

Intrinsic electrical transport properties of monolayer silicene and MoS₂ from first principlesXiaodong Li,¹ Jeffrey T. Mullen,² Zhenghe Jin,¹ Kostyantyn M. Borysenko,³ M. Buongiorno Nardelli,^{4,5} and Ki Wook Kim^{1,*}¹*Department of Electrical and Computer Engineering, North Carolina State University, Raleigh, North Carolina 27695-7911, USA*²*Department of Physics, North Carolina State University, Raleigh, North Carolina 27695-8202, USA*³*Department of Physics, Texas Southern University, Houston, Texas 77004, USA*⁴*Department of Physics and Department of Chemistry, University of North Texas, Denton, Texas 76203, USA*⁵*CSMD, Oak Ridge National Laboratory, Oak Ridge, Tennessee 37831, USA*

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The electron-phonon interaction and related transport properties are investigated in monolayer silicene and MoS₂ by using a density functional theory calculation combined with a full-band Monte Carlo analysis. In the case of silicene, the results illustrate that the out-of-plane acoustic phonon mode may play the dominant role unlike its close relative, graphene. The small energy of this phonon mode, originating from the weak sp^2 π bonding between Si atoms, contributes to the high scattering rate and significant degradation in electron transport. In MoS₂, the longitudinal acoustic phonons show the strongest interaction with electrons. The key factor in this material appears to be the Q valleys located between the Γ and K points in the first Brillouin zone as they introduce additional intervalley scattering. The analysis also reveals the potential impact of extrinsic screening by other carriers and/or adjacent materials. Finally, the effective deformation potential constants are extracted for all relevant intrinsic electron-phonon scattering processes in both materials.

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I. INTRODUCTION

Very recently, the attention to low-dimensional materials has expanded beyond the best-known example of this kind: graphene.¹⁻⁴ In particular, silicene⁵⁻⁸ and molybdenum disulfide⁹⁻¹² have gained much interest due to their unique properties in electronics, optoelectronics, and magnetics. Silicene is expected to share certain superior properties of graphene due to its structural similarity and the close position in the periodic table. More importantly, it is compatible with the current silicon-based technology and can be grown on a number of different substrates.¹³⁻¹⁵ On the other hand, atomically thin MoS₂ is a semiconductor with a finite band gap that ranges from approximately 1.3 to 1.9 eV depending on the thickness.⁹ It has been used as the channel material in field effect transistors with promising results.^{12,16} In addition, monolayer MoS₂ offers the possibilities of interesting spin and valley physics utilizing the strong spin-orbit coupling.¹⁷⁻²⁰

Characterization of electronic transport, particularly the intrinsic properties, is critical for assessing and understanding the potential significance of a material. In the case of silicene, many of the crucial parameters are presently unknown due to the brief history of this material. In comparison, notable advances have been made in MoS₂ lately. Experimental investigation of transistor characteristics claimed the channel mobilities ranging from ~ 200 cm²/Vs to ~ 1000 cm²/Vs at room temperature,^{12,16} while a theoretical study estimated an intrinsic phonon-limited value of ~ 410 cm²/Vs based on a first-principles calculation of electron-phonon interaction.²¹ However, questions remain regarding the intrinsic electron transport in MoS₂. For instance, those extracted from transistor current-voltage (I - V) measurements are indirect accounts and can be strongly affected by extrinsic factors, requiring caution as illustrated in the latest studies.²²⁻²⁴ Similarly, the latter work²¹ includes only the electronic states in the conduction band minima at the K points in the first Brillouin zone (FBZ); the impact of Q valleys located along the Γ - K symmetry

directions (which correspond to the minima of bulk MoS₂) were not considered. A detailed investigation is clearly called for.

In this paper, we examine intrinsic transport properties of monolayer silicene and MoS₂ by taking advantage of first-principles analysis and full-band Monte Carlo simulation. Along with the electronic band structure, the phonon spectra and electron-phonon coupling matrix elements are calculated for all phonon branches within the density functional theory (DFT) formalism.^{25,26} The obtained electron scattering rates are subsequently used in the Boltzmann transport equation to compute the intrinsic velocity-field characteristics with a full-band Monte Carlo treatment. The calculation results are compared with the available data in the literature and the key factors affecting electron transport in these materials elucidated. The investigation also provides the effective deformation potential constants extracted from the first-principles results.

II. THEORETICAL MODEL

Both monolayer silicene and MoS₂ are hexagonal crystals. To account for their delicate atomic structures accurately, the calculations are performed in the DFT framework, as it is implemented in the QUANTUM ESPRESSO package,²⁷ using ultrasoft pseudopotentials. A minimum of 35 Ry is used for the energy cutoff in the plane-wave expansion along with the charge truncation ~ 15 times larger. The generalized gradient approximation is used for the exchange-correlation potential for silicene, while the local density approximation is adopted for MoS₂. The momentum space is sampled on a $36 \times 36 \times 1$ Monkhorst-Pack grid with no offset (silicene) or on a $18 \times 18 \times 1$ grid (MoS₂) for electronic band calculation. The simulated cells are optimized in both cases until the atomic forces decrease to values less than 0.015 eV/Å.

Each phonon is treated as a perturbation of the self-consistent potential, created by all electrons and ions of the

system, within the linear response [i.e., density functional perturbation theory (DFPT)].²⁵ The calculation of the potential change due to this perturbation gives the value of the electron-phonon interaction matrix element²⁶:

$$\mathbf{g}_{\mathbf{q},\mathbf{k}}^{(i,j)v} = \sqrt{\frac{\hbar}{2M\omega_{v,\mathbf{q}}}} \langle j, \mathbf{k} + \mathbf{q} | \Delta V_{\mathbf{q},\text{SCF}}^v | i, \mathbf{k} \rangle, \quad (1)$$

where $|i, \mathbf{k}\rangle$ is the Bloch electron eigenstate with the wave vector \mathbf{k} , band index i , and energy $E_{i,\mathbf{k}}$; $\Delta V_{\mathbf{q},\text{SCF}}^v$ is the derivative of the self-consistent Kohn-Sham potential²⁵ with respect to atomic displacement associated with the phonon from branch v with the wave vector \mathbf{q} and frequency $\omega_{v,\mathbf{q}}$; and M is the atomic mass. Numerical calculations of lattice dynamics are conducted on a $18 \times 18 \times 1$ Monkhorst-Pack grid. Indices i, j are dropped hereinafter as only the first (lowest) conduction band is considered.

With the electron-phonon interaction matrix from the first-principles calculation, the scattering rate of an electron at state $|\mathbf{k}\rangle$ can be obtained by using Fermi's golden rule

$$\frac{1}{\tau_{\mathbf{k}}} = \frac{2\pi}{\hbar} \sum_{\mathbf{q},v} |\mathbf{g}_{\mathbf{q},\mathbf{k}}^v|^2 [N_{v,\mathbf{q}} \delta(E_{\mathbf{k}+\mathbf{q}} - \hbar\omega_{v,\mathbf{q}} - E_{\mathbf{k}}) + (N_{v,\mathbf{q}} + 1) \delta(E_{\mathbf{k}-\mathbf{q}} + \hbar\omega_{v,\mathbf{q}} - E_{\mathbf{k}})], \quad (2)$$

where $N_{v,\mathbf{q}} = [\exp(\hbar\omega_{v,\mathbf{q}}/k_B T) + 1]^{-1}$ is the phonon occupation number, k_B the Boltzmann constant, and T the temperature. As we are interested in the intrinsic scattering probability that is not limited to a specific carrier distribution (and thus, the Fermi level), our formulation assumes that all final electronic states are available for transition (i.e., nondegenerate) in the bands under consideration.

For transport properties, a Monte Carlo approach with full-band treatment is adopted. All of the details described above, including the wave-vector (\mathbf{k}, \mathbf{q}) dependence of the scattering matrix elements [e.g., Eq. (1)], are accounted for. This allows solution of the Boltzmann transport equation beyond the conventional relaxation time approximation.

III. RESULTS AND DISCUSSION

A. Monolayer silicene

Earlier first-principles studies have shown that the stable structure for monolayer silicene has a low-buckled configuration.^{4,6} While planar and high-buckled cases lead to imaginary phonon frequencies around the Γ point, indicating an unstable structure, the low-buckled construction provides well-separated phonon branches and positive frequencies. The origin of buckled geometry in silicene is the weakened π bonding of the electrons in the outer shell. Compared with graphene, which has very strong π bonding and planar geometry, the interatomic bonding distance is much larger in silicene, which decreases the overlap of p_z orbitals and dehybridizes the sp^2 states. Accordingly, the planar geometry can not be maintained. In our analysis, the lattice constant a is optimized to be 3.87 Å, with the buckling distance of 0.44 Å, in good agreement with Ref. 4.

Figure 1 shows the outcome of electronic and phononic band calculation in monolayer silicene. The Fermi velocity extracted from the Dirac cone is around 5.8×10^7 cm/s,

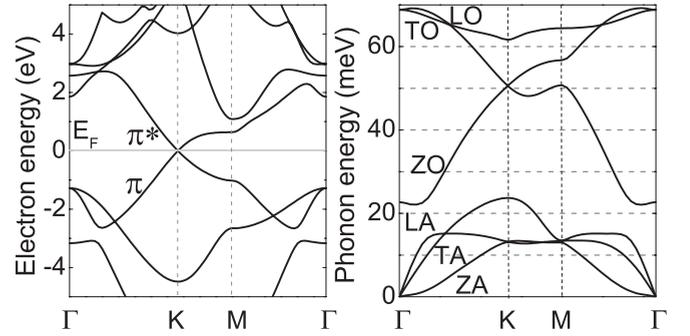


FIG. 1. Electronic and phononic band structures of monolayer silicene along the symmetry directions in the FBZ. The Dirac point serves as the reference of energy scale for electrons.

which is roughly one half of that in graphene [see bands π and π^* in Fig. 1(a)]. While this result is in agreement with other theoretical predictions,^{28,29} a value as high as 1×10^8 cm/s was also claimed in the literature.⁶ As for phonons in Fig. 1(b), six branches are identified with two atoms per unit cell. The transverse acoustic (TA) and longitudinal acoustic (LA) phonon dispersion relations are well approximated by sound velocities in the long-wavelength limit: $v_{\text{TA}} = 5.4 \times 10^5$ cm/s and $v_{\text{LA}} = 8.8 \times 10^5$ cm/s. Although the out-of-plane acoustic (ZA) phonon exhibits an approximate q^2 dependence near the center of the Brillouin zone, its sound velocity can also be estimated: $v_{\text{ZA}} = 6.3 \times 10^4$ cm/s. Interesting points to note in the phonon dispersion are the discontinuities in the frequency derivative of the highest optical branch that, similar to graphene, appear at the high-symmetry points Γ and K . These discontinuities are referred to as Kohn anomalies,^{28,30} induced by unusual screening of lattice vibrations by conduction electrons. Sharp cusps typically indicate strong electron-phonon coupling. Calculated phonon energies at the Γ , M , and K points in the FBZ are summarized in Table I.

The electron-phonon interaction matrix elements obtained for the electron at the Dirac point [i.e., $\mathbf{k} = (4\pi/3a, 0)$] are plotted in Fig. 2 as a function of phonon wave vector \mathbf{q} . Kohn anomalies, illustrated by the three peaks at three equivalent K points in the transverse optical (TO) mode and another at the zone center for the longitudinal optical (LO) branch, are not as distinct as those observed in graphene.²⁶ Overall, coupling of optical phonons with electrons appears to be relatively weak. In comparison, the acoustic phonons show much stronger interaction. Particularly striking is the large strength of ZA phonon coupling, unlike in graphene. Due to

TABLE I. Phonon energies (in units of meV) at the symmetry points for monolayer silicene.

Phonon modes	Γ	K	M
ZA	0	13.2	13.0
TA	0	23.7	13.4
LA	0	13.2	13.5
ZO	22.7	50.6	50.7
TO	68.8	50.6	56.7
LO	68.8	61.7	64.4

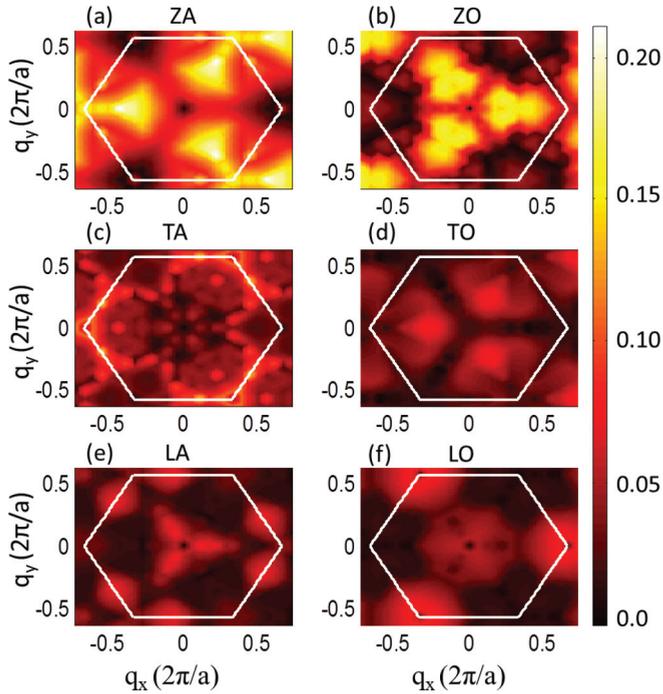


FIG. 2. (Color online) Electron-phonon interaction matrix elements $|G_{\mathbf{k}+\mathbf{q},\mathbf{k}}^v|$ (in units of eV) from the DFPT calculation in silicene for \mathbf{k} at the conduction band minimum K point [i.e., $(4\pi/3a, 0)$] as a function of phonon wave vector \mathbf{q} for all six modes v .

the buckled geometry (originating from the weak π bonding mentioned earlier), silicene does not maintain certain key characteristics of ideal planar lattice, especially the reflection symmetry with respect to the atomic plane. As such, the symmetry consideration, in which only the in-plane phonons can couple linearly to two-dimensional (2D) electrons,³¹ no longer applies. An increased role of ZA phonons is clearly expected.

The scattering rates calculated at room temperature ($T = 300$ K) are shown in Fig. 3. The result is plotted specifically for electrons with wave vector \mathbf{k} along the K - Γ direction. Since

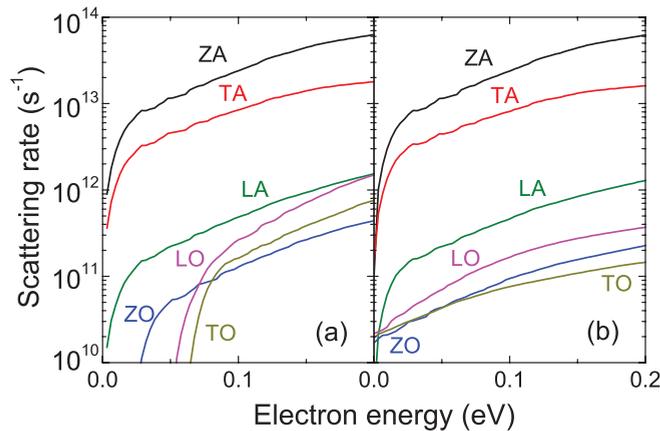


FIG. 3. (Color online) Electron scattering rates in silicene via (a) emission and (b) absorption of phonons calculated at room temperature. The electron wave vector \mathbf{k} is assumed to be along the K - Γ axis.

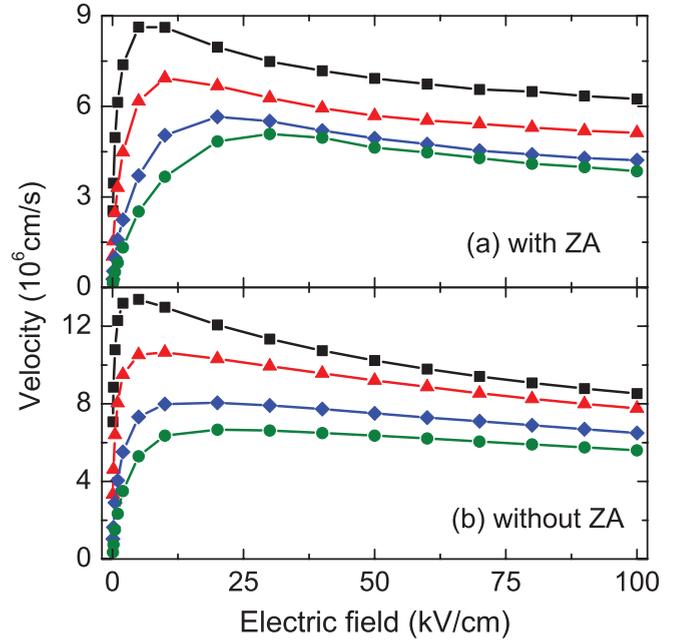


FIG. 4. (Color online) Drift velocity versus electric field in monolayer silicene obtained from a Monte Carlo simulation at different temperatures: 50 K (square), 100 K (triangle), 200 K (diamond), and 300 K (circle). The results in (a) consider the scattering by ZA phonons, while those in (b) do not.

the integration in Eq. (2) is over the entire FBZ, both intravalley ($K \rightarrow K$) and intervalley ($K \rightarrow K'$) transition events are included. As the interaction matrix elements illustrated above suggest, acoustic phonons have much larger scattering rates than optical modes. Specifically, the ZA branch provides the dominant contribution, which can be attributed to the observed large coupling strength as well as the small phonon energy near the zone center (i.e., a large occupation number $N_{ZA,\mathbf{q}}$). This also indicates that the scattering rates are very sensitive to the phonon energies (or, equivalently, the value of v_{ZA}). Since an accurate description of ZA dispersion in the long-wavelength limit requires a well-converged calculation with a sufficiently dense grid, care must be taken when evaluating accuracy of the data in the literature.^{6,29}

Figure 4(a) provides the drift velocity versus electric field at different temperatures obtained by full-band Monte Carlo simulations. The intrinsic mobility estimated from the figure is approximately $1200 \text{ cm}^2/\text{Vs}$ and the saturation velocity (defined at 100 kV/cm) $3.9 \times 10^6 \text{ cm/s}$ at 300 K . When the temperature decreases, both the mobility and the saturation velocity enhance due to the suppression of phonon excitation; the respective values at 50 K are $3.0 \times 10^4 \text{ cm}^2/\text{Vs}$ and $6.2 \times 10^6 \text{ cm/s}$. The drift velocities show a slight negative slope at high fields that becomes more pronounced at low temperatures. This phenomenon (i.e., the negative differential resistance) can be explained, at least in part, by the nonlinear band dispersion at high electron energies as in graphene.³²

The calculation results discussed above demonstrate the intrinsic properties of silicene. When this material is synthesized or placed on a substrate, however, additional scattering sources such as surface polar phonons and impurities must be considered, which could degrade the performance further.

A topic that may need additional attention is the role of ZA phonons in the presence of a supporting material. As recent measurement of graphene in-plane thermal conductivity attests,³³ even a weak binding between a 2D crystal and the substrate could dampen the ZA vibrations substantially. Moreover, it is reasonable to anticipate that the extent of this suppression would be dependent on the detailed interaction between two materials. Since ZA phonons provide the dominant role in the electron-phonon interaction in silicene, it (i.e., the damped vibration) could actually lead to sizable reduction in the scattering rate. To gauge the impact, transport characteristics are also studied without the ZA scattering. As shown in Fig. 4(b), the mobility experiences an increase of greater than threefold (3900 cm/s), while the saturation velocity goes up more modestly (5.6×10^6 cm/s). This estimate may be considered an upper limit for silicene on a substrate.

B. Monolayer MoS₂

In the present DFT calculation for monolayer MoS₂, the optimized lattice constant is 3.13 Å, consistent with other theoretical studies.^{21,34,35} Furthermore, this value is in good agreement with 3.15 Å determined experimentally in bulk MoS₂.³⁶ The resulting electronic and phononic band dispersion is depicted in Fig. 5. As shown, monolayer MoS₂ is a semiconductor with a direct gap of 1.86 eV at the *K* point, a number within a few percent from a recent measurement of 1.9 eV.⁹ Our calculation also predicts the presence of second energy minima only about 70 meV higher. These so-called *Q* valleys are located along the Γ -*K* axes at approximately the halfway points [e.g., $Q = (0.34 \times 2\pi/a, 0)$ versus $K = (4\pi/3a, 0)$]. At present, the energy separation E_{QK} between the *K* and *Q* valleys is unsettled with the estimates ranging from 50 to 200 meV.^{21,35} Since this is a crucial parameter for electron transport, a more extended discussion is provided later in the paper in relation to intervalley scattering. The band dispersion relations around the energy minima are nearly quadratic and can be well described by the effective mass approximation. For the *K* valleys (i.e., the energy minima at the equivalent *K* points), the extracted longitudinal and transverse effective masses are almost identical, $m_K^l = m_K^t = 0.50m_0$. On the other hand, the *Q* valleys yield $m_Q^l = 0.62m_0$,

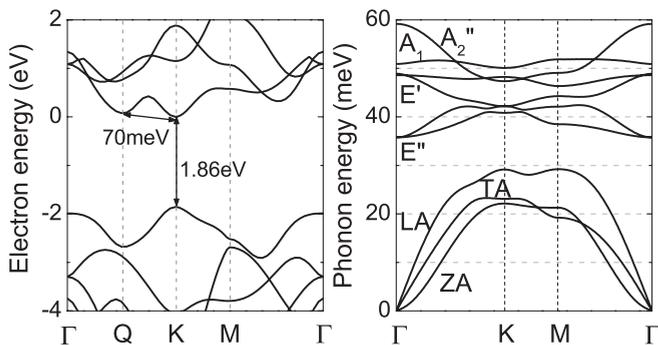


FIG. 5. Electronic and phononic band structures of monolayer MoS₂ along the symmetry directions in the FBZ. The conduction band minimum at the *K* point serves as the reference of energy scale for electrons.

TABLE II. Phonon energies (in units of meV) for TA, LA, TO(*E'*), LO(*E'*), and *A*₁ (or homopolar) modes at the Γ , *K*, *M*, and *Q* points in the FBZ of monolayer MoS₂.

Phonon modes	Γ	<i>K</i>	<i>M</i>	<i>Q</i>
TA	0	23.1	19.2	17.9
LA	0	29.1	29.2	23.6
TO(<i>E'</i>)	48.6	46.4	48.2	48.0
LO(<i>E'</i>)	48.9	42.2	44.3	44.2
<i>A</i> ₁	50.9	51.9	50.1	52.2

$m_Q^t = 1.0m_0$ with the longitudinal direction defined along the Γ -*K* axis. m_0 denotes the electron rest mass.

Monolayer MoS₂ has the symmetry of the point group D_{3h} , with nine branches of phonons. The irreducible representations associated with each phonon mode, together with the polarization (longitudinal or transverse), help denote all of the vibrational modes,^{34,37} as plotted in Fig. 5. The *E''* modes are degenerate at the Γ point. These two modes are the in-plane optical vibrations, with two *S* atoms moving out of phase and Mo atom static. The *E'* modes are polar LO and TO phonons, with Mo atom and two *S* atoms moving out of phase. Due to the coupling with the macroscopic electric field, there is LO-TO separation at the Γ point, which slightly lifts the LO(*E'*) mode upward on energy scale. A nonanalytical part is added to the dynamic matrix resulting in a small splitting of about 0.3 meV [not visible due to the energy scale of Fig. 5(b)]. The *A*₁ and *A*₂^{''} branches are two out-of-plane optical phonon modes. *A*₁ is also referred to as the homopolar mode, with two *S* atoms moving out of phase and the Mo atom static. In the *A*₂^{''} mode, the Mo atom and two *S* atoms vibrate out of phase. The three lowest branches are LA, TA, and ZA modes, with sound velocities of $v_{LA} = 6.6 \times 10^5$ cm/s, $v_{TA} = 4.3 \times 10^5$ cm/s. The phonon energies at different symmetric points are summarized in Table II.

Figures 6 and 7 show the electron-phonon interaction matrix elements for the initial electron state at $\mathbf{k} = K [= (4\pi/3a, 0)]$ and $\mathbf{k} = Q [\approx (2\pi/3a, 0)]$, respectively, for TA, LA, TO(*E'*), LO(*E'*), and *A*₁ (or homopolar) phonon modes. The contribution from the remaining four branches is found to be negligible due to the weak coupling. The matrix elements for $\mathbf{k} = K$ demonstrate a threefold-rotational symmetry (i.e., 120°), while those of $\mathbf{k} = Q$ show the reflection symmetry with respect to the q_x axis. As expected, the LO(*E'*) phonons near the Γ point possess the characteristics of Fröhlich coupling through the induced macroscopic electric field typical of polar materials. Since the relative electronic potential is not periodic in the long-wavelength limit,²⁵ DFPT does not yield a correct value to the electron-phonon interaction matrix. For an approximation, the matrix element of LO(*E'*) is interpolated at Γ by using the values from the nearby q points. This [i.e., LO(*E'*)] and *A*₁ are the only two modes that have nonzero scattering matrix as $\mathbf{q} \rightarrow 0$ (intervalley scattering); in the other three branches, the matrix elements only have first-order components $|\mathbf{g}_{\mathbf{q},\mathbf{k}}| \sim \mathbf{q}$, leading to $|\mathbf{g}_{\mathbf{q} \rightarrow 0, \mathbf{k}}| \rightarrow 0$. With regard to intervalley scattering that requires large- q phonons, a number of different transition processes are possible as shown in Figs. 6(f) and 7(f). For instance, Fig. 6 indicates strong electron-phonon interaction at the symmetry points *M* in the

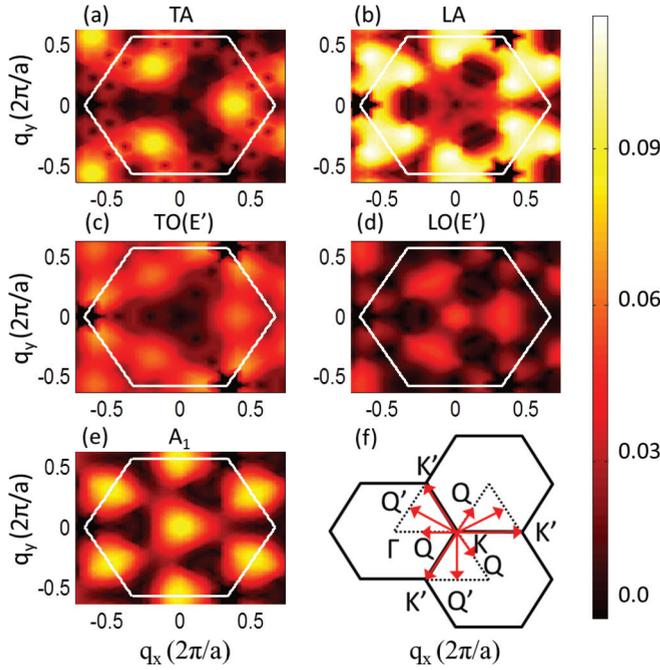


FIG. 6. (Color online) (a)–(e) Electron-phonon interaction matrix elements $|g_{\mathbf{k}+\mathbf{q},\mathbf{k}}^v|$ (in units of eV) from the DFPT calculation in MoS₂ for \mathbf{k} at the conduction band minimum K point [i.e., $(4\pi/3a, 0)$] as a function of phonon wave vector \mathbf{q} . Only the branches with significant contribution are plotted; i.e., TA, LA, TO(E'), LO(E'), and A_1 (or homopolar) modes. (f) Schematic illustration of *intervalley* scattering for electrons in the K valley.

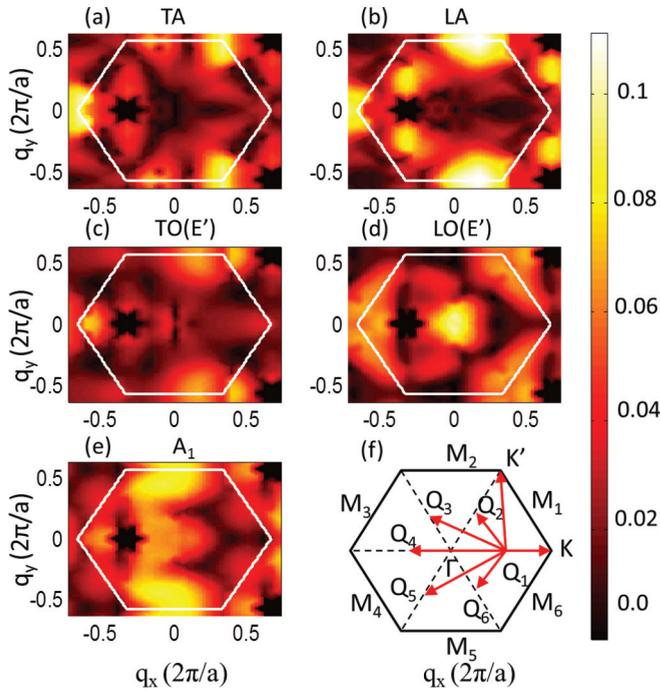


FIG. 7. (Color online) (a)–(e) Electron-phonon interaction matrix elements $|g_{\mathbf{k}+\mathbf{q},\mathbf{k}}^v|$ (in units of eV) from the DFPT calculation in MoS₂ for \mathbf{k} at the Q point [i.e., $Q_1 \approx (2\pi/3a, 0)$] as a function of phonon wave vector \mathbf{q} . Only the branches with significant contribution are plotted; i.e., TA, LA, TO(E'), LO(E'), and A_1 (or homopolar) modes. (f) Schematic illustration of *intervalley* scattering for electrons in the Q valleys.

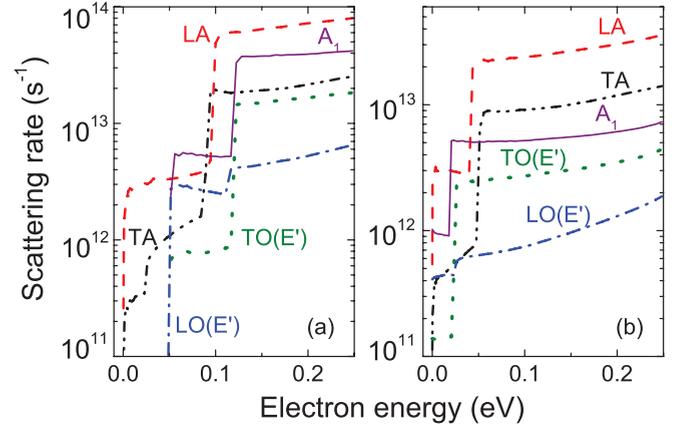


FIG. 8. (Color online) Scattering rates of K -valley electrons in MoS₂ via (a) emission and (b) absorption of phonons calculated at room temperature. The electron wave vector \mathbf{k} is assumed to be along the K - Γ axis.

phonon momentum space (denoted as $\mathbf{q} = M$ for simplicity) for all modes except LO(E'); these phonons can induce electron transition from K to Q' valleys. Another example is the phonons at $\mathbf{q} = K'$ for all five modes in Fig. 7, which can be associated with electron scattering from Q_1 to Q_4 .

The electron-phonon scattering rates are calculated as a function of electron energy using Fermi's golden rule. Figure 8 gives the rates for electrons in the K valleys at room temperature, while the result for Q -valley electrons is shown in Fig. 9. Similarly to silicene, the wave vector \mathbf{k} of the initial electronic state is chosen along the K - Γ or Q - Γ axis, respectively. As can be seen from the figures, the LA mode provides the largest scattering rates consistent with its large coupling strength. The discontinuities or steps in the curves represent either the onset of optical phonon emission or intervalley scattering. For instance, the abrupt increase observed in the rate of LA phonons at ~ 100 meV in Fig. 8(a) can be attributed to the above-mentioned strong $K \rightarrow Q'$ transition

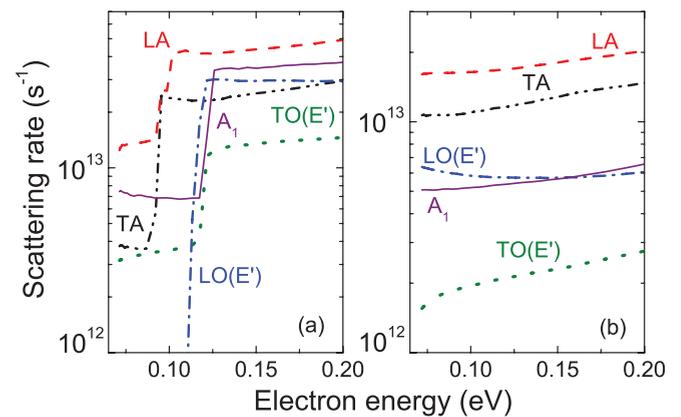


FIG. 9. (Color online) Scattering rates of Q -valley electrons in MoS₂ via (a) emission and (b) absorption of phonons calculated at room temperature. The Q - K separation energy E_{QK} ($=70$ meV) denotes the onset of curves as the K -valley minimum serves as the reference (zero) of energy scale. The electron wave vector \mathbf{k} is assumed to be along the Q - Γ axis.

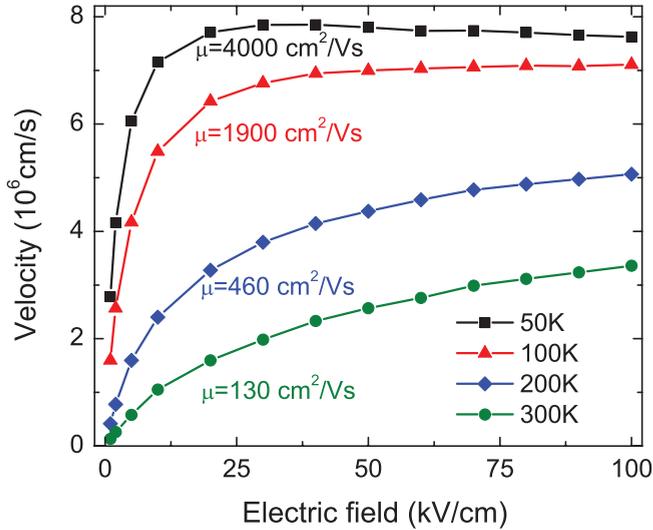


FIG. 10. (Color online) Drift velocity versus electric field in monolayer MoS₂ obtained from a Monte Carlo simulation at different temperatures with $E_{QK} = 70$ meV. When electron transfer to the Q valleys is not considered, the mobility increases to approximately 320 cm²/Vs at 300 K.

via emission of a LA phonon with $\mathbf{q} = M$. Since this phonon energy is approximately 30 meV (see Fig. 5), the final-state energy of ~ 70 meV indeed matches to the Q - K separation E_{QK} . Similarly, the onset of transition via absorption can be found around 40 meV in Fig. 8(b). The large density of states in the Q valleys (evident from the large effective masses) makes the contribution of this scattering even more prominent. If, on the other hand, all of the final states in the Q valleys are excluded, the total scattering rate for the K -valley electron reduces drastically to approximately 2×10^{13} s⁻¹, which is consistent with the prediction of an earlier first-principles calculation.²¹ The observed difference of an order of magnitude clearly illustrates the strong dependence of the scattering rates on E_{QK} . The inconsistency of this value in the recent publications^{21,35} adds difficulty to accurately evaluating the role of Q valleys.

Utilizing the scattering rates, the velocity versus field relation is obtained by a full-band Monte Carlo simulation at four different temperatures. As shown in Fig. 10, the extracted mobility decreases from 4000 cm²/Vs at 50 K to about 130 cm²/Vs at room temperature, while the saturation velocity changes from 7.6×10^6 to 3.4×10^6 cm/s. The small mobility and saturation velocity can be attributed to strong electron-phonon scattering as well as the heavy effective masses. With massive electrons that hinder acceleration and many states to scatter into (e.g., K , K' , Q , Q' valleys), this is an expected outcome.

Compared to a recent theoretical estimation²¹ of 410 cm²/Vs, however, our mobility is significantly smaller, requiring a careful analysis of the discrepancy. One factor that could yield at least a partial explanation is the issue surrounding the Q - K separation. With inconsistencies reported in several first-principles results on this sensitive quantity (see the discussion above), it is not unreasonable to imagine that our DFT calculation may have also experienced similar inaccuracies. If E_{QK} proves to be substantially larger

than the estimated 70 meV, then the Q valleys would have a limited influence on the low-field mobility and can be neglected in the calculation as in Ref. 21 (with 200 meV). In this case, our simulation estimates the K -valley dominated mobility of 320 cm²/Vs that is essentially in agreement with the earlier prediction (410 cm²/Vs).²¹ Clearly, both first-principles models produce a consistent picture of K -valley electron dynamics including intrinsic scattering with relevant phonon modes. The difference is the relative significance of Q valleys (e.g., 70 versus 200 meV). As such, further clarification of intrinsic mobility in monolayer MoS₂ may need to be preceded by accurate experimental determination of E_{QK} .

Even when the influence of Q valleys becomes negligible, a sizable disparity remains between the theoretically obtained mobility and the highest value claimed experimentally (e.g., ~ 1090 cm²/Vs).¹⁶ One reason could be inaccuracy in the measurement due to external artifacts as mentioned earlier. Indeed, the latest analysis revealed that the attempts based on the two-point method in the thin-film MoS₂ transistor channel ignore the coupling capacitance between top and bottom gates, leading to an overestimation of the mobility by approximately a factor of 14.^{22,23} When accounting for this effect, the recalibrated data from the experiments indicate mobility values substantially smaller than the theoretical prediction (< 130 – 410 cm²/Vs). A subsequent Hall measurement also yielded 63.7 cm²/Vs at 260 K.²⁴ Considering the influence of extrinsic scattering sources such as impurities, our calculation is actually consistent with the revised experimental estimate and provides a reasonable upper limit for the intrinsic mobility.

An additional point to note when interpreting the experimental data is that the screening by degenerate electrons can affect the bare scattering rates substantially (thus, the mobility) in a low-dimensional system.^{38,39} This effect can be included by renormalizing the electron-phonon interaction through the dielectric function; i.e., $\mathbf{g}_{\mathbf{q},\mathbf{k}}^v \rightarrow \mathbf{g}_{\mathbf{q},\mathbf{k}}^v/\epsilon(q)$. To gauge the potential significance in monolayer MoS₂, we adopt a simple model for the dielectric function based on Thomas-Fermi screening of only the K -valley electrons: i.e., $\epsilon(q) = 1 + q_{TF}/q$, where the screening wave vector $q_{TF} = 4m_{\kappa}e^2/\hbar^2\kappa$. Here, the factor of 4 accounts for the spin and valley degeneracies, e is the electron charge, and κ is the background dielectric constant. Subsequent calculation with a rough estimate of $\epsilon(q)$ shows that the scattering rates can experience a decrease of about an order of magnitude through screening. Thus, it is evident that the effect (i.e., screening) must be taken into account accurately when the carrier density becomes degenerate in MoS₂. A detailed analysis of $\epsilon(q)$ is, however, beyond the scope of this investigation as our focus is on the properties of intrinsic electron-phonon interaction (i.e., nondegenerate).

C. Deformation potential model

For practical applications, it would be convenient to approximate the *ab initio* results for electron-phonon coupling by a simple analytical model. Particularly useful in the present case is the deformation potential approximation. Under this treatment, the coupling matrix $\langle j, \mathbf{k} + \mathbf{q} | \Delta V_{\mathbf{q},\text{SCF}}^v | i, \mathbf{k} \rangle$ in Eq. (1) can be expressed in the first order ($D_1\mathbf{q}$), or in

the zeroth order (D_0).⁴⁰ The first-order deformation potential constant (D_1) is adopted to represent the coupling matrices for the acoustic phonon modes in the long-wavelength limit (i.e., intravalley scattering). In comparison, those involving the near zone-edge acoustic phonons (i.e., intervalley scattering) are treated by using the zeroth-order deformation potential (D_0) in a manner analogous to the optical modes. In the latter case (D_0), the phonon energy is assumed independent of the momentum for simplicity. The obtained analytical expressions of the scattering rates are then matched to the first-principles results by fitting the effective deformation potential constants.

For silicene, the *intervalley* scattering rate by acoustic mode v (=LA, TA, ZA) is obtained as

$$\frac{1}{\tau_{\mathbf{k},v}^{(1)}} \Big|_{\text{Si}} = \frac{D_1^2 k_B T}{\hbar^3 v_F^2 \rho v_v^2} E_{\mathbf{k}}. \quad (3)$$

Here, ρ is the mass density ($= 7.2 \times 10^{-8}$ g/cm²) and v_v denotes the sound velocity, for which we can take the value of $v_{ZA} = 6.3 \times 10^4$ cm/s, $v_{TA} = 5.4 \times 10^5$ cm/s, and $v_{LA} = 8.8 \times 10^5$ cm/s, respectively, as discussed earlier. On the other hand, the rate of optical phonon scattering (both *intervalley* and *intravalley* transitions) as well as the *intervalley* acoustic phonon scattering can be expressed by the following form:

$$\frac{1}{\tau_{\mathbf{k},v}^{(2)}} \Big|_{\text{Si}} = \frac{D_0^2}{2\hbar^2 v_F^2 \rho \omega_v} [(E_{\mathbf{k}} + \hbar\omega_v)N_v + (E_{\mathbf{k}} - \hbar\omega_v)(N_v + 1)\Theta(E_{\mathbf{k}} - \hbar\omega_v)], \quad (4)$$

where $\Theta(x)$ is the Heaviside step function and $N_v = [\exp(\hbar\omega_v/k_B T) + 1]^{-1}$ for phonon mode v . Since the phonon dispersion is treated constant in Eq. (4), the *intervalley* optical phonon scattering approximates the zone-center (i.e., Γ) phonon energies for $\hbar\omega_v$ ($v = \text{LO, TO, ZO}$). In the case of *intervalley* scattering via acoustic or optical phonons, $\hbar\omega_v$ takes the respective phonon energy at the zone-edge K point corresponding to electron transition $K \leftrightarrow K'$. The specific values used in the calculation can be found in Table I. When matched to the first-principles rates, the effective deformation potential constants can be extracted for each scattering process as summarized in Table III. A particularly interesting point to note from the result is that all three acoustic phonons show generally comparable values of D_1 and D_0 despite the large scattering rate of ZA mode (see also Fig. 3). Clearly, this mode (ZA) couples strongly with electrons but not enough to prevail over other acoustic branches; its dominant contribution is due mainly to the small phonon energy as discussed earlier.

TABLE III. Extracted deformation potential constants for electron-phonon interaction in silicene.

Phonon mode	Intravalley	Intervalley
ZA	2.0 eV	6.1×10^7 eV/cm
TA	8.7 eV	1.4×10^8 eV/cm
LA	3.2 eV	4.2×10^7 eV/cm
ZO	6.3×10^7 eV/cm	4.3×10^7 eV/cm
TO	1.8×10^8 eV/cm	1.4×10^8 eV/cm
LO	1.9×10^8 eV/cm	1.7×10^8 eV/cm

The electron-phonon scattering processes in MoS₂ are much more complicated as the deformation potentials need to be determined for both K - and Q -valley electrons. While feasible, it is not practically useful to define the effective interaction constants individually based on the mode of involved phonons and the transition types. Accordingly, we adopt a simplified description by combining the appropriate contributions into just two modes, acoustic and optical, respectively.

Using the effective mass approximation for the band structure near the valley minima, the scattering rate for *intervalley* acoustic phonon scattering (i.e., $K \rightarrow K$ or $Q_1 \rightarrow Q_1$ by both LA and TA phonons; see Figs. 6 and 7) is given by

$$\frac{1}{\tau_{\mathbf{k}}^{(1)}} \Big|_{\text{MoS}_2} = \frac{m_p^* D_1^2 k_B T}{\hbar^3 \rho v_s^2}, \quad (5)$$

where $\rho = 3.1 \times 10^{-7}$ g/cm² for MoS₂ and m_p^* is the density-of-states effective mass for the K or Q valley (final state), $m_p^* = \sqrt{m_p^i m_p^f}$ ($p = K, Q$). For electrons in the Q valleys, strictly speaking, an additional factor of $\Theta(E_{\mathbf{k}} - E_{QK})$ is multiplied to the left side of Eq. (5) to account for the nonzero energy at the bottom of the Q valley. By taking the sound velocity $v_s = v_{LA}$ ($= 6.6 \times 10^5$ cm/s), the value of D_1 is estimated to be 4.5 and 2.8 eV in the K and Q valleys, respectively.

The analytical expression that describes *intervalley* and *intervalley* optical phonon scattering as well as *intervalley* acoustic phonon scattering rate is obtained as

$$\frac{1}{\tau_{\mathbf{k},v}^{(2)}} \Big|_{\text{MoS}_2} = g_d \frac{m_p^* D_0^2}{2\hbar^2 \rho \omega_v} [N_v \Delta_1 + (N_v + 1) \Delta_2], \quad (6)$$

where g_d is the valley degeneracy for the final electron states, and Δ_1 and Δ_2 denote the onset of scattering for phonon absorption and emission, respectively. For instance, $\Delta_1 = 1$ and $\Delta_2 = \Theta(E_{\mathbf{k}} - \hbar\omega_v)$ for electron transitions between the K valleys, whereas $\Delta_1 = \Theta(E_{\mathbf{k}} - E_{QK})$ and $\Delta_2 = \Theta(E_{\mathbf{k}} - \hbar\omega_v - E_{QK})$ for transitions between the Q valleys. The factors corresponding to intervalley transfer between K and Q valleys can be constructed accordingly, where E_{QK} may be treated as a potentially adjustable parameter. Tables IV and V summarize the initial/final electron states, the phonon momentum that is involved (in the form of its location in the momentum space), and the extracted deformation potential constants for each transition process considered in the investigation. For a

TABLE IV. Extracted deformation potential constants for electron-phonon interaction in MoS₂ for electrons in the K valley [see also Fig. 6(f)].

Phonon momentum	Electron transition	Deformation potentials
Γ	$K \rightarrow K$	$D_1^{ac} = 4.5$ eV $D_0^{op} = 5.8 \times 10^8$ eV/cm $D_0^{ac} = 1.4 \times 10^8$ eV/cm
K'	$K \rightarrow K'$	$D_0^{op} = 2.0 \times 10^8$ eV/cm $D_0^{ac} = 9.3 \times 10^7$ eV/cm
Q'	$K \rightarrow Q$	$D_0^{op} = 1.9 \times 10^8$ eV/cm $D_0^{ac} = 4.4 \times 10^8$ eV/cm
M	$K \rightarrow Q'$	$D_0^{op} = 5.6 \times 10^8$ eV/cm

TABLE V. Extracted deformation potential constants for electron-phonon interaction in MoS₂ for electrons in the Q_1 valley [see also Fig. 7(f)]. Multiple equivalent valleys for the final state specify the degeneracy factor g_d larger than one in Eq. (6).

Phonon momentum	Electron transition	Deformation potentials
Γ	$Q_1 \rightarrow Q_1$	$D_1^{ac} = 2.8$ eV $D_0^{op} = 7.1 \times 10^8$ eV/cm $D_0^{ac} = 2.1 \times 10^8$ eV/cm
$Q_3(Q_5)$	$Q_1 \rightarrow Q_2(Q_6)$	$D_0^{op} = 4.8 \times 10^8$ eV/cm $D_0^{ac} = 2.0 \times 10^8$ eV/cm
$M_3(M_4)$	$Q_1 \rightarrow Q_3(Q_5)$	$D_0^{op} = 4.0 \times 10^8$ eV/cm $D_0^{ac} = 4.8 \times 10^8$ eV/cm
K'	$Q_1 \rightarrow Q_4$	$D_0^{op} = 6.5 \times 10^8$ eV/cm $D_0^{ac} = 1.5 \times 10^8$ eV/cm
Q_1	$Q_1 \rightarrow K$	$D_0^{op} = 2.4 \times 10^8$ eV/cm $D_0^{ac} = 4.4 \times 10^8$ eV/cm
$M_2(M_5)$	$Q_1 \rightarrow K'$	$D_0^{op} = 6.6 \times 10^8$ eV/cm

given phonon momentum, the actual value $\hbar\omega_v$ used in the analytical calculation is the average of the relevant phonon modes. Specifically, the acoustic (optical) phonon energy is obtained as the average of LA and TA [TO(E'), LO(E'), and A_1] modes at the respective symmetry points given in Table II. An additional point to note is that the estimate of D_0^{op} at the Γ point includes the effect of Fröhlich scattering by the LO(E') mode.²¹ While this is a mechanism physically distinct from the deformation potential interaction and must be handled separately, its impact is relatively modest, at least for the electrons in the K valley. Accordingly, the present treatment is considered adequate. Further simplification of the model may

also be possible judging from the narrow range of values in D_0^{ac} and D_0^{op} (mostly in the low to mid 10^8 eV/cm).

IV. SUMMARY

We have performed a first-principles calculation together with a full-band Monte Carlo analysis to examine electron-phonon interaction and the intrinsic transport properties in monolayer silicene and MoS₂. The results clearly elucidate the role of different branches as well as the intravalley/intervalley scattering. The predicted intrinsic mobility for silicene is approximately 1200 cm²/Vs, with saturation velocity of 3.9×10^6 cm/s at room temperature. In the case of MoS₂, the K -valley-dominated mobility gives approximately 320 cm²/Vs, while the intrinsic value reduces to about 130 cm²/Vs when the energy separation of 70 meV is used between the K and Q minima. The estimated saturation velocity is 3.7×10^6 cm/s. The investigation also illustrates the significance of extrinsic screening, particularly in numerical evaluation of transport characteristics. The extracted deformation potential constants may prove to be useful in further studies of these materials.

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