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Toward understanding the phase-selective growth mechanism of films and geometrically-shaped flakes of 2D MoTe₂[†]

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Two-dimensional (2D) molybdenum ditelluride (MoTe₂) is an interesting material for fundamental study and applications, due to its ability to exist in different polymorphs of 2H, 1T, and 1T', their phase change behavior, and unique electronic properties. Although much progress has been made in the growth of high-quality flakes and films of 2H and 1T'-MoTe₂ phases, phase-selective growth of all three phases remains a huge challenge, due to the lack of enough information on their growth mechanism. Herein, we present a novel approach to growing films and geometrical-shaped few-layer flakes of 2D 2H-, 1T-, and 1T'-MoTe₂ by atmospheric-pressure chemical vapor deposition (APCVD) and present a thorough understanding of the phase-selective growth mechanism by employing the concept of thermodynamics and chemical kinetics involved in the growth processes. Our approach involves optimization of growth parameters and understanding using thermodynamical software, HSC Chemistry. A lattice strain-mediated mechanism has been proposed to explain the phase selective growth of 2D MoTe₂, and different chemical kinetics-guided strategies have been developed to grow MoTe₂ flakes and films.

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Introduction

Among various members of two-dimensional (2D) transition metal dichalcogenides (TMDs), molybdenum ditelluride (MoTe₂) has sparked tremendous research interest in recent years, due to its structural variations and fascinating physical properties, such as existence of stable semiconducting 2H-phase to semi-metallic 1T and 1T'-phases,^{1,2} direct band gap similar to that of Si (~1.1 eV) in single and bilayer semiconducting 2H-phases,^{3,4} high carrier mobility of 4000 cm² V⁻¹ s⁻¹ and a giant magnetoresistance of 16 000% in a magnetic field of 14 T at 1.8 K,⁴ and room temperature phase change behavior from the 2H to 1T'-phase under applied strain⁵ and laser irradiation.⁶ Hence, 2D MoTe₂ is a promising material for several frontier applications, including electronics,^{3,4} optoelectronics,⁷ spintronics,⁸ switching devices,⁹ and thermoelectric devices.¹⁰ 2D MoTe₂-based devices can be realized if a scalable method for the production of high quality flakes and large area films of the material is developed. Chemical

vapor deposition (CVD) is a versatile and inexpensive method for the growth of flakes and large area films of 2D TMDs with high quality. However, the low chemical reactivity of transition metals with tellurium (Te) and the small electronegativity difference between them (for example, ≈0.3 eV between Mo and Te)¹ imposes a challenge for a straightforward synthesis of highly crystalline transition metal ditellurides by the CVD process. Hence, an in-depth understanding of the CVD reaction is needed to overcome such a complex nature of chemical reaction between the transition metals and Te.

Researchers have developed different strategies for the phase-controlled CVD growth of 2D MoTe₂. For example, Park *et al.*¹¹ successfully synthesized centimeter-scale 2H- and 1T'-MoTe₂ films by varying the Te partial pressure and quenching time, and Xu *et al.*¹² reported the synthesis of the single crystals of 2H- and 1T'-MoTe₂ in a two-step CVD process, where the 2D MoO₂ single crystals were grown first and were tellurized at different temperatures for growing different phases. Large area 2H- and 1T'-MoTe₂ films have also been controllably synthesized by adjusting the substrate location and the cooling rate after completion of the growth,¹³ and by tellurizing the Mo metal film and MoO₃ films,¹⁴ respectively. There are a few more reports^{15–18} that also demonstrated the phase-controlled growth of large area 2H- and 1T'-MoTe₂ films. Empante *et al.*² has reported a successful synthesis of 2H-, 1T-, and 1T'-MoTe₂ phases by varying the cooling rate after completion of the growth but has not demonstrated the controlled growth of films and flakes. 1T- phase of 2D MoTe₂, which has been known to be less stable

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The analysis chamber pressure is maintained below 5.0×10^{-9} mbar to avoid contamination of the surfaces to be analyzed. Current of Bi analysis beam was 14 nA over $500 \mu\text{m} \times 500 \mu\text{m}$ of area. Ion flux was 1.2×10^{12} ions per cm^2 for $100 \mu\text{m} \times 100 \mu\text{m}$ sample and 6.0×10^{13} ions per cm^2 for $100 \mu\text{m} \times 100 \mu\text{m}$ sample. Ion flux for sputtering was 6.8×10^{15} ions per cm^{-2} . For high lateral resolution mass spectral images acquired in this study, a Burst Alignment setting of 25 keV Bi^{3+} ion beam was used. The negative secondary ion mass spectra obtained were calibrated using C^- , O^- , OH^- , C_n^- . The positive secondary ion mass spectra were calibrated using C^+ , C_2H_3^+ , C_3H_5^+ , C_4H_7^+ .

Results and discussion

Fig. 1a shows a schematic of the CVD setup for the phase selective synthesis of 2D MoTe_2 films with 2D tellurene obtained as a byproduct material. The schematic for the phase selective synthesis of 2D MoTe_2 flakes or rings with 2D MoO_2 as a byproduct material has been presented in Fig. 1Sa, ESI.† Although the details of the chemical reactions of this CVD process are not known yet, the following reaction steps are expected to occur:

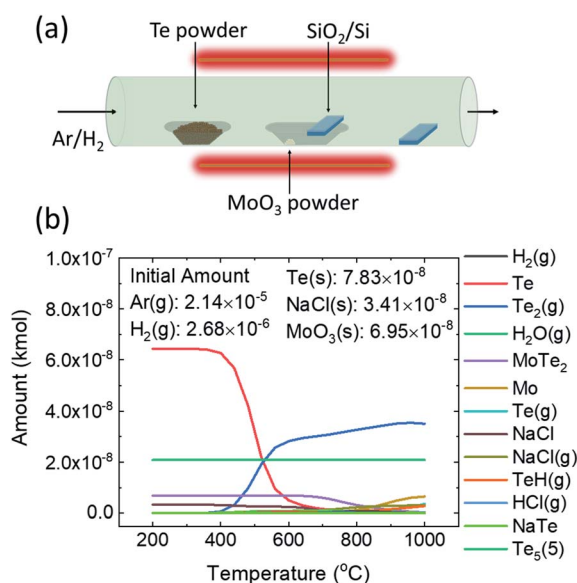
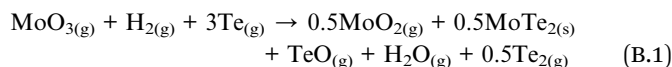
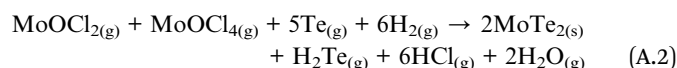
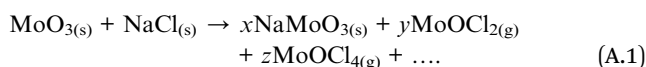


Fig. 1 (a) Schematic of the CVD setup for the growth of 2D MoTe_2 and byproduct 2D materials, MoO_2 and Te, and (b) thermodynamic equilibrium product compositions for the reaction of MoO_3 (6.95×10^{-8} kmol) with Te (7.83×10^{-8} kmol) for film growth, predicted by HSC Chemistry.



The reactions (A.1) and (A.2) have been written based on the previously suggested reaction mechanism¹⁹ for MoTe_2 synthesis, and the reactions (B.1) and (B.2) are the expected reactions written from our thermodynamic equilibrium product calculations. We suggest the additional reactions (B.1)–(B.2) for our CVD reactions based on the following reasoning: (1) HSC calculation shows that the Gibbs free energies are negative for these reactions, indicating that they are thermodynamically favorable, and (2) as predicted by the reaction (B.1), 2D MoO_2 crystals were deposited under Te deficient reaction condition.

Fig. 1b shows a typical graph showing thermodynamic equilibrium product compositions calculated with HSC software for the reaction of 10 mg of MoO_3 with 1 g of Te in the presence of promoter compound, NaCl, and gases H_2 and Ar to form the films of MoTe_2 . The corresponding graph obtained with smaller amounts of MoO_3 (1 mg) and Te (300 mg) for flakes or rings deposition has been presented in Fig. S1b, ESI.† From Fig. 1b, it is evident that the reaction leads to the formation of MoTe_2 even at a low temperature, 200 °C, but due to slow reaction kinetics, a lower growth temperature is not suitable. The equilibrium concentration of the product, MoTe_2 starts decreasing at 650 °C and almost vanishes at 900 °C. The effect of temperature on the reaction rate constant k is given by the Arrhenius equation, $k = Ae^{-E_a/RT}$,²⁰ where A is the pre-exponential factor, E_a is the activation energy, R is the molar gas constant, and T is the absolute temperature. Based on this equation, higher the temperature, faster is the reaction rate, which is generally intended for minimizing the total reaction time. However, the graph (Fig. 1b) shows that the formation of unintended byproducts, such as $\text{TeH}(\text{g})$, $\text{Te}_2(\text{g})$, $\text{Mo}(\text{s})$, and $\text{NaCl}(\text{g})$ increase after the temperature of 730 °C. Based on this thermodynamic equilibrium calculation, we chose 650 to 730 °C as the maximum deposition temperature for the growth of 2D MoTe_2 .

Fig. 2a, c, and e show that flakes of the different phases of MoTe_2 can exist in different geometrical structures, such as hexagonal for 2H-, hexagonal, triangular, and truncated-triangular (see Fig. S2, ESI†) for 1T-, and ring-like for 1T'-phases. The flakes were grown by controlling the amounts of the precursors to about 1 mg for MoO_3 and 200 mg for Te powder, and the SiO_2/Si substrates were placed upside down on the boat containing MoO_3 . For the film growth (see Fig. 2b, d, and f), the amounts of MoO_3 and Te powders were increased to about 10 mg and 500 mg, respectively, and the SiO_2/Si substrates were placed facing up on the boat. We note that despite a good controllability in film growth using the above-mentioned growth parameters and strategy, there remains a possibility of the formation of single crystals and/or streaks of the material in small areas, mostly near the edges of the substrates. However, films and flakes can be deposited with preference and with dominant morphology by following our growth parameters and strategies. Moreover, although it is known to be difficult to stabilize 1T- MoTe_2 ,² we successfully synthesized crystals and

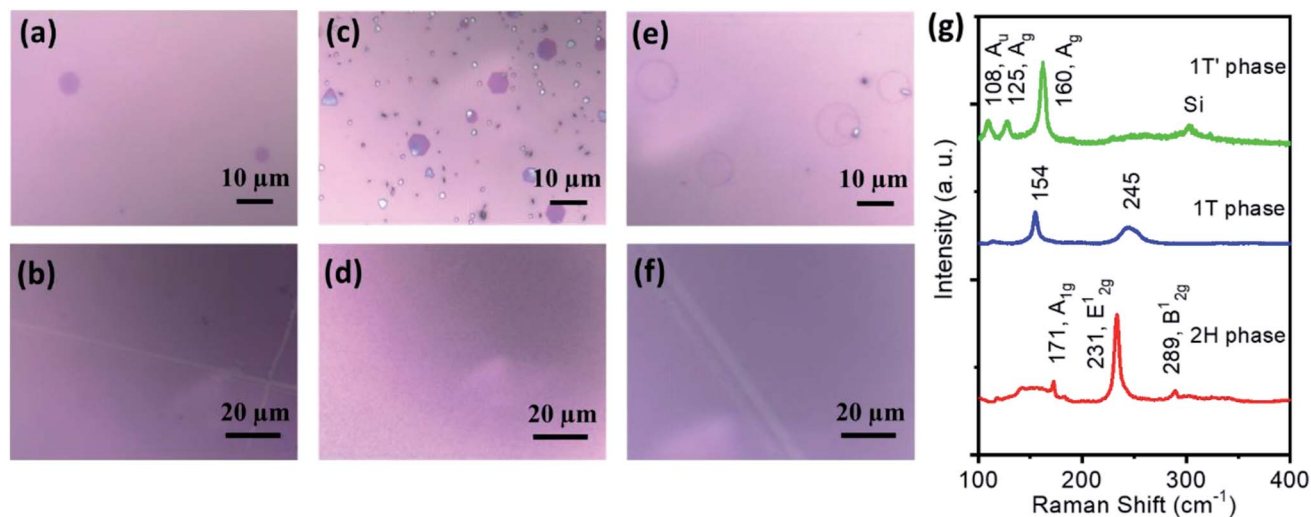


Fig. 2 Optical micrographs of the flakes and films of (a and b) 2H-phase, (c and d) 1T-phase, and (e and f) 1T'-phase of 2D MoTe₂. (g) Typical Raman spectra of as-synthesized 2H-, 1T-, and 1T'-2D MoTe₂.

films of this phase. The detailed synthesis mechanism will be discussed later.

Raman spectroscopy is the necessary and sufficient characterization technique for the identification of 2D materials and their phases. Fig. 2g depicts the typical Raman spectra of 2H-, 1T-, and 1T'-2D MoTe₂ phases for both flakes and films

morphologies. We observed that Raman spectra of flakes and films for each phase are similar in terms of full width at half maximum (FWHM) of their corresponding vibrational bands, indicating similar qualities of both morphologies. FWHM of the Raman bands in the Raman spectra collected on different flakes and within a film of the same phase also vary slightly, and the

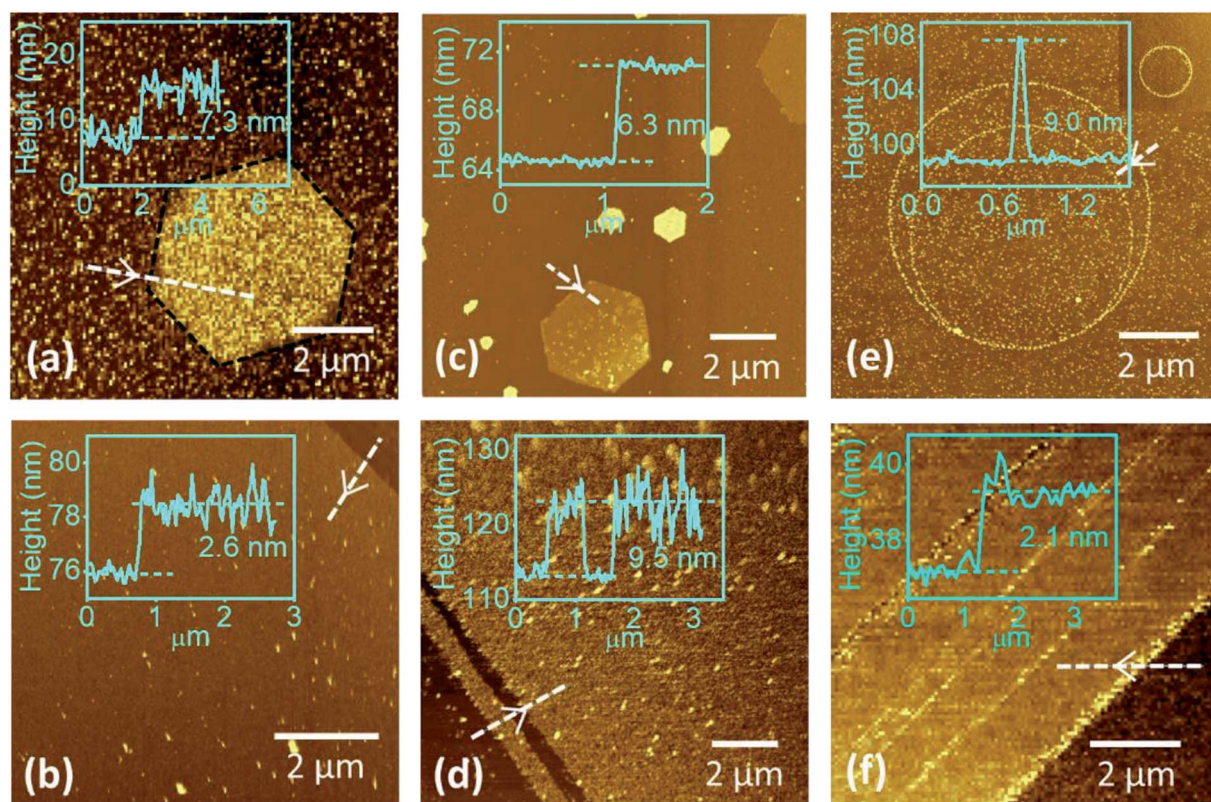


Fig. 3 Tapping-mode AFM images and corresponding height profile of the flakes and films of (a and b) 2H-, (c and d) 1T-, and (e and f) 1T'-MoTe₂. The dotted lines in (a) are drawn as a guide to show hexagonal shape of the 2H-phase crystal.



Raman spectra are not significantly different, which could be due to local temperature variation on the substrates or unintended particles formation on the flakes and films. Hence, we presented typical Raman spectra for both flakes and films morphologies to represent only the phases. The samples grown at 630 °C show peaks at 171, 231, and 289 cm^{-1} corresponding to A_{1g} , E_{2g}^1 , and B_{2g}^1 vibrational modes of 2H-MoTe₂, consistent with previous reports.^{2,4–6} The samples grown at 750 °C are confirmed to be 1T'-MoTe₂ as the Raman bands observed at 108, 125, and 160 cm^{-1} correspond to the A_u , A_g , and A_g , respectively.^{2–6,19,21} The samples grown with the two-step growth strategy show a Raman spectrum with a sharp peak at 154 and a broad peak at 254 cm^{-1} , which correspond to 1T-MoTe₂ with an octahedral crystal structure.² To our knowledge, this is the second report on the synthesis of 1T-MoTe₂, and the previous work² has shown a round structure for flakes, whereas our work shows perfect geometrically-shaped crystals, which suggests that the quality of 1T-MoTe₂ grown by our method could be higher. With higher amounts of MoO₃ and Te and SiO₂/Si substrate facing up on the boat, continuous films were obtained for 2H, 1T, and 1T' phases of 2D MoTe₂ as shown in Fig. 2b, d, and f, respectively.

Fig. 3a–f show the tapping-mode AFM images and corresponding height profiles of the flakes and films of 2H-, 1T-, and 1T'-MoTe₂. The images show that 2H-MoTe₂ flakes are formed with hexagonal shape, and 1T-MoTe₂ was formed with different geometrical shapes such as hexagon, triangle, and truncated triangle (see Fig. S2, ESI†). In contrast to 1T'-MoTe₂ flakes formed in various shapes such as rectangular,²² square,¹² star-like,²³ and irregular-shaped flakes^{1,19} reported previously, we obtained 1T' phase with ring-like structures in our growth condition. The detailed growth mechanism of ring-shaped 1T' phase of MoTe₂ remains to be explored. We hypothesize that the loop formation could be the result of a spontaneous polarization process, which originates from the binding of the positively and negatively charged surfaces on the opposite sides of belt-like MoTe₂ nanostructures. Fig. 3b shows the heights of the typical 2H-, 1T-, and 1T'-MoTe₂ films are ~2.6, ~9.5, and ~2.1 nm, which are equivalent to about 4, 15, and 3 layers, respectively. The measured root mean square (rms) roughness of the 2H-, 1T-, and 1T'-MoTe₂ films are ~0.8, ~3.0, and ~0.6 nm, respectively.

The bottom part of the Fig. 4a shows the Raman spectrum of the material grown at the downstream furnace edge obtained as a byproduct. The vibrational peaks appearing at 120 and 140 cm^{-1} resemble the longitudinal phonon mode (A_1) and the high frequency vibrational mode (E^2) of 2D tellurene.²⁴

The top part of the Fig. 4a shows the Raman spectrum of the byproduct obtained when Te was not well evaporated, *i.e.*, with the minimal amount of Te supply. The peaks appearing at 125, 202, 208, 228, 344, 361, 422, 458, and 496 cm^{-1} are the characteristic vibrational bands of monoclinic structure of 2D MoO₂.²⁵ From our observation, it can be understood that the deposition of the quasi-2D crystals of MoO₂ is due to the reduction of MoO₃ by H₂ gas, which is consistent with the previous reports.^{26,27} Fig. 4b and c show the AFM images of the

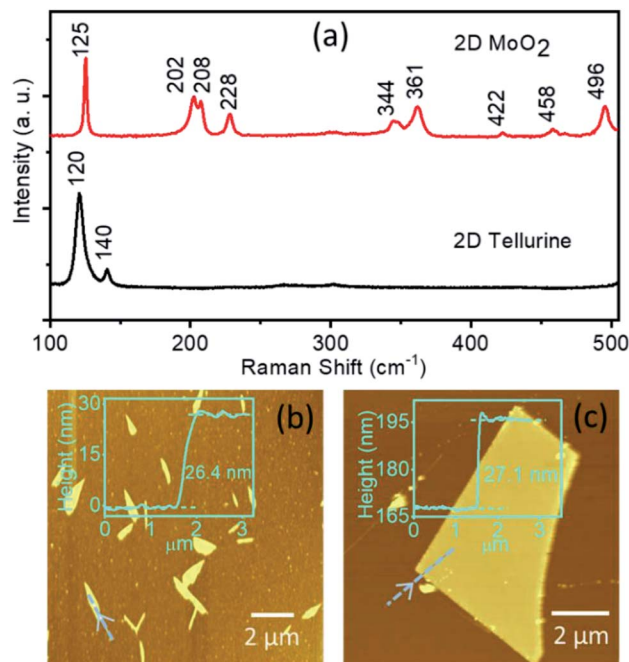


Fig. 4 (a) Raman spectra of 2D tellurene and MoO₂ showing characteristic vibrational bands. Tapping-mode AFM images and corresponding height profile of (b) tellurene nanobelts and (c) MoO₂ flakes.

2D tellurene nanobelts and MoO₂ flakes with a typical thickness of ~26.4, and ~27.1 nm, respectively.

XPS measurements were performed to analyze the elemental composition and chemical states of the synthesized samples, especially those that needed more characterization in addition to the Raman spectroscopy analysis and AFM measurements. We note that all the samples got oxidized to some extent when exposed to air for a few hours. During the course of various measurements, all samples were almost equally exposed to ambient, but 1T-MoTe₂ was not found oxidized more heavily than other phases and 2D tellurene, indicating a good stability of this phase. We sputtered each sample with Ar⁺ ion before analyzing for XPS. Fig. 5a shows the XPS survey spectra of the Ar⁺ ion-sputtered 1T-MoTe₂, 1T'-MoTe₂, and tellurene films. The survey spectra show Mo and Te peaks at ~232.8 eV and ~573.8 eV, with 4.1 and 3.4 atomic%, respectively for 1T-MoTe₂, and 3.3 and 2.9 atomic%, respectively, for 1T'-MoTe₂ films. A slightly smaller value of the Te content with respect to Mo is due to Ar⁺ ion sputtering, which removes the top layer consisting of mostly oxidized Te and adventitious carbons. For this reason, no or negligibly small carbon peaks are observed in the spectra. For comparison purpose, we analyzed the 1T-MoTe₂ film before Ar⁺ ion sputtering and observed Mo and Te atomic% as 1.9 and 4.1, respectively, with the Te to Mo atomic% ratio ~2 as expected (see Fig. S3, ESI†). The survey spectrum of the tellurene film, on the other hand, does not show Mo content, confirming the nanobelts were purely of Te. Since we did not observe any tellurene nanobelts or crystals on the samples placed in the central region of the furnace or a higher temperature region, it is apparent that Te vapor crystalizes at a relatively lower



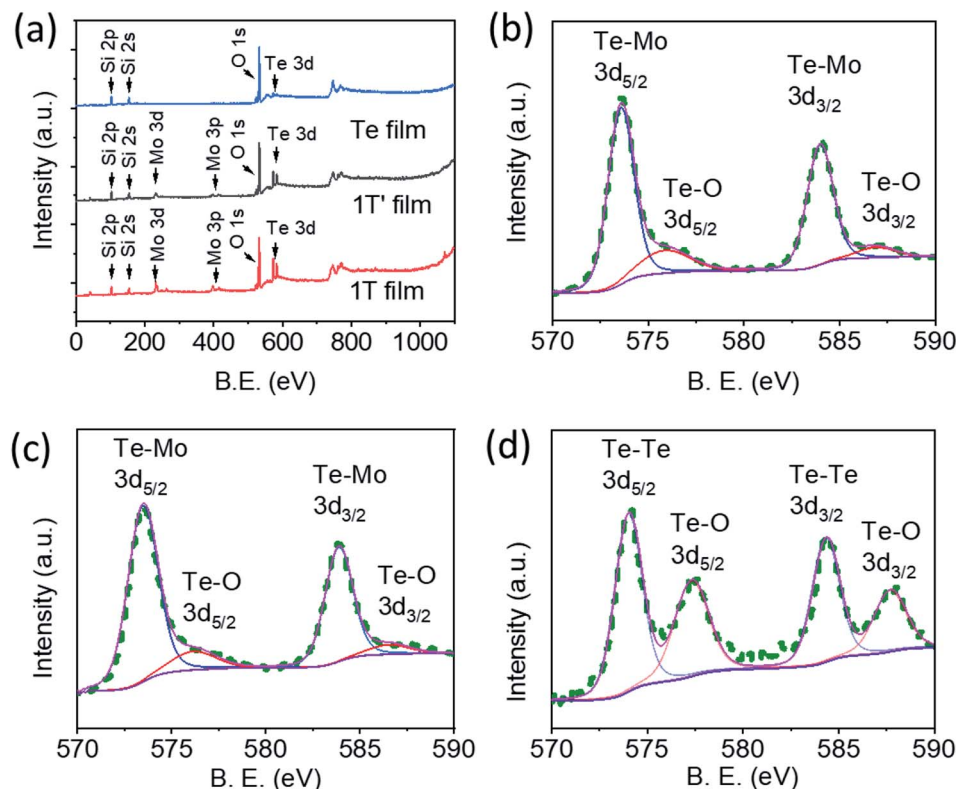


Fig. 5 (a) XPS survey spectra of 1T, 1T', and Te. High resolution peak of (b) Te 3d of 1T film, (c) Te 3d of 1T' film, (d) Te 3d of tellurene film.

temperature $\sim 500^\circ\text{C}$ and forms 2D tellurene nanostructures. Fig. 5b–d depict the high-resolution peaks of Te 3d region for 1T-MoTe₂, 1T'-MoTe₂, and tellurene films. The deconvolution of the Te peaks shows that a small amount of Te-O is still present in the samples even after the Ar ion sputtering.^{2,28} For 1T-MoTe₂

film, the high-resolution XPS peaks (Fig. 5b) of Te are observed at 573.4 (3d_{5/2}) and 583.7 (3d_{3/2}) due to Te-Mo bonding, and we did not observe a significant shift of the Te peaks in 1T'-MoTe₂ with respect to 1T'-MoTe₂. However, the Te peaks for tellurene nanobelts (Fig. 5d) appear at a slightly larger values of binding

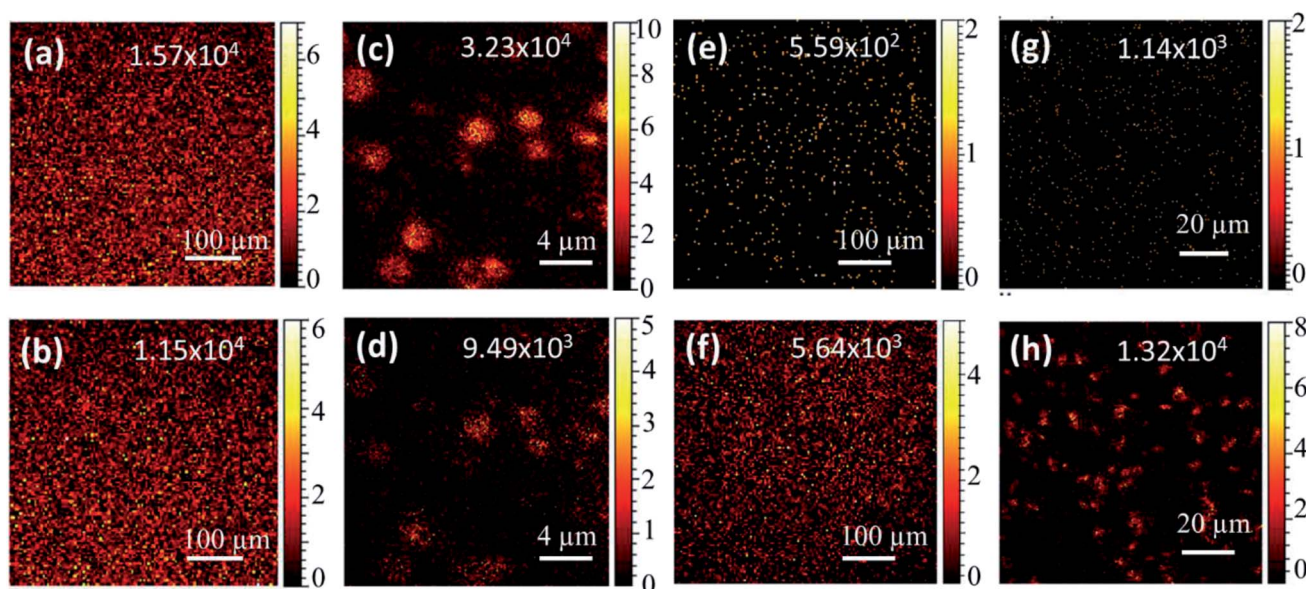


Fig. 6 Secondary ion ToF-SIMS color maps for (a and b) Mo and Te, respectively, in 1T film, (c and d) Mo and Te, respectively, in 1T flakes, (e and f) Mo and Te, respectively, in 2D Te film, and (g and h) Mo and Te, respectively, in 2D Te nanobelts.



energy (B. E.), 574.1 ($3d_{5/2}$) and 584.3 ($3d_{3/2}$) compared to that of Te-Mo peak for 1T-MoTe₂ or 1T'-MoTe₂ phases, which agrees with the report.²⁹

Elemental composition of the samples was further analyzed using ToF-SIMS. TOF-SIMS provides elemental, chemical state, and molecular information from surfaces of solid materials.³⁰ Secondary ion ToF-SIMS maps of Mo and Te collected on 500 $\mu\text{m} \times 500 \mu\text{m}$ area of 1T-MoTe₂ and tellurene films are presented in Fig. 6a, b, e and f, respectively, and the maps of 100 $\mu\text{m} \times 100 \mu\text{m}$ area of 1T-MoTe₂ flakes and tellurene nanobelts are presented in Fig. 6c, d, g and h, respectively.

The values in the inset show the respective secondary ion (Mo^+ and Te^+) counts recorded, but the values do not correspond to the stoichiometry of the compound material, MoTe₂. A much lower counts of Te^+ may be ascribed to larger ionization energy (9.01 eV) of Te^{3+} than that of Mo (7.09 eV)³² and the weak stability of Te^+ ions, due to a low electron affinity (1.97 eV).³³ For this reason, ToF-SIMS data alone are not ideal for determining stoichiometry of a compound. Hence, true amount of Te in the samples can be expected higher than what has been represented in Fig. 6b, d, f, and h. Secondary ion maps of Mo and Te for 1T-MoTe₂ flakes shown in Fig. 6c and d support that the flakes are comprised of Mo and Te. A higher-resolution maps of Mo and Te for 1T-MoTe₂ flakes deposited on SiO₂/Si have been presented in Fig. S4a and b, ESI† respectively. 2D tellurene growth on the substrates placed at the downstream furnace edge has also been confirmed by ToF-SIMS study. Secondary ion ToF-SIMS maps of Mo and Te in tellurene film deposited on SiO₂/Si shown in Fig. 6e and f clearly indicate that the Mo^+ ion counts is much smaller than that of Te^+ ions. We note that the actual amount of Te in 2D tellurene film should be much larger than the one detected in our experiment for the aforementioned reason. Similar observation has been made for the Mo and Te secondary ion maps for 2D tellurene nanobelts deposited on SiO₂/Si (Fig. 6g and h), confirming that the nanobelts are not a different polymorph of MoTe₂. Secondary ion maps of Si and SiHO (Fig. S4c and d, ESI†) further show a clearer picture of small and belt-like structures of 2D tellurene nanobelts.

We observed that under ambient pressure conditions, different growth temperatures and a two-step growth strategy can lead to the phase-selective stabilization of 2D MoTe₂. The phase selectivity may result from the lattice strain that influences the grown or growing crystals. Stress/strain-dependent phase transition rate in a material can be explained by the transition state theory (TST),³⁴ according to which the phase transition rate is given by the relation, $k(\sigma) \propto e^{-E_\sigma/k_B T}$, where σ is the stress acting on the growing material, E_σ is the barrier energy for phase transition, k_B is the Boltzmann constant, and T is the absolute temperature. In the case of our CVD process, the growth temperature is a key parameter, which however, is proportional to the strain developed on the grown or growing material arising from the thermal expansivity mismatch between MoTe₂ and the substrate SiO₂/Si. Hence, at a different growth temperature condition, the grown or growing material experiences different stress/strain. According to TST, a larger stress/strain (or larger temperature) is expected to lower the phase transition barrier energy,³⁴ facilitating the phase

transition of the material. For example, a lower strain on the grown or growing material at a lower growth temperature of 650 °C is not strong enough to lower the phase transition barrier energy, which stabilizes the 2H phase. On the other hand, a higher stress/strain at a higher growth temperature of 750 °C would lower the phase transition barrier energy significantly, facilitating the phase transition and making 1T'-MoTe₂ phase more thermodynamically favorable.⁴ This explanation is consistent with the observations made in previous reports,^{2,22} where 2H-MoTe₂ and 1T'-MoTe₂ were grown at lower and higher temperatures, respectively. Such a process of strain-mediated phase stabilization is also consistent with the report, where strain is found to have a strong influence on the phase change behavior of 2D MoTe₂, *i.e.*, 2H- to 1T'-MoTe₂ phase change occurs under the application of strain and reverse phase transition occurs under removal of the strain.³⁵ CVD growth of 1T-MoTe₂ phase under a two-step growth strategy may be ascribed to the sudden strain relaxation and additional phase stabilization time provided during the second step of the growth process. When the temperature of the furnace is lowered to 670 °C, the strain on the grown or growing material is reduced facilitating the transition to 1T-MoTe₂ phase, and the phase is stabilized when the furnace is maintained at 670 °C for 15 minutes. We performed several trials with a single step, *i.e.*, at 730 °C or at 670 °C, but none of the trials led to the growth of 1T-MoTe₂, which indicates that a two-step growth process was a key recipe to obtain this phase.

The controllable growth of flakes and films of 2H, 1T, and 1T' phases of 2D MoTe₂ may be understood considering the role of kinetic factors during the APCVD process. As mentioned above, it may be expected that the gaseous-phase reaction of MoO₃ and Te in the presence of NaCl and H₂ produces MoTe₂ molecules as active species. The gaseous active species then diffuse through the boundary layer, which is a thin and active reaction zone in contact with the substrate of the growing material,^{36,37} due to the concentration gradient between its two opposite regions (above and below the boundary layer in our case). The active species are adsorbed onto the surface of the substrate and nucleate for crystal growth. The other incoming gaseous active species interact with the edges of the growing crystals and are incorporated. Inactive gaseous species are taken away by the carrier gas, however, they are sometimes incorporated into the crystal as impurities or form unintended crystals. In the process, only a part of the active species that cross the boundary layer is incorporated into the crystals, and the rest return to the bulk of the gases. A lower amount, 1 mg of MoO₃ with 300 mg of Te supplies a much smaller concentration of the gaseous active species (C_g) than a larger amount, 10 mg of MoO₃ with 500 mg of Te, provided the same ramping rate and furnace temperature. Hence, the concentration of the gaseous active species that crosses the boundary layer and interacts with the growing crystals at their edges (C_s) is proportional to the amounts of the precursors. For such a condition, Flux of the active species diffusing to the substrate surface through boundary layer (ϕ_1) and flux of the unincorporated species returning from the substrate surface (ϕ_2) are given by:³⁸



$$\varnothing_1 = h_g (C_g - C_s) \quad (1)$$

$$\varnothing_2 = k_s C_s \quad (2)$$

where h_g and k_s are the gas phase mass transfer coefficient and surface reaction rate constant, respectively. At steady state, $\varnothing_1 = \varnothing_2$. Hence, C_s can be written as:

$$C_s = h_g C_g / (k_s + h_g) \quad (3)$$

Since we synthesized the flakes and films of each phase of 2D MoTe₂ at the same temperature, the temperature dependent quantity, k_s ,²⁰ can be treated constant for any amounts of precursors (e.g., 1 mg of MoO₃ and 300 mg of Te or 10 mg of MoO₃ and 1 g of Te). From eqn (3), it is apparent that the concentration of the gaseous active species that crosses the boundary layer and interacts with the growing crystals at their edges (C_s) is less for smaller amounts of the precursors. Furthermore, the mass transport coefficient is dependent upon the thickness of the boundary layer, δ , as $h_g = D_g/\delta$,³⁹ where D_g is the gas diffusion coefficient. According to the Blasius model, the thickness of the laminar boundary layer (δ) is inversely proportional to the square root of the gas flow speed.⁴⁰ Since the gas speed can be expected less below than above the SiO₂/Si substrate placed with SiO₂ surface facing down on the boat, h_g is smaller for such a substrate orientation. Eqn (3) suggests that C_s is smaller as both factors C_g and h_g are smaller, when a smaller amount of the precursors were used and the SiO₂/Si substrate placed facing down on the ceramic boat. Hence, this growth strategy leads to a smaller nucleation density resulting in the growth of flakes or ring morphologies of the 2D MoTe₂. On the other hand, when larger amounts of the precursors were used, and the materials were grown on the SiO₂/Si substrate placed facing up on the boat, both the quantities C_g and h_g are larger, making C_s much larger. As a result, this strategy leads to a much larger nucleation density, promoting the film growth of 2D MoTe₂.

Conclusions

In summary, we demonstrated a facile approach for selective growth of few-layer flakes and films of 2H-, 1T-, and 1T'-MoTe₂ by APCVD and presented a thorough understanding of the growth mechanism. A thermodynamical software, HSC Chemistry was used to optimize the growth parameters and understand the role of thermodynamics for materials growth. Lattice strain-mediated growth mechanism has been proposed to explain the phase selective growth of 2D MoTe₂, and the strategies for selective growth for flakes or films have been developed based on the role of chemical kinetics. We also suggested the facile routes for the synthesis of 2D MoO₂ and tellurene nanocrystals, based on the observation of these nanocrystals deposited as byproducts during our synthesis. Our investigation imparts knowledge on the growth strategies to overcome the complexity of the CVD reactions and will be useful for growing not only MoTe₂ but also other layered 2D materials.

Conflicts of interest

There are no conflicts to declare.

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