


 Cite this: *Nanoscale*, 2020, **12**, 11448

Received 14th March 2020,

Accepted 16th May 2020

DOI: 10.1039/d0nr02084b

rsc.li/nanoscale

Transition from Schottky to Ohmic contacts in Janus MoSSe/germanene heterostructures†

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The performance of an electronic device based on a two-dimensional material is strongly affected by the contact with the metallic electrodes. In this article, we study the electronic properties of two-dimensional MoSSe in contact with a germanene electrode by first-principles calculations. The results show that the contact characteristics are significantly different for the two sides of MoSSe. Notably, for both sides in-plane tensile strain induces a transition from Schottky to Ohmic behavior. Increasing the thickness of MoSSe also leads to an Ohmic contact. We propose an effective route to high performance MoSSe electronic devices.

Introduction

Semiconductor/metal heterojunctions are of critical importance for electronic devices. In this context, van der Waals heterostructures consisting of stacked two-dimensional materials have opened up a new paradigm to design the interface properties.^{1–5} As an Ohmic contact with low resistance is the ultimate goal to achieve high performance devices,⁶ the Schottky barrier height (SBH; energy barrier that must be overcome by charge carriers to cross the interface; positive value indicating formation of a Schottky instead of an Ohmic contact) becomes a key parameter.^{7,8}

Two-dimensional Janus MoSSe has first been prepared by Lu and coworkers in 2017.⁹ Similar to MoS₂, its band gap and carrier mobility show a distinct thickness-dependence.¹⁰ On the other hand, the intrinsic dipole of MoSSe (structural symmetry of MoS₂ is broken) modifies the electronic properties.¹¹ A number of theoretical studies have shown that two-dimensional Janus materials perform well in gas sensors,^{12,13} photocatalytic water splitting,^{14–16} field-effect transistors,¹⁷ and solar cells.^{18,19} However, practical use of MoSSe in a device will

require a contact with metal electrodes to enable injection of carriers.²⁰ For this reason, we investigate in the present work van der Waals heterostructures constructed from MoSSe (considering both the S and Se sides) and germanene (Ge). Ge is chosen as electrode, because it is a Dirac metal with perfect lattice match to MoSSe.²¹ Employing first-principles calculations, we explore the electronic properties of the heterostructures, focusing on the SBH and band alignment, which are critical for the device performance. We analyze how Ge interacts with the different sides of MoSSe and determine the resulting contact characteristics.

Computational methods

All the calculations are performed using the Vienna *Ab initio* Simulation Package.²² The generalized gradient approximation (GGA) of Perdew–Burke–Ernzerhof²³ is applied to the exchange correlation functional and the DFT-D3 correction²⁴ is used to account for the van der Waals interaction. The plane-wave cutoff energy is set to 500 eV and a Monkhorst–Pack²⁵ 9 × 9 × 1 k-mesh is utilized for Brillouin-zone integration (5 × 5 × 1 in the structural relaxation). The convergence criteria are set to 0.01 eV Å⁻¹ for the maximal residual force and 10⁻⁶ eV for the total energy. Combining a 4 × 4 × 1 supercell of Ge with 32 germanium atoms and a 5 × 5 × 1 supercell of MoSSe with 75 atoms leads to a small lattice mismatch of less than 0.5%. In each heterostructure a vacuum layer of 15 Å thickness is added for ensuring that there is no interaction between periodic images in the out-of-plane direction.

Results and discussion

We first investigate the geometries of Ge and monolayer MoSSe, obtaining the lattice parameters 4.04 and 3.25 Å after optimization, respectively, in good agreement with previous studies.^{26,27} In contrast to MoS₂, MoSSe has an intrinsic dipole pointing from Se to S with a calculated dipole moment of $D =$

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†Electronic supplementary information (ESI) available. See DOI: 10.1039/d0nr02084b

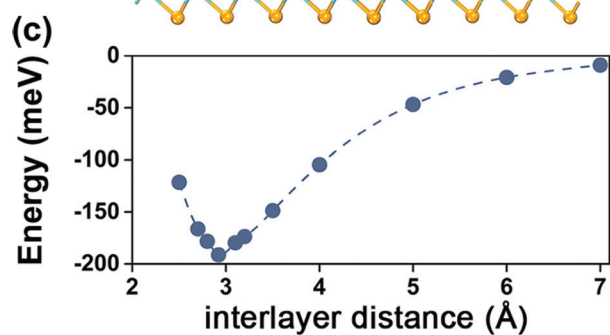
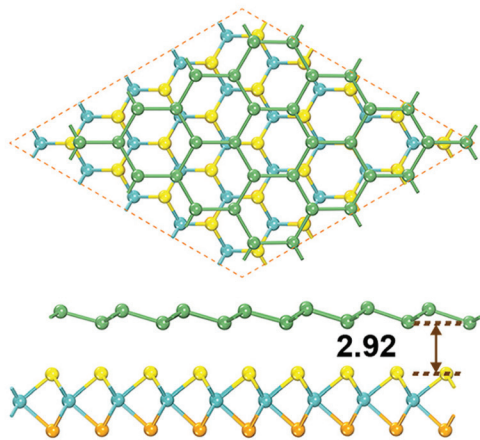


0.25 Debye, which is also consistent with previous results.²⁸ Both the S and Se sides of MoSSe can be in contact with Ge to form a heterostructure, namely, SeMoS/Ge with S–Ge interface and SMOSe/Ge with Se–Ge interface. To verify stability of the heterostructures, the binding energy

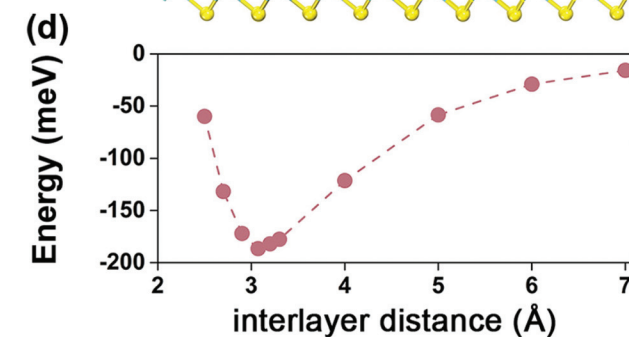
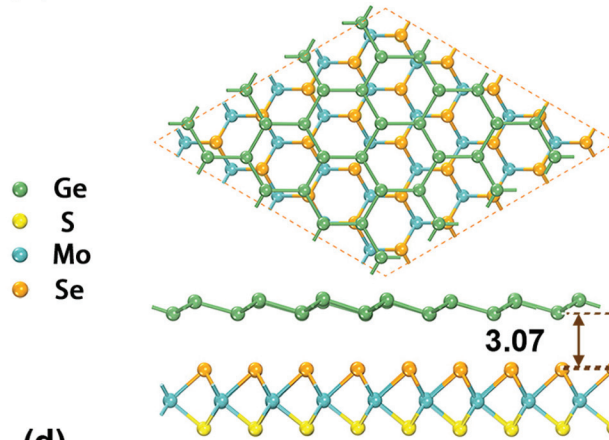
$$E_b = [E_{\text{Ge}} + E_{\text{MoSSe}} - E_{\text{hetero}}]/N \quad (1)$$

is calculated, where E_{Ge} , E_{MoSSe} , and E_{hetero} are the total energies of Ge (consisting of N atoms), MoSSe, and the heterostructure. By configuration scanning (see Fig. S1† for the considered stacking configurations), the energetically favorable heterostructures are determined. They are shown in Fig. 1(a and b). We obtain equilibrium interlayer distances of 2.92 Å for SeMoS/Ge and 3.07 Å for SMOSe/Ge, see Fig. 1(c and d).

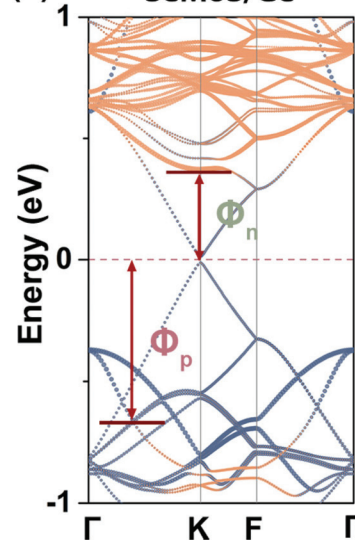
(a) SeMoS/Ge



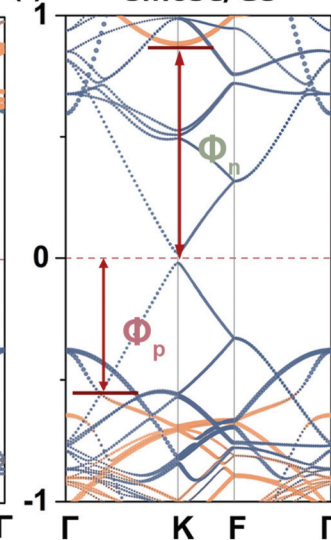
(b) SMOSe/Ge



(e) SeMoS/Ge



(f) SMOSe/Ge



(g)

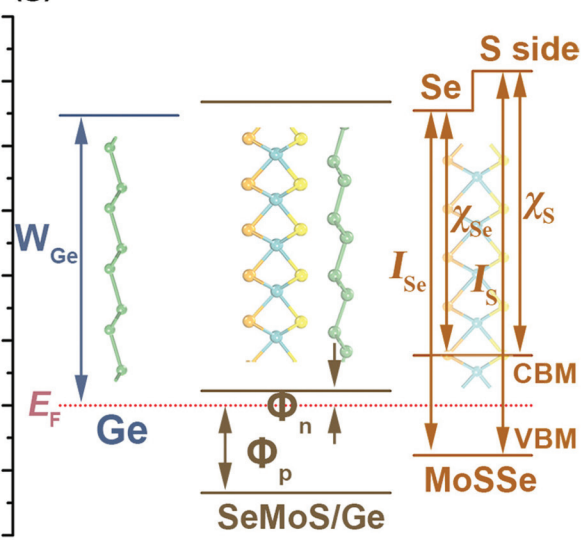


Fig. 1 Top and side views of (a) SeMoS/Ge and (b) SMOSe/Ge. The dependence of the binding energy on the interlayer distance is shown in (c) and (d). Weighted band structures are given in (e) and (f) with orange and blue colors indicating the contributions of Ge and MoSSe, respectively. Corresponding densities of states are shown in Fig. S5.† (g) Energy band diagram of Ge, monolayer MoSSe, and SeMoS/Ge.



The corresponding weighted band structures are shown in Fig. 1(e and f). Orange and blue colors represent the contributions of MoS₂ and Ge, respectively. Comparing to the pristine component materials (Fig. S2[†]), the band structures of the heterostructures are more than simple superpositions due to hybridization of the atomic orbitals at the interfaces. The effects of spin-orbital coupling have been tested carefully for MoS₂ and Ge, showing no significant influence on the band dispersions (Fig. S3[†]). Therefore, spin-orbit coupling is not taken into account for the heterostructures. According to Fig. 1 (e and f), a band gap of 25 and 41 meV opens at the Dirac point for SeMoS/Ge and SMOSe/Ge, respectively. Interestingly, the band gap of the semiconducting layer (measured from the highest valence to the lowest conduction state dominated by MoS₂) in SeMoS/Ge is 1.00 eV, *i.e.*, smaller than in the case of pristine MoS₂ (1.55 eV), while in SMOSe/Ge it is 1.45 eV. This

trend does not reflect the typical behavior of two-dimensional heterostructures²⁹ due to the fact that the strong hybridization seen in Fig. 1(e and f) between the component layers is unusual.^{30–32}

Our results show that an n-type Schottky contact is formed in SeMoS/Ge and a p-type Schottky contact in SMOSe/Ge. The SBHs of the heterostructures are given by³³

$$\begin{aligned}\Phi_n &= W_{\text{SeMoS/Ge}} - \chi_S, \Phi_p = I_S - W_{\text{SeMoS/Ge}}, \\ \Phi_n &= W_{\text{SMoSe/Ge}} - \chi_{\text{Se}}, \Phi_p = I_{\text{Se}} - W_{\text{SMoSe/Ge}},\end{aligned}\quad (2)$$

where $W_{\text{SeMoS/Ge}}$ and $W_{\text{SMoSe/Ge}}$ are the work functions. Moreover, $\chi_{\text{S/Se}}$ and $I_{\text{S/Se}}$ denote the electron affinity and ionization potential of the S/Se side of MoS₂, respectively. In Fig. 1(g) we show for SeMoS/Ge (S side in contact with Ge), the energy band diagram at large interlayer distance (no interaction

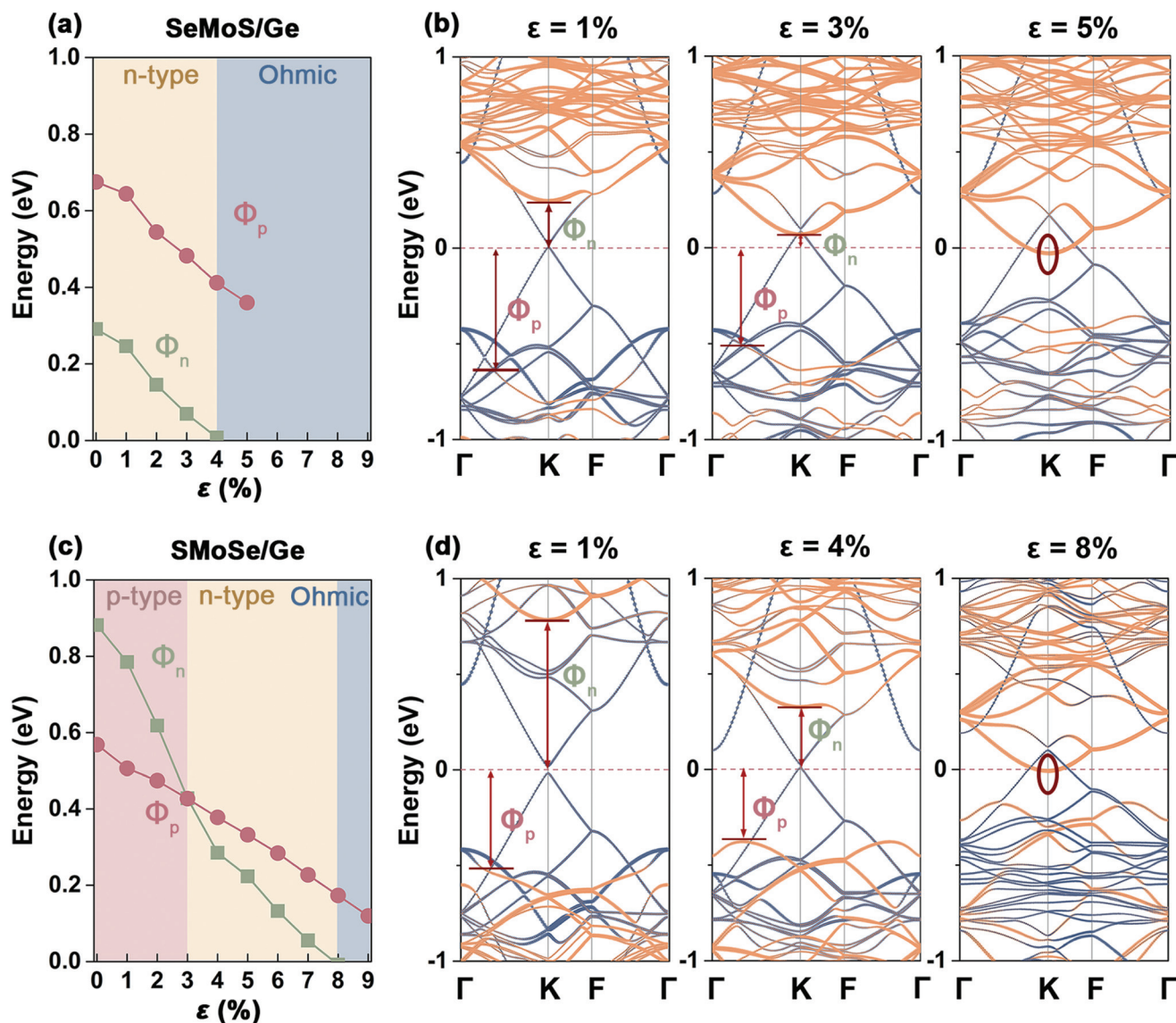


Fig. 2 SBHs of (a) SeMoS/Ge and (c) SMOSe/Ge as functions of strain. Weighted band structures are given in (b) and (d) for different strains with orange and blue colors indicating the contributions of Ge and MoS₂, respectively. Corresponding densities of states are shown in Fig. S5.[†]



between the components), *i.e.*, there is no effect of charge transfer. A quantitative calculation at an interlayer distance of 7 Å results in $W_{\text{SeMoS/Ge}} = 4.68$ eV, $I_S = 5.93$ eV, and $\chi_S = 4.38$ eV. Therefore, $\Phi_n = 0.30$ eV and $\Phi_p = 1.25$ eV, consistent with the values measured in the band structures (Fig. S4;† 0.23 and 1.34 eV).

A tunable SBH is essential to improve the transition of charge carriers between the two components of a heterostructure. To this aim, we explore the effect of tensile strain (ϵ), finding that for increasing ϵ the interaction between MoS₂ and Ge intensifies. According to Fig. 2(a), both Φ_n and Φ_p of SeMoS/Ge decrease. The n-type SBH vanishes at $\epsilon \sim 4\%$, *i.e.*, the contact becomes Ohmic. Band structures at different strains are shown in Fig. 2(b), demonstrating that the conduction band minimum of MoS₂ shifts dramatically to lower energy (relative to the Fermi energy) for increasing ϵ , while the Dirac point of Ge shifts to higher energy similar to isolated Ge.³⁴ A more complicated situation is encountered in SMOSe/Ge, see Fig. 2(c). Faster decrease of Φ_n than Φ_p results in an

n-type SBH for strain above $\epsilon \sim 3\%$ and an Ohmic contact is achieved at $\epsilon \sim 8\%$. According to the band structures in Fig. 2(d), similar to SeMoS/Ge, for increasing ϵ the Dirac point of Ge shifts to higher energy. We have confirmed our conclusions for the non-local van der Waals method (otherwise same computational setup),^{35–38} finding that the interlayer distance increases significantly from 2.92 Å to 3.50 Å for SeMoS/Ge and from 3.07 Å to 3.67 Å for SMOSe/Ge, while Φ_n (0.36 eV instead of 0.29 eV for SeMoS/Ge; 0.89 eV instead of 0.88 eV for SMOSe/Ge) and Φ_p (1.02 eV instead of 0.67 eV for SeMoS/Ge; 0.71 eV instead of 0.57 eV for SMOSe/Ge) increase only slightly and the contact type is not modified.

To obtain insight into the Schottky-to-Ohmic transition, the strain-dependence of the electrostatic potential is addressed in Fig. 3(a and e) (vacuum level set to zero energy). While the shortened interlayer distance of the heterostructures under tensile strain results in minor shifts of the peaks, the heights of the peaks decrease gradually. We observe in Fig. 3(b and f) that the charge transfer from Ge to MoS₂ (positive Δq) gener-

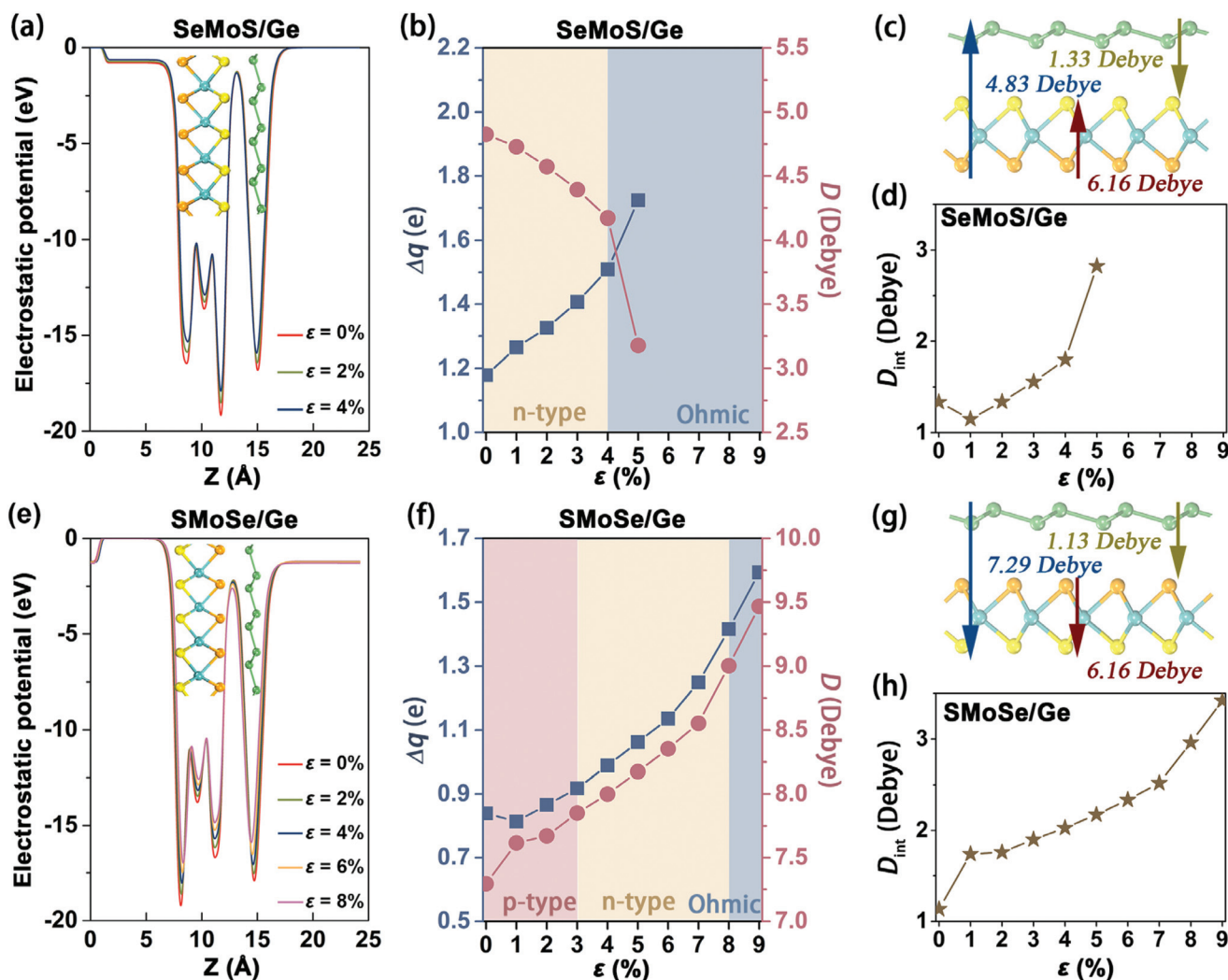


Fig. 3 Electrostatic potentials of (a) SeMoS/Ge and (e) SMOSe/Ge for different strains. Corresponding charge transfers and dipole moments as functions of strain are given in (b) and (f), dipole moments without strain in (c) and (g), and interface dipole moments as functions of strain in (d) and (h).



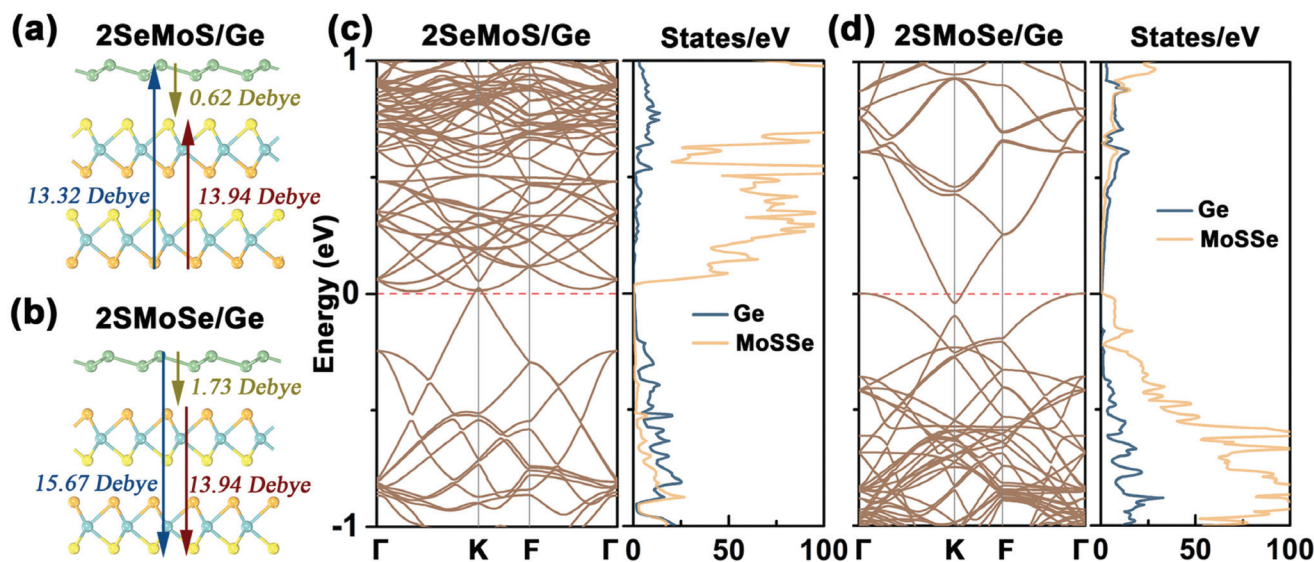


Fig. 4 Dipole moments of (a) 2SeMoS/Ge and (b) 2SMoSe/Ge. Corresponding band structures and densities of states are shown in (c) and (d).

ally increases under tensile strain (faster for higher strain). The value is larger for SeMoS/Ge than SMoSe/Ge due to the fact that S is more electronegative than Se. In addition, Fig. 3(b and f) shows that the total dipole moment points from Se to S ($D > 0$) for both heterostructures, being larger for SMoSe/Ge than SeMoS/Ge. We have $D = D_0 + D_{\text{int}}$, where D_0 is the intrinsic dipole moment of MoS₂ (6.16 Debye for the $5 \times 5 \times 1$ supercell) and D_{int} is the interlayer dipole moment (1.33 and 1.13 Debye for SeMoS/Ge and SMoSe/Ge without strain, see Fig. 3(c and g)). D of SeMoS/Ge decreases under tensile strain, see Fig. 3(b), whereas D of SMoSe/Ge increases, see Fig. 3(f). Though the direction of D is different in the two heterostructures (pointing from MoS₂ to Ge for SeMoS/Ge and from Ge to MoS₂ for SMoSe/Ge), D_{int} always points from Ge to MoS₂. D_{int} increases under tensile strain, as Δq increases, see Fig. 3(d and h). At $D_{\text{int}} = 2.82$ Debye for SeMoS/Ge and $D_{\text{int}} = 2.96$ Debye for SMoSe/Ge an Ohmic contact is achieved. Therefore, D_{int} controls the SBH.

As not only monolayer MoS₂ but also bilayer MoS₂ has an intrinsic dipole, we next address heterostructures of bilayer MoS₂ with Ge, see Fig. 4(a and b). As expected, D_0 increases in the case of bilayer MoS₂ (13.94 Debye for the $5 \times 5 \times 1$ supercell) as compared to the case of monolayer MoS₂. While D_{int} still points from Ge to MoS₂, it decreases to 0.62 Debye for 2SeMoS/Ge and increases to 1.73 Debye for 2SMoSe/Ge. The charge transfer is still directed from Ge to MoS₂ with values of 1.15 e for 2SeMoS/Ge and 0.78 e for 2SMoSe/Ge, which are similar to the case of monolayer MoS₂. According to Fig. 4(c and d), both heterostructures have a gapped Dirac cone near the Fermi energy, shifted as expected from the dipole (built-in electric field³⁹). We obtain for 2SeMoS/Ge $\Phi_n = 0.01$ eV, *i.e.*, an almost Ohmic contact, while for 2SMoSe/Ge the bands of the semiconducting layer touch the Fermi energy, *i.e.*, an Ohmic contact is formed.

Conclusion

We have systematically studied heterostructures of two-dimensional Janus MoSSe with germanene (Ge). An n-type Schottky contact is found to be formed for SeMoS/Ge and a p-type Schottky contact for SMoSe/Ge. A transition from Schottky to Ohmic behavior occurs under tensile strain ($\sim 4\%$ for SeMoS/Ge, $\sim 8\%$ for SMoSe/Ge), which is explained by modifications of the interface dipole. Of great interest from a technological perspective is the observation that an increased thickness of MoSSe (bilayer sufficient) directly results in an Ohmic contact. Our investigation thus demonstrates that Ge is an ideal electrode for contacting two-dimensional Janus MoSSe in electronic devices.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

The research reported in this publication was supported by funding from King Abdullah University of Science and Technology (KAUST).

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