ROYAL SOCIETY
OF CHEMISTRY

COMMUNICATION

Check for updates

Defect-engineered room temperature negative differential resistance in monolayer MoS$_2$ transistors†

Wen-Hao Chang,[a] Chun-I Lu,[a] Tilo H. Yang, [ID][a] Shu-Ting Yang,[a]
Kristan Bryan Simbulan, [ID][ab] Chih-Pin Lin,[c] Shang-Hsien Hsieh,[d]
Jyun-Hong Chen,[e] Kai-Shin Li,[e] Chia-Hao Chen, [ID][df] Tuo-Hung Hou,[c]
Ting-Hua Lu[a] and Yann-Wen Lan [ID]*[a]

The negative differential resistance (NDR) effect has been widely investigated for the development of various electronic devices. Apart from traditional semiconductor-based devices, two-dimensional (2D) transition metal dichalcogenide (TMD)-based field-effect transistors (FETs) have also recently exhibited NDR behavior in several of their heterostructures. However, to observe NDR in the form of monolayer MoS$_2$, theoretical prediction has revealed that the material should be more profoundly affected by sulfur (S) vacancy defects. In this work, monolayer MoS$_2$ FETs with a specific amount of S-vacancy defects are fabricated using three approaches, namely chemical treatment (KOH solution), physical treatment (electron beam bombardment), and as-grown MoS$_2$. Based on systematic studies on the correlation of the S-vacancies with both the device's electron transport characteristics and spectroscopic analysis, the NDR has been clearly observed in the defect-engineered monolayer MoS$_2$ FETs with an S-vacancy (V_S) amount of $\sim 5 \pm 0.5\%$. Consequently, stable NDR behavior can be observed at room temperature, and its peak-to-valley ratio can also be effectively modulated *via* the gate electric field and light intensity. Through these results, it is envisioned that more electronic applications based on defect-engineered layered TMDs will emerge in the near future.

New concepts

In this research, defect-engineered monolayer MoS$_2$ field-effect transistors are fabricated using physical and chemical treatments. Based on the results of electrical characteristics and spectroscopic analysis to determine the amount of sulfur vacancy defects, the negative differential resistance (NDR) can be reliably obtained in the monolayer MoS$_2$ FETs with a sulfur vacancy amount of $\sim 5 \pm 0.5\%$. Furthermore, this NDR behavior is stable at room temperature, and its peak-to-valley ratio can be effectively modulated *via* the gate electric field and light intensity.

Introduction

Since the discovery of the Esaki tunneling diode,[1,2] the negative differential resistance (NDR) effect has been widely applied in

[a] *Department of Physics, National Taiwan Normal University, Taipei 116, Taiwan.*
   E-mail: ywlan@ntnu.edu.tw
[b] *Department of Mathematics and Physics, University of Santo Tomas, Manila 1008, Philippines*
[c] *Department of Electronics Engineering & Institute of Electronics, National Yang Ming Chiao Tung University, Hsinchu 300, Taiwan*
[d] *National Synchrotron Radiation Research Center, Hsinchu 300, Taiwan*
[e] *Taiwan Semiconductor Research Institute, National Applied Research Laboratories, Hsinchu 300, Taiwan*
[f] *Department of Electrophysics, National Yang Ming Chiao Tung University, Hsinchu 300, Taiwan*
† Electronic supplementary information (ESI) available. See DOI: https://doi.org/10.1039/d2nh00396a

designing various devices such as logic switches,[3] computer memory,[4–6] and high-frequency oscillators.[7,8] In addition to typical semiconductor-based tunneling diodes, ultra-thin field-effect transistors (FETs), fabricated with low dimensional material channels, such as carbon nanotubes,[9,10] graphene[11–13] or graphite,[14] have also been reported to exhibit NDR behavior. Likewise, vertically-integrated two-dimensional (2D) layered van der Waals (vdW) heterostructures that involve the recently well-studied 2D transition metal dichalcogenides (TMDs), among others, possess the NDR effect and have been documented in several material arrangements, including MoS$_2$-MoTe$_2$,[15] MoS$_2$-WSe$_2$,[16,17] and MoS$_2$-WSe$_2$-graphene.[18] However, to simplify the fabrication process and to have more practical applications in the future, it is better to possess the NDR effect at room temperature using a single type of material *via* a simple method. According to theoretical calculations in monolayer 2D materials, the NDR effect is predicted to exist in the presence of, and be more profoundly affected by, either deformations or defects (such as S-vacancies in MoS$_2$) rather than under defect-free conditions.[19] Therefore, defect engineering in 2D materials using several techniques, including molecular modulation,[20] oxygen plasma,[21] ion irradiation,[22–24] or electron bombardment,[25,26] have been proposed to achieve NDR behavior.

In this work, initially, two post-treatment approaches are used, namely soaking in alkaline solution (KOH) and electron beam bombardment, to determine the amount of defects in

monolayer MoS_2 that would cause NDR behavior. To evaluate the correlation of the percentage of S-vacancy defects with the electrical characteristics and spectroscopic analysis, we perform electrical measurements, and carry out Raman, photoluminescence (PL), and X-ray photoemission spectroscopy (XPS) analysis on monolayer MoS_2 that contains defects. Based on the results of the abovementioned approaches, the NDR is observed in the defect-engineered MoS_2 with an S-vacancy V_S amount of ~ 4.5–5% at room temperature. The amount of S-vacancies is determined from the atomic S/Mo ratio in the experimental XPS data.[24] Accordingly, using the direct growth method with a specific amount of S-vacancies, the as-grown monolayer MoS_2 FETs clearly show stable NDR behavior that can be actively modulated *via* the gate electric field and light intensity. Density-functional theory (DFT) simulations were also conducted to confirm the occurrence of the NDR phenomenon at the said amount of S-vacancies in monolayer MoS_2. Therein, the band structure will have additional defect states formed in the bandgap, which will assist the electron tunneling. With the NDR effect successfully observed in FETs formed using defect-containing monolayer MoS_2, given a specific amount of S-vacancy defects, we believe that defect engineering on 2D materials could attract more attention in exploring diverse applications for quantum technologies in the future.

Results and discussion

Fig. 1a presents a schematic of defect-engineered monolayer MoS_2, where the red dashed circles indicate S-vacancies. The defect-containing monolayer MoS_2 with a specific amount of

S-vacancies can be created using the three approaches highlighted in this work, in particularly, (a) chemical treatment with KOH, (b) electron beam bombardment, and (c) control of the S/Mo ratio during the chemical vapor deposition (CVD) synthesis process. In order to figure out the amount of S-vacancies that can cause NDR behavior, we first study the PL and Raman spectra (excitation with a 532 nm laser) of the KOH-treated monolayer MoS_2 at various soaking times as shown, respectively, in Fig. 1b and c. The interesting dependencies of the optical properties of KOH-treated MoS_2 on the soaking time are manifested by the exponentially decreasing behavior of the PL intensity of the A excitons (blue squares) and the blueshift in the peak energy position (red dots) as the soaking time is increased, as shown in Fig. 1d.[22,24] The blueshifting of the A exciton peak can be attributed to the defect-induced strain on the monolayer MoS_2.[27] To compare the Raman shifts, the spectra are normalized to the intensity of the A_{1g} peak, as presented in Fig. 1c. Consistent with previously reported results, the separation between the E_{2g} and A_{1g} modes of pristine monolayer MoS_2 is ~ 18.76 cm^{-1} at room temperature, as presented in Fig. S1a of the ESI.† [28] Fig. 1e demonstrates the Raman shift as a function of the KOH soaking time, and shows that the E_{2g} peak (in-plane mode) redshifts and the A_{1g} peak (out-of-plane mode) blueshifts. A comparison of the overall spectra, including the silicon peak for all conditions, is given to address concerns about the influence of the measurement setup. It turns out that the Si Raman peak measured in our system remains at the same position, as demonstrated in Fig. S1b of the ESI.† The redshifting of the E_{2g} peak and the blueshifting of the A_{1g} peak can also be attributed to defect-induced strain.[27,29,30] These phenomena are different from the
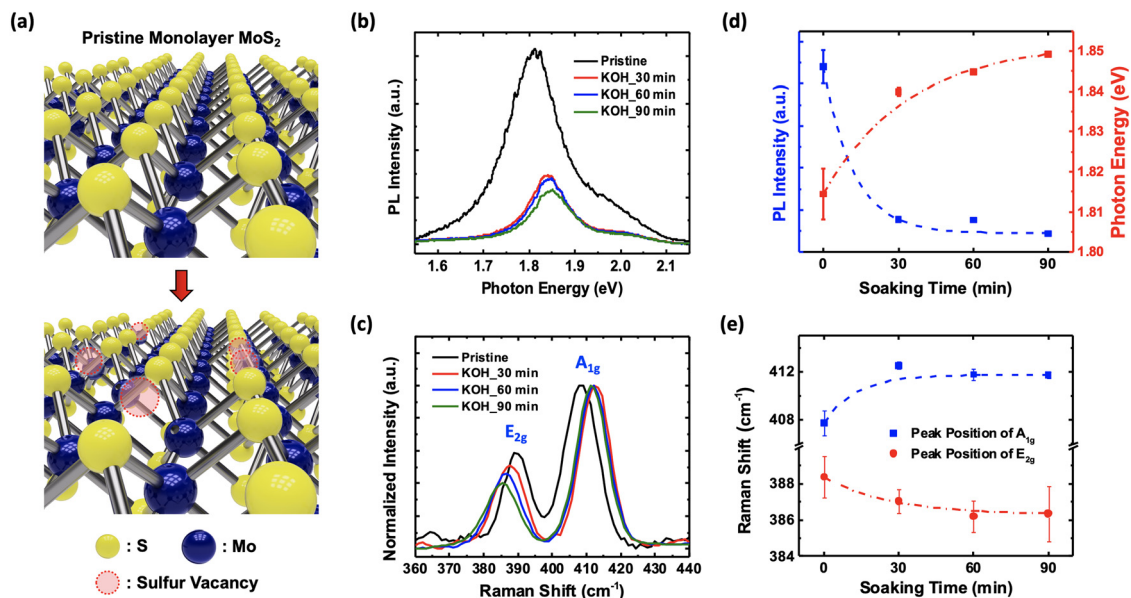

Fig. 1 (a) Schematic illustrations of the pristine and defect-containing monolayer MoS_2 that contains S-vacancies indicated by red dashed circles. (b and c) Photoluminescence (PL; b) and Raman (c) spectra of monolayer MoS_2 treated with KOH for varying soaking times. (d) PL intensity and photon energy of the A exciton as a function of soaking time, showing a decrease in intensity (blue squares) and a blue shift (red dots), respectively, with increasing soaking time. (e) Raman shift as a function of soaking time, displaying the redshifting of the E_{2g} mode and the blueshifting of the A_{1g} mode. The error bars for three samples are shown in (d) and (e).

electron doping effect, which may result in an almost unaffected E_{2g} peak and a redshifted A_{1g} peak.[31] Similar defects that affect the Raman shifts have also been discussed in monolayer MoS_2 treated using ion[23,24] or electron[26] irradiation.

To evaluate the percentage of S-vacancies in the KOH-treated MoS_2 at various soaking times, we exhibit the evolution of the XPS spectrum in the Mo 3d (left panel) and S 2p (right panel) regions, recorded from pristine MoS_2 (bottom panel) to the 90 minute-treated (top panel) samples as demonstrated in Fig. 2a. The additional peaks of Mo $3d_{3/2}$ and Mo $3d_{5/2}$ that appear at slightly lower binding energy values for the 90 minute treatment (top panel) correspond to the defect-containing MoS_2 with S-vacancies.[32–34] For the semiquantitative analysis of the S/Mo ratio, the XPS spectra are fitted with Lorentzian functions to acquire the area underneath the curves and the S/Mo ratio is normalized to 2, assuming that the S-vacancy in pristine MoS_2 is negligible. By following the equation proposed by Bertolazzi *et al.*,[24] the S-vacancy (V_S) amount can be expressed as $V_S = 1 - r/2$, where r is the S/Mo ratio. Fig. 2b exhibits a decrease in the normalized S/Mo ratio (left y-axis) and an increase in V_S (right y-axis) with increasing soaking time. With these observed relationships, it can be said that, in Fig. 2c, the peak positions of both Mo $3d_{5/2}$ and S $2p_{3/2}$ shift to lower binding energies with an increasing amount of S-vacancies, resulting in metallic-like behavior.[22] Accordingly, to further verify whether the aforementioned defect-induced phenomena also exist in the other two defect-engineered techniques, we also performed XPS experiments on the electron-beam-bombarded MoS_2. After testing several electron beam bombardment conditions (as displayed in Fig. S2, ESI†), we irradiated monolayer MoS_2 at varying dosages under a fixed electron acceleration voltage

($V_{acc} = 15$ kV) and current ($I = 0.49$ nA) as presented in Fig. S3 of the ESI.† The XPS results of the electron-beam-bombarded MoS_2 are consistent with the results observed in the chemically-treated MoS_2, wherein a decreasing normalized S/Mo ratio is accompanied by a lowering binding energy as the e-beam dosage is increased (more details can be seen in Fig. S4, ESI†).

Based on the aforementioned results that include PL, Raman and XPS, defect-engineered MoS_2 FETs with an appropriate amount of S-vacancies can be controlled and adopted to observe NDR behavior using either chemical or physical treatments. Fig. 3a–c present the I_d–V_g characteristics of three MoS_2 FET devices, which are chemically treated with KOH in 0, 15 and 30 minute durations, respectively, corresponding to S-vacancies from 0 to 4.5%. The optical microscope (OM) images of the measured devices are shown in the insets. The I_d–V_g curves progressively become less gate dependent when the duration of the treatment is lengthened, indicating a transition from semi-conducting to metallic-like properties. This result can be attributed to the defect states induced by the S-vacancies, which may shift the Fermi level toward the conduction band edge, resulting in metallic-like behavior.[35,36] Accordingly, Fig. 3d–f display the corresponding I_d–V_g characteristics, respectively, where the NDR effect can be seen clearly in Fig. 3f. It turns out that with a specific amount of S-vacancies (~4.5%), similar results can also be observed in another 30 minute KOH-treated device presented in Fig. S5 (ESI†). The inset shown in Fig. 3f displays the shifting of the NDR peak positions, which is represented by the correlating bias voltage *versus* the applied gate voltage. It shows the linear dependence of the NDR peak positions to the gate voltage in the range of −60 V to +40 V. The gate voltage could possibly shift the Fermi level and align it
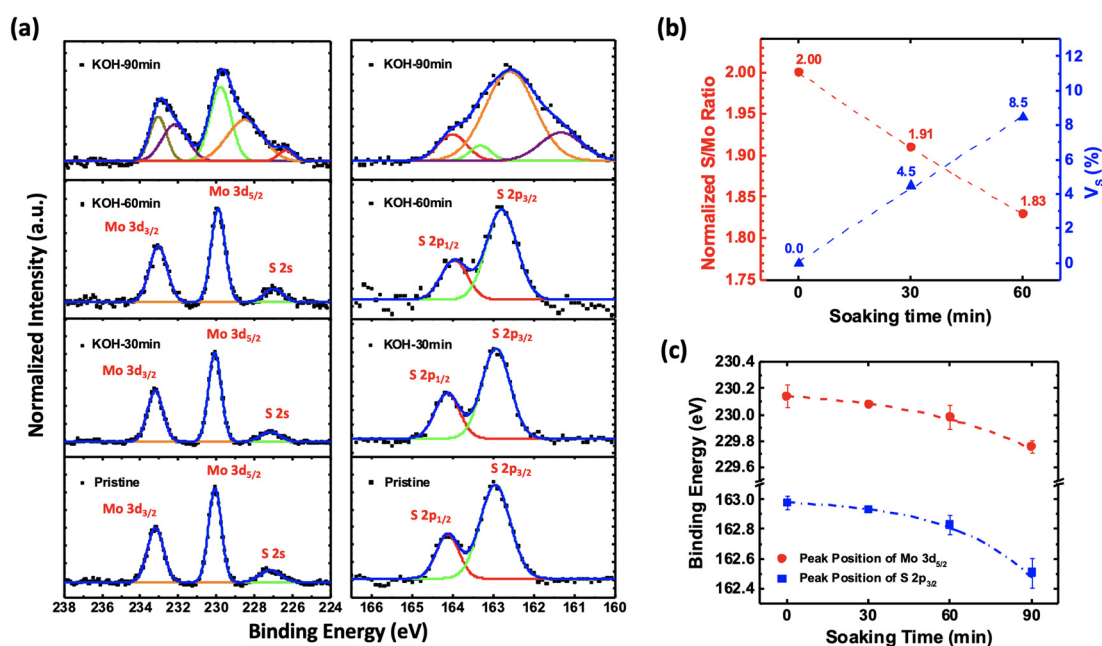

Fig. 2 (a) X-ray photoemission spectroscopy (XPS) of Mo 3d (left panel) and S 2p (right panel) at various KOH soaking times. (b) Normalized S/Mo ratio and S–vacancy (V_S) as a function of soaking time showing an increasing percentage of V_S with increasing soaking time. (c) Peak positions of Mo $3d_{5/2}$ and S $2p_{3/2}$ shift to lower binding energy values with increasing soaking time.
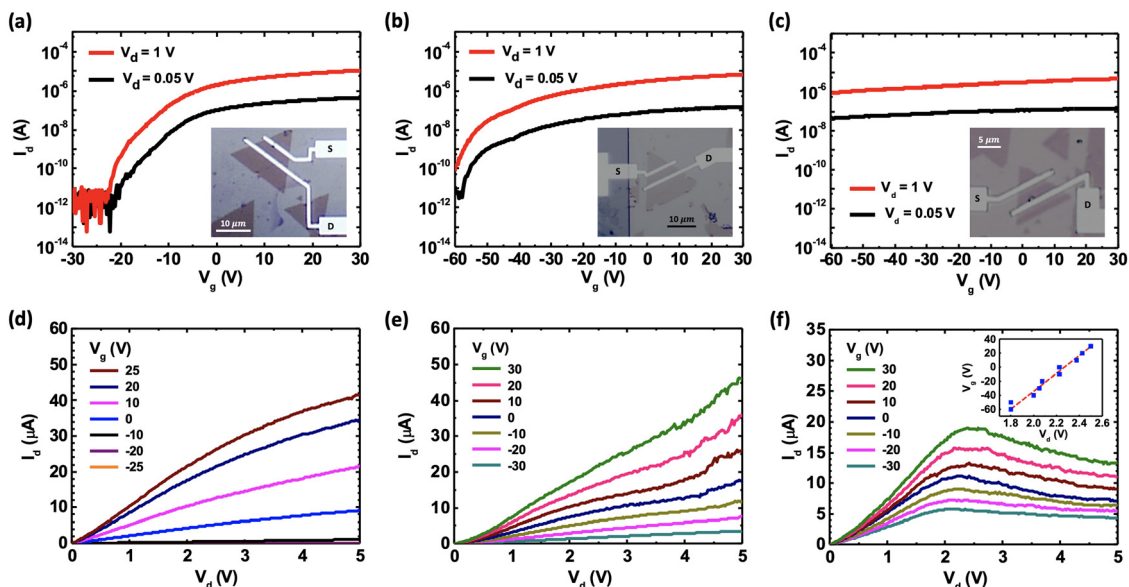
Fig. 3 (a–c) I_d–V_g characteristics of the MoS$_2$ FET devices showing an evolution from semiconducting to metallic-like behavior with increasing KOH soaking time (0, 15 and 30 minutes, respectively). The insets show the OM images of the respective measured devices. (d–f) Corresponding I_d–V_g characteristics demonstrating typical n-type semiconducting behavior. The NDR effect appears in the metallic-like MoS$_2$ device as shown in (f), in which the inset displays the peak positions, represented by their corresponding bias voltage *versus* gate voltage showing a linear correlation.

with the defect/quantum states, causing more tunneling electrons, while the increasing bias voltage may generate more diffusing holes. Consequently, the competition between electrons and holes could result in a shifting of the NDR peak position with increasing V_g.[14,17,37] To prove that the S-vacancies in defect-containing MoS$_2$ could result in defect states that can further affect the device's electron transport, we also performed electrical measurements on devices bombarded with different electron beam dosages and observed the resulting NDR behavior at a dosage of $\sim 5 \times 10^5$ μC cm^{-2}, which corresponds to V_S $\approx 5\%$ (determined from Fig. S4b, ESI†), as demonstrated in Fig. S6 (ESI†). However, the electrical signal becomes noisy due to the severe damage caused by the physical electron beam bombardment.

In order to minimize the noisy signal caused by the damage caused by the electron beam bombardment, using as-grown monolayer MoS$_2$ with specific amounts of S-vacancies is an appropriate method to achieve stable NDR behavior. We thus synthesized such MoS$_2$ flakes with a particular range of S/Mo ratios using the CVD method under low S supply conditions.[38] Accordingly, the as-grown monolayer MoS$_2$ samples with different amounts of S-vacancies, from $\sim 1\%$ to $\sim 6\%$, along with their corresponding XPS results are shown in Fig. S7 (ESI†). Based on the aforementioned results of NDR behavior under the specified S-vacancy conditions, we intentionally fabricated such a MoS$_2$ device with $V_S \approx 5\%$, which was precisely determined through its XPS data shown in Fig. S8 (ESI†). In this device, the observation of NDR can be seen in the I_d–V_g curves at different gate voltages without light illumination, as shown in Fig. 4a, while a clearer NDR performance is perceived at a fixed gate voltage $V_g = +10$ V with different light illumination power values as presented in Fig. 4b. A larger peak to valley

ratio is observed since more majority carriers can be induced in the n-type monolayer MoS$_2$ at either $V_g = +10$ V or at higher light illumination powers. More details of the device's electrical characteristics at different light power values and gate voltages can be seen in Fig. S9 (ESI†). Additionally, another as-grown MoS$_2$ FET with $V_S \approx 4.5\%$, identically confirmed by XPS data, also exhibits NDR behavior, as presented in Fig. S10 (ESI†). Consequently, by growing to achieve a certain range of S-vacancies ($\sim 5 \pm 0.5\%$), the NDR behavior can be reproduced in another MoS$_2$ device, as displayed in Fig. S11 (ESI†). Since the competition between the field-dependent carrier density and the drift velocity determines whether or not the NDR can be observed, we speculate that the defect states at the interface between MoS$_2$ and the electrodes affect the contact resistance, which could further dominate the tunneling mechanism.[14,39] However, the hugely different current level appearing in the two types of device, namely the KOH-treated MoS$_2$ and the as-grown MoS$_2$, which correspond to μA and pA current levels, respectively, is probably due to either the channel resistance or contact issues. It is obvious that the devices with a shorter channel length of ~ 2 μm have a higher current level, as shown in Fig. 3a–c, than that with a longer channel length of ~ 65 μm, as presented in Fig. 4a. Furthermore, according to ref. 40 and our recent experimental results, using semimetallic bismuth (Bi) instead of titanium (Ti) as the buffer layer of the electrodes can achieve almost ohmic contact behavior. Therefore, we believe that the devices designed and fabricated with the shorter channel length and Bi-MoS$_2$ contacts can help to reach a higher current level for approaching the requirement of real applications.

In order to figure out the possible mechanism, the density of states (DOS) of monolayer MoS$_2$ with S-vacancies is evaluated
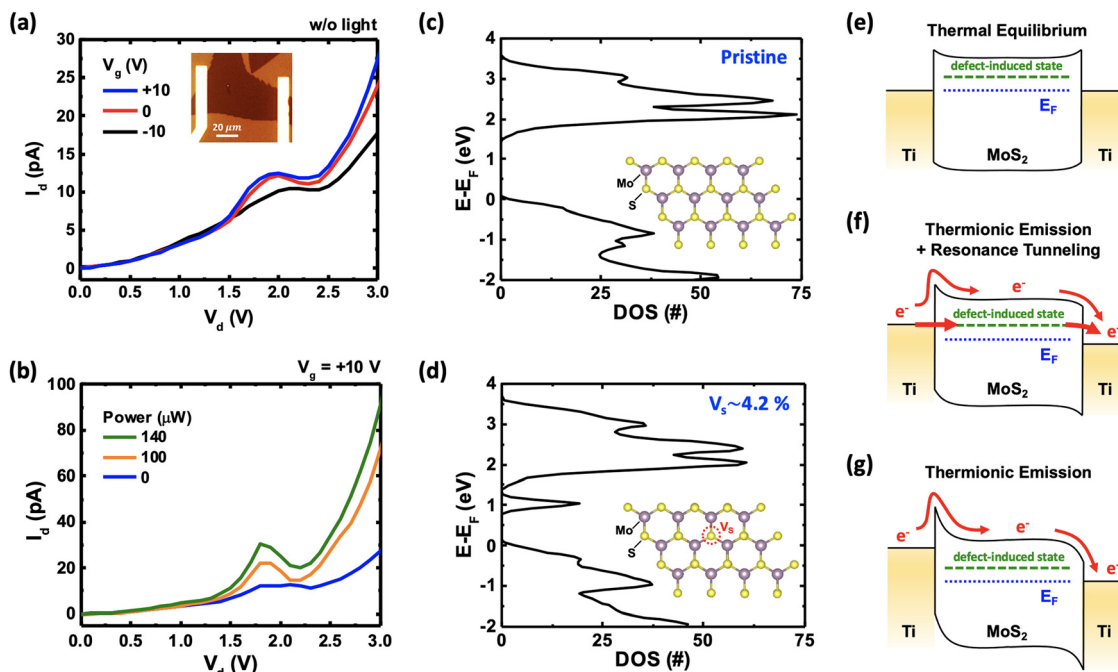
Fig. 4 (a) I_d–V_d curves at different gate voltages without light illumination. The inset presents the OM image of the measured device. (b) I_d–V_d curves illuminated with different light power values at a fixed gate voltage V_g = +10 V. (c and d) Density of states (DOS) of pristine monolayer MoS_2 and that with $V_S \approx 4.2\%$, respectively. (e–g) Schematics of band diagrams for different stages of the NDR mechanism.

and a density-functional theory (DFT) simulation is performed using the Vienna *ab initio* simulation package (VASP).[41,42] Fig. 4c and d present the simulated DOS of monolayer MoS_2 for the pristine sample and that with $V_S \approx 4.2\%$, respectively, in which an additional defect-induced DOS can clearly be seen in the band gap as illustrated in Fig. 4d. A similar defect-induced DOS is obtained for V_S from ~4 to 6%, and it could be responsible for the observation of the NDR behavior. More details about the simulated DOS of MoS_2 from pristine condition up to that with $V_S \approx 12.6\%$ are presented in Fig. S12 (ESI†), wherein the energy bands move toward to the Fermi level and the number of defect states increases in the bandgap with increasing S-vacancies.[30] Consequently, the increasing number of defect states helps more electrons to tunnel through the barrier causing the defect-containing MoS_2 FETs to exhibit metallic-like properties. To clearly understand the potential mechanism for NDR behavior, the band diagrams are proposed as demonstrated in Fig. 4e–g. Fig. 4e is the band diagram at thermal equilibrium conditions for MoS_2 with an appropriate number of S-vacancies contacted with the metal of Ti, in which an additional defect-induced state is located in the band gap. When a small bias voltage (V_d) is applied to the device with a suitable gate voltage (V_g), the defect-induced state will be aligned to the Fermi level of Ti. Subsequently, the current (I_d) will increase due to both the thermionic emission and resonance tunneling mechanisms as presented in Fig. 4f. However, if the bias voltage (V_d) is increased slightly further, the defect-induced state is not aligned with the Fermi level of Ti, and only thermionic emission contributes to the electron transport as demonstrated in Fig. 4g, leading to the decreased current

forming an NDR peak as shown in Fig. 4a and b. Therefore, with an appropriate amount of S-vacancies, the NDR behavior can be theoretically predicted and experimentally observed.

Conclusions

The NDR effect is observed in monolayer MoS_2 FETs *via* defect engineering approaches, including chemical treatment with KOH, electron beam bombardment, and as-grown MoS_2 with a specific amount of S-vacancies. Based on systematic studies for the correlation of the S-vacancies with both the electrical characteristics and spectroscopic analysis (PL, Raman and XPS), the NDR effect has been clearly observed at room temperature in defect-engineered monolayer MoS_2 FETs with $V_S \approx 5 \pm 0.5\%$. Moreover, stable NDR behavior can also be effectively modulated *via* the gate electric field and light intensity. We believe that these results may provide an alternative route to develop more electronic applications based on defect-containing monolayer TMDs in the future.

Methods

Material synthesis

MoS_2 was synthesized on a sapphire substrate *via* the chemical vapor deposition (CVD) method at 190 °C. Initially, molybdenum trioxide (MoO_3) powder was placed in a quartz boat located at the heating zone center of the furnace, and the sapphire substrate was placed next to the quartz boat. Sulfur (S) powder was prepared in another quartz boat at the upstream side of the furnace. An argon

gas flow was then used to bring both the S and MoO_3 vapors towards the sapphire substrate.

Device fabrication

After synthesizing the monolayer MoS_2, PMMA was coated on the MoS_2 film at a spin speed of 4000 rpm for 55 seconds. The sapphire/MoS_2/PMMA substrate was subsequently immersed in an ammonia solution for one to two hours to separate the MoS_2/PMMA film from the sapphire substrate.[43,44] After transferring the MoS_2/PMMA film onto the Si/SiO_2 substrate, the PMMA protection layer was removed using acetone. The sample was then immersed in an N-methyl-2-pyrrolidone solution for wet cleaning to remove residues or contamination on the MoS_2 surface.[45] A new layer of PMMA was coated on the Si/SiO_2/MoS_2 substrate at a spin speed of 4000 rpm for 55 seconds for the e-beam lithography process. Finally, the Ti (20 nm)/Au (50 nm) electrodes were deposited using a thermal evaporator.

XPS experiments

The XPS data for KOH-treated MoS_2 were recorded using a Thermo Fisher Scientific Theta Probe at the Taiwan Semiconductor Research Institute. Monochromatic anode X-rays (1486.6 eV photon energy) were focused down to ~ 15 μm. Additionally, the XPS data for e-beam bombarded MoS_2 and as-grown MoS_2 were obtained using the scanning photoelectron microscopy (SPEM) endstation at beamline 09A1 of the National Synchrotron Radiation Research Center's Taiwan Light Source. Monochromatic soft X-rays (400 eV photon energy) were focused down to ~ 200 nm using Fresnel-zone-plate-based focusing optics. By raster-scanning the sample relative to the focused beam and collecting the photoelectrons, SPEM can produce the surface elemental distribution image. Based on the SPEM image, we moved the focused beam to a specific location of interest to record the high-resolution photoelectron spectra of the microscopic area. All samples were annealed at about 450 K for 8 hours prior to SPEM measurement. The measurements were recorded under ultra-high vacuum (UHV) conditions near 300 K.

Density functional theory simulations

The density of states (DOS) for monolayer MoS_2 with different S-vacancy concentrations was investigated through density-functional theory (DFT) simulations using the Vienna *ab initio* simulation package (VASP). A $3 \times 4 \times 1$ supercell of MoS_2 with a vacuum layer of 18 Å was used as a good approximation for the isolated 2D slab. An $11 \times 11 \times 1$ *k*-points and a 500 eV cut-off energy were selected in all supercells. The relaxed lattice constant of monolayer MoS_2 without sulfur vacancies is 3.18 Å. The DOS of MoS_2 with sulfur vacancies ranging from 4.2% to 12.6% was then calculated by removing one to three sulfur atoms in the $3 \times 4 \times 1$ supercell.

Author contributions

In this work, W.-H. Chang, T. H. Yang and S.-T. Yang performed the device fabrication and data analysis. The XPS experiments

Conflicts of interest

There are no conflicts of interest to declare.

Acknowledgements

References

1 L. Esaki, *Phys. Rev.*, 1958, **109**, 603–604.

2 L. Esaki and R. Tsu, *IBM J. Res. Dev.*, 1970, **14**, 61–65.

3 M. Herzig, M. Weiher, A. Ascoli, R. Tetzlaff, T. Mikolajick and S. Slesazeck, *J. Phys. D: Appl. Phys.*, 2019, **52**, 325104.

4 S. Chen, P. B. Griffin and J. D. Plummer, *IEEE Trans. Electron Devices*, 2009, **56**, 634–640.

5 S. Wang, A. Pan, C. Grezes, P. K. Amiri, K. L. Wang, C. O. Chui and P. Gupta, *IEEE Trans. Electron Devices*, 2017, **64**, 4084–4090.

6 K.-H. Kim, H.-Y. Park, J. Shim, G. Shin, M. Andreev, J. Koo, G. Yoo, K. Jung, K. Heo, Y. Lee, H.-Y. Yu, K. R. Kim, J. H. Cho, S. Lee and J.-H. Park, *Nanoscale Horiz.*, 2020, **5**, 654–662.

7 R. J. Hwu, A. Djuandi and S. C. Lee, *IEEE Trans. Microwave Theory Tech.*, 1993, **41**, 890–893.

8 V. Ulansky, A. Raza and H. Oun, *Electronics*, 2019, **8**, 409.

9 F. Léonard and J. Tersoff, *Phys. Rev. Lett.*, 2000, **85**, 4767–4770.

10 M. Rinkiö, A. Johansson, V. Kotimäki and P. Törmä, *ACS Nano*, 2010, **4**, 3356–3362.

11 Y. Wu, D. B. Farmer, W. Zhu, S.-J. Han, C. D. Dimitrakopoulos, A. A. Bol, P. Avouris and Y.-M. Lin, *ACS Nano*, 2012, **6**, 2610–2616.

12 P. Sharma, L. S. Bernard, A. Bazigos, A. Magrez and A. M. Ionescu, *ACS Nano*, 2015, **9**, 620–625.

13 S. Rathi, I. Lee, M. Kang, D. Lim, Y. Lee, S. Yamacli, H.-I. Joh, S. Kim, S.-W. Kim, S. J. Yun, S. Choi and G.-H. Kim, *Sci. Rep.*, 2018, **8**, 7144.

14 S.-T. Yang, T. H. Yang, C.-I. Lu, W.-H. Chang, K. B. Simbulan and Y.-W. Lan, *Carbon*, 2021, **176**, 440–445.

15 N. T. Duong, S. Bang, S. M. Lee, D. X. Dang, D. H. Kuem, J. Lee, M. S. Jeong and S. C. Lim, *Nanoscale*, 2018, **10**, 12322–12329.

16 T. Roy, M. Tosun, X. Cao, H. Fang, D.-H. Lien, P. Zhao, Y.-Z. Chen, Y.-L. Chueh, J. Guo and A. Javey, *ACS Nano*, 2015, **9**, 2071–2079.

17 C.-Y. Lin, X. Zhu, S.-H. Tsai, S.-P. Tsai, S. Lei, Y. Shi, L.-J. Li, S.-J. Huang, W.-F. Wu, W.-K. Yeh, Y.-K. Su, K. L. Wang and Y.-W. Lan, *ACS Nano*, 2017, **11**, 11015–11023.

18 Y.-C. Lin, R. K. Ghosh, R. Addou, N. Lu, S. M. Eichfeld, H. Zhu, M.-Y. Li, X. Peng, M. J. Kim, L.-J. Li, R. M. Wallace, S. Datta and J. A. Robinson, *Nat. Commun.*, 2015, **6**, 7311.

19 A. Sengupta and S. Mahapatra, *J. Appl. Phys.*, 2013, **114**, 194513.

20 K. Cho, M. Min, T.-Y. Kim, H. Jeong, J. Pak, J.-K. Kim, J. Jang, S. J. Yun, Y. H. Lee, W.-K. Hong and T. Lee, *ACS Nano*, 2015, **9**, 8044–8053.

21 M. R. Islam, N. Kang, U. Bhanu, H. P. Paudel, M. Erementchouk, L. Tetard, M. N. Leuenberger and S. I. Khondaker, *Nanoscale*, 2014, **6**, 10033–10039.

22 Q. Ma, P. M. Odenthal, J. Mann, D. Le, C. S. Wang, Y. Zhu, T. Chen, D. Sun, K. Yamaguchi, T. Tran, M. Wurch, J. L. McKinley, J. Wyrick, K. Magnone, T. F. Heinz, T. S. Rahman, R. Kawakami and L. Bartels, *J. Phys.: Condens. Matter*, 2013, **25**, 252201.

23 S. Mignuzzi, A. J. Pollard, N. Bonini, B. Brennan, I. S. Gilmore, M. A. Pimenta, D. Richards and D. Roy, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2015, **91**, 195411.

24 S. Bertolazzi, S. Bonacchi, G. Nan, A. Pershin, D. Beljonne and P. Samorì, *Adv. Mater.*, 2017, **29**, 1606760.

25 H.-P. Komsa, J. Kotakoski, S. Kurasch, O. Lehtinen, U. Kaiser and A. V. Krasheninnikov, *Phys. Rev. Lett.*, 2012, **109**, 035503.

26 W. M. Parkin, A. Balan, L. Liang, P. M. Das, M. Lamparski, C. H. Naylor, J. A. Rodríguez-Manzo, A. T. C. Johnson, V. Meunier and M. Drndić, *ACS Nano*, 2016, **10**, 4134–4142.

27 E. Scalise, M. Houssa, G. Pourtois, V. V. Afanas'ev and A. Stesmans, *Phys. E*, 2014, **56**, 416–421.

28 H. Li, Q. Zhang, C. C. R. Yap, B. K. Tay, T. H. T. Edwin, A. Olivier and D. Baillargeat, *Adv. Funct. Mater.*, 2012, **22**, 1385–1390.

29 C. Rice, R. J. Young, R. Zan, U. Bangert, D. Wolverson, T. Georgiou, R. Jalil and K. S. Novoselov, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2013, **87**, 081307.

30 H. Li, C. Tsai, A. L. Koh, L. Cai, A. W. Contryman, A. H. Fragapane, J. Zhao, H. S. Han, H. C. Manoharan, F. Abild-Pedersen, J. K. Nørskov and X. Zheng, *Nat. Mater.*, 2016, **15**, 48–53.

31 B. Chakraborty, A. Bera, D. V. S. Muthu, S. Bhowmick, U. V. Waghmare and A. K. Sood, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2012, **85**, 161403.

32 I. S. Kim, V. K. Sangwan, D. Jariwala, J. D. Wood, S. Park, K.-S. Chen, F. Shi, F. Ruiz-Zepeda, A. Ponce, M. Jose-Yacaman, V. P. Dravid, T. J. Marks, M. C. Hersam and L. J. Lauhon, *ACS Nano*, 2014, **8**, 10551–10558.

33 K. C. Kwon, C. Kim, Q. V. Le, S. Gim, J.-M. Jeon, J. Y. Ham, J.-L. Lee, H. W. Jang and S. Y. Kim, *ACS Nano*, 2015, **9**, 4146–4155.

34 D. M. Sim, M. Kim, S. Yim, M.-J. Choi, J. Choi, S. Yoo and Y. S. Jung, *ACS Nano*, 2015, **9**, 12115–12123.

35 S. McDonnell, R. Addou, C. Buie, R. M. Wallace and C. L. Hinkle, *ACS Nano*, 2014, **8**, 2880–2888.

36 C.-P. Lin, P.-C. Chen, J.-H. Huang, C.-T. Lin, D. Wang, W.-T. Lin, C.-C. Cheng, C.-J. Su, Y.-W. Lan and T.-H. Hou, *ACS Appl. Electron. Mater.*, 2019, **1**, 684–691.

37 G. P. Lansbergen, R. Rahman, C. J. Wellard, I. Woo, J. Caro, N. Collaert, S. Biesemans, G. Klimeck, L. C. L. Hollenberg and S. Rogge, *Nat. Phys.*, 2008, **4**, 656–661.

38 J. Chen, X. Zhao, G. Grinblat, Z. Chen, S. J. R. Tan, W. Fu, Z. Ding, I. Abdelwahab, Y. Li, D. Geng, Y. Liu, K. Leng, B. Liu, W. Liu, W. Tang, S. A. Maier, S. J. Pennycook and K. P. Loh, *Adv. Mater.*, 2018, **30**, 1704674.

39 P. X. Tran, *J. Electron. Mater.*, 2018, **47**, 5905–5912.

40 P.-C. Shen, C. Su, Y. Lin, A.-S. Chou, C.-C. Cheng, J.-H. Park, M.-H. Chiu, A.-Y. Lu, H.-L. Tang, M. M. Tavakoli, G. Pitner, X. Ji, Z. Cai, N. Mao, J. Wang, V. Tung, J. Li, J. Bokor, A. Zettl, C.-I. Wu, T. Palacios, L.-J. Li and J. Kong, *Nature*, 2021, **593**, 211–217.

41 J. Hafner, *J. Comput. Chem.*, 2008, **29**, 2044–2078.

42 Y. Tsai and Y. Li, *IEEE Trans. Electron Devices*, 2018, **65**, 733–738.

43 C.-P. Lin, P.-S. Liu, L. Lyu, M. Li, C.-C. Cheng, T.-H. Lee, W. Chang, L. Li and T. Hou, N-type Doping Effect of Transferred MoS_2 and WSe_2 Monolayer, Extended Abstracts of the 2015 International Conference on Solid State Devices and Materials, Sapporo, 2015, pp. 642–643.

44 K. B. C. Simbulan, P. C. Chen, Y. Y. Lin and Y. W. Lan, *J. Visualized Exp.*, 2018, **138**, e57885.

45 P.-C. Chen, C.-P. Lin, C.-J. Hong, C.-H. Yang, Y.-Y. Lin, M.-Y. Li, L.-J. Li, T.-Y. Yu, C.-J. Su, K.-S. Li, Y.-L. Zhong, T.-H. Hou and Y.-W. Lan, *Nano Res.*, 2019, **12**, 303–308.