Near-ideal van der Waals rectifiers based on all-two-dimensional Schottky junctions

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The applications of any two-dimensional (2D) semiconductor devices cannot bypass the control of metal-semiconductor interfaces, which can be severely affected by complex Fermi pinning effects and defect states. Here, we report a near-ideal rectifier in the all-2D Schottky junctions composed of the 2D metal 1T'-MoTe2 and the semiconducting monolayer MoS2. We show that the van der Waals integration of the two 2D materials can efficiently address the severe Fermi pinning effect generated by conventional metals, leading to increased Schottky barrier height. Furthermore, by healing original atom-vacancies and reducing the intrinsic defect doping in MoS2, the Schottky barrier width can be effectively enlarged by 59%. The 1T'-MoTe2/healed-MoS2 rectifier exhibits a near-unity ideality factor of ~1.6, a rectifying ratio of >5 × 10^5, and high external quantum efficiency exceeding 20%. Finally, we generalize the barrier optimization strategy to other Schottky junctions, defining an alternative solution to enhance the performance of 2D-material-based electronic devices.

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Metal-semiconductor interfaces are crucial to the operations and applications of every semiconductor devices, and are at the basis of Schottky diodes. With excellent on-off switching characteristics, Schottky diodes often replace p–n diodes in many essential components of modern electronics and optoelectronics. Recently, without the constraint limits of lattice matching and processing compatibility requirements of conventional bonded heterojunctions, two-dimensional (2D) van der Waals (vdWs) heterojunctions have attracted widespread interests. To date, the traditional merits of Schottky diodes are not reproduced in 2D material-based Schottky diodes but are replaced by many deficiencies, such as excessive reverse leakage current (i.e., large power consumption), low current rectifying ratio, and poor ideality factor. This is attributed to the composition effect of the low Schottky barrier height and the tiny Schottky barrier width in 2D Schottky diodes in Fig. 1a, which are originated respectively from the strong Fermi pinning effect of the metal–semiconductor junctions and the ultrathin body thickness.

Both achieving the designed Schottky barrier height and precisely controlling the barrier width are crucial for realizing the full potential of Schottky diodes. For Schottky barrier height, transferring metals or employing 2D metals can build weak Fermi pinned Schottky junctions to roughly satisfy the personalized barrier height. The 2D metals (e.g., 1T′-MoTe2, 1T′-WTe2, 1T′-PtSe2, and 2H-NbSe2), by contrast, have more potential in creating high-grade Schottky junctions because their Schottky barrier heights strictly follow the trend of the Schottky–Mott model. Since 2D metals can form the weaker chemical bonding interface with 2D vdWs semiconductors and efficiently suppress the metal-induced gap states in conventional Schottky junctions. The metal-induced gap states are one of the important sources of the Fermi pinning effect.

On the other hand, the tunneling currents widely exist in the 2D semiconductor-based Schottky junctions, which is highly related to its tiny Schottky barrier widths and will form the large leakage currents to cause the high power consumptions of diodes. To date, the adjustments of the Schottky barrier widths often rely on surface absorption chemical doping, which is environment unstable and inefficient in controlling the electronic structures. Lattice defect healing is an optimal strategy to efficiently and stably decrease the intrinsic defect doping of ultrathin 2D materials.

Here, we report a near-ideal Schottky diode in 1T′-MoTe2/MoS2 Schottky junction composed of the metallic 1T′-MoTe2 and the semiconducting monolayer MoS2 in Fig. 1b. We show constructing all 2D Schottky junctions greatly reduce the Fermi pinning effect in the metal–semiconductor interfaces. Furthermore, we firstly demonstrate that by healing original defects, shrinking the intrinsic defect doping in monolayer MoS2 can effectively enlarge the Schottky barrier width to significantly suppress the reverse tunneling leakage current in Fig. 1c. After the defect healing in monolayer MoS2, the 1T′-MoTe2/healed-MoS2 Schottky diode shows a near-unity ideality factor of ~1.6, a high rectifying ratio of >5 × 10^5 in monolayer semiconductor-based Schottky diodes to date, and high external quantum efficiency of >20%. The Schottky barrier optimization strategy is generally applicable to other Schottky junctions, including Pd/MoS2, 1T′-MoTe2/WS2, 1T-PtSe2/MoS2, and 1T-PtSe2/WS2. This work provides an alternative solution for improving the performance of 2D material-based Schottky junctions.

Results

Basic characterizations of the Schottky junctions. The idea and strategy of the Schottky junction optimization are shown in Fig. 1a–c. The detailed differences between the general and optimized Schottky junctions by the conventional and 2D metals are summarized in Fig. 1a, b. A monolayer MoS2 transistor with symmetric 1T′-MoTe2 electrodes in Fig. 2a was built to characterize the Schottky barrier height between monolayer MoS2 and 1T′-MoTe2. The monolayer MoS2 and few-layer 1T′-MoTe2 are chemical vapor deposition.
deposition (CVD) grown and mechanically exfoliated (‘Methods’), respectively. The 1H-MoS2 has a trigonal prismatic structure and lacks inversion symmetry; while the 1T′-MoTe2 has a distorted octahedral structure and inversion symmetry in Fig. 2b. The intensities of the several modes (e.g., 194 cm\(^{-1}\)) exhibit a four-lobed shape with four maximum-intensity angles at ~10°, ~100°, ~190°, and ~280° in Fig. 2c and Supplementary Fig. 1. This data conform to the polarization dependence of the Raman signals of 1T′-MoTe2 was measured under the atmosphere environment and may be slightly different from the true vacuum value. According to the Schottky–Mott rule, the Pd/MoS2 Schottky junction should possess a larger Schottky barrier height than the 1T′-MoTe2/MoS2 Schottky junction. However, the Schottky barrier height of the 1T′-MoTe2/MoS2 Schottky junction is significantly higher than the reported values of ~0.12 eV of the Pd/MoS2 Schottky junction in Fig. 2e (ref. 2). The fundamental reason for causing the opposite result is that unlike 3D metal Pd, the 2D metal 1T′-MoTe2 can suppress the strong Fermi pinning effect at the metal-semiconductor interface. In reality, the Schottky barrier heights between most 3D metals and monolayer MoS2 had seriously deviated from the Schottky–Mott rule22,23, resulting in the significantly lower Schottky barrier of 0–0.3 eV than the theoretical values in Fig. 2c. Therefore, followed by the formula (1), the ideal value of the slope \( S = \left| d\Phi_{SB}/d\Phi_M \right| \) in Fig. 2e is 1. However, the S value of the Schottky barrier height between 3D metals and monolayer MoS2 is only 0.09 which is much lower than the ideal value, indicating the Fermi levels of the MoS2 in these metal-semiconductor interfaces are severely pinned.

Characterization of Schottky barrier height. Kelvin probe force microscopy (KPFM) was used to characterize the work function of the metallic 1T′-MoTe2 nanosheet as ~4.83 eV (Supplementary Fig. 3c, d). The Schottky barrier height is measured as ~501 ± 26 meV in Fig. 2b by the variable temperature study (Supplementary Fig. 4), and is the same as the largest Schottky barrier height of 0.5 eV in the reported all 2D Schottky junctions32. In the Schottky–Mott rule, the Schottky barrier height in the Schottky junction is defined as32,

\[
\Phi_{SB} = \Phi_M - \chi_S
\]

where \(\Phi_M\) and \(\chi_S\) are the work function of the metal electrode and the electron affinity of the n-type semiconductors, respectively. The \(\chi_S\) of monolayer MoS2 previously reported is ~4.3 eV (ref. 28,32), implying the theoretical Schottky barrier height \(\Phi_{SB}\) of the 1T′-MoTe2/MoS2 Schottky junction should be ~0.53 eV. The experimental and theoretical values are close, suggesting that our constructed 1T′-MoTe2/MoS2 Schottky junctions are excellent and largely follow the Schottky–Mott rule.

The work function of Pd is ~5.3 eV (ref. 2), which is larger than that of ~4.83 eV of 1T′-MoTe2. It should be noted that this work function of 1T′-MoTe2 was measured under the atmosphere environment and may be slightly different from the true vacuum value. The gunmetal dash line and points are the calculated SBH from the ideal Schottky–Mott model. The blue solid line is fitted line of experimental data. 

Fig. 2 The 1T′-MoTe2/MoS2 Schottky junction with large Schottky barrier height and weak Fermi pinning effect. a Schematic diagram (top) and optical micrograph (bottom) of a monolayer MoS2 FET with symmetric 1T′-MoTe2 electrodes. b Crystal structures of the 1H-MoS2 (left) and 1T′-MoTe2 (right) monolayers. c Angle-resolved Raman spectra of the mechanically exfoliated 1T′-MoTe2 flake. d Barrier heights of the 1T′-MoTe2/MoS2 Schottky junction as a function of gate voltage obtained from Supplementary Fig. 4. The Schottky barrier height is extracted under a flat band gate voltage condition, which is responsible for the start of deviations from the linear behavior. e Comparisons of the extracted Schottky barrier height (SBH) between monolayer MoS2 and the electrodes of evaporated 3D metals (extracted from ref. 2) and the 2D metal 1T′-MoTe2. The gunmetal dash line and points are the calculated SBH from the ideal Schottky–Mott model. The blue solid line is fitted line of experimental data. 

f, g Band diagrams (f) and output curves (g) of the monolayer MoS2 diodes with the Pd and 1T′-MoTe2 electrodes. All gate voltages are 0 V.
The Fermi pinning effect is highly related to the defect-induced gap states of the semiconductors, the additional defects and chemical disorders involved by device fabrications, and the metal-induced gap states. The benefits of using 2D metals in suppressing the Fermi pinning effect are as follows. Firstly, lattice structure damages of the channel materials from the 3D metal evaporation are avoided. Secondly, some chemical disorders in the interface will not be involved by various solvents used in lithography processes. Thirdly, unlike 3D metals, 2D metals can suppress the metal-induced gap states formed from the decaying metallic wavefunction of the 3D metal itself.

To explore the dependence of the Schottky barrier height on the performance of 2D semiconductor-based diodes, we compared the rectifying performance of the Pd/MoS2 and 1T-′MoTe2/MoS2 diodes. According to the aforementioned discussions, the intrinsic electron concentration is significantly depressed by the treatment operation details and healing mechanism are also presented later (Methods). To further improve the performance among monolayer 2D semiconductor-based Schottky diodes reported values of 4 × 10^5 at VD = ± 2 V and VG = -40 V in Fig. 3d. This ratio is the best performance among monolayer 2D semiconductor-based Schottky diodes reported to date (Supplementary Table 1). The gate-dependent output curves and transfer curves of the Schottky junction before and after the SVSH are presented in Supplementary Fig. 8. To minimize the performance degradation from transferring monolayer MoS2 in the device fabrication, another device design has been abandoned, in which the MoS2 is top on the metallic 1T-′MoTe2. Though the abandoned device design can achieve the SV healing of the MoS2 above the 1T-′MoTe2, the performance degradations from transferring MoS2 may offset the benefits of defect healing.

To further evaluate the rectifying performance of the 1T-′MoTe2/MoS2 Schottky junction diode, an ideality factor (n) was estimated at a small forward bias (here is 0.02–0.35 V, Fig. 3c) by fitting to the Schottky diode equation:

$$I_D = I_S \exp \left( \frac{V_D}{nV_T} \right) - 1$$

where ID, IS, VD, and VT denote drain current, reverse leakage current, drain voltage, and thermal voltage, respectively. As the gate voltage sweep from positive to negative in Fig. 3e, the ideality factor derived from the parameters of the fitting equation gradually approaches the perfect limit value of 1 and gain ~1.6 at VG = -40 V (the fitted lines of other gate voltages in Supplementary Fig. 8a, b), indicating that the Schottky junction probably reaches a near-ideal diode behavior under the dual effect of the atom-vacancy healing and gate doping. This similar performance improvement effect can be reproduced in dozens of devices. Besides, with gate voltage sweeping from positive to negative, both the rectifying ratios and ideal factors will gain the best performance at some negative gate voltage and then start to degrade. This performance transition may be related to the trade-off between the resistances of the bare Schottky junction (consist of region A, B1, and B2 in Fig. 3f) and the series resistances of the MoS2 channel (region B3) of the Schottky junction dominates the total resistance of the diode above the transition voltage. While the interference series resistance of the MoS2 channel will gradually increase its proportion in total resistance and replace the Schottky junction to dominate the diode after crossing the transition voltage.

To remove the interferences from electrode contacts of the MoS2/Cr and 1T-′MoTe2/Cr interfaces, both MoS2 and MoTe2 transistors before and after the PEDOT:PSS treatment was also fully characterized. The nearly linear current–voltage relationships in the output curves of the Cr/MoS2–Cr transistors before and after the SVSH are observed in Supplementary Fig. 7b, c, indicating that the Ohmic contacts of the MoS2/Cr interfaces have not been adversely affected. It should be noted that the Cr/ MoS2 interfaces are not truly Ohmic contacts, and its Schottky barrier heights before and after the SVSH are largely maintained at ~0.1 eV (Supplementary Fig. 9). The Schottky
barriers of the Cr/MoS\textsubscript{2} interfaces are believed to be highly related to the formation of CrS\textsubscript{x} (ref. 37). On the other hand, compared to the untreated one, the treated 1T\textsuperscript{′}-MoTe\textsubscript{2} transistor still maintains a linear current–voltage relationship and constant conductance in Supplementary Fig. 7f–h, indicating that the variations on the 1T\textsuperscript{′}-MoTe\textsubscript{2} side from the acid treatment are almost negligible. Therefore, the main action object of the PEDOT:PSS treatment is the metal–semiconductor interface of the 1T\textsuperscript{′}-MoTe\textsubscript{2}/MoS\textsubscript{2} Schottky junction, rather than the other electrode contacts.

**Fig. 3** Giant optimization of the Schottky junction through thickening the Schottky barrier width. a Schematic diagram (top) and optical micrograph (bottom) of 1T\textsuperscript{′}-MoTe\textsubscript{2}/MoS\textsubscript{2} Schottky junctions in which the monolayer MoS\textsubscript{2} serves as the transport channel, p-Si is the control gate, SiO\textsubscript{2} layer is the gate dielectric, the 1T\textsuperscript{′}-MoTe\textsubscript{2} is the bias drain electrode, and the Cr/Au is grounded. b Schematic diagram of the atomic structure of the 1T\textsuperscript{′}-MoTe\textsubscript{2}/MoS\textsubscript{2} Schottky junctions before and after the SVSH. c Schematic diagram of the atomic structure of the 1T\textsuperscript{′}-MoTe\textsubscript{2}/MoS\textsubscript{2} Schottky junctions before and after the SVSH. d, e Gate-dependent current rectifying ratios (d) and ideality factors (e) extracted from Supplementary Fig. 8 before and after the SVSH. f Device schematic diagrams (top) and band structure diagrams (bottom) before and after the SVSH. The $E_F$, $E_C$, and $E_V$ are Fermi level, conduction band, and valence band, respectively. The A, B1, B2, and B3 regions refer to the metal, contacted MoS\textsubscript{2}, depletion exposed MoS\textsubscript{2}, and pure channel MoS\textsubscript{2}. The yellow and blue gradient areas mean the depletion regions of the Schottky junctions. It should be noted that to simplify the band structure, the tunneling barriers originated from the vdWs gap between 1T\textsuperscript{′}-MoTe\textsubscript{2} and MoS\textsubscript{2} are intentionally removed in the whole manuscript. g Band structure diagrams of the two-state devices at forward and reverse biased voltages. The green and red arrows (solid/dashed: main/almost-impossible style) indicate the thermionic emission current and thermally assisted tunneling current, respectively. The movement direction of an electron during reverse (forward) biased voltage is from the MoTe\textsubscript{2} to the MoS\textsubscript{2} (MoS\textsubscript{2} to MoTe\textsubscript{2}).
While reducing the electron doping of the SVs of the covered MoS$_2$ (region B2) in Fig. 3f only will efficiently enlarge the depletion width of the Schottky barrier but will not significantly alter the Schottky barrier height $\Phi_{SB}$. The variable temperature transport studies show the Schottky barrier heights $\Phi_{SB}$ before and after treatment are calculated as $501 \pm 26$ and $510 \pm 19$ meV in Supplementary Fig. 4, suggesting the SVSH effect can not largely alter the Schottky barrier height $\Phi_{SB}$. On the other hand, if the Schottky barrier height $\Phi_{SB}$ is considered to be significantly enhanced by the SVSH effect, both the forward and reverse biased currents will be significantly reduced. However, the forward-biased currents are nearly unchanged or even increased in the partial negative voltage range in Supplementary Fig. 8c right. Besides, previous studies showed that not doping the covered region B1 but doping the exposed region B2 (Fig. 3f) could also significantly modulate the Schottky barrier width. In reality, even if the monolayer MoS$_2$ of the covered region B1 (Fig. 3f) was healed, the improvement effect induced by the SVSH on the rectifying behavior of the 1T’-MoTe$_2$/MoS$_2$ Schottky diode would not be significantly changed (Supplementary Fig. 10).

**Improvement mechanism of the rectifying performance.** So how does enlarging the Schottky barrier width improve the rectifying behavior? Based on this band structure transformation of the Schottky junction in Fig. 3f, the depletion region width of the as-prepared Schottky junction under reverse bias is too narrow to easily generate large tunneling current Fig. 3g. When the Schottky barrier height remains roughly constant in Fig. 3g, enlarging the Schottky barrier width can transform the charge injection style under reverse bias from thermionic emission to thermionic field emission (also called thermally assisted tunneling). Thus, different from the 1T’-MoTe$_2$/as-prepared MoS$_2$ Schottky junction, the reverse leakage currents of the optimized Schottky junctions are significantly suppressed. But in the forward bias, an electron only needs to pass through the Schottky barrier with a small built-in potential to reach the metal electrode 1T’-MoTe$_2$ in Fig. 3g bottom. On the other hand, as forward bias is added, the resistance of the bare Schottky junction decreases, the negative effect of the series resistance of the MoS$_2$ channel (region B3) becomes comparable. To put it simply, the total resistance of the 1T’-MoTe$_2$/healed MoS$_2$ Schottky junction in the forward bias can be presumably assumed to be divided into the large MoS$_2$ series resistance (region B3) and small bare Schottky junction resistance (regions A, B1, and B2). While the total resistance of the 1T’-MoTe$_2$/as-prepared MoS$_2$ Schottky junction is the sum of the small MoS$_2$ series resistance and large bare Schottky junction resistance in Fig. 3g bottom-left. In this case, the total resistance of the two Schottky junctions in the forward bias may be close in the most 1T’-MoTe$_2$/MoS$_2$ Schottky junctions, which has been confirmed by the two similar forward currents in Fig. 3c and Supplementary Fig. 8d. Based on the performance survey of the dozens of Schottky junctions, the differences between the two forward currents of the as-prepared and healed devices are fluctuated by the MoS$_2$ channel length. In a word, the decline of $E_P$ induced by the SVSH can greatly reduce the reverse leakage current, roughly maintain the forward current, and finally improve the rectifying ratio and ideal factor.

**Optimization of other Schottky junctions.** Except for the 1T’-MoTe$_2$/MoS$_2$ Schottky diode, the performance optimization strategy induced by the vacancy healing can also be generalized to other MoS$_2$-based diodes. The monolayer MoS$_2$ Schottky diodes with the Pd and Cr asymmetric metal electrodes after the solution treatment can only exhibit a maximum rectifying ratio of $\sim 2.9 \times 10^3$ in Fig. 4a, which are considerably worse than that of the 1T’-MoTe$_2$/healed MoS$_2$ Schottky junctions in Fig. 3d.

The poor performance of the Pd/MoS$_2$ Schottky junction can be partially attributed to its small Schottky barrier height of $\sim 0.12$ eV (ref. 2), which is lower than the value of $\sim 0.5$ eV of the 1T’-MoTe$_2$/healed MoS$_2$ Schottky junction in Fig. 2d. Nevertheless, this optimized performance of the Pd/MoS$_2$ Schottky junction is much higher than that (1~100) of similar MoS$_2$-based Schottky junctions with evaporated asymmetric metal electrodes (Supplementary Table 1). As mentioned before, the Pd/MoS$_2$ Schottky junctions may be suffered from the severe Fermi pinning effect in the interface between the evaporated Pd electrode and the monolayer MoS$_2$. Thus, the origin of the superior rectifying behaviors of the 1T’-MoTe$_2$/healed-MoS$_2$ Schottky junctions can be attributed to the dual effect with the large Schottky barrier height and the spacious Schottky barrier width.

Similar to the 1T’-MoTe$_2$/MoS$_2$ and Pd/MoS$_2$ Schottky junction, the rectifying ratios of the 1T’-PtSe$_2$/MoS$_2$ also were systematically enhanced by the SVSH in Fig. 4b. Due to the similar layered structure between transition metal dichalcogenides (TMDCs) materials, the acid-induced vacancy healing effect has been confirmed to apply to other TMDCs semiconductors. Thus, we also generalize the performance optimization strategy to other TMDCs-based Schottky junctions (Supplementary Fig. 11 and Fig. 4c, d.). Similarly, the vacancy healing effect can also increase the rectifying ratios of the 1T’-MoTe$_2$/WS$_2$ and 1T’-PtSe$_2$/WS$_2$ Schottky junctions, which means that their Schottky barrier widths may be greatly enhanced. It should be noted that similar to the 1T’-MoTe$_2$ film, the 1T’-PtSe$_2$ transistor still maintains a linear current–voltage relationship and constant metallic conductance in Supplementary Fig. 12.

**Photoresponse characterization of the Schottky junctions.** Next, we focus on the photoresponse of the 1T’-MoTe$_2$/MoS$_2$ Schottky junctions before and after the SVSH and evaluate its performance in photovoltaic detectors in the visible wavelength range. Figure 5a shows that regardless of the as-prepared device or the healed device, the short-circuit current $I_{SC}$ and open-circuit voltage $V_{OC}$ show a monotonic increase with the increase of laser power density. Under the same laser power density, both the $I_{SC}$ and $V_{OC}$ are significantly increased by the SVSH effect. As a comparison, the obvious photovoltaic response is not observed in the as-prepared and healed MoS$_2$ transistors. Similarly, the interface charge separation of the 1T’-MoTe$_2$/MoS$_2$ Schottky junction has been confirmed by the PL spectra (Supplementary Fig. 14). With the power intensity increasing, the $I_{SC}$ linearly increases, and the $V_{OC}$ will gradually enter the saturated state (Fig. 5b). In the 1T’-MoTe$_2$/healed-MoS$_2$ Schottky junction, a high $I_{SC}$ of $\sim 22.2$ nA and a large $V_{OC}$ of $\sim 0.19$ V are obtained under the power density of $8.23$ mW/mm$^2$ (junction area $\sim 671$ m$^2$). Under the maximum laser power density, the output electrical power density is increased by about 16 times from $\sim 16.7$ to $\sim 270.7$ mW/cm$^2$ through the SVSH strategy in Fig. 5c.

To evaluate the performance improvement of the MoTe$_2$/MoS$_2$ photodetector, the photoresponsivity and external quantum efficiency (EQE) are systematically characterized. The photoresponsivity is defined as $R = I_{ph}/(P_{in} A)$, where $I_{ph}$ is the photocurrent, $P_{in}$ is the incident laser power density, and $A$ is the junction area of the 1T’-MoTe$_2$/MoS$_2$ diode. EQE is the ratio of collected charge carriers to incident photons and can be calculated by EQE = ($I_{ph} h c)/(e \lambda P_{in} A)$, where $h$, $c$, and $\lambda$ are the Planck’s constant, the speed of light, and the wavelength of the incident laser, respectively. At laser power density of $\sim 1.9$ mW/mm$^2$ (the laser spot size is $\sim 2.25$ m$^2$), the
photoresponsivities of the as-prepared and healed devices are \~59/~88 mA/W and EQE of \~13/~20% in Fig. 5d. This improvement in photoresponse performance can be attributed to the SVSH effect which greatly enlarges the Schottky barrier width, meaning that the photoresponse area is increased in Fig. 5e, f. The second possible reason is that the sub-bandgap defect states of SVs will trap the photo-generated electrons and promote non-radiative recombination in Fig. 5e, leading to lower $I_{SC}$ and $V_{OC}$ (ref. 41). Thirdly, a neglected but reliable reason is that the removal of electron doping induced by the SVSH effect greatly improves the light absorption efficiency of monolayer MoS$_2$ to form more photocarriers 42,43. The electron doping removal induced by gate doping also can enhance the $I_{SC}$ and $V_{OC}$ of the MoS$_2$ Schottky junction2, while our improvement strategy of defect healing is gate-free and doesn’t rely on the external gate voltage.

Discussion

In conclusion, a near-ideal 1T'–MoTe$_2$/MoS$_2$ Schottky diode with a large Schottky barrier height was successfully fabricated by stacking the 2D metallic 1T'-MoTe$_2$ and the semiconducting monolayer MoS$_2$. We systemically demonstrate that besides the large Schottky barrier height, thickening the tiny Schottky barrier width can further improve their performance of 2D Schottky junctions by healing the original defects in the 2D semiconductor counterparts. We further confirm that this Schottky barrier optimization strategy is generally applicable to other Schottky diodes, including 1T-PtSe$_2$/MoS$_2$, 1T'-MoTe$_2$/WS$_2$, and Pd/MoS$_2$. This work provides a proof-of-concept performance optimization strategy for building high-performance Schottky diodes.

Methods

Growth of monolayer MoS$_2$. The monolayer MoS$_2$ films were grown by the atmospheric pressure oxygen-assisted CVD method28,35. MoO$_3$ (Sigma-Aldrich, \geq 99.5% purity) and sulfur (Sigma-Aldrich, \geq 99.5% purity) were applied as the precursor and reactant materials, respectively. MoO$_3$ powder (10 mg) was placed in a quartz boat at the center of a single zone furnace (Tianjin Zhonghuan Electrical Furnace Co., Ltd.). A 2 × 2 cm$^2$ Si substrates with 300 nm SiO$_2$ (from Silicon Valley Microelectronics, Inc.) were put face down at top of the MoO$_3$ powder. S powder was heated to 170 °C by the home-made heating belt and carried through a high-purity Ar flow of 500 sccm. The experiments were implemented at a reaction temperature of 850 °C for 30 min. When the temperature reached the set value of 850 °C, the oxygen of \~1.2 sccm was introduced as the auxiliary growth gas and it was kept for 30 min. Finally, the monolayer MoS$_2$ samples were cooled naturally under S vapor to minimize SVs.

Thermal CVD growth of WS$_2$. The WS$_2$ films were grown by the typical thermal CVD process44. To prepare the WS$_2$ flake single crystals, an alumina boat loaded with WS$_2$ (Alfa Aesar, 99.8% purity) powder (3–5 g), was placed into the middle heating zone of the single zone furnace (from Therm craft XST-2-0-12-MINI-240). The Si substrates (with 300 nm SiO$_2$) were put face down at top of the MoO$_3$ powder. S powder was heated to 170 °C by the home-made heating belt and carried through a high-purity Ar flow of 500 sccm. The experiments were implemented at a reaction temperature of 850 °C for 30 min. When the temperature reached the set value of 850 °C, the oxygen of \~1.2 sccm was introduced as the auxiliary growth gas and it was kept for 30 min. Finally, the monolayer MoS$_2$ samples were cooled naturally under S vapor to minimize SVs.

Material preparation and device construction. The metal phase 1T'-MoTe$_2$ and 1T-PtSe$_2$ films were obtained by mechanical exfoliation of the 1T'-MoTe$_2$ (from Hefei Kejing Co.) and 1T-PtSe$_2$ (from 2D Semiconductors) bulks. The devices were constructed by stacking CVD-grown monolayer MoS$_2$ and exfoliated 1T'-MoTe$_2$ flakes with the accurate transfer platform (Metatest, E1-T). We used a standard electron beam lithography (EBL) to pattern source/drain contacts of MoS$_2$ and 1T'-MoTe$_2$.
MoTe₂ FETs, followed by thermal beam evaporation of 10/50-nm-thick Cr/Au electrodes and lift-off. The metallization was performed by a thermal evaporation system (KYKY Technology), and the thermal evaporation pressure was 1–5 × 10⁻⁵ Pa. Especially, our electrode contact optimization for CVD-grown monolayer MoS₂ FETs was the mild Argon plasma etching which can remove residual polymethyl methacrylate (PMMA, A8, Microchem Inc.) and produce a small amount of SVs. The plasma etching parameters (from Tailong Electronics RIE 100) are the radiofrequency (13.56 MHz) power of 5 W, the Ar flow of 10 sccm, and the treatment duration of 35–45 s. Both the ~15/45 nm Pd/Au electrodes and the ~15/45 nm Cr/Au electrodes in the Schottky junction diodes of the asymmetric metal electrodes were deposited by thermal evaporation.

Acid solution treatment for sulfur vacancy self-healing (SVSH). Firstly, the MoS₂ sample was immersed in the undiluted poly(3,4-ethylenedioxythiophene):poly(4-styrenesulfonate) (PEDOT:PSS) solution (Sigma-Aldrich, 1.0 wt%) at room temperature, after standing for 5 min, and then immersed in plenty of deionized (DI) water to wash the water-soluble PEDOT:PSS solution for 30 min. The quality of the DI water, which was used to rinse the samples (~0.8 × 0.8 cm² Si substrates), is >200 + >200 mL to the twice cleaning. Further, the residual deionized water was dried with nitrogen flow. Both Raman spectra and gas chromatography–mass spectrometry (GC–MS) can prove that there is no residual PEDOT:PSS in the 1T′-MoTe₂/MoS₂ Schottky junctions before (Supplementary Figs. 5b, c top). After these processes, the STEM images are converted into the plain text signal, in which the surface contaminations (Supplementary Fig. 5b, c bottom) by using an Origin Pro 9.0. It should be noted that to reduce the measurement error of the 1T′-MoTe₂ film from strong laser irradiation, both the PMMA package and the weak laser power density (<5 mW/μm²) are simultaneously arranged [45].

**AFM and KPFM measurement.** The AFM and KPFM images were obtained via the Bruker Icon AFM system. An AFM tip (SCM-PIT-V2, Bruker Nano Inc., USA) was selected to probe the topography and thickness of the samples under the peak-force working mode, while the AM-KPFM measurements were measured under the tapping mode via the same tip of SCM-PIT-V2 with a resonant frequency of 75 kHz, a tip radius of 25 nm, a Pt–Ir coated conductive tip, a tip-sample height of 100 nm, and the ac bias of 0.8 V on a grounded substrate. These tests were performed in an atmospheric environment.

**Calculation of Schottky barrier width W.** The Schottky barrier width W of the Schottky junction can be obtained by Eq. (2) [4]. We use a relative dielectric constant of 3.3 for MoS₂ (ref. [46]). The Φ_M of 1T′-MoTe₂ and the Φ_S of 1H-MoS₂ before and after the SVSH effect can be extracted from Supplementary Fig. 3c, d and Supplementary Fig. 6e. The d_Mo is the thickness of 0.65 nm of monolayer MoS₂. Thus, the Schottky barrier widths W before and after the SVSH effect are ~2.9 and ~4.6 nm, respectively.

**STEM measurement.** STEM samples were prepared using a standard PMMA-based transfer, where the samples were spin coated and soaked in a KOH solution (5 wt%). Subsequently, acetone (AR, from RCI Labscan Limited) and isopropyl alcohol (AR, from Aladdin) were used to clean the TEM grid, allowing for the suspension of the films in the solution and the transfer to TEM grids. All STEM samples were baked at 160 °C for 5 h under vacuum (the pressure was <1 Pa) before the microscopy experiment and characterized on a FEI-ARM200F TEM operating at 80 kV. The STEM images were prepared by partially filtering out the direct spot in the FFT of the image, which helps to remove the contrast generated by the surface contaminations (Supplementary Fig. 5b, c top). After these filtering processes, the STEM images are converted into the plain text signal, in which the signal of the Mo atoms was accurately eliminated by a home-made program. Finally, the program-processed data was produced into the filtered STEM images (Supplementary Fig. 5b, c bottom) by using an Origin Pro 9.0. It should be noted that there is no residual PEDOT:PSS in the 1T′-MoTe₂/MoS₂ Schottky junctions.

Fig. 5 Photoreponse optimization of the 1T′-MoTe₂/MoS₂ Schottky junction photodiodes. **a** Output curves of the 1T′-MoTe₂/healed MoS₂ and 1T′-MoTe₂/as-prepared MoS₂ (inset) Schottky junctions in dark and under 532 nm light irradiation with various laser power densities (P_laser). The gate voltages before and after the SVSH are zero. **b** Short circuit current (I_sc) and open-circuit voltage (V_OC) as a function of P_laser extracted from (a). **c** Output electrical power of the as-prepared (left) and healed (right) devices as a function of drain-source voltage. **d** P_laser-dependent photoresponsivities (red) and EQE (blue) at zero bias. **e** f Band structure diagrams of the 1T′-MoTe₂/MoS₂ Schottky junctions before (e) and after (f) the SVSH under light irradiation. The purple and hollow circles represent electrons and holes, respectively. The red dashed arrow refers to the electron movement direction. The yellow and blue gradient areas refer to the depletion regions of the Schottky junctions. W_light-eff refers to the efficient light absorption width of the Schottky junction.
that to reduce the interference of ex-situ measurement, the CVD-grown monolayer MoS₂ samples in the STEM measurement were originated from the same SiO₂/Si substrate, and the pass energy for the narrow scan was 30 eV. The base pressure was about \(6.5 \times 10^{-10}\) mbar. The analyzer of the spectrometer was calibrated using sputter cleaned Au, Cu, and Ag foils, as is outlined in the GB/T 19500-2004 standard. In this calibration method, the standard calibrated binding energies of the Au, Cu, and Ag standard samples were \(\text{Au 4f}_{7/2} = 84.0 \pm 0.1\) eV, \(\text{Cu 2p}_{3/2} = 932.7 \pm 0.1\) eV, and \(\text{Ag 3d}_{3/2} = 368.3 \pm 0.1\) eV, respectively. To compensate for sample charging, the XPS spectra were calibrated using the Si 2p⁹/₂ peak of the SiO₂ substrate at 103.6 eV as a reference. For XPS peak analysis and deconvolution, the software Avantage 4.15 was employed, where a mixed Gaussian–Lorentzian function and an active Shirley background were used for peak fitting. The S/Mo ratios were determined from the integrated areas of the S 2p and Mo 3d peaks factored by their corresponding relative sensitivity factors of 0.668 and 3.321. The error in the S/Mo ratios was obtained from the peak fitting residuals.

**UPD**

UPD curves were obtained in an ultrahigh vacuum chamber using a helium lamp source emitting (AXIS ULTRA DLD). The ultraviolet light source used was a He I (21.22 eV) that has not been monochromated. The basic vacuum of the analysis chamber during the UPS analysis and testing of this equipment was \(3.0 \times 10^{-8}\) Torr. The biased voltage during the test was \(\sim 9\) V.

**GC-MS measurement.** GC was carried out using an Agilent 7890A/5975C instrument with an HP-5MS column (30 m x 0.25 mm x 0.25 μm). The temperatures of both inlet and detection were set at 300 °C. The initial temperature of the oven was set at 40 °C and kept for 1 min, and then programmed from 30 to 250 °C with a heating rate of 10 °C/min, and finally retained at 250 °C for additional 5 min. This last increase was to clean the column from any residues. The MS was set as the positive ion model with the mass-to-charge ranging from 20 to 1024. The preparation process of the sample as follows: Firstly, after the 1T–MoTe₂/MoS₂ Schottky diodes were constructed on the SiO₂/Si substrate, the Si substrate was immersed in the absolute ethanol (AR, from Honeywell International Inc.) solution of 3 mL for 2 h. Secondly, remove and blow-dry with nitrogen. After the sample was soaked in the PEDOT:PSS solution for 5 min, was rinsed it twice and after the PEDOT:PSS treatment were extracted 1 μL for the GC-MS test.

**Electrical measurement.** The electrical characteristics were implemented by a semiconductor analysis system (Keithley 4200 and B2902 A). Electrical measurements were performed in a vacuum probe station (Lakeshore TTP4) with liquid nitrogen and base pressure \(\sim 10^{-5}\) Torr.

**Data availability**

All data supporting this study and its findings are available within the article and its Supplementary Information or from the corresponding author upon reasonable request.

**Code availability**

The codes that support the findings of this study are available from the corresponding author upon reasonable request.

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Author contributions
X.Z., Z.Z., and Y.Z. initiated and supervised the project. L.G., Y.L., and B.L. deposited MoS2 monolayers by CVD. X.Z. and B.L. performed device fabrication, data collection, and analysis. J.D., H.Y., M.H., and L.G. assisted in carrying out the film fabrication and characterizations. J.X., Y.L., and L.G. assisted in the device performance measurements. Q.L., Z.K., and B.L. performed part of the Raman, PL, UPS, and XPS characterizations. X.Z. and H.Y. finished the GC-MS, AFM, and KPFM test. L.G. and X.L. carried out STEM experiments. X.Z., Z.Z., and Y.Z. co-wrote the manuscript. All authors discussed the results and commented on the manuscript.

Competing interests
The authors declare no competing interests.

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