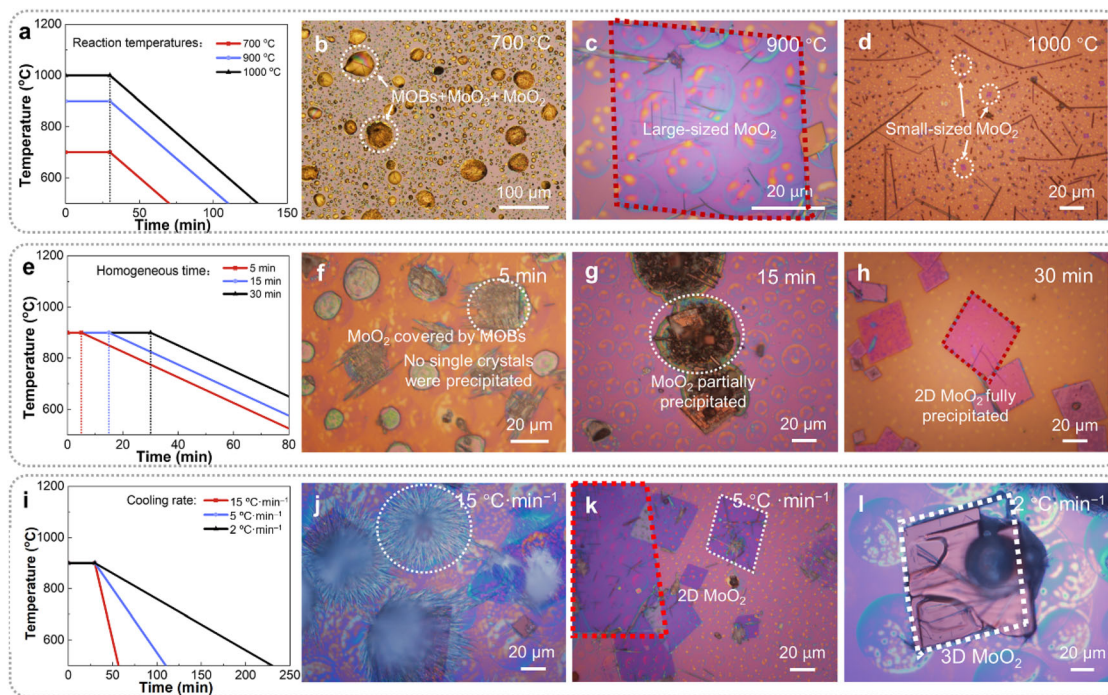


Fig. 4 Influence of key parameters on preparation of 2D MoO₂ by flux method: **a** time–temperature curves of experiments investigating effect of reaction temperature on products; typical OM images of samples maintained at **b** 700, **c** 900 and **d** 1000 °C for 30 min before cooling to room temperature at 5 °C·min⁻¹; **e** time–temperature curves of experiments investigating effect of homogeneous time on products; typical OM images of samples held at 900 °C for **f** 5, **g** 15 and **h** 30 min and then cooled to room temperature at 5 °C·min⁻¹; **i** time–temperature curves of experiments investigating effect of cooling rate on products; typical OM images of samples after holding at 900 °C for 30 min and then cooling to room temperature at **j** 15, **k** 5 and **l** 2 °C·min⁻¹

method is generally applicable to bulk single crystals. However, the thickness of the target crystal and the cooling rate are highly related, and the key to growing 2D crystals is to control their cooling rate [45]. The pivotal factors influencing the reaction encompass temperature, homogeneous time and cooling rate. We selected the readily available KCl to dissolve the precursor, MoO₃ [16, 17, 20, 46, 47].

We first conducted experiments to illustrate the effect of temperature. Thermogravimetric analysis suggests that KCl and MoO₃ start to react at 510 °C (Fig. S5). The temperature range in our experiment was thus set from 700 to 1000 °C. Results were recorded at each temperature (maintained for 30 min before cooling to room temperature at 5 °C·min⁻¹). The corresponding time–temperature curve is shown in Fig. 4a. The typical OM image for the experiment is depicted in Fig. 4b–d, which contains distributed circular particles when cooled from a starting temperature of 700 °C (Fig. 4b), indicating a condensation process. The particles are composed of potassium molybdenum oxide bronzes (MOBs), i.e., K₂Mo₃O₁₀, K₂Mo₂O₇, K₂MoO₄ and MoO₃ based on SEM, Raman and energy dispersive X-ray spectrometry (EDS) analyses in Fig. S6a–c [48–51]. Although no clear MoO₂ grains can be found and

confirmed, and neither does MoO₂ exhibit a 2D morphology, several Raman peaks of MoO₂ are present at 126, 229 and 367 cm⁻¹ [52]. We speculate that the Raman signal of MoO₂ comes from its spontaneous nucleation during the cooling process [26, 53]. A control experiment without salt reveals that there are only a few MoO₃ sheets on the substrate (Fig. S6d–f). Next, it was discovered that 2D MoO₂ flakes were obtained at 900 °C as shown in Figs. 4c, S7. Moreover, different from the typical growth substrate of 2D materials, many bubbles are exhibited on the surface beneath every MoO₂ flake (Fig. S8). These are attributed to the reaction of KCl and MoO₃ to produce a mixed solution containing MOBs. Most MOBs melted at 900 °C, (K₂Mo₂O₇ melted at ~ 560 °C, K₂MoO₄ at ~ 900 °C and MoO₃ at ~ 795 °C) [36, 48]. For the 1000 °C-sample, only a few MoO₂ flakes with a size of just a few microns could be found on the surface, as displayed in Fig. 4d. Consequently, we can conclude that 900 °C is the most suitable temperature for preparing the large-area 2D MoO₂ flakes in our flux growth method.

After determining the holding temperature of 2D MoO₂, further investigation of the homogenization process of 2D MoO₂ prepared by flux growth is essential to understand the whole growth process. Because flux growth

is achieved by dissolving the precursor with flux to form a saturated solution, after which the target crystals precipitate out of the supersaturated solution during the cooling process, the saturated solution is prepared by holding the composition and flux of the desired crystals at a temperature slightly above the saturation temperature for long enough to form a complete solution, a process known as homogenization [45].

We designed a series of experiments that only altered the holding times at 900 °C to explore the effect of the homogenization process on the resulting products. The corresponding time–temperature curve is shown in Fig. 4e. We present OM images of the products prepared under different conditions, where the reaction times were kept at 900 °C for 5, 15, 30 and 60 min, respectively, followed by cooling to room temperature at 5 °C·min⁻¹. After holding at 900 °C for 5 min, the corresponding Raman analysis was conducted as shown in Fig. S9a. The result shows that the particles are made up of K₂Mo₃O₁₀, K₂Mo₂O₇, K₂MoO₄, MoO₃ and MoO₂, too. As can be surmised, with a holding time of only 5 min, the saturated solution of MoO₂ may not have formed yet and the characteristic peak position of MoO₂ still did not appear, so no single crystals were precipitated (Fig. 4f). After holding for 15 min and dropping to room temperature, as shown in Fig. 4g, some of the MoO₂ had separated from the round particles, indicating that the homogenization process had started, and MoO₂ partially precipitated from the supersaturated solution. Then, after homogenization at this temperature for 30 min, MoO₂ single crystals were formed in Fig. 4h, demonstrating that holding at 900 °C for 30 min permits the homogenization process to be finished and complete 2D MoO₂ single crystals to precipitate. Ultimately, as the homogenization time extends to 60 min, conspicuous circular formations emerge on the substrate, accompanied by a central darkening (Fig. S9b). Notably, the substrate manifests a discernible degree of impairment, devoid of discernible 2D crystals. This occurrence may be attributed to the protracted presence of a corrosive solution containing MOB [54], stemming from an overly prolonged homogenization interval, causing substrate degradation. This deduction underscores the optimal homogenization period of 30 min under our experimental regimen.

It is crucial to continue looking at the cooling rate after obtaining the saturated homogenous solution of 2D MoO₂. The cooling rate affects the thickness of crystals from flux growth [45]. Therefore, the key to expanding the flux method, which was previously limited to preparing bulk single crystals, to 2D non-layered materials is to control its cooling rate. To illustrate the effect of cooling rate on samples, we designed a series of experiments, varying only the cooling rate of the 900 °C-grown samples. The corresponding time–temperature curve is shown in Fig. 4i.

OM images of thus-prepared products precipitated by different cooling rates of 15, 5 and 2 °C·min⁻¹ are illustrated in Fig. 4j–l. As shown in Fig. 4j, if the cooling rate is as rapid as 15 °C·min⁻¹, MoO₂ is obtained with a large number of MOB atop. However, the quality of the MoO₂ crystals will be compromised by the rapid cooling rate [24]. When the rate is 5 °C·min⁻¹, well-shaped 2D MoO₂ crystals are produced, with only a few MOB present (Fig. 4k). The slow cooling rate of 2 °C·min⁻¹ yields 3D MoO₂, as shown in Fig. 4l. Therefore, in the flux method, non-layered 2D MoO₂ with good crystal quality can only be obtained at a suitable cooling rate of around 5 °C·min⁻¹.

This precipitation process has been demonstrated by the presence of particulate residues on the surface of the 2D MoO₂ flakes, which were examined by SEM, Raman and EDS. As can be seen in Fig. 5a, several hexagonal-shaped particles are neatly aligned, and EDS and Raman results assist in confirming the presence of K₂Mo₂O₇, K₂Mo₃O₁₀ and K₂MoO₄ on the surface of the MoO₂ as illustrated in Fig. 5b, c. Moreover, SEM and EDS were randomly conducted around MoO₂ flakes as shown in Fig. S9c–f, it can be distinctly observed that there are particles underneath each MoO₂ flake which are confirmed by EDS result to be most likely potassium MOB. It's worth noting that these residues are all soluble in water and can be removed by using deionized water, as shown in Fig. S10. Based on these findings, it can be assumed that after the homogenization process is complete, the majority of the MOB and MoO₃ are converted to MoO₂, allowing the MoO₂ to precipitate out during the cooling process, with a small amount of residual MOB being present around the MoO₂ single crystal. Next, the products were collected and analyzed by XPS. Three groups of the fitted curve of Mo 3d (3d_{5/2} and 3d_{3/2}) are shown in Fig. 5d. The curves at 229.2 and 232.2, 230.9 and 234.2 eV are assigned to the oxidation state of Mo⁴⁺, which is expected from the stoichiometry of MoO₂ and MoO_xCl_y, while the third group is 232.9 and 235.9 eV, corresponding to Mo⁶⁺, which together with the signal from K 2p can be explained by the presence of MOB (Fig. 5e). The signals from Cl–Mo bonds in Cl 2p confirm the existence of the MoO_xCl_y (Fig. 5f) [39, 55, 56]. Thus, the existence of MOB and MoO_xCl_y in the preparation process can be confirmed from the experimental results.

Based on the results, a growth model for the flux-grown MoO₂ can be proposed as shown in Fig. 5g. MoO₃ and KCl start to form MOB at 510 °C. A saturated homogeneous solution (mixed by the MOB and MoO₃) is formed after holding at 900 °C for thirty minutes, after which the highly crystalline 2D MoO₂ precipitates from the supersaturated solution at the cooling rate at ~ 5 °C·min⁻¹. Without the addition of flux, MoO₃

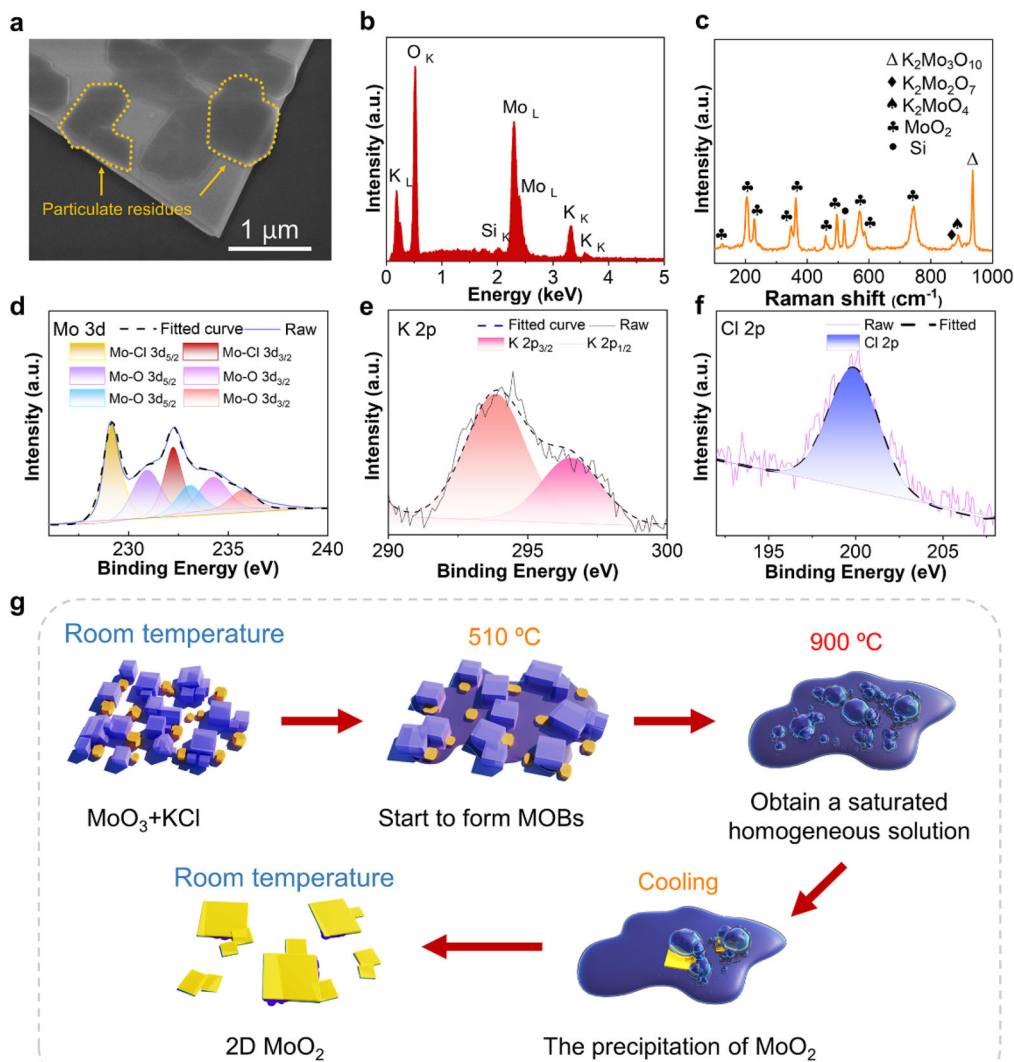
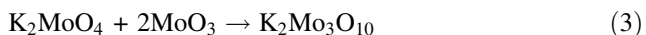
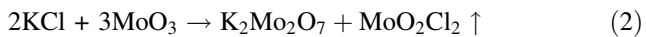
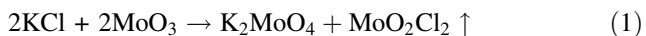


Fig. 5 Mechanism of 2D MoO₂ obtained by flux method: **a** typical enlarged SEM images of flux-grown MoO₂ after holding at 900 °C for 30 min and cooled to room temperature at 5 °C·min⁻¹; corresponding **b** EDS and **c** Raman spectra; XPS spectra of **d** Mo 3d, **e** K 2p and **f** Cl 2p of sample; **g** schematic illustration of 2D MoO₂ obtained by flux method

powder yields MoO₃ flakes at ~ 700 °C, followed by 3D MoO₂ formation at ~ 900 °C.

The possible reaction routes we propose are as follows [39, 57–59]:



Finally, the 2D MoO₂ is precipitated from a supersaturated solution, which contains K₂Mo₂O₇, K₂Mo₃O₁₀ and K₂MoO₄ and MoO₃.

3.3 Electrical properties of 2D MoO₂

To evaluate the electrical performance of the materials, we calculated the density of states (DOS) of the bulk and 2D

MoO₂ films with O-terminated (100) surfaces. It is found that the MoO₂ crystal is metallic and the DOS has a low valley at the Fermi level (Fig. 6a, d), which is similar to the previous calculations [56, 60]. The DOS of two layers and four layers O-terminated MoO₂ (100) surfaces is very close to that of MoO₂ crystal, both being metallic with DOSs near the Fermi level (Fig. 6b, c, e, f). Figure 6e, f exhibits relatively large peaks in DOS at E_F , suggesting electronic instability, which may be beneficial for electron transport. The experimental results show that there is no shift in the in-plane current (I)-voltage (V) dependence when the gate voltage is modulated from -40 to 40 V as shown in Figure S11. Figure 6g depicts the I - V dependence of the MoO₂ device. The insets are the schematic illustration and OM image of a typical device made in this work. We measured the sheet resistance of several devices and plotted them as histograms as shown in Fig. 6h, where the

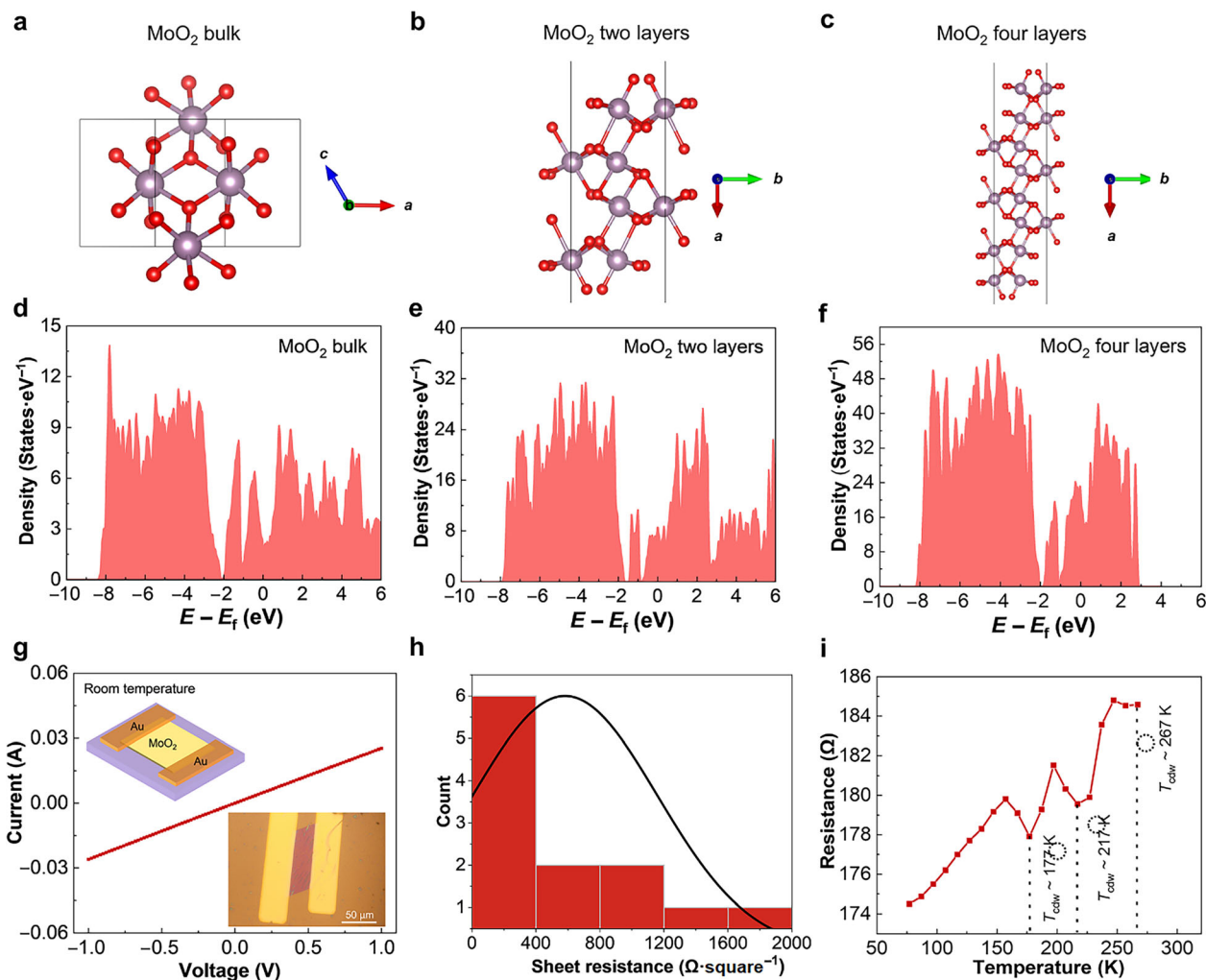


Fig. 6 Electrical properties of 2D MoO₂: structure of **a** MoO₂ bulk crystal, **b** two layers and **c** four layers of O-terminated MoO₂ (100) flakes; DOSs of **d** bulk MoO₂ crystal, **e** two layers thick MoO₂ flake and **f** four layers thick MoO₂ flake with O-terminated (100) surfaces; **g** *I*-*V* dependence for MoO₂ device at room temperature and (insets) schematic illustration and OM image of flux-grown MoO₂ device; **h** histogram of measured sheet resistance distribution of flux-grown MoO₂ devices; **i** resistance vs. temperature of flux-grown MoO₂ devices

value could be as low as $99 \Omega\text{-square}^{-1}$, indicating the good conductivity of 2D MoO₂, which may be ascribed to the high crystallinity that turns into less scattering. To further determine the temperature-dependent electrical properties, the resistivity was measured as a function of temperature. In Fig. 6i, kinks are observed at ~ 177 , 217 and 267 K. The exact origin of the kinks is still under investigation. However, it is reasonable to attribute them to the metal–semiconductor transition caused by the so-called charge density wave (CDW) instabilities. CDW is a periodic modulation of electron density, which is accompanied by periodic lattice distortions and is mostly observed in low-dimensional materials [61–65]. In fact, as early as the 1960s, the behavior has been demonstrated experimentally in a variety of quasi-low-dimensional compounds, such as MOBs [61]. For example, the blue bronzes (K_{0.3}MoO₃),

which is one of the MOBs, are metallic at ambient temperatures but exhibit a metal-to-semiconductor transition at $T_P = 180$ K [66]. The layered purple bronze K_{0.9}Mo₆O₁₇ has a triple-Q CDW phase below $T_c = \sim 120$ K [67]. Furthermore, it has been reported that doping MoO₂ with K⁺ can induce superconductivity and anomalous metallic conductivity [68], where two phase transitions have been observed in the compound, one at ~ 54 K and another near room temperature. An explanation as to why the CDW phase transition arises in certain low-dimensional materials has been provided by Pouget et al. [69]. Typically, in an energy band of a 2D metal, when the Fermi surfaces in *k*-space are two parallel lines, one line can be superposed to the other line by a translational vector. In this case, the Fermi surface is said to be nested by the vector. Here, the occupied and unoccupied band levels (near the Fermi

surface) related by the nesting vector are roughly degenerate. Nevertheless, under a certain perturbation, a small energy gap at the Fermi level can be introduced by virtue of CDW caused by the interaction between those band levels [69]. For a realistic 2D metal with several partially overlapping energy bands, even if the Fermi surface nesting does not exist at a first glance, “hidden nesting” may still happen as long as the band structures fulfill certain requirements, which could anyway lead to CDW related metal-to-semiconductor transition [69]. Detailed discussion is out of the scope of this paper. We note that in electrical measurement, it may be reflected by kinks in resistance–temperature curves where metal-to-semiconductor phase transitions occur, such as the phenomenon observed in Fig. 6i. The three kinks appearing in Fig. 6i may be due to different CDW phase transitions happening at different temperatures.

4 Conclusion

In conclusion, we have showcased the efficacy of flux growth in synthesizing 2D MoO₂ flakes via CVD. This method presents notable benefits in achieving reduced thickness and expanded dimensions when contrasted with traditional approaches. Additionally, the resulting 2D MoO₂ shows high crystallinity and good conductivity and exhibits the CDW phenomenon in the measurement of resistivity as a function of temperatures. We demonstrate that the prevailing MSAM in the 2D community is in fact the flux method. We have conducted a systematic investigation into the impact of temperature, homogeneous time and cooling rate on growth products. This aims to extend the application of the flux method, commonly employed for synthesizing bulk single crystals, to non-layered 2DMOs and complete the understanding of the MSAM for the preparation of non-layered materials. We anticipate that by further refinement, thinner and larger areas of 2D non-layered materials prepared by the flux method can be achieved, with the possibility of extending to layered materials, assisting in the further exploration of their properties and applications.

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Declarations

Conflict of interest The authors declare that they have no conflict of interest

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