Facile hydrothermal synthesis of layered 1T′ MoTe₂ nanotubes as robust hydrogen evolution electrocatalysts

Yuxi Lei¹,²*, Xuefeng Xiao¹,², Tianpeng Ma¹,², Weiyin Li¹,², Huan Zhang¹,² and Chao Ma¹,²

¹School of Electrical and Information Engineering, North Minzu University, Yinchuan, China, ²The Key Laboratory of Physics and Photonic Information Functional Materials, North Minzu University, Yinchuan, China

*CORRESPONDENCE
Yuxi Lei,
Lei_yuxi@nun.edu.cn

KEYWORDS
MoTe₂ nanotubes, layered nanostructure, electrocatalysts, hydrogen evolution reaction, hydrothermal method

1 Introduction

Layered transition metal dichalcogenides (TMDs), such as molybdenum ditelluride (MoTe₂), have attracted much attention because of their novel structure-related physicochemical properties. In particular, semi-metallic-phase MoTe₂ (1T′) is considered as a competitive candidate for low-cost electrocatalysts for water splitting. However, there are few reports on the simple hydrothermal synthesis of MoTe₂ nanostructures compared with other layered TMDs. In this study, a facile one-step hydrothermal process was developed for the fabrication of layered MoTe₂, in which uniform nanotubes with a few layers of 1T′ MoTe₂ were fabricated at a lower temperature for the first time. The as-obtained MoTe₂ nanotubes were fully characterized using different techniques, which revealed their structure and indicated the presence of layered 1T′ nanocrystals. The efficient activity of MoTe₂ nanotubes for the electrocatalytic hydrogen evolution reaction (HER) in 0.5 M H₂SO₄ was demonstrated by the small Tafel slope of 54 mV/dec⁻¹ and endurance ability, which is attributed to the abundant active sites and remarkable conductivity of 1T′ MoTe₂ with a few-layer feature. This provides a facile method for the design and construction of efficient layered MoTe₂ based electrocatalysts.
(Wang et al., 2012) and electrochemistry (Chia et al., 2015), and has been well-studied in WS₂, MoS₂ and MoSe₂ based 2D nanostructures in recent years (Bonaccorso et al., 2015; Zhang et al., 2016; Manzeli et al., 2017; Jha et al., 2019; Han et al., 2020; Singh et al., 2022; Singh et al., 2022). In particular, with the demand for effective catalysts for clean energy hydrogen production, TMDs have been widely studied and are considered to be as one of the most outstanding electrocatalysts for the hydrogen evolution reaction (HER) (Jin et al., 2018; Fu et al., 2021; Zhang et al., 2021; Li et al., 2022). TMDs nanostructures possess the capability of tuning catalytic active sites in comparison with their bulk counterparts. Importantly, they represent earth-abundant and low-cost alternatives to noble metal Pt groups (Chandrasekaran et al., 2021; Qiao et al., 2018; Bhat and Nagaraja, 2019; McGlynn et al., 2019). Importantly, they represent earth-abundant and low-cost alternatives to noble metal Pt groups (Chandrasekaran et al., 2021; Qiao et al., 2018; Bhat and Nagaraja, 2019; McGlynn et al., 2019).

Recently, van der Waals (vdW) layered TMDs MoTe₂ has been widely reported for its potential HER performance with enhanced activity (Jinbong et al., 2017; McGlynn et al., 2018; Qiao et al., 2018; Bhat and Nagaraja, 2019; McGlynn et al., 2019; Zazpe et al., 2021). MoTe₂ has two typical stable phases: a hexagonal structure and a distorted monoclinic structure, also known as 2H- and 1T*-MoTe₂, respectively (Manchanda et al., 2020; Singh et al., 2020). 1T*-MoTe₂ exhibits semi-metallic characteristics and high conductivity, which favor efficient electrocatalytic activity. Seok et al. investigated the enhanced HER driving Peierls-type lattice distortion in 1T*-MoTe₂ single crystals prepared using a solid-state method (Jinbong et al., 2017). McGlynn et al. studied the rapid electrochemical activation of 1T*-MoTe₂ nanocrystals for HER, which originates from the adsorption of H onto Te sites on the surface of 1T*-MoTe₂ (McGlynn et al., 2019). Mao et al. (2021) synthesized MoTe₂ nanowires using a mesoporous silica SBA-15 template via a nanocasting method and investigated their HER performances. Besides, Lu et al. (2020) prepared 1T'-MoTe₂ on carbon cloth via direct vapor deposition and demonstrated enhanced and stable HER performance in 1 M H₂SO₄. Zazpe et al. (2021) grew 2D flaky MoTe₂ nanosheets on TiO₂ nanotubes via atomic layer deposition for excellent pollutant photodegradation and HER properties. The aforementioned MoTe₂ catalysts were prepared mainly using a high-temperature solid-state method or chemical vapor deposition. Sun et al. (2016) synthesized 1T'-MoTe₂ nanoflowers using a solution method at a relatively low temperature of 300°C. Liu et al. (2017) prepared 1T'-MoTe₂ nanosheets using a colloidal chemical strategy at 320°C. However, the synthesis of MoTe₂ nanostructures with layered ultrathin sheets using a simple hydrothermal process at low temperatures has rarely been reported.

In this study, we prepared MoTe₂ nanotubes with ultrathin nanosheets by a facile one-step hydrothermal process for the first time. The growth of MoTe₂ nanosheets was derived from a template of Te nanorods, which were reduced rapidly from the TeO₂ precursor via a moderate reductant of ascorbic acid, forming a tubular hierarchical structure with a diameter of 200 nm. The MoTe₂ nanotubes demonstrated efficient and stable HER performance, exhibiting a lower overpotential of ~374 mV at ~100 mA cm⁻² and a Tafel slope of 54 mV dec⁻¹, which are better than those of another typical nanostructure of MoTe₂ nanoparticles. The improved electrocatalytic activity originates from the effective active sites in the nanosheet structure and high conductivity of 1T'-MoTe₂. This facile synthetic strategy is expected to be helpful in the design of novel MoTe₂-based electrocatalysts with 3D hierarchical nanostructures.

2 Experimental section

2.1 Synthesis of MoTe₂ nanostructures

The MoTe₂ nanotubes were synthesized via a one-step hydrothermal method using the following procedure. Tellurium dioxide (8 mmol, TeO₂, 5N, Aladdin Reagent) was added to a 100 ml transparent solution containing 1.2 g ascorbic acid (C₆H₈O₆, 99%, Sinopharm), and 40 ml ethanolamine (C₉H₁₇NO, 99%, Macklin Reagent) was added to the above suspension under constant stirring to form a light-yellow transparent solution. The configured solution was transferred to a 200 ml Hastelloy autoclave and mixed with 4 mmol molybdenum hexacarbonyl [Mo(CO)₆, 98%, Macklin Reagent] that was directly added. The autoclave was heated to 180°C with a rate of 3°C/min and kept 20 h under constant stirring at 200 rpm. After the reaction, the autoclave was naturally cooled to room temperature, and the black precipitate was collected and washed three times with deionized water and ethanol. Finally, the product was desiccated in an oven at 60°C. The experimental process was schematic illustrated by Supplementary Figure S1.

For comparison, another MoTe₂ nanostructure was prepared. MoTe₂ nanoparticles were prepared using a similar procedure but with a different Te source without the addition of ascorbic acid. Te powder (2 mmol, Te, 5N, Aladdin Reagent) and 1 mmol Mo(CO)₆ were directly added to a 200 ml Hastelloy autoclave containing 110 ml deionized water under constant stirring, and then 10 ml hydrazine hydrate aqueous solution (H₂N₂·xH₂O, 85%, Sinopharm) was dropped into the above suspension. The autoclave was heated to 200°C at a rate of 3°C/min and maintained at 200 rpm for 20 h with constant stirring. The second step was the same as that used for the MoTe₂ nanotubes.

2.2 Characterizations

The phase and crystal structures of the as-prepared samples were analyzed using X-ray powder diffraction
(XRD) patterns measured on an X-ray diffractometer (Rigaku SmartlabSE) with Cu Kα radiation (λ = 1.5406 Å) at a scanning rate of 8°/min in the 2θ range of 10°C–80°C. The morphologies of the samples were observed using field-emission scanning electron microscopy (FESEM, ZEISS SIGMA 500) at an accelerating voltage of 10 kV. The crystalline characteristics of the MoTe2 nanostructure, dark-field scanning transmission electron microscopy (STEM), and energy-dispersive X-ray spectroscopy (EDX) mapping were performed using field-emission transmission electron microscopy (FETEM, JEOL JEM 2100F, 200 kV). X-ray photoelectron spectroscopy (XPS) was performed on an ESCALAB Xi+ X-ray photoelectron spectrometer (Thermo Fisher Scientific Ltd.) using Al-Kα radiation. The Raman spectrum was collected by Raman spectroscopy (Horiba Ltd.) using a 532 nm laser. The Brunauer-Emmett-Teller (BET) method was used to calculate the c surface areas of the samples via N2 adsorption-desorption isotherms using a Micromeritics ASAP 2460 instrument.

2.3 Electrochemical measurements

All electrochemical measurements were carried out using an electrochemical workstation (CHI650E, Shanghai CH Instruments) with a three-electrode system in 0.5 M H2SO4 (unless otherwise stated). A graphite rod and Ag/AgCl were used as counter and reference electrodes, respectively. For the working electrode, the samples (5 mg) were added to 1 ml of isopropanol containing 0.25% NaCl under ultrasonication for 1 h, and 20 μl of the ink was coated onto the L-type glassy carbon electrode (5 mm in diameter) and dried naturally. All the potentials were converted to the reversible hydrogen electrode (RHE) using the following equation: $E_{\text{RHE}} = E_{\text{SHE}} + 0.059 \log \left( \frac{c_{H^+}}{c_{H^2O}} \right) + 0.242$, where $E_{\text{RHE}}$ is the standard potential of the reference electrode versus the standard hydrogen electrode (SHE). Linear sweep voltammetry (LSV) curves were obtained in the selected potential range at a scan rate of 5 mV/s without iR compensation. The Tafel slope was fitted using the equation $\eta = a + b \log j$, where $\eta$ is the overpotential, $a$ is a constant and $b$ is the Tafel slope. Cyclic voltammetry (CV) data were collected at scan rates of 10 mV/s, 20 mV/s, 40 mV/s, 60 mV/s, 80 mV/s, and 100 mV/s in selected potential ranges to estimate the electrochemical active surface area (ECSA) and double-layer capacitance ($C_{dl}$). Electrochemical impedance spectra (EIS) were obtained in the frequency range of 0.1 Hz–106 Hz with an amplitude of 10 mV. Chronoamperometry was performed for 20 h at a potential of −0.2 V (vs. RHE).

3 Results and discussion

3.1 Morphology and phase

The phase and morphology of the as-synthesized MoTe2 powder obtained by a facile ascorbic acid (VC)-assisted hydrothermal method were characterized by XRD, SEM, and TEM. Figure 1A shows the collected XRD pattern, the diffraction peaks at 12°C, 29°C, 32°C, and 51°C match the (0 0 2), (2 0 1), (1 1 2), and (3 1 0) planes of 1T’ MoTe2 (JCPDS No. 71–2,157), and the low intensity and broadness of the diffraction peak (0 0 2) are related to a few sheets of the MoTe2 nanostructure and poor crystallinity by the hydrothermal route, which is similar to that of a few MoS2 nanosheets (Lei et al., 2017b; Liu et al., 2019). No other impurity peaks were detected, confirming that MoTe2 nanocrystals were synthesized. A typical TEM image (Figure 1B) indicates distinctly uniform tubular nanosheets on the microstructure of the products. A typical SEM image in Figure 1C and the corresponding high-resolution SEM image (Figure 1D) indicate that the nanotubes were decorated with ultrathin sheets on their surface, and the nanosheets interlaced with each other, forming a hierarchical structure. An individual nanotube with a diameter of approximately 200 nm and a length of several micrometers. Uniform MoTe2 nanotubes were synthesized using a facile hydrothermal method.

The microstructural components and crystalline characteristics of the as-synthesized MoTe2 nanotubes were further investigated using HRTEM analysis. As shown in Figure 2A, an individual nanotube manifested a distinct tubular structure with a diameter of 200 nm and length of approximately 2 μm, on which thin sheets were uniformly distributed. The clear lattice stripes with a spacing of 0.71 nm are assigned to the (0 0 2) plane of the monoclinic MoTe2 crystal from the HRTEM image in Figure 2B, which is slightly larger than that of the standard MoTe2 crystal; the curved lattice fringe represents the characteristics of the layered TMDs nanosheets obtained by the wet chemical method (Lei et al., 2018; Wang et al., 2021, Zarpe et al., 2021). STEM-EDX mapping performed on a selected individual MoTe2 nanotube revealed a uniform and homogeneous elemental distribution of Te and Mo in the integrated MoTe2 nanotube (Figure 2C). The STEM-EDX spectrum and analytical results obtained for the entire nanotube are shown in Figure 2D, and are close to the stoichiometric proportion of the MoTe2 compound.

The growth of MoTe2 nanotubes using a facile hydrothermal method was explored using controlled experiments under different conditions. The additive agent VC plays a key role in the formation of uniform tubular MoTe2 nanosheets. VC can reduce TeO2 to Te nanorods in the precursor solution (Xi et al., 2006; Lei et al., 2017a) and act as a template/seed to grow MoTe2 nanosheets. Uniform MoTe2 nanotubes were obtained when an appropriate amount of VC was used (Figure 1; Supplementary Figure S2D); otherwise, irregular Te impurities and non-uniform
MoTe₂ microtubes with rough surfaces were obtained (Supplementary Figures S2,3).

To investigate the facile hydrothermal route for the synthesis of MoTe₂ nanoparticles, using Te powder as the Te source and hydrazine hydrate as the reducing reagent under different reaction conditions. Figure 3A shows the XRD pattern of the as-prepared sample, where the broad hump diffraction peaks around 12°C, 32°C, and 51°C are ascribed to the (0 0 2), (1 1 2), and (3 1 0) planes of 1T′ MoTe₂ (JCPDS No. 71–2157), respectively, which is consistent with the characteristics of MoTe₂ nanotubes. Notably, a small amount of incompletely reacted Te impurities is detected. From the low-resolution SEM image (Figure 3B), it can be seen that irregular particles aggregated on a large rod (Te seed). The high-resolution SEM images in Figures 3C,D indicate that the size of the fine particles is approximately dozens of nanometers. In addition to the nanoparticle morphology, there are large quantities of amorphous sheets, forming a porous structure. The EDX spectrum and analytical results (insets of Figures 3C,D) collected from the corresponding objects are close to the stoichiometric ratio of MoTe₂. Then, only irregular MoTe₂ nanoparticles were obtained using the as-designed process but with uniform layered nanosheets.

3.2 XPS analysis

The elemental compositions and surface chemical states of the as-prepared MoTe₂ nanocrystals were investigated using XPS. In the high-resolution Mo 3d spectrum, as illustrated in Figure 4A, the peaks at 572.8 eV and 583.2 eV correspond to Te 3d₃/₂ and Te 3d₅/₂, respectively, which are ascribed to the typical peaks of Te-Mo bonds (Guo and Wang, 2018; Li et al., 2020). The other two satellite peaks at 576.3 eV and 586.7 eV originate from the Te-O bond on sample exposed to ambient atmosphere, which is commonly detected in nanostructured tellurides (Luxa et al., 2017; Anantharaj et al., 2018). As for the Mo 3d (Figure 4B), the peaks at 227.8 and 231.1 eV should be ascribed to Mo 3d₅/₂ and Mo 3d₃/₂, similarly two observed peaks at 232.5 eV and 235.6 eV related to Mo-O bonds due to dynamic surface states. These findings are consistent with those of previous studies (Zazpe et al., 2021; Wang et al., 2021). Figure 4C shows the Raman spectrum of the as-synthesized MoTe₂ nanotubes under a 532 nm laser, where the peaks situated at 115 cm⁻¹, 135 cm⁻¹, and 158 cm⁻¹ correspond to the A₅ modes. Notably, the absence of a peak at approximately 230 cm⁻¹ which is the phase-specific Raman peak of 2H-MoTe₂, indicating 1T′-MoTe₂ was obtained using a facile solution-phase synthetic method.

The nitrogen adsorption–desorption isotherm was employed to calculate the specific surface area of the samples using the BET method, as shown in Supplementary Figure S4, which can be identified as type IV with hysteresis loops, confirming typical nanocrystalline structures. The specific area of the MoTe₂ nanoparticles was approximately 87 m²/g, which is significantly larger than that of the MoTe₂ nanotubes, possibly because of its fine particles and mesoporous structure.
3.3 Electrochemical performance

The electrocatalytic activity of the as-prepared MoTe₂ nanocrystals for the hydrogen evolution reaction (HER) was investigated using electrochemical measurements and compared with that of commercial Pt/C. Figure 5A shows the linear sweep voltammetry (LSV) curves in a 0.5 M H₂SO₄ electrolyte. The as-prepared MoTe₂ nanotubes exhibited lower overpotentials of $-317$ mV and $-349$ mV at current densities of $-10$ mA·cm⁻² and $-100$ mA·cm⁻², respectively, which were much smaller than those of MoTe₂ nanoparticles and MoTe₂ nanowires (Mao et al., 2021). The MoTe₂ nanotubes obtained with different amounts of VC (except for 0 VC) showed similar catalytic performance (Supplementary Figure S5). It can be seen from the LSV curve that the current density increased rapidly with increasing potential in the MoTe₂ nanotubes. The HER performance of MoTe₂ related electrocatalysts reported in recent literature is listed in Supplementary Table S1.

Tafel plots and their slopes were obtained using the Tafel equation for the kinetic investigation of HER, as shown in Figure 5B. The Tafel slope of the MoTe₂ nanotubes was $54$ mV·dec⁻¹, which was even lower than that of the commercial Pt/C catalyst ($61$ mV·dec⁻¹). A slight decrease in the Tafel slope ($67$ mV·dec⁻¹) was observed for MoTe₂ nanoparticles. The HER kinetics of MoTe₂ nanotubes can be inferred from the Volmer-Heyrovsky mechanism (Watzele et al., 2018). The catalytic kinetics were further revealed by Nyquist plots from the electrochemical impedance spectra (EIS) shown in Figure 5C. The results indicate that the MoTe₂ nanotubes exhibit a small charge-transfer resistance ($R_{ct}$) in comparison with that of the MoTe₂ nanoparticles, which is consistent with the LSV results. The electrochemical surface area (ECSA) of the MoTe₂ nanotubes was further evaluated from the double-layer capacitances ($C_{dl}$) measured by CV curves at different scan rates in the potential range of $0.55$ V–$0.65$ V (Figure 5D). The $C_{dl}$ of MoTe₂ nanotubes is linear fitting to $2.62$ mF·cm⁻¹.

The endurance and long-term stability of the electrocatalysts were investigated. Figure 6A depicts the HER performance of the two MoTe₂ nanocrystals exposed to a $1.0$ M H₂SO₄ electrolyte. The MoTe₂ nanotubes exhibited much lower overpotentials at a certain current density than MoTe₂ nanoparticles. The potential slightly increased with the current density in the MoTe₂
nanotubes, indicating an endurable stability of the HER activity even in a strong alkaline electrolyte. Figure 6B shows a curve of current density with time (i - t) under a fixed overpotential of −0.2 V by using chronopotentiometry measurements. It can be seen that the current density was distinctly increased in the first two hours, which is similar with recent reports about the gradual electrochemical activation of MoTe₂ nanocrystals (McGlynn et al., 2019; Zazpe et al., 2021). Subsequently, the current density remained stable over time. Cyclic voltammetry measurements were conducted to assess stability. As shown in Figure 6C, the polarization curve of the MoTe₂ nanotubes reveals a decrease in the overpotential (~30 mV at a current density of −10 mA·cm⁻²) after 100 cycles and a negligible change after 20 h of chronopotentiometry measurements, which is consistent with the i-t result. SEM and XPS were performed to evaluate the structural and chemical stability of the MoTe₂ nanotubes.
Supplementary Figure S6). The SEM images reveal no obvious changes in the morphological characteristics of the tubular nanosheets. No obvious change in the position of the peaks was observed in the XPS high-resolution Te 3d and Mo 3d spectra. Notably, there is an obvious decrease in the intensity of the Te-O peaks, suggesting that the electrochemical activation would originated from the Te sites of the layered MoTe2 nanosheets in the HER process, which is in line with previous reports (Zazpe et al., 2021; McGlynn et al., 2019).

4 Conclusion

In this work, two typical MoTe2 nanocrystals were prepared by low-temperature hydrothermal methods. Uniform MoTe2 nanotubes with hierarchical few-layer nanosheets were synthesized for the first time. An appropriate amount of VC contributed to the formation of tubular nanosheet structures. The as-synthesized MoTe2 nanotubes exhibited efficient performance in the hydrogen evolution reaction with a Tafel
slope of 54 mV·dec⁻¹. A relatively gradual electrochemical activation and durable HER stability were also observed. The uniform nanosheet structure improved the electrochemical active sites and facilitated HER. The development of sheet-like MoTe₂ nanocrystals using facile synthetic methods is expected to be helpful for the promising applications of layered MoTe₂.

Data availability statement

The original contributions presented in the study are included in the article/Supplementary Material, further inquiries can be directed to the corresponding author.

Author contributions

Material preparation, data collection, and analysis were performed as previously described by Lei et al. YL wrote the first draft of the manuscript. XX supervised and provided funding for this study. TM, HZ, and CM participated in the discussion of results. All authors have proofread and refined the manuscript.

Funding

This research was supported by the Natural Science Foundation of Ningxia (Grant Nos. 2018AAC03118, 2020AAC03241 and 2020AAC03239), the Scientific Research Project of Ningxia Higher Education Institutions (No. NGY2018-144), the National Natural Science Foundation of China (Grant No. 61965001), the Ningxia Province Key Research and Development Program (Grant No. 2021BEE03005) and the Ningxia New Solid Electronic Materials and Devices Research and Development Innovation Team (No. 2020CXTDLX12).

Acknowledgments

The authors would like to thank the Key Laboratory of North Minzu University (Physics and Photoelectric Information Functional Materials).

Conflict of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Publisher’s note

All claims expressed in this article are solely those of the authors and do not necessarily represent those of their affiliated organizations, or those of the publisher, the editors and the reviewers. Any product that may be evaluated in this article, or claim that may be made by its manufacturer, is not guaranteed or endorsed by the publisher.

Supplementary Material

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fchem.2022.1005782/full#supplementary-material

References

Anantharaj, S., Karthick, K., and Kundu, S. (2018). NiTe₂ nanowire outperforms Pt/C in high-rate hydrogen evolution at extreme pH conditions. Inorg. Chem. 57, 3082–3096. doi:10.1021/acs.inorgchem.7b02947

Bhat, K. S., and Nagaraja, H. S. (2019). Performance evaluation of molybdenum dichalcogenide (MoX₂; X= S, Se, Te) nanostructures for hydrogen evolution reaction. Int. J. Hydrogen Energy 44, 17878–17886. doi:10.1016/j.ijhydene.2019.05.179

Bonacorso, F., Colombo, L., Yu, G., Stoller, M., Touzini, V., Ferrari, A. C., et al. (2015). 2D materials. Graphene, related two-dimensional crystals, and hybrid systems for energy conversion and storage. Science 347, 1246501. doi:10.1126/science.1246501

Chandrasekaran, S., Yao, L., Deng, L., Bowen, C., Zhang, Y., Chen, S., et al. (2019). Recent advances in metal sulfides. From controlled fabrication to electrocatalytic, photocatalytic and photoelectrochemical water splitting and beyond. Chem. Soc. Rev. 48, 4178–4280. doi:10.1039/c8cs00664d

Chia, X., Eng, A. Y. S., Ambrosi, A., Tan, S. M., and Pumera, M. (2015). Electrochemistry of nanostructured layered transition-metal dichalcogenides. Chem. Rev. 115, 11941–11966. doi:10.1021/acs.chemrev.5b00287

Fu, Q., Han, J., Wang, X., Xu, P., Yao, T., Zhong, J., et al. (2021). 2D transition metal dichalcogenides: Design, modulation, and challenges in electrocatalysis. Adv. Mat. 33, 1907818. doi:10.1002/adma.201907818

Guo, Y., Park, T., Yi, J. W., Henzie, J., Kim, J., Wang, Z., et al. (2019). Nanosarchitectonics for transition-metal-sulfide-based electrolystas for water splitting. Adv. Mat. 31, 1807134. doi:10.1002/adma.201807134

Han, W., Liu, Z., Pan, Y., Guo, G., Zou, J., Xia, Y., et al. (2020). Designing champion nanostructures of tungsten dichalcogenides for electrocatalytic hydrogen evolution. Adv. Mat. 32, 2002584. doi:10.1002/adma.202002584

Jha, R. K., D’Costa, J. V., Salduja, N., and Bhat, N. (2019). MoSe₂ nanoflakes based chemiresistive sensors for ppb-level hydrogen sulfide gas detection. Sensors Actuators B Chem. 297, 126667. doi:10.1016/j.snb.2019.126667

Ji, J., and Choi, J. H. (2022). Recent progress in 2D hybrid heterostructures from transition metal dichalcogenides and organic layers: Properties and applications in energy and optoelectronics fields. Nanoscale 14, 10648–10689. doi:10.1039/ d2nr01358d

Jin, H., Guo, C., Liu, X., Liu, J., Vasileff, A., Jiao, Y., et al. (2018). Emerging two-dimensional nanomaterials for electrocatalysis. Chem. Rev. 118, 6337–6408. doi:10.1021/acs.chemrev.7b00689
Enhanced photocatalytic properties of novel 3D MoS$_2$/Na$_0.9$Mg$_{0.45}$Ti$_{3.55}$O$_8$ nanorods and diversification CoTe nanorods for water-splitting.

Lei, Y., Wang, J., Zhou, J., Hassan, Q. U., and Wang, J. (2017b). One-step synthesis of NiTe$_2$ nanorods coated with few-layers MoS$_2$ for enhancing photocatalytic activity. Nanotechnology 28, 495602. doi:10.1088/1361-6528/aa94ae

Manchanda, P., Kumar, P., and Dev, P. (2020). Thickness dependence of hydrogen-induced phase transition in MoTe$_2$. Phys. Rev. B 101, 144104. doi:10.1103/physrevb.101.144104

Zazpe, R., Sopha, H., Charvot, J., Krumpolec, R., Rodriguez-Pereira, J., Zazpe, R., Sopha, H., Charvot, J., Krumpolec, R., Rodriguez-Pereira, J., Mglynn, J. C., Cascallana-Mat As, I., Fraser, J. P., Roger, I., Mcallister, J., Miras, E. K., et al. (2019). The rapid electrochemical activation of MoTe$_2$ for the hydrogen evolution reaction. Nat. Commun. 10, 4916. doi:10.1038/s41467-019-12831-0

Qian, X., Liu, J., Fu, L., and Li, J. (2014). Quantum spin Hall effect in two-dimensional transition metal dichalcogenides. Science 346, 1344–1347. doi:10.1126/science.1256815

Qiao, H., Huang, Z., Liu, S., Liu, Y., Li, J., and Qi, X. (2018). Liquid-exfoliated molybdenum telluride nanosheets with superior electrocatalytic hydrogen evolution performances. Ceram. Int. 44, 21205–21209. doi:10.1016/j.ceramint.2018.08.166

Singh, S., Deb, J., Sarkar, U., and Sharma, S. (2022a). MoSe$_2$/multilayer carbon nanotube composite for ammonia sensing in natural humid environment. J. Hazard Mater. 435, 128821. doi:10.1016/j.jhazmat.2022.128821

Singh, S., Deb, J., Singh, J. V., Sarkar, U., and Sharma, S. (2022b). Highly selective ethyl mercaptan sensing using a MoTe$_2$/SnO$_2$ composite at room temperature. ACS Appl. Interfaces 14, 23916–23927. doi:10.1021/acs.ajc.2c01512

Mcglynn, J. C., Dankworth, T., Kienle, L., Bandeira, N. A. G., Fraser, J. P., Gibson, E. K., et al. (2019). The rapid electrochemical activation of MoTe$_2$ for the hydrogen evolution reaction. Nat. Commun. 10, 4916. doi:10.1038/s41467-019-12831-0

Khan, M. A., and Leuenberger, M. N. (2018). Optoelectronics with single layer group-VIB transition metal dichalcogenides. Nanophotonics 7, 1589–1600. doi:10.1515/nanoph-2018-0041

Lei, Y.-X., Miao, N.-X., Zhou, J.-P., Hassan, Q. U., and Wang, J.-Z. (2017a). Novel magnetic properties of CoTe nanorods and diversified CoTe$_2$ nanostuctures obtained at different Na$_2$O$_2$ concentrations. Sci. Technol. Adv. Mater. 18, 325–333. doi:10.1088/1468-9971.2017.13.17218

Lei, Y., Wang, J., Zhou, J., Guo, Z., and Hassan, Q. U. (2018). Fabrication and enhanced photocatalytic properties of novel 3D Mo$_3$S$_8$/Na$_3$Mg$_{10}$Ti$_{31}$O$_{78}$ heterostructures. Appl. Surf. Sci. 427, 733–741. doi:10.1016/j.apsusc.2017.08.169

Ni$_x$CoTe$_2$/graphene ternary heterostructures. Nanotechnology 28, 495602. doi:10.1088/1361-6528/aa94ae

Li, C., Liu, M., Ding, H., He, L., Wang, E., Wang, B., et al. (2020). A lightly Fe-doped (Ni$_{30}$S$_{70}$/Mo$_{50}$)carbon nanotube hybrid electrocatalyst film with laser-drilled micropores for stabilized overall water splitting and pH-universal hydrogen evolution reaction. J. Mat. Chem. A 8, 17527–17536. doi:10.1039/d0ta04586a

Li, X.-Y., Zhu, S.-J., Wang, Y.-L., Lian, T., Yang, X.-Y., Ye, C.-F., et al. (2022). Synergistic regulation of S vacancy of Mo$_y$S$_{3-y}$based materials for highly efficient electrocatalytic hydrogen evolution. Front. Chem. 10, 915408. doi:10.3389/fchem.2022.915408

Li, Y., Liu, C., Geng, F., Liu, Y., Yang, S., Zhang, S., et al. (2019). 2H- and 1T-mixed phase few-layer MoS$_2$ as a superior to Pt co-catalyst coated on TiO$_2$ nanorod arrays for photocatalytic hydrogen evolution. Appl. Catal. B Environ. 241, 236–245. doi:10.1016/j.apcatb.2018.09.040

Liu, D., Ren, X., Ren, L., Xu, W., Liu, S., Liu, Y., et al. (2020). Direct vapor deposition growth of 1T MoTe$_2$ on carbon cloth for electrocatalytic hydrogen evolution reactions. ACS Appl. Energy Mater. 3, 3212–3219. doi:10.1021/acsaem.9b01589

Liu, C., Duan, J., Shen, C., Wang, J., Zhang, J., et al. (2017). Layered transition-metal ditellurides in electrocatalytic water splitting. J. Mat. Chem. A 5, 2070–2092. doi:10.1039/c5ta01855d

Liu, M., Wang, Z., Liu, J., Wei, G., Du, J., Li, Y., et al. (2017). Synthesis of few-layer 1T-MoTe$_2$ ultrathin nanosheets for high-performance pseudocapacitors. J. Mat. Chem. A 5, 1035–1042. doi:10.1039/c7ta0260e

Liu, Y., Li, Y., Peng, F., Liu, Y., Yang, S., Zhang, S., et al. (2019). 2H- and 1T-mixed phase few-layer MoS$_2$ as a superior to Pt co-catalyst coated on TiO$_2$ nanorod arrays for photocatalytic hydrogen evolution. Appl. Catal. B Environ. 241, 236–245. doi:10.1016/j.apcatb.2018.09.040

Ma, J., Zhou, L., Li, Y., Tan, Y., Chai, K., Shu, Y., et al. (2021). Synthesis of MoTe$_2$ nanowire as an efficient hydrogen evolution reaction material. Mater. Lett. 290, 124971. doi:10.1016/j.matlet.2021.124971

Mas-Ballester, R., Gomez-Narváez, C., Gomez-Herrero, J., and Zamora, F. (2011). 2D materials: To graphene and beyond. Nanoscale 3, 20–30. doi:10.1039/c0nr00323a

Meyglyn, J. C., Cascallana-Mat As, I., Fraser, J. P., Roger, I., Mcallister, J., Miras, E. K., et al. (2018). Molybdenum ditelluride rendered into an efficient and stable photocatalyst for the hydrogen evolution reaction by polymorphic control. Energy Technol. 6, 345–350. doi:10.1002/ente.201700489

Meyglyn, J. C., Dankworth, T., Kienle, L., Bandeira, N. A. G., Fraser, J. P., Gibson, E. K., et al. (2019). The rapid electrochemical activation of MoTe$_2$ for the hydrogen evolution reaction. Nat. Commun. 10, 4916. doi:10.1038/s41467-019-12831-0

Zarpe, R., Scala, H., Chavot, J., Kumpoles, R., Rodriguez-Pereira, J., Michalíčka, I., et al. (2021). 2D MoTe$_2$ nanosheets by atomic layer deposition: Excellent photo- electrocatalytic properties. Appl. Mater. Today 23, 101017. doi:10.1016/j.apmt.2021.101017

Zhang, G., Liu, H., Lu, J., and Li, J. (2016). Two-dimensional layered Mo$_3$S$_2$: Rational design, properties and electrochemical applications. Energy Environ. Sci. 9, 1190–1209. doi:10.1039/c6ee03761a

Zhang, X., Jia, F., and Song, S. (2021). Recent advances in structural engineering of molybdenum disulfide for electrocatalytic hydrogen evolution reaction. Chem. Eng. J. 405, 127013. doi:10.1016/j.cej.2020.127013