

Tailoring Quantum Tunneling in a Vanadium-Doped WSe₂/SnSe₂ Heterostructure

Sidi Fan, Seok Joon Yun, Woo Jong Yu, and Young Hee Lee*

2D van der Waals layered heterostructures allow for a variety of energy band offsets, which help in developing valuable multifunctional devices. However, p–n diodes, which are typical and versatile, are still limited by the material choice due to the fixed band structures. Here, the vanadium dopant concentration is modulated in monolayer WSe₂ via chemical vapor deposition to demonstrate tunable multifunctional quantum tunneling diodes by vertically stacking SnSe₂ layers at room temperature. This is implemented by substituting tungsten atoms with vanadium atoms in WSe₂ to provoke the p-type doping effect in order to efficiently modulate the Fermi level. The precise control of the vanadium doping concentration is the key to achieving the desired quantum tunneling diode behaviors by tuning the proper band alignment for charge transfer across the heterostructure. By constructing a p–n diode for p-type V-doped WSe₂ and heavily degenerate n-type SnSe₂, the type-II band alignment at low V-doping concentration is clearly shown, which evolves into the type-III broken-gap alignment at heavy V-doping concentration to reveal a variety of diode behaviors such as forward diode, backward diode, negative differential resistance, and ohmic resistance.

2D van der Waals (vdW) layered semiconductors possess various bandgaps ranging from 0.4 to 2.0 eV and different electron affinities, establishing predetermined heterojunctions of type-I (straddling gap), type-II (staggered gap), or type-III (broken gap) band alignments.^[1–4] Accordingly, a variety of functional devices can be developed by taking advantage of p–n junctions with different band alignments, which are the elementary

building blocks for numerous applications in diodes, transistors, photodetectors, and solar cells.^[5–9] In addition, the charge transfer originating from the thermionic emission or quantum tunneling is affected by the band bending near the heterojunction, which relies on the Fermi level difference between materials.

Previously, carrier transport properties through a heterojunction appear to be in principle material-dependent behaviors. Recent reports show the possibility to integrate multiple functions in a certain vdW heterostructure; for example, black phosphorus (BP) of different thicknesses are employed to provide a tunable Fermi level for the desired band bending.^[10,11] Nevertheless, BP suffers from sensitivity in air.^[12] Besides, the use of mechanically exfoliated BP is not tenable for controlling the exact flake thickness, which directly determines its Fermi level, let alone realize accessible integration.

Chemical vapor deposition (CVD) technique is a promising approach in a wafer-scale synthesis for industrial applications. Moreover, introducing the dopant during CVD growth is an effective way to change the carrier density in the material and thus modulate the position of the Fermi level.^[13] Here, we report a reliable and repeatable method to synthesize vanadium (V) substituted WSe₂ monolayer by CVD and demonstrate multifunctional p–n diode behaviors in V-doped WSe₂/SnSe₂ heterostructures. A liquid precursor with tungsten host and vanadium dopant atoms is adopted in our approach with only two zones for the precise control of V-doping concentration in WSe₂. The coverage of the grown-WSe₂ flakes is increased up to 90% at a high liquid precursor concentration (Figure S1, Supporting Information). In addition to the p-type V-doped WSe₂ monolayer, multilayer SnSe₂ as an n-type material is introduced to construct the p–n junction. We observe the diverse p–n diode behaviors in V-WSe₂/SnSe₂ devices with various V-doping concentrations at room temperature, including quantum tunneling p–n diodes for the forward rectification, backward rectification, negative differential resistance (NDR), and ohmic resistance.

Figure 1a,b is the schematics for synthesizing V-doped WSe₂ monolayer with the CVD approach. Liquid precursor containing solutions of ammonium metatungstate (AMT: W-precursor) and ammonium metavanadate (AMV: V-precursor) together with a promoter of alkali metal (NaOH) and iodixanol is spin-casted on SiO₂/Si substrate and then introduced into a two-zone furnace CVD for the selenization

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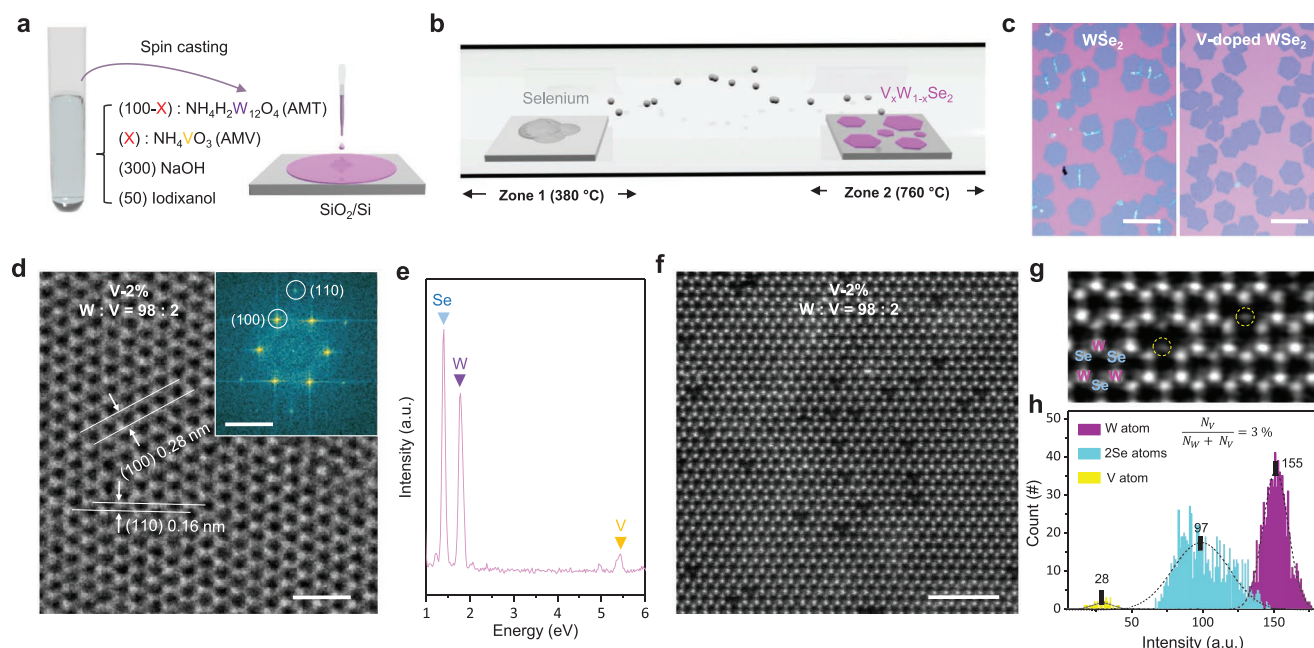


Figure 1. Synthesis and atomic structure of monolayer V-doped WSe₂. a) Schematic for preparation process of water-based precursor. The precursor containing sodium metal (promoter), ammonium tungstate (W-precursor), and ammonium vanadate (V-precursor) are spin-casted onto SiO₂/Si wafer. b) Two-zone furnace CVD process for growing V-doped WSe₂. The precursor-coated substrate is introduced into the CVD chamber and is followed by selenization. c) Optical images of pure and V-doped WSe₂. The monolayer (0.8 nm) was retained up to V-10% doping. The scale bar is 150 μm. d,e) High-resolution TEM image (d) and EDS spectrum (e) of V-2% WSe₂. The scale bar is 1 nm. f,g) Unfiltered (f) and Gaussian-filtered (g) ADF-STEM images of V-2% WSe₂. After the filtering process, the substituted V atoms in W sites are clearly seen. The scale bar is 2 nm. h) The number of W, Se, and V atoms with the STEM intensity. The scale bar is 1 nm.

process. The molar ratio of AMT to AMV in the liquid precursor is precisely controlled using a micropipette.

The hexagonal shape of flakes is preserved in V-doped WSe₂ samples (up to V-10%), similar to the pristine WSe₂ flakes (Figure 1c; Figure S2, Supporting Information). The V-10% sample shows a uniform contrast in the entire flake area with a height profile of 0.8 nm, confirmed by atomic force microscope (AFM). The monolayer is maintained up to V-10% doping (Figure S2, Supporting Information), at which multilayers are partially spotted. The highly crystalline atomic structure of V-doped WSe₂ monolayer (V-2%) is observed without noticeable defect sites from the high-resolution transmission electron microscopy (HRTEM) image with selective-area electron-diffraction pattern (Figure 1d). The interlayer distances of the (100) and (110) planes are identified to be 0.28 and 0.16 nm, respectively, in consistent with the reported values in WSe₂.^[14]

To explore V-incorporation in WSe₂, we further performed energy-dispersive X-ray (EDX) spectroscopy and annular dark field scanning TEM (ADF-STEM). The chemical elements of V atom in addition to W and Se atoms were clearly discernible in EDX spectrum (Figure 1e) and EDX mapping (Figure S3, Supporting Information). Unlike the HRTEM image in Figure 1d, numerous dark spots are clearly distinguished in ADF-STEM image (Figure 1f). The intensity of STEM is proportional to the atomic number (AN) by $a^*(AN)^x$, where a is constant and x ranges from 1.5 to 2.5.^[15] Therefore, the lowest atomic number of V (AN = 23) atoms are seen as the dark spots compared with brightest W (AN = 74) and Se (AN = 34) atoms. After Gaussian filtering (Figure 1g), V atoms are clearly seen by substituting

W atoms. The average intensities of W (155), 2Se (97), and V (28) atoms are well fitted to $a^*(AN)^x$ ($a = 0.29$, $x = 1.55$), indicating that the dark spots are indeed V atoms (Figure 1h). Se-vacancies are also presented in Figure S4 of the Supporting Information with a vacancy density of $1.7 \times 10^{13} \text{ cm}^{-2}$.

Figure 2a demonstrates the Raman spectra of V-doped WSe₂ with different V-doping concentrations (Figure S5, Supporting Information). The Raman shift near 250 cm^{-1} is ascribed to the mixed modes of E_{2g}^1 (strain) with A_{1g} (charge transfer) and is red-shifted to 3 cm^{-1} at 4% V-doping concentration. Such a red-shift is dominantly ascribed to the A_{1g} mode by the p-doping effect of V atoms on WSe₂.^[16] The 2LA(M) mode appears around 260 cm^{-1} and is gradually developed as V-doping concentration increases, reflecting the emergent crystalline disorders due to the randomly distributed V atoms.^[17] The photoluminescence (PL) is further conducted to study the p-doping effect of V-substitution to WSe₂ (Figure 2b). The asymmetric PL emission near 1.66 eV is reminiscent of the positive trions in intrinsic p-doped WSe₂ (trion binding energy $\approx 30 \text{ meV}$).^[18] As V-doping concentration increases, the PL intensity is reduced due to charge (hole) screening and the position is red-shifted by the developed positive trions.^[19]

To investigate V-doping effect on the electrical performance, we fabricated field effect transistors (FETs) using V-doped WSe₂ with Pd contact electrodes for efficient hole transport (Figure 2c). The pristine WSe₂-FET shows ambipolar transfer characteristics at a drain bias of 0.5 V. This prevails the p-type behavior as V-doping concentration increases. The threshold voltage in the hole region is shifted to the positive

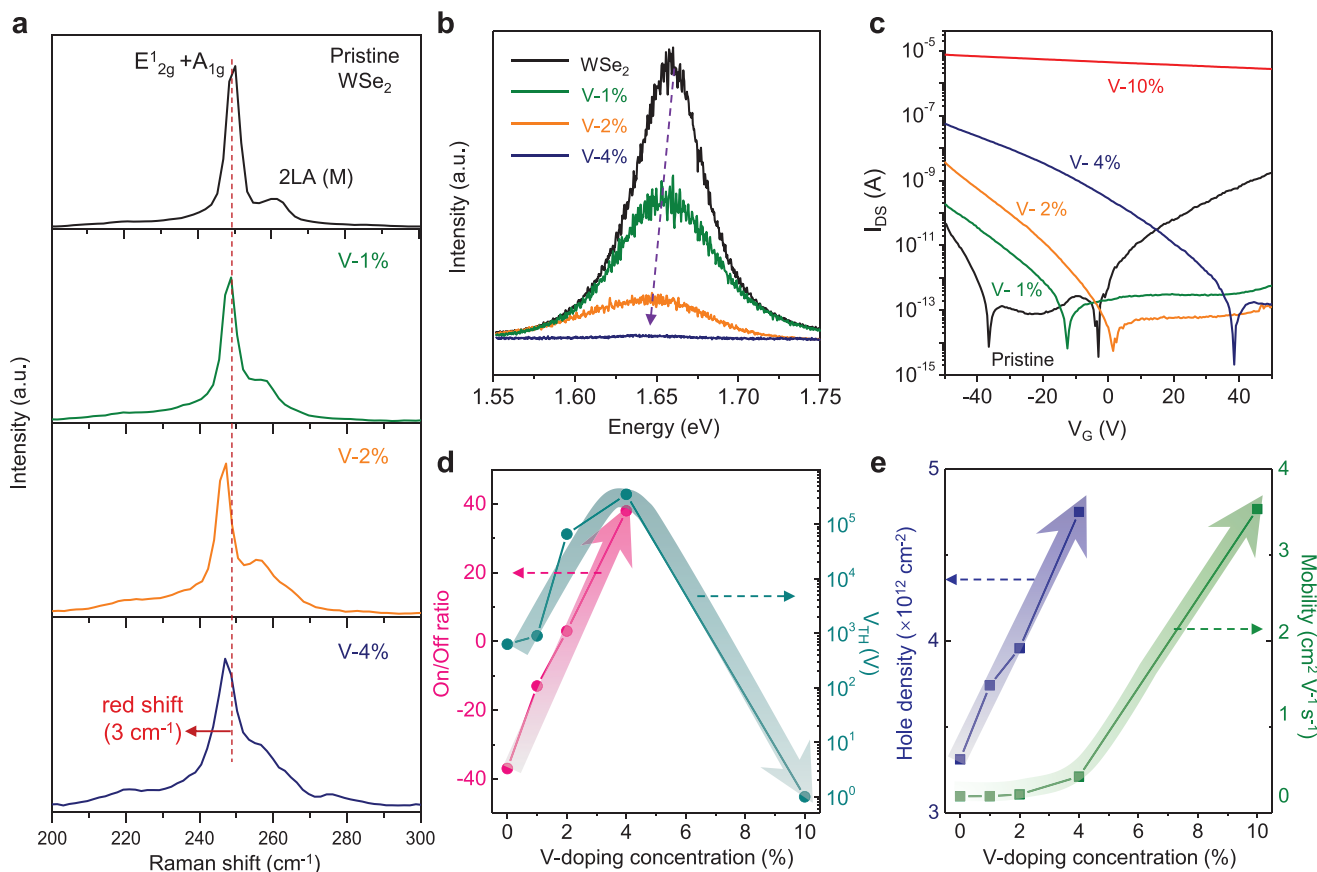


Figure 2. p-Doping effect of V-substitution to WSe₂. a,b) Raman (a) and PL (b) spectra of V-doped WSe₂ monolayer at an excitation wavelength of 532 nm in terms of V-doping concentration. c) I_{DS} - V_G curves of V-doped WSe₂ transistors under $V_{DS} = 0.5$ V. d,e) On/off ratio (I_{L-50V}/I_{off}) and threshold voltage (d) field-effect hole mobility and intrinsic hole carrier concentration (e) as a function of V-doping concentration.

side, while the on/off current ratio reaches to $\approx 10^5$ at V-4% doping (Figure 2d). At a high V-doping concentration of 10%, the FET shows a heavily degenerate behavior with a negligible current modulation within the entire range of gate voltages (-50 to 50 V). Moreover, the hole mobility and carrier density increase as a function of V-doping concentration (Figure 2e; for the detail calculation, see Note S1 and S2, and Figure S6, Supporting Information).

We next construct two-terminal p-n diodes by vertically stacking the grown V-doped WSe₂ as a p-type and the exfoliated SnSe₂ as an n-type, as shown schematically (Figure 3a). The source and drain electrodes of Pd are deposited on the V-WSe₂/SnSe₂ heterostructure (Figure 3b; for the detail fabrication process, see Figure S7, Supporting Information).^[20,21] The thickness of SnSe₂ flakes in all diodes were 11–17 nm, as confirmed by AFM for the controlled experiments (Figure S8, Supporting Information). The Fermi level of the V-doped WSe₂ is modulated with V-doping concentration. Meanwhile, the multilayer SnSe₂ FET device reveals a heavily degenerate n-type behavior by the high n-type current of $\approx 100 \mu\text{A } \mu\text{m}^{-1}$ (Figure S8, Supporting Information). As V-doping concentration increases, the Fermi level is shifted toward the valence band edge up to V-4% and further into the valence band at V-10%, implying a near broken-gap band alignment with n-type SnSe₂ (Figure 3c).^[22]

We now demonstrate a variety of diode behaviors in V-WSe₂/SnSe₂ devices by varying the V-doping concentration at room temperature. The diode device at V-1% (Figure 3d) as well as the pristine WSe₂ device (Figure S9, Supporting Information) shows a typical forward rectifying diode behavior. The Fermi level of V-1% or pristine WSe₂ sample locates near the middle of the bandgap, allowing for a large current flow from WSe₂ to SnSe₂ driven by the forward bias while suppressing the reverse current. At V-2% doping, the Fermi level downshifts closer to the valence band. An asymmetric ambipolar transistor is manifested, preferring a backward rectifying diode behavior (Figure 3e). A more prominent device is constructed at V-4% doping. NDR behavior is clearly observed in the forward bias region from 2.8 to 5.0 V with a peak-to-valley ratio of 1.6 (Figure 3f). When the Fermi level of n-SnSe₂ is located above the valence band edge of V-doped WSe₂, the current drops gradually with the reduction in electron band-to-band tunneling (BTBT). At a high V-doping concentration (V-10%), the V-doped WSe₂ turns into the degenerate state with the Fermi level getting inside the valence band, which is confirmed by the subtle gating effect in the entire range of gate bias (Figure S10, Supporting Information). In such a case, the diode behaves as an ohmic resistance due to the presence of both degenerate n-SnSe₂ and degenerate p-WSe₂ (Figure 3g). The contact issues of resistance are excluded

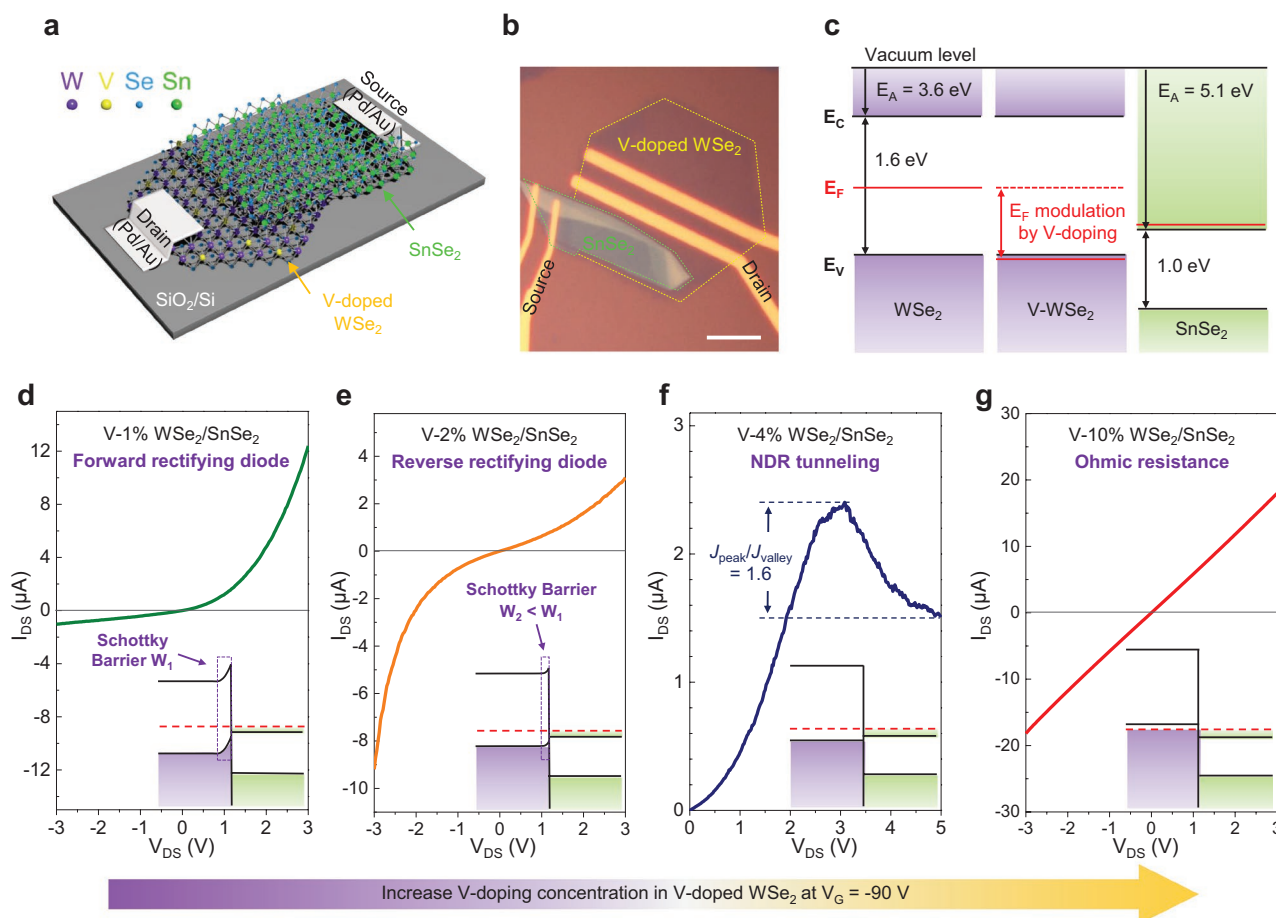


Figure 3. Device structure and electrical performance of two-terminal diodes from CVD-synthesized V-doped WSe₂ and exfoliated SnSe₂. a) Schematic showing p-type V-WSe₂ at the bottom and n-type SnSe₂ on top, with both Pd contacts. b) Optical image of a representative device of V-2%. The scale bar is 10 μm . c) Band structures of V-WSe₂ and SnSe₂ with their electron affinities and bandgaps, suggesting near broken-gap alignment in the heterostructures. The Fermi levels are estimated from the threshold voltages. d–g) $I_{\text{DS}}-V_{\text{DS}}$ characteristics (linear scale at room temperature) with diverse diode behaviors corresponding to different V-doping concentrations in WSe₂. The inset figures are the band alignments simulated by Poisson's equation when SnSe₂ was brought into contact with V-doped WSe₂. All the devices are gated at -90 V.

here owing to the near-ohmic contact in V-doped WSe₂/Pd junctions (Figure S11, Supporting Information).^[10]

The band bending of alignments in heterostructures occurs in planar V-doped WSe₂ rather than the vertical V-WSe₂/SnSe₂ junction due to the thin thickness of monolayer V-WSe₂^[9,23] and low carrier density while the V-doping concentration is low. In view of the SnSe₂, which is degenerate and appears in metal, the band bending can be negligible. According to Poisson's equation, band alignments in the V-WSe₂/SnSe₂ heterojunctions are calculated upon different V-doping concentrations in V-doped WSe₂ (insets in Figure 3d–g; the details are provided in Note S3 and Figure S12, Supporting Information). The band bending (depletion region width) in the V-WSe₂ at low V-doping concentration are much longer than the atomic thickness of monolayer V-WSe₂, inducing the bands bend along the planar direction in V-WSe₂. The initial type II band alignment is preserved at low V-concentration (up to V-4%). Further increase V-concentration to V-10%, the width of the depletion layer decreases, eventually obtains a type III broken-gap alignment, implying the evolution of the carrier transport mechanism.

The aforementioned appealing device characteristics modulated by V-doping concentration are further improved for high device efficiency via electrostatic doping. The $I_{\text{DS}}-V_{\text{DS}}$ output curves are measured at various gate biases from -30 to -110 V (Figure 4). In addition to the p-type V-doping in WSe₂, the hole doping by the negative gate bias is applied to boost the efficiency without altering the device characteristics.^[24,25] The Fermi level in heavily n-doped SnSe₂ is strongly pinned. In the forward rectifying diode at 1% V-WSe₂ (Figure 4a), a high current is achieved at a large negative gate bias; however, it yields a high rectification ratio ($I_{3\text{V}}/I_{-3\text{V}}$) of 135 at $V_{\text{G}} = -30$ V, benefiting from the suppression of minority carrier drift in the off-state compared to that at $V_{\text{G}} = -110$ V. For the backward rectifying diode, the rectification ratio ($I_{-3\text{V}}/I_{3\text{V}}$) increases up to 2.8 at $V_{\text{G}} = -110$ V (Figure 4b). Meanwhile, the peak-to-valley ratio in NDR diode rises up to 2.3 at $V_{\text{G}} = -110$ V (Figure 4c). Similar device characteristics are obtained from other four devices with the corresponding V-doping concentration (Figure S13, Supporting Information). Figure 4d summarizes the improved device performances in terms of the rectification ratio (forward and backward) and peak-to-valley ratio with the gate bias

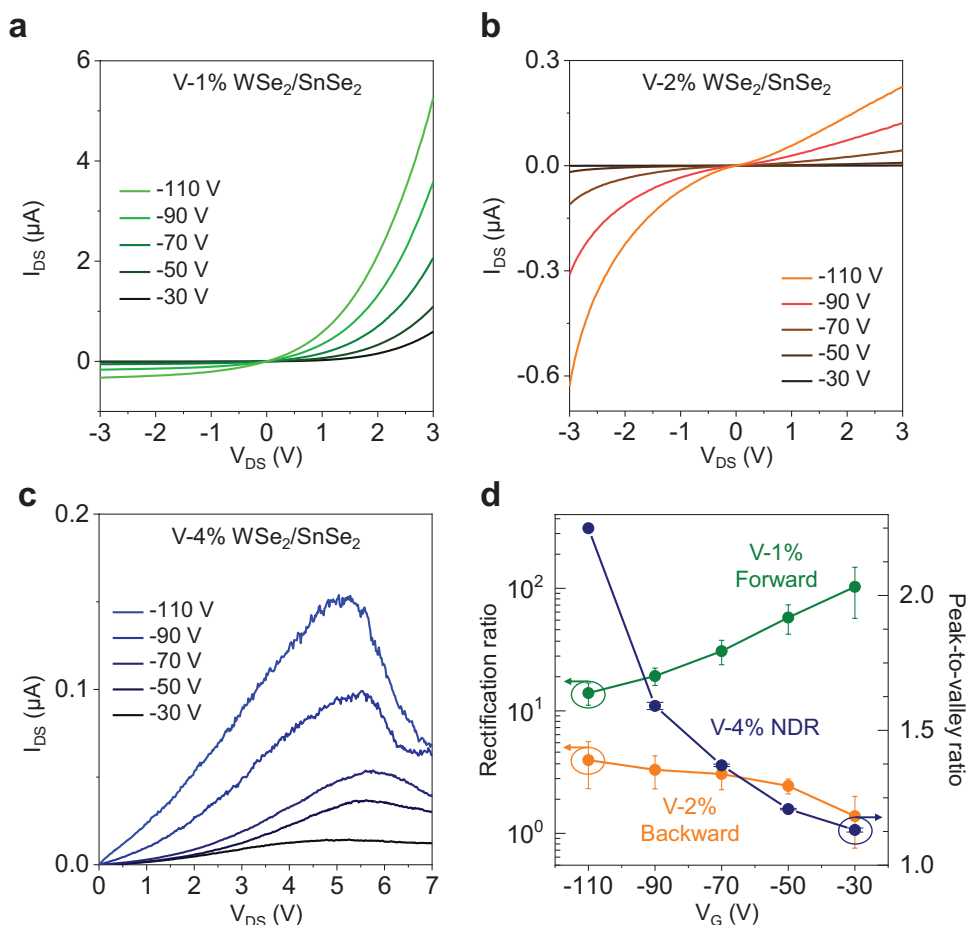


Figure 4. Gate-tunable behaviors of V-doped WSe₂/SnSe₂ devices. a–c) I_{DS} – V_{DS} output curves of the forward rectifying diode, backward rectifying diode, and NDR tunneling under various gate biases. d) Device rectification ratio (I_{3V}/I_{-3V} in the forward direction, and I_{-3V}/I_{3V} in the backward direction) and NDR peak-to-valley ratio as a function of gate bias with reasonable error bars, demonstrating the reproducibility.

modulation. The reproducibility of our devices is ensured with reasonable error bars. As the negative gate bias increases, the rectification ratio degrades in the forward direction but is improved in the backward diode. The peak-to-valley ratio is inversely proportional to the gate bias with a maximum value of 2.3, especially with 4% V-WSe₂, higher than the value of 1.8 obtained from the WSe₂/SnSe₂ without h-BN insulating layer.^[26] This is ascribed to the increase in available energy states for tunneling, modulated by the high V-doping concentration.

Diverse diode behaviors indicate different carrier transport mechanisms. In the pristine WSe₂ or V-1% device, there is a depletion region in WSe₂ nearby the heterointerfaces (Figure S12a, Supporting Information). Under a positive drain bias (Figure 5a), the energy bands of WSe₂ shift down, thinning the barrier width. The majority electrons residing in the SnSe₂ conduction band, therefore, overcome the triangular barrier and migrate to the other side, generating a forward current. This process is dominated by the thermionic Fowler–Nordheim (FN) tunneling or FN tunneling, which can be modeled as^[27]

$$I(V) = \frac{Aq^3 m V^2}{8\pi h \phi W^2 m^*} \exp\left[\frac{-8\pi\sqrt{2m^*} \phi^{3/2} W}{3hqV}\right] \quad (1)$$

where A is the effective contact area, q is the unit charge, h is the Planck's constant, m and m^* are the electron free mass and effective mass, respectively, and W and ϕ are the barrier width and height, respectively. Based on the derivation, $1/V$ follows $\ln(I/V^2)$ in the relationship of negative linear correlation. The $\ln(I/V^2)-1/V$ plot (Figure S14, Supporting Information) is calculated from Figure 3d, confirming FN tunneling mechanism with the negative slope at a high drain bias (near zero).^[28] When $V_{DS} < 0$, the minority carrier drift starts to dominate the current flowing from SnSe₂ to WSe₂. Since the intrinsic concentration of minority carriers is limited and the diffusion length stays identical, the reverse current is small and tends to saturate, showing a forward rectifying behavior.

The Fermi level in V-2% WSe₂ locates closer to its valence band edge compared to that in pristine or V-1% sample because of the V-doping effect, and further approaches at a high negative gate bias. When $V_{DS} > 0$, the SnSe₂ conduction band aligns with the forbidden band of WSe₂, resulting in the forward FN tunneling current as in pristine WSe₂ or V-1% device (upper panel in Figure 5b). The depletion region in V-2% WSe₂ is much thinner but the built-in potential shows $\phi_b < \phi_a$ (marked in upper panels in Figure 5b and Figure 5a, respectively) as indicated in the calculated band alignment (Figure S12b,

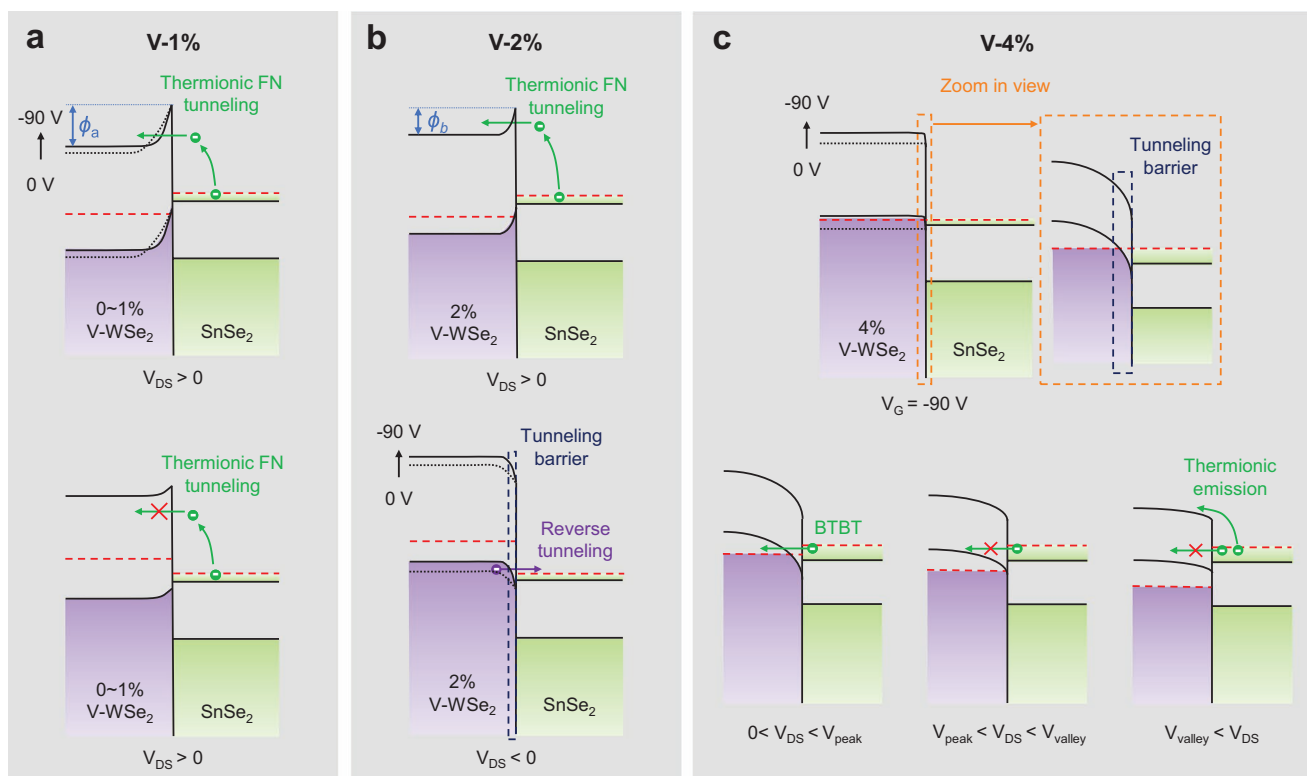


Figure 5. Band alignments in various V-doped $\text{WSe}_2/\text{SnSe}_2$ devices. a) In the pristine WSe_2 and V-1% devices, driven by a positive drain bias, the majority electrons thermally excited from the SnSe_2 conduction band dominate the carrier transport via FN tunneling, generating a forward rectifying current. ϕ_a is the built-in potential. b) In the V-2% device, when $V_{\text{DS}} < 0$, electrons tunnel from the WSe_2 valence band into the empty conduction band of SnSe_2 by the band-to-band tunneling, exhibiting a backward rectifying behavior. ϕ_b is the built-in potential. c) In the V-4% WSe_2 , the Fermi level is expected to enter its valence band at a high negative gate due to the strong accumulation of holes. A tunneling channel is thereby established in the device for the band-to-band tunneling. Electrons are transferred across the tunneling barrier at the interface with an NDR behavior in the positive drain bias region.

Supporting Information), leading to a smaller forward current. Meanwhile, the energy bands of V- WSe_2 move upward when $V_{\text{DS}} < 0$, which allows for electrons from the WSe_2 valence band heading to the empty states in the SnSe_2 conduction band, generating a backward BTBT current. In this case, the thinner tunneling barrier promotes the efficient band-to-band tunneling in backward, whereas the lower built-in potential reduces the forward thermionic current, forming the backward rectifying behavior.

As V-4% WSe_2 -FET shows a strong p-type property, the original Fermi level is expected to locate near its valence band edge (Figure S12c, Supporting Information). Gated by a high negative bias (Figure 5c), the electrostatic doping pushes the Fermi level into the valence band due to the strong accumulation of holes, leaving large amounts of empty states. Driven by a small positive drain bias ($0 < V_{\text{DS}} < V_{\text{peak}}$), electrons from the SnSe_2 conduction band tunnel into the WSe_2 valence band owing to the overlap of equivalent energy states, and thus the forward tunneling current is generated. This current then reaches its peak when the SnSe_2 Fermi level matches the WSe_2 valence band edge, where the populations of equivalent states in two sides reach a maximum. A further increase in V_{DS} induces the reduction in the tunneling current, namely, NDR behavior. The upper part of the SnSe_2 conduction band aligns with the forbidden band of WSe_2 , thereby prohibiting the BTBT of electrons, and the overlap region shrinks continuously until

vanishes. Normally, I_{DS} would pick up after the valley point and be dominated by the thermionic emission. However, this part is absent in some of our devices, which is probably because of the tunneling barrier or large hysteresis. Most of the drain bias drops at the wide interface barrier, which results in the appearance of peak current at a high V_{DS} . In addition, the large hysteresis in the V-4% WSe_2 transistor (Figure S15a, Supporting Information) indicates the carrier trapping or scattering, which severely impedes the majority carrier diffusion. Apart from this, device with apparent I_{DS} pick-up behavior is provided in Figure S15b of the Supporting Information with a peak-to-valley ratio of 1.4.

In conclusion, we have shown the tunable quantum tunneling in the V-doped $\text{WSe}_2/\text{SnSe}_2$ heterostructure by varying the doping concentration at room temperature. A liquid precursor containing transition metal and vanadium dopant atoms is used in our approach to synthesize V- WSe_2 via CVD. The precise control of the doping concentration efficiently modulates the Fermi level position, achieving the p-type doping effect. This method induces the substitution of W by V, endowing a proper band bending for the desired band alignment, which determines the charge transfer across the heterostructure. We theoretically probed the variation in the band alignment upon to different V-doping concentration, where the intrinsic type-II band alignment is preserved at low concentration, and evolves into the type-III broken-gap alignment by heavily doping.

Hence, the realization of diverse functions with the forward rectifying diode, backward rectifying diode, NDR diode, and ohmic resistance has been illustrated. They therefore suggest the promising opportunities to be applied for future integration in the scaled electronics.

Experimental Section

Preparation of Liquid Precursor: The precursor solution was prepared by mixing four types of water-based solutions: (i) tungsten precursor (ammonium metatungstate: $\text{NH}_4\text{H}_2\text{W}_{12}\text{O}_4$), (ii) vanadium precursor (ammonium metavanadate: NH_4VO_3), (iii) promoter (sodium hydroxide: NaOH), and (iv) medium solution (iodixanol). The V-doping concentration in V-doped WSe_2 was adjusted by mixing the molar ratio of tungsten to vanadium precursor. The mixed solution was then coated onto SiO_2/Si wafer by spin-casting at 3000 rpm for 1 min.

Synthesis of V-Doped WSe_2 by CVD: An atmospheric CVD system with two-zone furnaces was used for separately controlling the temperature of the selenium source and reaction zone. Here, 0.5 g of selenium (Sigma, 209643) was introduced, while the precursor-coated substrate was placed in the reaction zone. For the growth of V-doped WSe_2 , the selenium zone was heated up to 400 °C at a rate of 50 °C per min. At the same time, the substrate zone was set to 760 °C. Nitrogen gas of 500 sccm and hydrogen gas of 25 sccm were injected during CVD.

TEM Measurement: TEM and ADF-STEM images were taken by a probe aberration-corrected JEM ARM 200F machine, operated at 80 kV for high-resolution TEM observations. The imaging time was set within 1 min under a high-magnification STEM mode to avoid beam damage on the monolayer V-doped WSe_2 samples.

Device Fabrication: The fabrication of the V- $\text{WSe}_2/\text{SnSe}_2$ devices consists of three steps: (i) pristine WSe_2 or V- WSe_2 flakes transfer, (ii) exfoliation of SnSe_2 flakes and dry transfer in the glovebox, and (iii) electrode metallization. First, the CVD grown WSe_2 flakes were transferred onto a 300 nm SiO_2/Si substrate by wet-etching using hydrofluoric acid. Second, the SnSe_2 flakes were exfoliated using a Scotch tape onto a polypropylene carbonate (PPC)/300 nm SiO_2/Si substrate, visualizing the thin and large-size flakes. The desired SnSe_2 flake along with the PPC film was then mechanically peeled off from the silicon substrate and attached onto a polydimethylsiloxane stamp. Next, SnSe_2 and WSe_2 were brought together, followed by poly methyl methacrylate spin-coating prior to performing electrode metallization for protection. All the manipulations of SnSe_2 flakes were performed in the glovebox. Finally, the source (S) and drain (D) contacts (Pd/Au of 5/50 nm) for both WSe_2 and SnSe_2 were patterned by e-beam lithography prior to metal deposition. The entire fabrication process was schematically illustrated in Figure S7 of the Supporting Information.

Device Characterization: Electrical measurements were conducted by a probe station equipped with source/measurement units (Keithley 4200 and Agilent B2900A) at the high vacuum (10^{-6} Torr). AFM mapping analyses were recorded in a SPA400 (SEIKO) system. NT-MDT Raman & PL spectroscopy (532 nm laser) were used to study the samples.

Supporting Information

Supporting Information is available from the Wiley Online Library or from the author.

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Conflict of Interest

The authors declare no conflict of interest.

Keywords

2D tunneling heterojunctions, chemical vapor deposition, functional diodes, tin diselenide, vanadium-doped tungsten diselenide

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