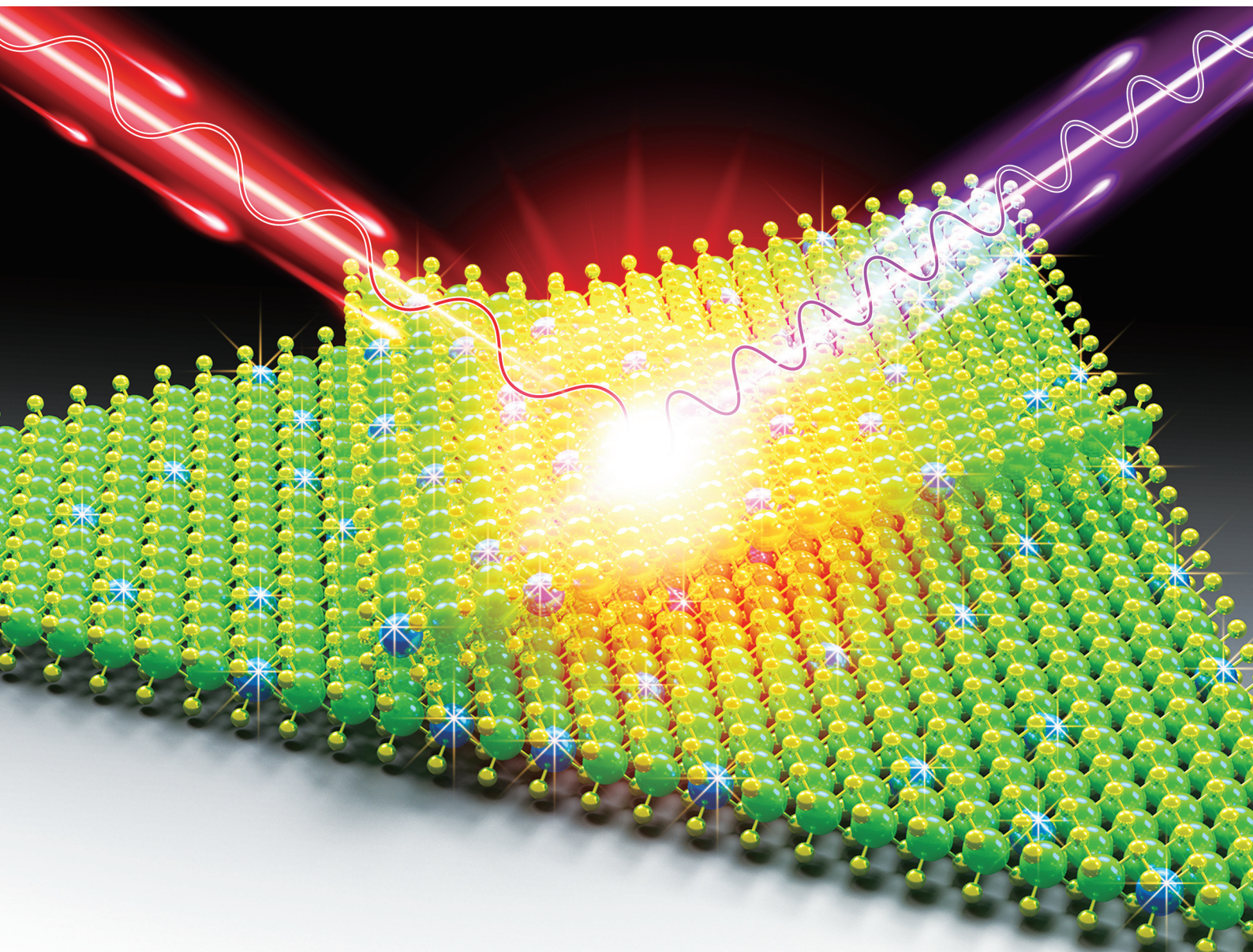


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Inversion symmetry broken in 2H phase vanadium-doped molybdenum disulfide†

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Two-dimensional (2D) layered transition metal dichalcogenides (TMDs) have received much attention in nonlinear optical applications due to their unique crystal structures and second harmonic generation (SHG) efficiency. However, SHG signals in TMDs show a layer-dependent behavior, consistent with the presence (absence) of inversion symmetry in even-layer (odd-layer) of TMDs. Herein, we synthesized monolayer and bilayer 2H and 3R phase vanadium (V)-doped MoS₂ crystal. Raman spectroscopy, XPS, and STEM were used to identify the chemical composition and crystalline structure of as-grown nanoflakes. SHG measurement was used to research the symmetry of V-doped MoS₂ crystals with different stacking orders. Significantly, the SHG efficiency in bilayer 2H phase V-doped MoS₂ is equivalent to the 3R phase, indicating an inversion symmetry broken lattice structure caused by the *in situ* V substitute for Mo sites. This study will be conducive to promote the development of promising nonlinear optical devices based on 2D material.

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Introduction

Two-dimensional (2D) transition metal dichalcogenides (TMDs), represented by MoS₂, WS₂, WSe₂, MoTe₂, have gained enhanced research interest in recent years. Many excellent physical properties were observed in atomically thin few layers or monolayer TMDs, such as high Young modulus,^{1–3} excellent transistor mobility and on-off ratio,⁴ as well as piezoelectricity,^{5,6} superconductivity^{7,8} and nonlinear Hall effect.⁹ These make the TMDs an ideal candidate for next-

generation optoelectronic device.^{10,11} Typically, 2D TMDs possess three crystalline structure, trigonal prismatic 3R, 2H phase and octahedral 1T phase, respectively.^{12,13} 3R polytype has AA stacking order of adjacent layers in same crystalline orientation with a slight in-plane shift. Based on the AA stacking order, 3R phase crystals exist with parallel in-plane nonlinear dipoles due to the absence of inverse symmetry.^{14–16} Thus, 3R TMDs possess effective second-order nonlinear susceptibility ($\chi^{(2)}$) and phase-matched second harmonic generation (SHG) from monolayer to bulk crystal regardless of layer numbers. Distinguished from 3R polytype, 2H polytype TMDs have AB stacking order of adjacent layers in opposite crystalline orientation, leading to centrosymmetry (non-centrosymmetry) in even (odd) numbered layers.^{5,17,18} As a result, 2H TMDs present a layer-dependent SHG due to the broken or recovery of inverse symmetry, which limits the application of 2D TMDs in the nonlinear optical devices since most 2D TMDs acquired by CVD synthesis or mechanical exfoliation from bulk crystal is 2H phase.

Currently, many strategies have been introduced to break the inverse symmetry of 2D TMDs and change the lattice structure,^{19–23} such as tuning by electric field or defects,^{24,25} strain regulation,⁹ twisted TMDs *via* artificial stacking,^{26,27} and controllable 3R phase crystal growth.^{14,15,28} Yu *et al.* measured a strong SHG signal in bilayer 2H WSe₂ by applying an electrical gate field.²⁴ Further bond-charge model analysis indicated that accumulated sub-monolayer screening charge in the tungsten plane resulting in a nonuniform field distribution. The nonlinear Hall effect was observed in monolayer WSe₂ by

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‡These authors contribute equal to this work.

applying uniaxial strain to break C_{3v} symmetry.⁹ Apart from applying external condition, precise regulation of growth factor during 2D materials preparation process is a direct way to achieve crystalline phase control. Chen *et al.* fabricated the 3R-stacked MoS₂ by homoepitaxial growth, and the flakes performed a vigorous SHG intensity by a factor of 6 compared with that of monolayer MoS₂.¹⁵ Although 3R MoTe₂, WS₂ and WSe₂ have been synthesized by CVD or PVD route and exhibited considerable SHG in even layers, the large-area and high-quality synthesis of a wide range of 2D TMDs remain big challenge up to now.¹³

Recently, atomic doping or alloy has been a common means in regulating electric and magnetic properties,^{7,29,30} and phase controlled-synthesis of 2D material.^{31,32} V-doped MoS₂ with different concentrations has been reported in tuning the electric conductance from n-type to p-type.²⁹ However, the nonlinear optical effect of 2D V-doped MoS₂ has not been investigated systematically before. Polarization angle-dependent SHG of monolayer Re_xMo_{1-x}S₂ ($0 < x < 0.16$) demonstrated that a slight lattice strain emerged in 2H MoS₂ due to Re substitute for Mo sites.³³ It provides a promising possibility to change the symmetry of pristine 2D material by atomic doping.

In this work, we synthesized V-doped MoS₂ flakes *via* CVD growth using MoO₃ and V₂O₅ precursors. SHG measurement was employed to investigate the doping effect of bilayer flakes with different stacking orders for 2H phase AB stacking and 3R phase AA stacking. We find significant SHG efficiency in bilayer 2H phase V-doped MoS₂ flakes compared with that of 3R phase, confirming a broken inverse symmetry regardless of crystalline orientation and layer numbers. We believe the significant result will promote the application of 2D materials in the nonlinear optoelectronic device. This work also provides a

potentially valuable strategy for the design of 2D optical crystals with broken inversion symmetry.

Experimental section

CVD synthesis of V-doped MoS₂

V-Doped MoS₂ crystals were synthesized by chemical vapor deposition (CVD) under standard atmospheric pressure. Fig. 1a illustrates the synthesis process of V-doped MoS₂ in this work. MoO₃ (Aladdin, 99.999%), V₂O₅ (Aldrich, 99.99%) and S powder (Aladdin, 99.95%) were used as source materials for the synthesis of V-doped MoS₂ crystal. A powder mixture of 2 mg NaCl, 4 mg MoO₃ and 4 mg V₂O₅ in an aluminium oxide boat was placed in the center of the quartz tube. Especially, the source MoO₃ and V₂O₅ powder were mixed uniformly in the mortar in advance. 100 mg S powder was placed at the upstream side of the furnace. The grown substrate SiO₂/Si was placed on downstream of the aluminium oxide boat with ~1 cm distance. The furnace was heated to the growing temperature 800 °C with a ramp rate of 35 °C min⁻¹ and the growth time was 5 minutes. After that, the furnace was decreased to room temperature naturally. Argon (Ar) with a flow rate of 80 sccm (standard cubic centimeters per minute) was used as the carrier gas.

Characterization

Optical imaging was taken by ZEISS Primotech microscope. AFM topography and nanoindentation experiments were performed by Bruker Dimension Icon system. PL spectra and Raman mapping characterizations were carried out with the excitation laser of 532 nm using a WITEC alpha 300R Raman system. The system was first calibrated with the Raman peak of Si at 520 cm⁻¹ before the Raman measurement of V-doped MoS₂ flakes. The PMMA-assisted wet-transfer method was employed to prepare the HADDF-STEM sample. STEM imaging was performed by FEI Themis system at 200 kV.

SHG measurement of the 2D V-doped MoS₂ flakes was carried out in a home-built vertical microscope setup with the reflection geometry. A fiber-based pulsed laser with a central wavelength at 1550 nm, a repetition rate of 18.5 MHz, and a pulse width of 8.8 ps was used as the fundamental pump radiation. A microscope objective lens (50×) with a numerical aperture of 0.75 was employed to focus the pump laser into a spot size of about 2 μm on the sample. The second harmonic signal scattered from the sample was collected by the same objective lens. In the signal collection path, a dichroic mirror was used to filter out the pump laser from the SHG signal, which was finally analyzed and detected by a spectrometer mounted with a cooled silicon charge-coupled device camera.

Results and discussion

Monolayer or bilayer V-doped MoS₂ flakes were synthesized by the salt-assisted CVD method, as shown in Fig. 1(a). Briefly,



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manufacturing technology for highly-integrated flexible smart sensors. His group is devoted to developing in situ and ex situ characterization techniques to reveal the sensing and failure mechanism for flexible materials and flexible sensors at the nanoscale.

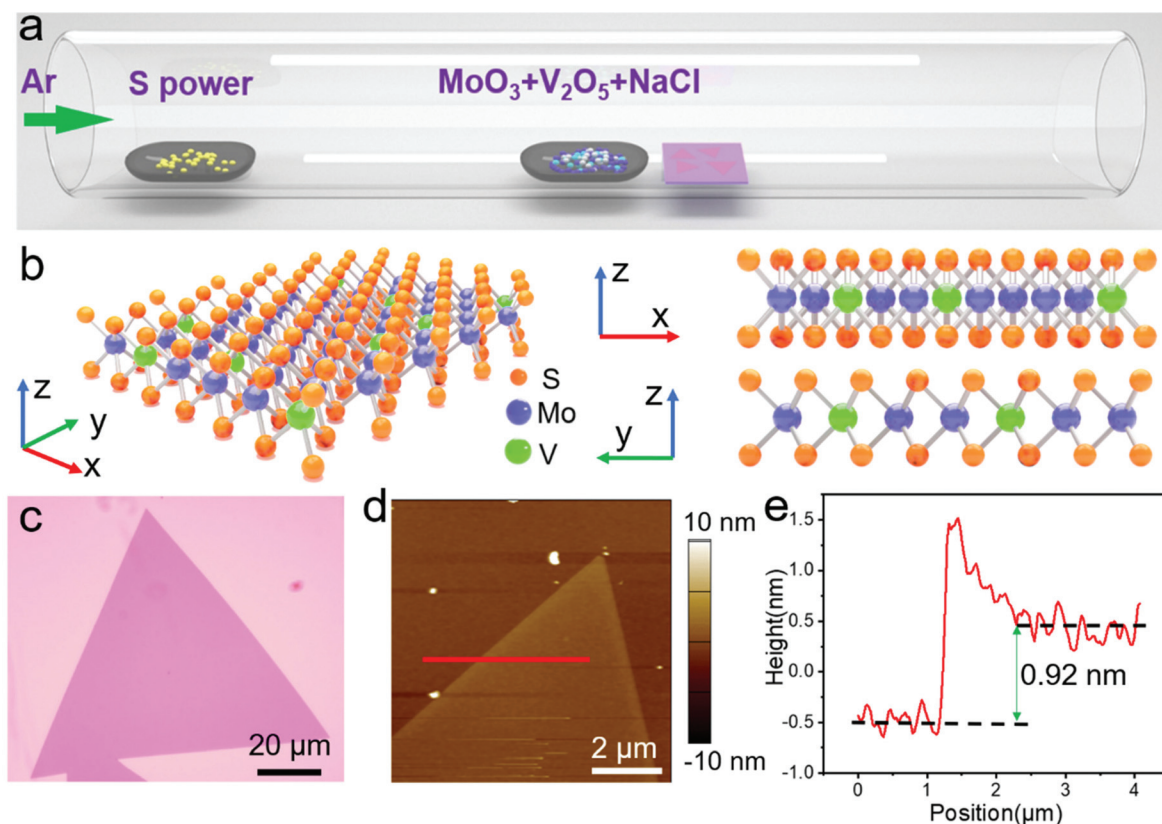


Fig. 1 (a) Schematic of synthesis of V-doped MoS₂ by CVD. (b) Atom structure and side view of V-doped MoS₂. (c) Optical image of a V-doped MoS₂ single crystal. (d and e) AFM topography and thickness measurement of V-doped MoS₂.

the mixed MoO₃ powder, V₂O₅ powder and NaCl particles were placed in the center of the heating zone. The S powder was placed in the upstream. More detailed information can be found in the Experimental section. The atomic structure of monolayer V-doped MoS₂ crystal was schematically shown in Fig. 1(b). The left panel in Fig. 1(b) gives its three-dimensional atomic structure, which is similar as the structure of 1H phase MoS₂. The difference is that in the center layer of the sandwich, partial Mo atoms were replaced by V atoms. The side view from *x*-*z* direction and *y*-*z* direction in the right panel gives a clear sight. The optical image of the growing V-doped MoS₂ flake was shown in Fig. 1(c). A single triangle crystal was observed with lateral size of about 90 μm. AFM measurement was used to characterize the topography and thickness of the flakes. Its corresponding result was shown in Fig. 1(d). The step height of 0.92 nm between the SiO₂/Si substrate and the sample measured by line scan indicates a monolayer flake.

Raman spectroscopy and X-ray photoelectron spectroscopy (XPS) characterization of the 2D V-doped MoS₂ sheets were collected and presented in Fig. 2. Raman spectra of pristine monolayer MoS₂ (black curve) and V-doped MoS₂ (red curve) flakes were displayed in Fig. 2(a). Typical Raman characteristic peaks were observed from the pristine MoS₂. The E_{2g}¹ mode at 384.5 cm⁻¹ refers to the in-plane vibration of Mo and S atoms. The A_{1g} mode at 403 cm⁻¹ refers to the out-of-plane vibration of S atom. While with the doping of V atoms in MoS₂ lattice,

significant differences appear at the Raman spectra. Multiple peaks at 158 cm⁻¹, 188 cm⁻¹, 226 cm⁻¹, 325 cm⁻¹, 352 cm⁻¹ were observed in the V-doped MoS₂, which was coincident with the previous report.^{29,30} In particular, the peak at 226 cm⁻¹ is the characteristic peak of disordered MoS₂, which could be caused by the V atoms induced broken of lattice symmetry. In order to find out the distribution of V dopant, Raman intensity mapping of V-doped MoS₂ flake was further carried out. As shown in Fig. 2(b), the upper left panel is an optical image of the sample, and the selected region is marked by a white frame. The homogeneous and uniform Raman intensity mapping of both E_{2g}¹ mode at 384 cm⁻¹, A_{1g} mode at 403 cm⁻¹ and characteristic mode of disordered MoS₂ at 226 cm⁻¹ reveal the V atoms are uniformly doped throughout the entire nanoflakes. XPS measurement further demonstrates the existence of S, Mo, and V elements. Fig. 2(c-e) give the narrow spectrum of S 2p, Mo 3d, and V 2p state, respectively. The spectra of S 2p core level binding energy have two strong peaks at 161.3 eV and 162.5 eV, corresponding to S 2p_{3/2} and S 2p_{1/2}. And the peaks of Mo 3d core level state at about 228.4 eV and 231.6 eV, corresponding to Mo 3d_{5/2} and Mo 3d_{3/2}, respectively. The bonding states of S and Mo elements in the V-doped MoS₂ alloy exhibit a slight energy shift compared with that of pristine MoS₂ in the previous report.^{34,35} This can be explained by the effect of the nearby doped V atom. The red curve in Fig. 2(e) is the Gaussian fit of the signal of V 2p orbital split.

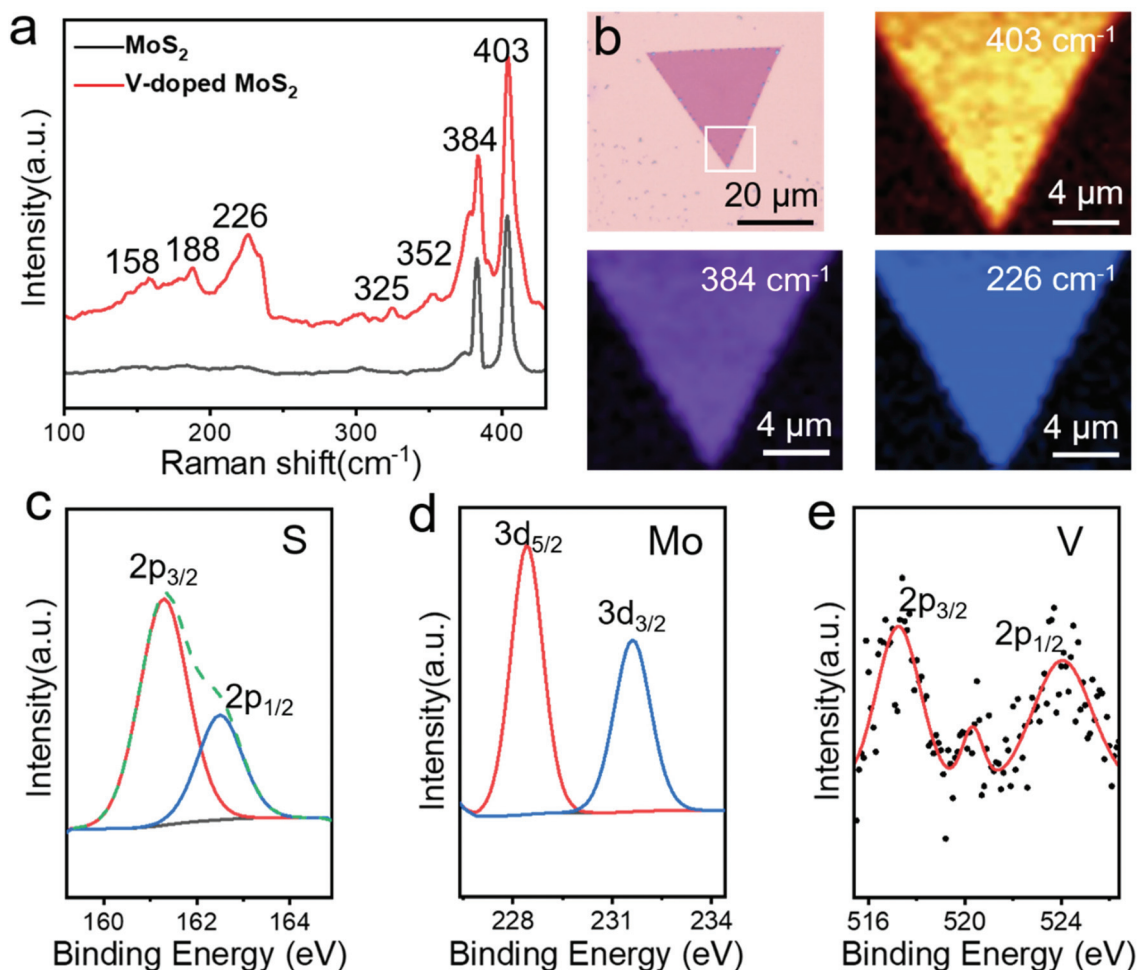


Fig. 2 Raman and XPS measurement of V-doped MoS₂. (a) Raman spectra of pristine MoS₂ and V-doped MoS₂. (b) Optical image of V-doped MoS₂ and its corresponding Raman intensity mapping images at 403 cm⁻¹, 384 cm⁻¹ and 226 cm⁻¹. (c–e) XPS narrow spectra of the S 2p, Mo 3d, and V 2p orbitals, respectively.

Two strong peaks located at about 517.2 eV (2p_{3/2}) and 524.0 eV (2p_{1/2}), which is consistent with the original chemical state of V atom in a hexagonal structure, demonstrating the successful *in situ* replacement of Mo atom in MoS₂ lattice.³⁶ As shown in Fig. 2(e), the experimental result presented by black points is not so flat, indicating a low concentration doping of V atoms in the sample.

To assess the quality of the as-synthesized V-doped MoS₂ flakes precisely, high-angle annular dark-field scanning transmission electron microscopy (HAADF-STEM), selected area electron diffraction (SAED) and energy dispersive X-ray spectroscopy (EDS) were employed to explore the crystalline structure and chemical composition of the sample. Fig. 3(a) is a low magnification STEM image of a monolayer V-doped MoS₂ flake transferred on a copper screen. The edge of the nanofilm has been lapped over during the transfer process. The SAED pattern (Fig. 3(b)) taken from the flake exhibits regular hexagonally arranged diffraction spots, which can be fitted to the hexagonal lattice structure of MoS₂. It means that no lattice mismatch generation with the doping of V atoms in MoS₂. The

high-resolution STEM imaging gives obvious evidence of *in situ* substitute between Mo and V atoms further. The aberration correction annular dark-field (ADF)-STEM imaging can provide the Z-contrast with atomic resolution, which is proportional to the quadratic square of the atomic number (*Z*). So, the Mo (*Z* = 42) site is bright and V (*Z* = 23) site is dark in the ADF-STEM imaging (Fig. 3(c)).

As a contrast, STEM imaging of pristine MoS₂ flakes was also characterized and no dark field was found in the MoS₂ lattice (Fig. S1†). Red arrows marked the substitute of Mo sites by V atoms. The calculated doping concentration of V is ~8% by counting the number of Mo and V atoms in the image, which is the highest value of V-doping in MoS₂ using V₂O₅ as the precursor. An enlarged HAADF-STEM image (Fig. 3(d)) and its corresponding atomic structure (Fig. 3(e)) show a detailed atom distribution. The intensity line profile marked by the highlighted red frame in Fig. 3(d) is shown in Fig. 3(f). The four stronger intensity peaks are Mo sites and the weak peak in the center is V site. To identify the dark field is V substitute other than atomic defect, EDS spectroscopy (Fig. 3(g)) was

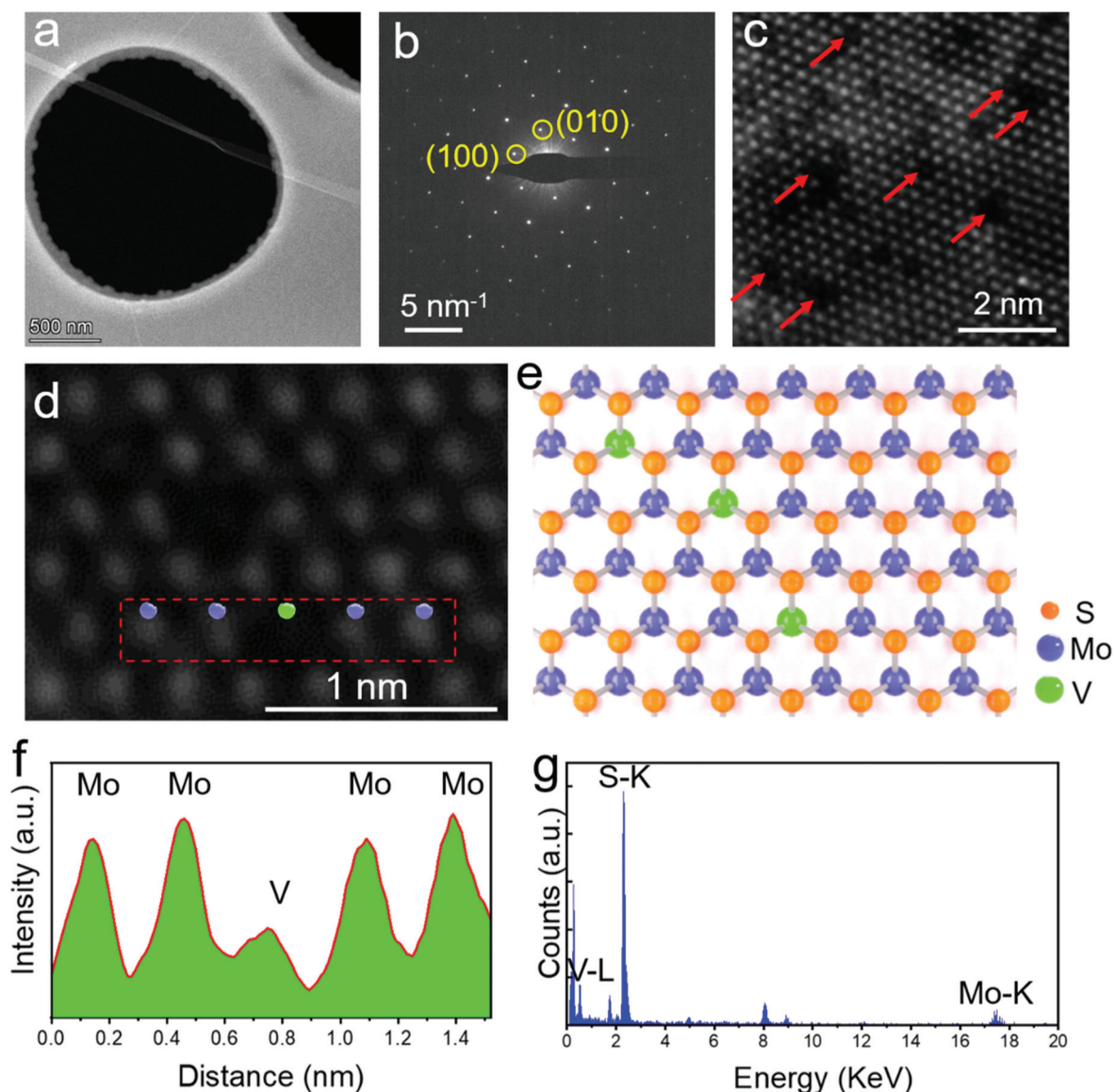


Fig. 3 HADDF-STEM characterization of monolayer V-doped MoS₂. (a) Low magnification image of transferred V-doped MoS₂ nanofilm on grid copper. (b) SAED pattern of H-phase V-doped MoS₂. (c) HADDF-STEM image of V-doped MoS₂. Partial V substitutes for Mo sites are marked by red arrow; (d and e) enlarged STEM image and its corresponding atomic distribution of Mo (blue), V (green) and S (yellow). (f) Z-Contrast intensity profile along the highlighted red dashed box in (d). (g) EDS spectra of V-doped MoS₂ flakes, confirming the coexistence of Mo, V and S elements.

applied to investigate the chemical composition of the sample. All the expected characteristic peaks of S-K at 2.31 keV, Mo-K at 17.48 keV and V-L at 0.51 keV certified their existence.³⁷

Except for most monolayer V-doped MoS₂ nanoflakes, twisted bilayer nanoflakes with different angles were also observed in SiO₂/Si substrate. Here, we measured the SHG response of monolayer, bilayer AA-stacking and AB-stacking pristine MoS₂ and V-doped MoS₂ flakes to verify the crystal character. Fig. 4(a) is a schematic diagram of SHG process of 2D crystal under a pump excitation ($\lambda = 1550$ nm). We measured the SHG intensity of a monolayer flake under various excitation power (2 mW–10 mW). First, all the expected single peak has emerged at 775 nm, which is exactly at the

doubled frequency of the fundamental 1550 nm excitation laser. Second, the function of SHG intensity and excitation power was presented by a double logarithmic plot (Fig. S2†). The slope of 1.9 affirmed the second-order nonlinear optical process. The side view of the atomic structure of bilayer 3R phase and 2H phase MoS₂ was shown in Fig. 4(b). The 3R phase has AA stacking of different layers being in the same orientation with a slight in-plane shift. The S atoms and Mo atoms of different layers do not align on top of each other precisely, ensuring the expected broken inversion symmetry. Comparing with 3R stacking, the 2H phase MoS₂ has AB stacking of bottom layer and top layer being in the opposite orientation with inversion symmetry.^{28,38} Fig. 4(c) and (e) display

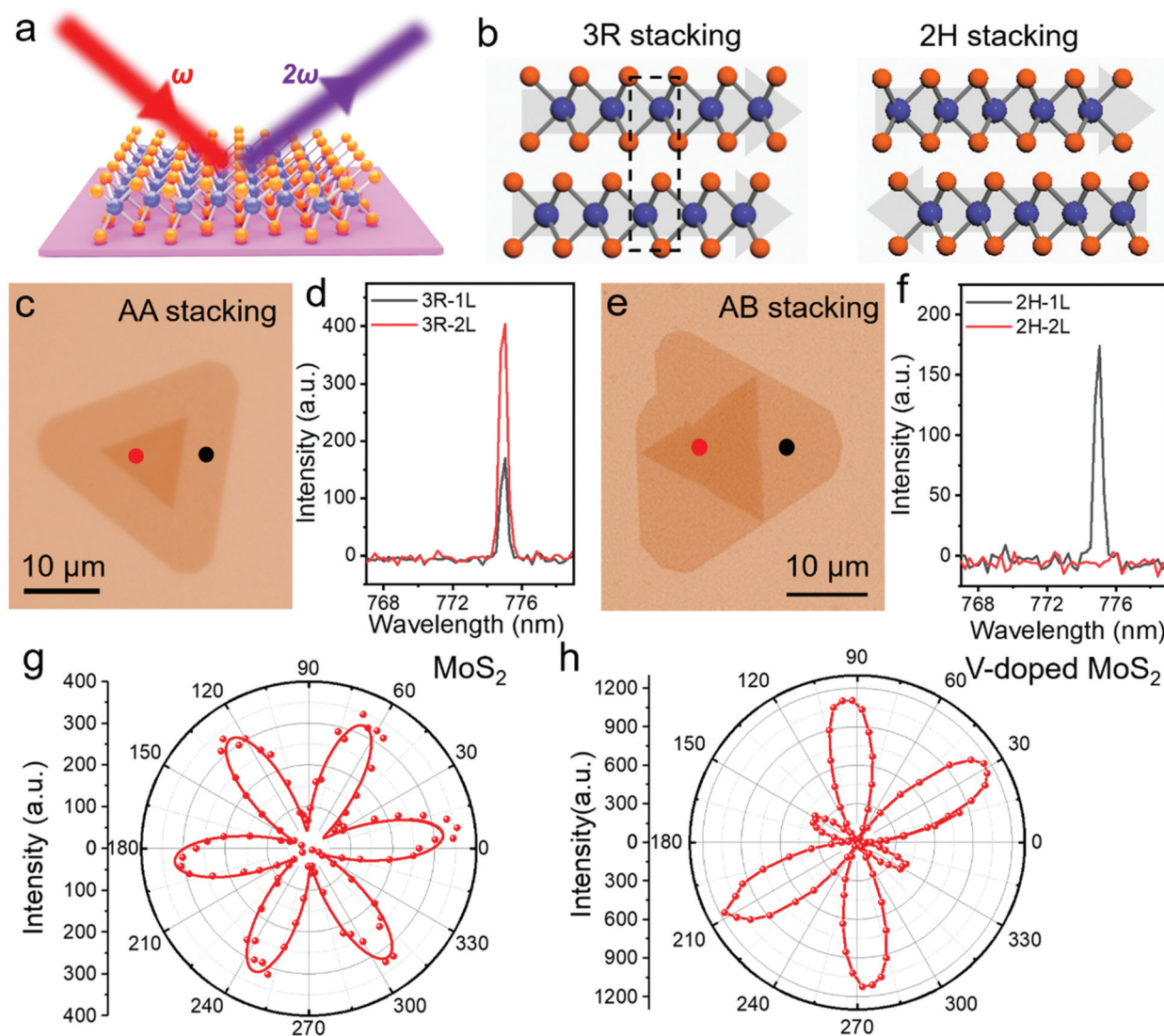


Fig. 4 (a) Schematic diagram of SHG process. (b) Side view of bilayer 3R and 2H phase MoS₂ with different stacking order. (c and e) Optical images of MoS₂ with 3R and 2H stacking. (d and f) Layer number-dependent SHG response 3R and 2H phase V-doped MoS₂. (g and h) Angular-dependent SHG intensity of monolayer MoS₂ and V-doped MoS₂.

the optical microscopy images of pristine AA-stacking and AB-stacking MoS₂ flakes with different stacking orders. As shown in Fig. 4(d), an increased SHG intensity was observed in bilayer AA-stacking MoS₂. As a contrast, no SHG response was observed in bilayer AB-stacking MoS₂ (Fig. 4(f)). Generally, the SHG disappeared in even layers of 2H phase MX₂ (MoS₂, WS₂, WSe₂, etc.) due to the recovered inverse symmetry, which has been widely reported before.^{38,39}

Polarization angle-dependent SHG was carried out to explore the crystalline structural characteristics. The monolayer MoS₂ presents a typical six-fold symmetric pattern (Fig. 4(g)). However, the six-fold pattern of monolayer V-doped MoS₂ was nonuniform shaped, with four folds enlarged compared with the two shrunken folds (Fig. 4(h)). Obviously, the situation changed with the doping of V atom in MoS₂ lattice. We further measured the SHG response of bilayer V-doped

MoS₂ flakes with different stacking orders. Fig. 5(a) and (d) display the optical microscopy images of as-synthesized AA-stacking and AB-stacking V-doped MoS₂ flakes. Their corresponding Raman spectra were given in Fig. S3.† Both AB-stacking and AA-stacking flakes grown on the uniform SiO₂/Si substrate show consistent optical contrast, indicating they possess the same number of layers. AFM measurements were carried out to identify the thickness and layers, and the results of an AA-stacking flake demonstrate the center was bilayer (Fig. S4†). We characterized the SHG mapping (Fig. 5(b)) of a flake with bilayer AA-stacking in the center and monolayer in the edge, highlighted by red dashed box in Fig. 5(a). It is apparent that the center section presented stronger SHG response than the edge according to the image's contrast. The quantitative SHG intensities of 3R stack monolayer and bilayer are presented in Fig. 5(c). To our surprise, the bilayer AB-stacking 2H phase

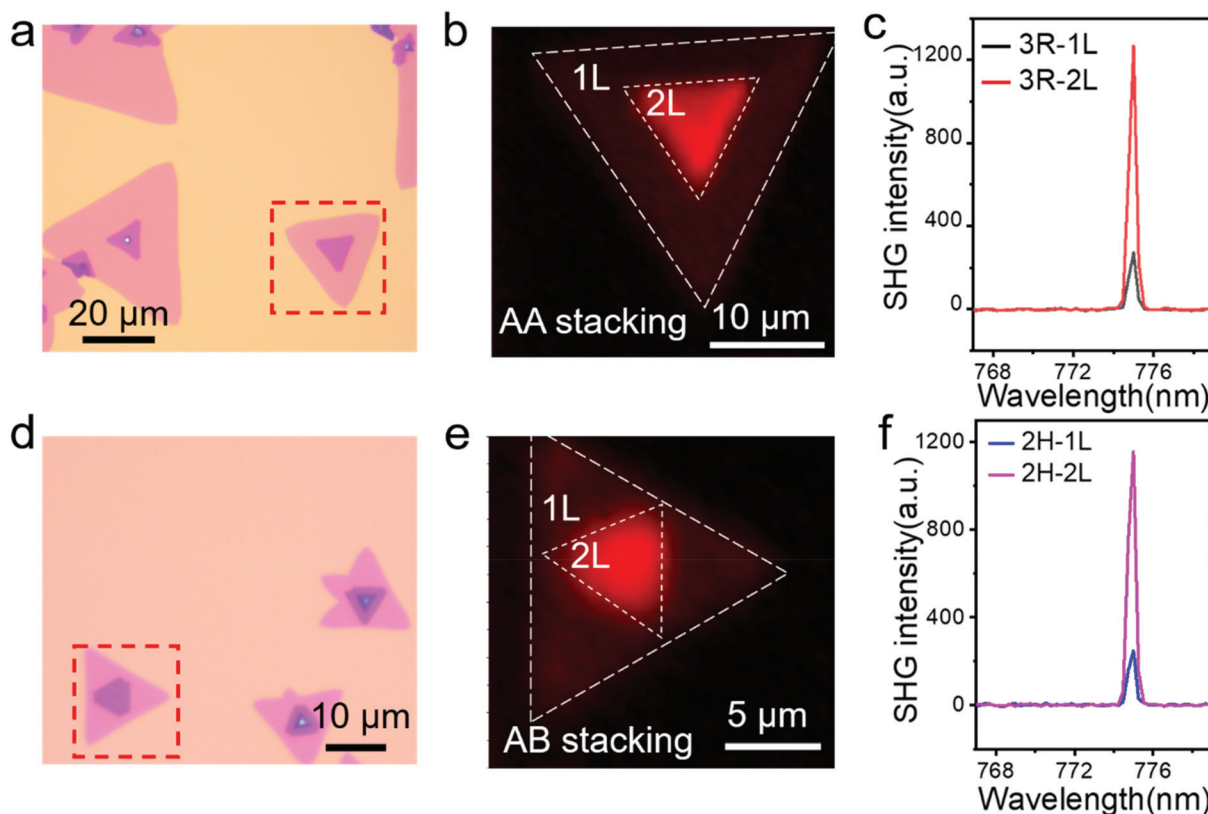


Fig. 5 SHG mapping of bilayer 3R and 2H phase V-doped MoS₂. (a and d) Optical images of V-doped MoS₂ with 3R and 2H stacking. (b and e) SHG intensity mapping of V-doped MoS₂ flakes from (a and d), marked by red dashed boxes. (c and f) Layer number-dependent SHG response 3R and 2H phase V-doped MoS₂.

flakes also generate a strong SHG signal (Fig. 5(e)) and the intensity is almost same as that from 3R bilayer (Fig. 5(f)), demonstrating a breakage of inverse symmetry in 2H phase V-doped MoS₂. This was originally from the random distribution of V-substitute for Mo sites, which causes a slight lattice strain to break the symmetric structure.³³

Conclusions

In summary, we synthesized the V-doped MoS₂ flakes using V₂O₅ precursor *via* CVD method. The crystalline structure was determined by STEM characterization. SHG measurement was further used to investigate the inverse symmetry of 2H and 3R phase V-doped MoS₂ with different stacking orders. The nearly consistent SHG response in bilayer 2H and 3R phase V-doped MoS₂ indicated the broken inversion symmetry regardless of stacking order and layer numbers. The doping 2D crystal provides a profitable strategy to extend the range of atomically thin optical material with second-order nonlinearity.

Conflicts of interest

There is no conflict to declare.

Acknowledgements

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