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Correlation of grain orientations and the thickness of gradient MoS₂ films†

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In this study, we synthesized gradient MoS₂ films with a home-made suspended mask and characterized them by transmission electron microscopy (TEM) and Raman spectroscopy. The advantage of using gradient films is to simultaneously produce numerous samples under the same growth condition but with different thicknesses. The cross-sectional TEM images and their Fourier transform spectra revealed the thickness dependency of the grain orientations for synthetic MoS₂ films. Combining the TEM results and the data of Raman A_{1g} and E_{2g}¹ peaks, we found the correlation between the grain orientation and the A_{1g}/E_{2g}¹ peak area ratio. We demonstrated the potential of using the non-polarized Raman Spectroscopy to characterize the grain structures of synthetic MoS₂ films.

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Introduction

Transition metal dichalcogenides (TMDs) have attracted considerable interest in the past decade because of their two-dimensional (2-D) characteristics and various unique properties. TMDs cover a wide range of materials, including semiconductors (MoS₂ and WSe₂), semimetals (PtSe₂) and ferromagnets (VSe₂).^{1–4} Among many, MoS₂ is the most extensively investigated TMD in terms of device applications and material synthesis. It can be fabricated into various kinds of devices such as transistors,¹ electroluminescence devices,⁵ photodetectors,⁶ gas sensors,⁷ hydrogen-evolution-reaction catalysts,⁸ memristors,⁹ and memtransistors.¹⁰ Various methods for the scalable synthesis of MoS₂ films have been developed, including chemical vapor deposition (CVD),^{11,12} atomic layer deposition (ALD),¹³ molecular beam epitaxy (MBE),¹⁴ and magnetron sputtering.¹⁵ Precursors such as Mo, MoO₂, MoO₃, MoCl₅, S, H₂S, and (NH₄)₂MoS₄ were successfully demonstrated for their capability of forming MoS₂ from single layer to multi-layers.^{11–17} In contrast to the micro-sized monolayer flakes, the synthetic MoS₂ film is typically nanocrystalline with grain sizes ranging from tens to hundreds of nanometers. Different grain orientations lead to different applications. Horizontally oriented (in-plane) grains are suitable for

electronic devices, while vertically oriented (out-of-plane) grains are efficient for electrocatalysts.

From the precision point of view, transmission electron microscopy (TEM) is effective to probe the MoS₂ grain structures, providing an important guideline for material design.^{18–20} For example,¹⁸ TEM results showed that MoS₂ has a vertical growth mode after sulfurizing the Mo precursor of 5 nm at 550 °C; but it became horizontally oriented after annealing at 800 °C for 8 h. Some TEM studies reported the effects of precursor thickness¹⁹ and sulfurization time²⁰ on the orientations of MoS₂ or the growth mechanism of monolayer MoS₂ flakes. However, TEM characterization is a destructive and time-consuming method.

On the other hand, the optical Raman spectroscopy is a fast and non-destructive tool to characterize 2-D materials. The Raman spectrum of MoS₂ contains two major peaks, E_{2g}¹ and A_{1g}, which represent the in-plane shear vibration and out-of-plane breathing vibration of the molecules, respectively.²¹ Qualitatively, the frequency difference between E_{2g}¹ and A_{1g} is related to the layer number of MoS₂ crystals²² and the A_{1g}/E_{2g}¹ intensity ratio is sensitive to the orientation of MoS₂.^{18,19} Although the theoretical simulation showed that the predominant orientation of grains and the distribution of orientation angles can be evaluated from the A_{1g}/E_{2g}¹ peak area ratio, it is hard to experimentally control or identify the angular distribution of grains (not just horizontal and vertical). The polarized Raman study²³ reported that the A_{1g}/E_{2g}¹ ratio increased by a factor of 2 in MoS₂ with 75% of grains being vertically grown, in comparison with that of horizontally grown MoS₂.²³ However, the relationship between the A_{1g}/E_{2g}¹ ratio and the angular distribution of grains has not been experimentally established yet. Since the information of grain orientation is extremely important to the film-device applications, we conducted

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† Electronic supplementary information (ESI) available: S1: design of the gradient deposition, S2: *t*_{Mo} and *t*_{MoS₂} calculation, S3: analysis procedures of Raman spectroscopy, and S4: theoretical simulation of the intensity *versus* θ (PDF). See DOI: 10.1039/d1ra05982c



a parallel study on the TEM images and Raman spectra of the gradient MoS₂ films, with an attempt to find the correlation between the A_{1g}/E_{2g}¹ ratio and the angular distribution of grains. The advantage of using gradient films is to simultaneously produce numerous samples with various thicknesses.

Results and discussion

We fabricated two pieces of gradient MoS₂ films under the same conditions but with different patterns (Fig. 1a and b). The Mo precursor layers of MoS₂ were produced by the gradient deposition method adopted in a previous study.²⁴ The sulfurization processes are described in the Methods section. We measured Raman spectra on nineteen successive sites of one gradient MoS₂ film with “cross” pattern (Fig. 1a) and took cross-sectional TEM images on five MoS₂ samples cut out from another gradient film with “squared” pattern (Fig. 1b). The thickness of the Mo precursor (t_{Mo}) on the measured site was calculated by a process described in S1 and S2.†

Ten out of nineteen Raman spectra after the baseline removal are shown in Fig. 2a–e, with two spectra as a pair to correspond the thickness of the TEM sample. All Raman spectra display two peaks around 380 and 405 cm⁻², representing the E_{2g}¹ and A_{1g} mode of MoS₂, respectively.²² Cross-sectional TEM images of five MoS₂ samples are shown in Fig. 2f–j. In Fig. 2f (for the sample of $t_{\text{Mo}} = 0.88$ nm), the TEM image did not show any trace of MoS₂. However, the Raman spectra of $t_{\text{Mo}} = 0.75$ and 0.93 nm exhibit E_{2g}¹ and A_{1g} peaks, suggesting the existence of MoS₂ in the sample of $t_{\text{Mo}} = 0.88$ nm. The reason for MoS₂ not being found in the TEM image is probably because the MoS₂ film is too thin and destroyed during the sputtering of the Pt layer, which is the protective layer to prevent the damage of dual-beam FIB (focused ion beam). Except in Fig. 2f, MoS₂ textures with various grain orientations are clearly seen. The corresponding thicknesses of four MoS₂ samples (t_{MoS_2}) were determined from TEM image (Fig. 2g–j) and they have a polynomial relation with t_{Mo} , as shown in Fig. 2k. Note that the t_{Mo} values of five TEM samples ($t_{\text{Mo}} = 0.88, 1.55, 2.21, 2.88,$ and 3.54 nm) fall between the t_{Mo} values of ten Raman samples ($t_{\text{Mo}} = 0.75, 0.93, 1.46, 1.64, 2.17, 2.35, 2.71, 2.89, 3.42,$ and 3.60 nm).

The magnified and aligned TEM images are displayed in Fig. 3a–d with their Fourier transform being shown in Fig. 3e–h,

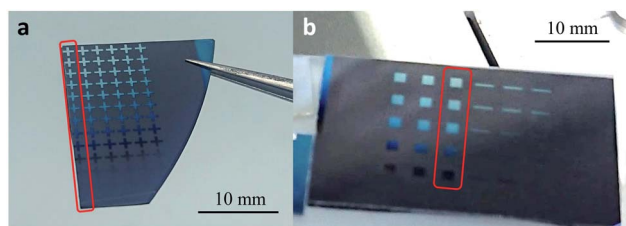


Fig. 1 (a) MoS₂ gradient film for Raman measurements. The nineteen measurement sites are on the “cross” of film within the area of the red frame. The positions of sites and the calculation of t_{Mo} and t_{MoS_2} are described in Table S2-2.† (b) Five square samples were cut from the gradient MoS₂ film (marked by red frame) for the experiments of FIB and TEM. The calculation of t_{Mo} was described in Table S2-1.†

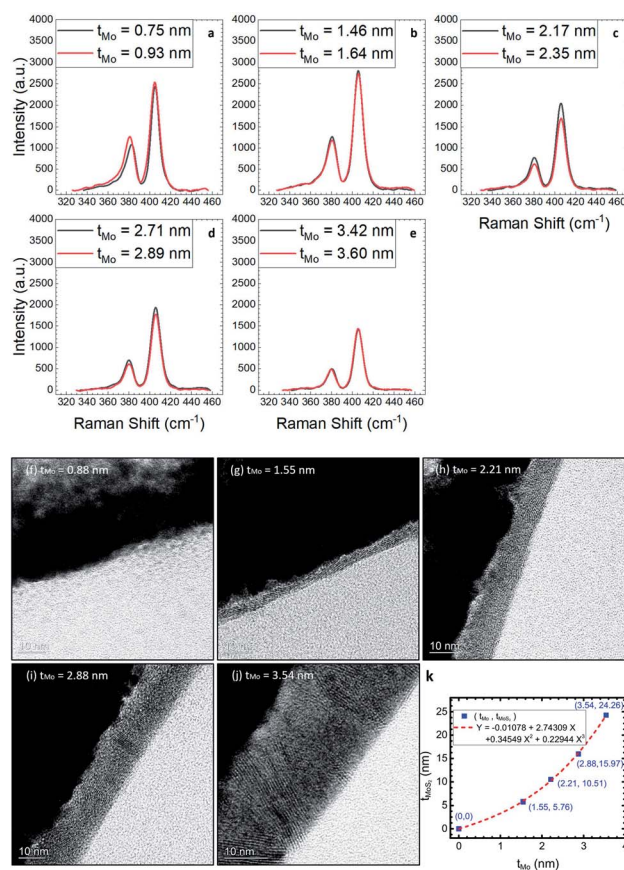


Fig. 2 (a)–(e) Ten Raman spectra of MoS₂ films with two spectra in one graph to correspond the thickness of TEM sample. (f)–(j) Cross-section TEM images of five MoS₂ samples. In the TEM images, the black region is Pt films (left), the dark grey region is MoS₂ (middle), and the light grey region is the amorphous SiO₂ (right). (k) t_{MoS_2} versus t_{Mo} for the TEM samples of (g)–(j). The paired numbers in each sample mark their X and Y positions. The origin is also shown as the boundary condition of the polynomial fitting.

respectively. The pattern and distance of bright spots in Fig. 3e–h provide the information of grain orientations. Fig. 3e has a two-fold symmetry of a pair-spot, indicating two kinds of predominant orientations with a tilted angle smaller than 20° with respect to the substrate surface. Fig. 3f shows a two-fold

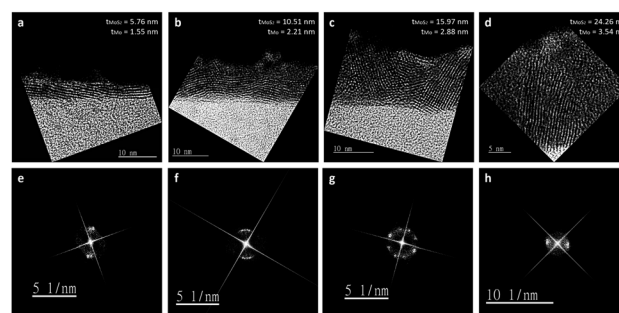


Fig. 3 (a)–(d) Magnified cross-section TEM images of four MoS₂ films. (e)–(h) The corresponding Fourier transform of (a)–(d).



symmetry of an arc structure, indicating multiple grains with different orientations confined within an angle smaller than 30° . Fig. 3g has eight major spots, indicating predominant four kinds of grains with different orientations covering from 0 to 90° . Fig. 3h shows a two-fold symmetry of a big spot, indicating the preferred orientations are within 20° . It is noted that the relative coordination of bright spots changes 90° from Fig. 3e–h due to the change in grain orientation from horizontal to vertical.

Next, we analyze the nineteen Raman spectra after the baseline removal and Lorentzian fitting (described in S3†). The Raman shifts of E_{2g}^1 and A_{1g} peaks, frequency difference between the two peaks, intensity, normalized intensity, peak widths, and peak area are plotted against t_{MoS_2} in Fig. 4a–f respectively. The t_{MoS_2} dependences of E_{2g}^1 and A_{1g} Raman shift are similar but with opposite trends (see Fig. 4a). They both show a dramatic variation at small thickness ($t_{\text{MoS}_2} < 5.46$ nm), and then reach a constant value. The Raman shift of ($A_{1g} - E_{2g}^1$) increases first with t_{MoS_2} increasing, and then saturates around 26.5 cm^{-1} with $t_{\text{MoS}_2} \geq 5.46$ nm (see Fig. 4b). The above thickness dependencies of Raman shifts are very similar with that found in layer-number dependency of MoS_2 single crystal.²² In Fig. 4c, the thickness dependencies of A_{1g} and E_{2g}^1 intensity have a similar characteristic, having a decreasing trend but with some oscillations. This decreasing trend is attributed to the optical field enhancement of Si as found in the MoS_2 single

crystal.²² We normalized the Raman peak intensity of MoS_2 with the intensity of Si (I_{Si}), as shown in Fig. 4d. Compared with Fig. 4c, the oscillation behavior becomes more profound and the decreasing trend becomes less significant. The oscillation behavior can be attributed to the optical interference of excitation and the emitted light in association with the interfaces of $\text{MoS}_2/\text{SiO}_2$ and SiO_2/Si .²⁶ In Fig. 4e, the peak-width of A_{1g} increases slightly, while that of E_{2g}^1 decreases about 27% with an increase in t_{MoS_2} , and the two widths become comparable for $t_{\text{MoS}_2} \geq 9.95$ nm (see Fig. 4e). The change in the E_{2g}^1 peak-width may be attributed to the variation of force constants associated with the structural change.²² We therefore plotted the peak areas of A_{1g} and E_{2g}^1 to include the variation of peak-width, as shown in Fig. 4f. In addition, the peak area ratio of A_{1g} over E_{2g}^1 is calculated to remove the effects of Si-substrate and interfacial interferences.

The A_{1g}/E_{2g}^1 area ratio against t_{MoS_2} is shown in Fig. 5a. Based on the slope of curve, the data were classified into 4 regions: (I) a near constant low value of A_{1g}/E_{2g}^1 ratio (~ 1.87), (II) a rising trend, (III) without clear trends and (IV) a linear relationship. These four regions can be correlated with TEM patterns in Fig. 3e–h in terms of the grain orientation and the MoS_2 thickness. Here, we defined the angle between the c -axis of grain and the substrate surface normal as θ . Region I is for t_{MoS_2} of $2.4 \text{ nm} < t_{\text{MoS}_2} < 5.5 \text{ nm}$ and with predominant horizontal grains

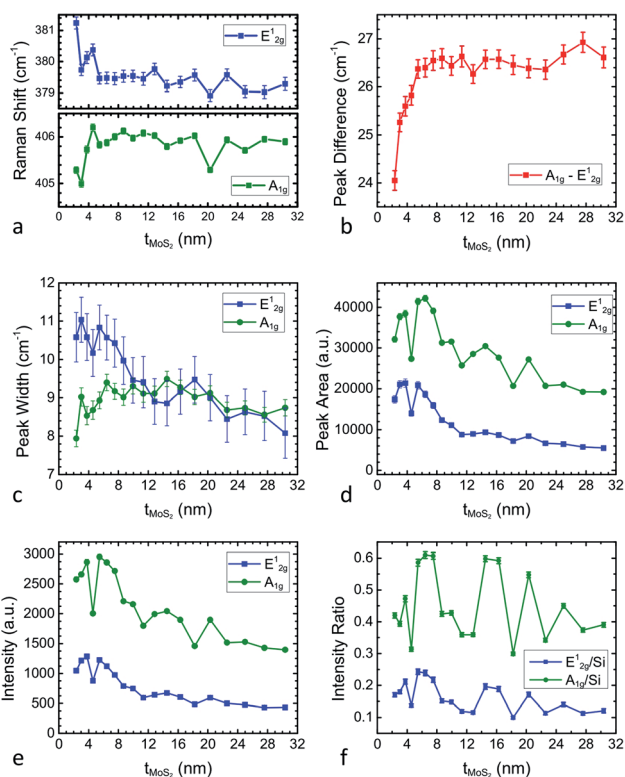


Fig. 4 The t_{MoS_2} -dependencies of (a) A_{1g} and E_{2g}^1 Raman shifts, (b) ($A_{1g} - E_{2g}^1$) Raman shift, (c) intensity of A_{1g} and E_{2g}^1 , (d) normalized intensity of A_{1g} and E_{2g}^1 with the intensity of silicon, (e) peak width of A_{1g} and E_{2g}^1 , and (f) peak area of A_{1g} and E_{2g}^1 .

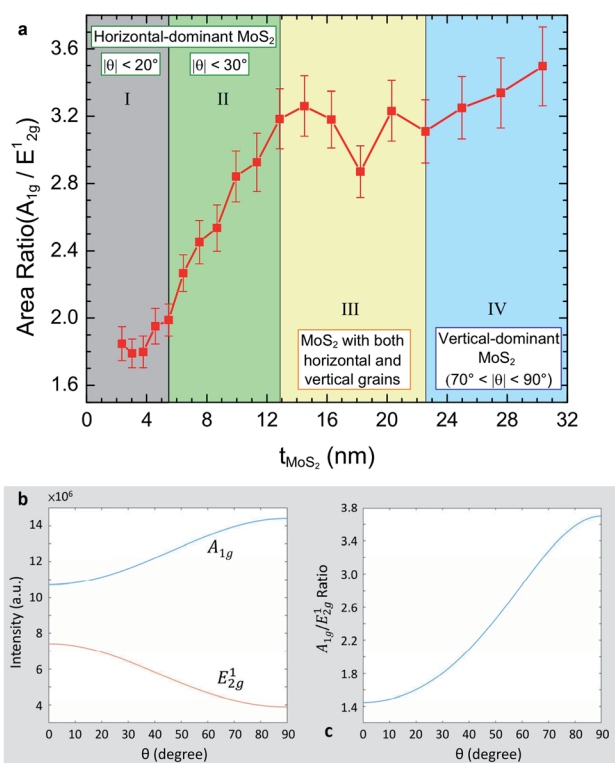


Fig. 5 (a) The diagram of the A_{1g}/E_{2g}^1 area ratio versus t_{MoS_2} with four distinguishable regions, marked by different colors. The angular distribution shown in the insets are estimations based on TEM images and their corresponding Fourier transform. (b) Simulation of the intensity of A_{1g} and E_{2g}^1 mode from $\theta = 0^\circ$ to 90° . (c) Simulation of the ratio of A_{1g}/E_{2g}^1 from $\theta = 0^\circ$ to 90° .



of $|\theta| < 20^\circ$, as found from Fig. 3e. Region II is for $5.5 \text{ nm} < t_{\text{MoS}_2} < 12.9 \text{ nm}$ and with predominant horizontal grains of $|\theta| < 30^\circ$ (Fig. 3f). Region III is for $12.9 \text{ nm} < t_{\text{MoS}_2} < 22.6 \text{ nm}$ with both horizontal and vertical grains (Fig. 3g). Region IV is for $22.6 < t_{\text{MoS}_2} < 30.4$ with predominant vertical grains of $70^\circ < |\theta| < 90^\circ$ (Fig. 3h). Overall, the A_{1g}/E_{2g}^1 area ratio enhances with the increase in the tilt angle of grain by a factor of 2.2 from the lowest value (1.70) to the highest value (3.73).

To theoretically explain the above-mentioned diagram, we adopted a method used in previous work^{23,25} to simulate the intensity of A_{1g} and E_{2g}^1 , and the ratio of A_{1g}/E_{2g}^1 by considering the contributions from the electric field of incident/scattered light, the numerical aperture of the objective lens (NA), and the Raman tensor matrices of A_{1g} and E_{2g}^1 (eqn (1) and (2)).

The total scattering intensity (denoted as I in the following passages) of a vibrational mode can be calculated as shown below:

$$I = \iint_{V\Omega} |\vec{E}_{\text{in}} R_{xyz} \vec{E}_{\text{sc}}|^2 d\Omega dV$$

$$= (R_{xx}^2 A + R_{xy}^2 A + R_{xz}^2 B) C_x$$

$$+ (R_{yx}^2 A + R_{yy}^2 A + R_{yz}^2 B) C_y$$

$$+ (R_{zx}^2 A + R_{zy}^2 A + R_{zz}^2 B) C_z \quad (1)$$

\vec{E}_{sc} and \vec{E}_{in} : vectors describing the electric field of the scattering laser and incident laser.

C_x, C_y, C_z : coefficients proportional to the integrated intensities of $E_{\text{in},x}, E_{\text{in},y}$ and $E_{\text{in},z}$.

A and B : coefficients related to NA.

R_{ij} ($i = x, y, z; j = x, y, z$): the elements of R_{xyz} matrix.

$$R_{xyz} = R_1 R_2 R_3 R_{xyz} R_3^T R_2^T R_1^T$$

$$R_{xyz, A_{1g}} = \begin{pmatrix} 195 & 0 & 0 \\ 0 & 195 & 0 \\ 0 & 0 & 276 \end{pmatrix}$$

$$R_{xyz, E_{2g}^1, x} = \begin{pmatrix} 125 & 0 & 0 \\ 0 & -125 & 0 \\ 0 & 0 & 0 \end{pmatrix} \quad (2)$$

$$R_{xyz, E_{2g}^1, y} = \begin{pmatrix} 0 & -125 & 0 \\ -125 & 0 & 0 \\ 0 & 0 & 0 \end{pmatrix}$$

R_1, R_2 and R_3 are the rotation matrix elements of the Euler angle (α, θ, γ) system (see S4† for details). (A, B) = (8.593, 5.111) for the numerical aperture of objective lens (NA) to be 0.9, and (C_x, C_y, C_z) = (13.722, 0.082, 2.192) for an X -polarized incident laser.^{23,25} In our simulation, we use (C_x, C_y, C_z) = (13.722 + 0.082, 13.722 + 0.082, 2×2.192) = (13.804, 13.804, 4.384) for the non-polarized laser beam, and considered $\alpha = 0$ to 2π and $\gamma = 0$ to 2π for each value of θ .

Fig. 5b is the simulation of angle dependences of A_{1g} and E_{2g}^1 intensity, showing that A_{1g} increases and E_{2g}^1 decreases with θ increasing from 0° to 90° . The two simulation curves cannot be compared with the data in Fig. 4d because of the interference effect being dominant over the angular effect in the actual Raman scattering measurement. The interference effect can be removed by normalized the A_{1g} intensity to the E_{2g}^1 intensity. Fig. 5c is the simulation of the angular-dependent A_{1g}/E_{2g}^1 ratio, showing a monotonically enhancement from 1.447 ($\theta = 0^\circ$) to 3.697 ($\theta = 90^\circ$), which is a factor of 2.5. The simulation range is for the grain orientation varying from zero to 90° and the predicted enhancement factor of the A_{1g}/E_{2g}^1 ratio is close to our experimental value (1.70 to 3.73 as shown in Fig. 5a). Therefore, it is evident that the enhancement of the A_{1g}/E_{2g}^1 ratio with an increase in the MoS_2 thickness originates from the increase of θ . However, the simulation curve in Fig. 5c does not completely match with the diagram of the A_{1g}/E_{2g}^1 ratio *versus* thickness in Fig. 5a owing to the fact that the simulation is based on a simple model that all grains have the same size and with coherent angles from zero to 90° . In reality, our synthetic MoS_2 films consisted of the grains with different sizes and specific angles. Therefore, further theoretical consideration and experiment design are urged to clarify the influences of different structural factors on the A_{1g}/E_{2g}^1 ratio for 2-D materials.

Conclusions

In conclusion, we used the method of gradient synthesis to simultaneously produce numerous MoS_2 film samples with different thicknesses and characterized them with TEM and non-polarized Raman spectroscopy. We demonstrated that it is possible to make MoS_2 films of various thicknesses from 2 to 32 nm at the same time. The Fourier transform patterns of TEM images provide the information of grain orientation, which can be correlated with the parameters of the A_{1g}/E_{2g}^1 peak area ratio extracted from non-polarized Raman spectra. We found a significant enhancement factor of the A_{1g}/E_{2g}^1 ratio (2.2 by experiment; 2.55 by simulation) with a change of grain orientation by 90° , suggesting that the non-polarized Raman spectra can be an effective probe to determine the grain-orientations of synthetic multi-grained MoS_2 films. The methodology in this study may be applicable to other TMD materials; thus, it benefits a fast development of large-scale production of TMD devices.

Methods

Sample preparation

Molybdenum films were deposited by a typical e-beam evaporator on the Si/SiO₂ (80 nm) substrate. The deposition rate is 0.1 \AA s^{-1} measured by the thickness monitor. The details for the gradient deposition system and the sample thickness can be found in S1 and S2.†

Si/SiO₂/Mo was placed into the quartz tube equipped with a high temperature furnace. After vacuuming the tube to a pressure lower than 5×10^{-2} torr, 1% H₂S/Ar mixed gas flowed into the tube at a rate of 400 sccm till the pressure



reached 3.6 torr. The heating process follows three steps: (i) from room temperature to 750 °C in 150 min (+4.8 °C min⁻¹), (ii) at 750 °C for 60 min, and (iii) from 750 °C to room temperature in 150 min (-4.8 °C min⁻¹). The flow rate of H₂/Ar mixed gas was kept the same during the whole heating process.

Raman measurements and analysis

Raman spectroscopy is equipped by a 532 nm laser and a computer controlled XYZ stage with the smallest step in X, Y direction being 1 μm. The numerical aperture of the objective is 0.9 and the size of the laser spot is around 6.25 μm². Details for the site positions can be found in Table S2-2.† Details for the baseline removal and the fitting procedures can be found in S3.†

FIB and TEM

A layer of 20 nm Pt was sputtered on the rectangular MoS₂ film (Fig. 1b) as a protection layer, and then the substrate was cut and thinned by FIB (FEI Helios 600i) for cross-sectional TEM samples. The samples were characterized by the 300 kV TEM (FEI Tecnai G2 F30). Details about the thickness of MoS₂ samples can be found in Table S2-1.†

Author contributions

H.-P. Chang conducted the design of the gradient deposition system, sample synthesis, Raman measurement, simulation, data analysis and first draft writing. M. Hofmann proposed the idea of gradient deposition and provided the e-beam evaporator. Y.-P. Hsieh provided the raw materials, CVD system and Raman Spectroscopy. Y. S. Chen assisted the data analysis. J. G. Lin proposed the idea of combined characterization of TEM and Raman spectra, and provided the major research funding.

Conflicts of interest

There are no conflicts to declare.

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