

Hole mobility of strained GaN from first principles

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Nitride semiconductors are ubiquitous in optoelectronic devices such as LEDs and Blu-Ray optical disks. A major limitation for further adoption of GaN in power electronics is its low hole mobility. In order to address this challenge, here we investigate the phonon-limited mobility of wurtzite GaN using the *ab initio* Boltzmann transport formalism, including all electron-phonon scattering processes, spin-orbit coupling, and many-body quasiparticle band structures. We demonstrate that the mobility is dominated by acoustic deformation-potential scattering, and we predict that the hole mobility can significantly be increased by lifting the split-off hole states above the light and heavy holes. This can be achieved by reversing the sign of the crystal-field splitting via strain or via coherent excitation of the A_1 optical phonon through ultrafast infrared optical pulses.

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I. INTRODUCTION

Wurtzite GaN plays a key role in solid state lighting due its wide emission spectrum, high efficiency, and scalable manufacturing [1]. GaN has a high breakdown field, high thermal conductivity, and high electron mobility, and is therefore an excellent candidate for high-power electronic devices [2–5] and radio-frequency electronics [6]. GaN also exhibits high Seebeck coefficient and excellent temperature stability, which makes it a prime candidate for high-temperature thermoelectric applications [7–9]. It can also be used for thermal neutron and gamma radiation detection [10]. More generally, group-III nitrides can be engineered to form continuous alloys with a tunable band gap from 6.2 eV (AlN) through 3.4 eV (GaN) to 0.7 eV (InN) [1], finding applications in blue [11] and green [12] lasers, photodetectors [13], and light-emitting diodes [14]. Recently, a GaN/NbN semiconductor/superconductor heterojunction was achieved through epitaxial growth, paving the way for superconducting qubits [15]. Widespread adoption of GaN for applications such as complementary metal-oxide-semiconductor (CMOS) and high-power conversion devices is hindered by the low hole mobility of GaN. In fact, the room-temperature hole mobility of GaN does not exceed 40 cm^2/Vs [16–22]. In comparison, electron mobilities as high as 1265 cm^2/Vs have been reported in the bulk, and exceeding 2000 cm^2/Vs in two-dimensional (2D) electron gases [23]. It is therefore important to find practical ways to increase the hole mobility in this semiconductor.

Transport properties in wurtzite GaN have been investigated theoretically decades ago by Ilegems and Montgomery [24], taking into account conduction band nonparabolicity, deformation potential, piezoelectric acoustic-phonon scattering, and polar-optical phonon scattering. More recently, analytical models based on experimental results have been developed to

accurately describe low-field carrier mobilities in a wide temperature and doping range [25–27], and the electron-phonon scattering rates of GaN were computed from first-principles using the EPW software [28].

In a recent work [29], we clarified the atomic-scale mechanisms that are responsible for the low hole mobilities in GaN using the state-of-the-art *ab initio* Boltzmann transport formalism, and we discussed strategies to significantly increase the hole mobility in wurtzite GaN. We showed that the origin of the low *hole* mobility lies in the scattering of carriers in the light-hole (lh) and heavy-hole (hh) bands predominantly by long-wavelength longitudinal-acoustic phonons. Using this understanding, we predicted that the hole mobility could significantly be enhanced if the split-off hole band (sh) could be raised above the lh and hh bands. Such band inversion can be achieved by reversing the sign of the crystal-field splitting via uniaxial compressive or biaxial tensile strain. In this paper, we give details on the computational parameters used in Ref. [29], convergence studies, phase diagram, and we discuss various strategies to perform momentum integration in the calculation of mobility.

The paper is organized as follows. In Sec. II, we compute the GaN phase diagram and show that the wurtzite phase is thermodynamically stable for the range of pressures and temperatures investigated. We also discuss ground-state properties of unstrained GaN. We then discuss in Sec. III the electronic structure including many-body quasiparticle correction, the electron and hole effective masses, as well as spin-orbit and crystal-field splitting. In Sec. IV, we briefly present the linearized Boltzmann transport formalism to compute the mobility of GaN, the associated convergence studies, the band velocity, and the Hall factor. We compare our theoretical results to experiment and analyze the origin of the low hole mobility in unstrained GaN. In Sec. V, we analyze the elastic properties, phonon dispersion and electronic band structure of GaN under biaxial and uniaxial strain. We also show how to increase the hole mobility via biaxial tensile or uniaxial

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compressive strain. Finally, in Sec. VI, we discuss how to realize high-mobility *p*-type GaN in experiments. We draw our conclusions in Sec. VII.

II. GALLIUM NITRIDE PHASE DIAGRAM AND GROUND-STATE PROPERTIES

In this section we present the computed phase diagram of bulk GaN, highlighting the approximations and computational parameters used. Such a diagram is important to make sure that no phase change occurs within the range of applied strains and temperature investigated in this study. We then focus on the stable wurtzite phase and discuss the optimized structure.

GaN can form the following allotropes: (i) wurtzite $P6_3mc$, (ii) zinc-blende $F\bar{4}3m$, and (iii) rocksalt $Fm\bar{3}m$. The wurtzite phase is the naturally occurring phase, while the zinc-blende phase has been stabilized experimentally by epitaxial growth on cubic GaAs [001] surfaces [30]; the rocksalt phase can be obtained under high pressure [31].

We use the fully relativistic norm-conserving Perdew-Zunger [32] parametrization of the local density approximation (LDA) to density functional theory, and the Perdew-Burke-Ernzerhof (PBE) [33] generalized gradient approximation. The pseudopotentials are generated using the ONCVSPcode [34] and optimized via the PSEUDODOJO initiative [35]. The semicore $3s$, $3p$, and $3d$ electrons of Ga are explicitly described, as *GW* quasiparticle corrections are sensitive to semicore states. The electron wave functions are expanded in a plane-wave basis set with kinetic energy cutoff of 120 Ry, and the Brillouin zone is sampled using a homogeneous Γ -centered $6 \times 6 \times 6$ mesh.

Within the quasiharmonic approximation [37], the Helmholtz free energy of a crystal is given by [38]

$$F(T, V) = U(V) + F^{\text{vib}}(T, V) + F^{\text{el}}(T, V), \quad (1)$$

where U is the static (clamped-ion) energy at 0 K, F^{vib} is the contribution due to lattice vibrations, and F^{el} is the energy due to electronic thermal excitations. We rely on the adiabatic approximation to treat each term independently. The vibrational Helmholtz free energy per cell is given in the harmonic approximation by [38]

$$F^{\text{vib}}(T, V) = \frac{1}{2N} \sum_{\mathbf{q}, \nu} \hbar \omega_{\mathbf{q}, \nu}(V) + \frac{k_B T}{N} \sum_{\mathbf{q}, \nu} \ln \left[1 - \exp \left(\frac{-\hbar \omega_{\mathbf{q}, \nu}(V)}{k_B T} \right) \right], \quad (2)$$

where N is the number of $\mathbf{q}$ points, the first term is the contribution to the zero-point energy, and the second term is the phonon contribution at finite temperature. F^{el} can be neglected as the band gap is much larger than thermal energies.

The energy minimum of $U(V) + F^{\text{vib}}(T, V)$ at a given temperature corresponds to zero pressure and gives the variation of volume with temperature due to thermal expansion. To perform these calculations, we use the QUANTUM ESPRESSO [39] and THERMO_PW codes [40]. The phonon frequencies were computed using the PBE pseudopotentials, without spin-orbit coupling (SOC), at 11 different volumes. The resulting energies were fitted using the Murnaghan equation of

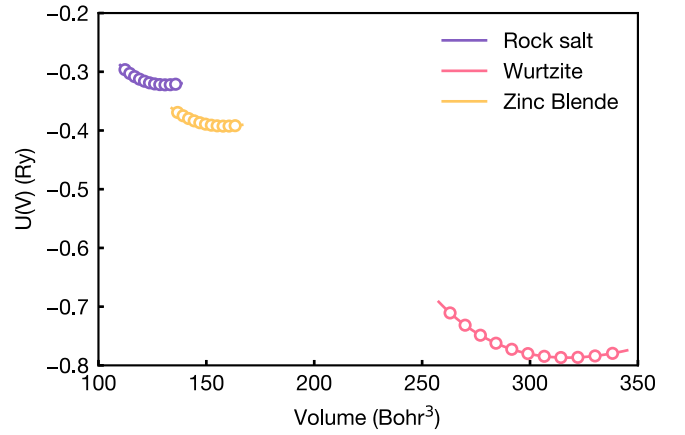


FIG. 1. Total energy versus volume at zero temperature for the rocksalt, wurtzite, and zinc-blende phases of GaN.

state [41]. We used a $6 \times 6 \times 6$ $\mathbf{q}$ -point grid for the phonons. In order to obtain accurate dielectric permittivity tensors and Born effective charges, we employed a much denser, shifted Monkhorst-Pack grid with $16 \times 16 \times 16$ $\mathbf{k}$ points. To compute phonon dispersion relations, we applied the crystal acoustic sum rule [42,43]. The calculated dependence of the static energy $U(V)$ on the volume is shown in Fig. 1.

The complete phase diagram can be obtained by comparing the Gibbs free energy of the various allotropes at each temperature and pressure. The Gibbs free energy $G(T, P)$ can be obtained from the Helmholtz free energy as

$$G(T, P) = F(T, V) + PV, \quad (3)$$

where the pressure is obtained by computing the first-order derivative of the Helmholtz free energy, with respect to volume at fixed temperature:

$$P = - \left. \frac{\partial F}{\partial V} \right|_T. \quad (4)$$

The Gibbs free energy was computed using the python toolkit PHASEGO [44]. The resulting pressure-temperature phase diagram is given in Fig. 2, where the light gray rectangle represents the maximum stress (9.8 GPa at 2% strain) and temperature (500 K) investigated in this paper. We see that the wurtzite structure is always the thermodynamically stable phase in the strain and temperature considered in this study. We note that our calculated *ab initio* phase diagram overestimates the coordinates of the triple point (10 GPa, 4700 K) with respect to experiment (6.2 GPa, 2300 K) [36], however, this error does not affect the region of interest for this study.

Our optimized lattice parameters of wurtzite GaN are $a = 5.961(6.081)$ bohrs and $c = 9.716(9.9049)$ bohrs, and the internal parameter is $u = 0.376(0.377)$ in LDA (PBE). As expected, the experimentally measured parameters, $a = 6.026$ bohrs and $c = 9.800$ bohrs [45], fall in-between the LDA and PBE data. All subsequent calculations of electron band structures, phonon dispersion relations, and electron-phonon interactions are performed using these optimized lattice parameters.

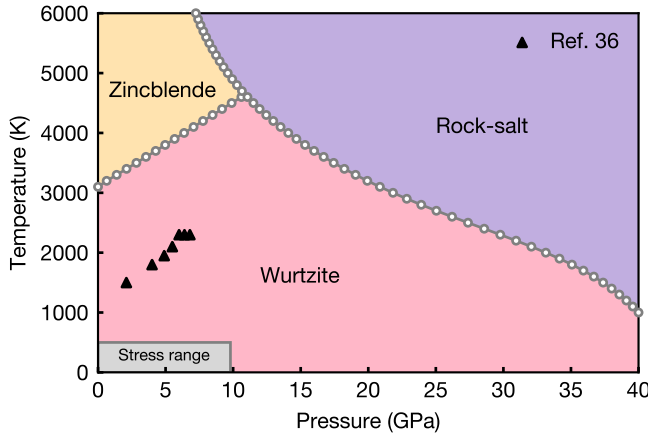


FIG. 2. Phase diagram of GaN. The phonon frequencies are computed using the PBE functional, without spin-orbit coupling, at 11 different volumes. The experimental values (black triangles) are from Ref. [36]. The light gray rectangle represents the maximum strain (9.8 GPa at 2% strain) and temperature (500 K) investigated in this paper.

III. GW QUASIPARTICLE CORRECTIONS

The electronic band structure computed within the LDA is presented in Fig. 3(a) for states close to the band edge along the high-symmetry directions Γ -A (which is the direction parallel to the c axis, denoted with a $\parallel$ symbol) and Γ -M (which is the perpendicular direction, denoted with a $\perp$ symbol). Due to the wurtzite symmetry, the in-plane Γ -K direction is equivalent to the Γ -M direction, and therefore is not shown.

The calculated direct band gap is 2.14 eV, strongly underestimating the measured value of 3.5 eV [52,62]. To overcome

this shortcoming, we calculated the GW quasiparticle band structures of wurtzite GaN within the many-body G_0W_0 approximation including SOC, as implemented into the YAMBO code [63]. We used a plane-wave kinetic energy cutoff of 120 Ry for the exchange self-energy and 29 Ry for the polarizability. In addition, we included 1500 bands, a plasma energy of 16.5 eV for the plasmon pole [64], and a $6 \times 6 \times 6$ Γ -centered Brillouin-zone grid. We employed the band extrapolation scheme of Ref. [65] to speed up convergence with the number of empty states. We obtained a corrected band gap of 3.41 eV, much closer to the experimental one.

As discussed in Ref. [66], the accuracy of the effective masses is improved by using a self-consistent, $\mathbf{k}$ -dependent scissor shift. This approximation, which we call $G_0W_0 + \Delta_{\mathbf{k}}$, yields a wider band gap of 3.85 eV. This value is in agreement with the gaps 3.24 and 3.81 eV obtained in previous calculations [49,50]. The calculated $LDA + G_0W_0 + \Delta_{\mathbf{k}}$ band structure for unstrained GaN is reported in Fig. 3(b). Throughout this paper, GW -corrected band structures were obtained via Wannier interpolation [67], using 20 Wannier functions for the ground-state structure and the structures with 1% strain, and 28 Wannier functions for the structures with 2% strain (to be discussed below). In contrast, we note that the PBE functional yields much too small a band gap, even after $G_0W_0 + \Delta_{\mathbf{k}}$ corrections (2.94 eV). The various band gaps and their comparison to previous calculations and experiment are summarized in Table I. We note that the theoretical values reported in Table I do not account for the zero-point renormalization, which has been calculated to be -150 meV for zinc-blende GaN [68].

As shown in Fig. 3, the conduction band bottom of GaN is singly degenerated, while at the valence band top we have a lh and a hh, which are split into doublets by SOC as

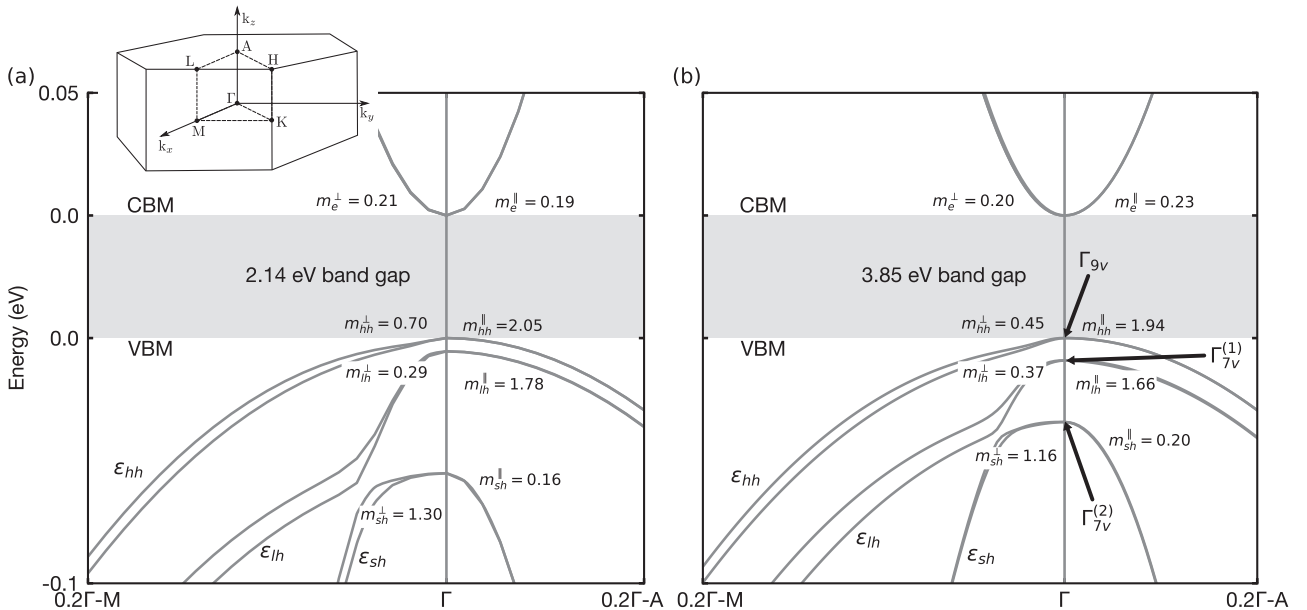


FIG. 3. Electronic band structure of wurtzite GaN using (a) the LDA functional in the optimized ground-state LDA structure, and (b) quasiparticle $G_0W_0 + \Delta_{\mathbf{k}}$ calculation. We indicate the effective masses at the zone center, obtained from the second derivatives of the band energy with respect to the wave vector along the Γ -M and Γ -A directions, respectively. The band gap is off scale for clarity. We indicate the naming convention for the three topmost eigenstates at Γ . The energy levels have been aligned to the band edges. A schematic of the Brillouin zone of wurtzite GaN is given in the upper left corner.

TABLE I. Comparison between our calculated band gap E_g , spin-orbit splitting Δ_{so} , crystal-field splitting Δ_{cf} , and effective masses of wurzite GaN with earlier theory and experiment. QSGW stands for quasiparticle self-consistent approach [59], 0.8 Σ -QSGW is an empirical hybrid method with 20% LDA self-energy [60], and OEPx stands for exact-exchange optimized effective potential [49]. [‡] Calculation done using the quasicubic approximation. The bold values are recommended and are used throughout this paper.

	Energy gaps							
	E_g	Δ_{so}	Δ_{cf}					
This work	(eV)	(meV)	(meV)					
PBE	1.74	9	39					
PBE+ G_0W_0	2.60	11	26					
PBE+ $G_0W_0+\Delta_{\mathbf{k}}$	2.94	13	22					
LDA	2.14	8	53					
LDA+ G_0W_0	3.41	12	35					
LDA+ $G_0W_0+\Delta_{\mathbf{k}}$	3.85	14	30					
Previous work								
LDA [46]		13	42					
LDA [47]		12	37					
LDA [48]		16	36					
LDA [49]	1.78		49					
OEPx+ G_0W_0 [49]	3.24		34					
QSGW [50]	3.81							
0.8 Σ -QSGW [50]	3.42							
0.8 Σ -QSGW [51]	3.60	5 ‡	18 ‡					
0.8 Σ -QSGW [51]	3.60	14	12					
Experiment [52]	3.47							
Experiment [53]	3.47	11	22					
Experiment [54]		18	10					
Experiment [55]		12	16					
Experiment [56]	3.51	17	25					
Experiment [57]		19	10					
	Effective masses							
	$m_{hh}^\parallel$	$m_{lh}^\parallel$	$m_{sh}^\parallel$	$m_{hh}^\perp$	$m_{lh}^\perp$	$m_{sh}^\perp$	$m_e^\parallel$	$m_e^\perp$
This work								
LDA	2.05	1.78	0.16	0.70	0.29	1.30	0.19	0.21
LDA+ G_0W_0	1.98	1.68	0.18	0.58	0.33	1.29	0.22	0.19
LDA+ $G_0W_0+\Delta_{\mathbf{k}}$	1.94	1.66	0.20	0.45	0.37	1.16	0.23	0.20
Previous work								
Theory [51]	1.85	0.55	0.20	0.69	0.50	0.80	0.20	0.22
Theory ^a [49]	1.88	0.92	0.19	0.33	0.36	1.27	0.19	0.21
Theory ^b [49]	1.88	0.37	0.26	0.33	0.49	0.65	0.19	0.21
Theory [48]	2.00	1.19	0.17	0.34	0.35	1.27	0.19	0.23
Theory [47]	2.03	1.25	0.15	0.33	0.34	1.22	0.17	0.19
Experiment [57]	1.76	0.42	0.30	0.35	0.51	0.68		
Experiment [58]							0.22	0.24

^aCalculation using $\Delta_{so} = 16$ meV and $\Delta_{cf} = 25$ meV from Ref. [61].

^bCalculation using $\Delta_{so} = 19$ meV and $\Delta_{cf} = 10$ meV from Ref. [57].

we move from the Γ to the M point of the Brillouin zone. We also have a split-off hole resulting from crystal-field splitting. In order to determine the spin-orbit splitting Δ_{so} and the crystal-field splitting Δ_{cf} , we employ the quasicubic model of Refs. [46,69] for the triplet of states Γ_{9v} , $\Gamma_{7v}^{(1)}$, and $\Gamma_{7v}^{(2)}$ at the valence band top, with energies ε_{hh} , ε_{lh} , and ε_{sh} ,

respectively:

$$\varepsilon_{hh} = \frac{1}{2}(\Delta_{so} + \Delta_{cf}), \quad (5)$$

$$\varepsilon_{lh}, \varepsilon_{sh} = \pm \frac{1}{2} \sqrt{(\Delta_{so} + \Delta_{cf})^2 - \frac{8}{3} \Delta_{so} \Delta_{cf}}. \quad (6)$$

Having checked that $\Delta_{so} \ll \Delta_{cf}$, we can simplify these expressions as

$$\Delta_{so} = \frac{3}{2}(\varepsilon_{hh} - \varepsilon_{lh}), \quad (7)$$

$$\Delta_{cf} = \varepsilon_{lh} - \varepsilon_{sh} + \frac{\varepsilon_{hh} - \varepsilon_{lh}}{2}. \quad (8)$$

Using the last two equations, we determine the spin-orbit splitting and the crystal-field splitting from the calculated band structure energies ε_{hh} , ε_{lh} , and ε_{sh} and systematically report them in Table I. As seen in Fig. 3, the effect of G_0W_0 and the self-consistent scissor is to increase the spin-orbit splitting and decrease the crystal-field splitting. The same effect is observed using either the LDA or PBE exchange-correlation functional. In our LDA+ $G_0W_0+\Delta_k$ calculations for unstrained GaN we find $\Delta_{so} = 14$ meV and $\Delta_{cf} = 30$ meV, in the range of experimental values $\Delta_{so} = 11$ –19 meV and $\Delta_{cf} = 10$ –25 meV [47,53,55–57,70]. We note that LDA tends to slightly overestimate the crystal-field splitting with respect to experiment, in line with previous theoretical findings [46–50]. In contrast, PBE yields slightly smaller values for the crystal-field splitting, but since the band gap is strongly underestimated, we proceed with LDA for the remainder of the paper.

Using the parabolic band approximation, our calculated effective masses with quasiparticle and scissor-shift corrections are $m_e^{\perp/\parallel} = 0.20/0.23 m_e$, $m_{hh}^{\perp/\parallel} = 0.45/1.94 m_e$, $m_{lh}^{\perp/\parallel} = 0.37/1.66 m_e$, and $m_{sh}^{\perp/\parallel} = 1.16/0.2 m_e$, respectively. These values are in reasonable agreement with experimental data ranging from $0.30 m_e$ to $2.03 m_e$ [57,71–74] for holes, and in good agreement with $0.2 m_e$ [75] for the electrons. A detailed comparison with previously computed effective masses and experimental masses is given in Table I. The largest discrepancy with respect to experiment are the overestimated $m_{lh}^{\parallel}$ and $m_{sh}^{\perp}$ effective masses, resulting from an overestimation of the crystal-field splitting in the LDA. This effect was already reported in Ref. [49]. Indeed, as seen in Table I, their $m_{lh}^{\parallel}$ and $m_{sh}^{\perp}$ effective masses decrease from 0.92 to 0.37 and from 1.27 to 0.65 when using the calculated $\Delta_{cf} = 25$ meV from Ref. [61] or the experimental value of $\Delta_{cf} = 10$ meV from Ref. [57]. Overall, we find that increasing the level of theory (from LDA to LDA+ G_0W_0 to LDA+ $G_0W_0+\Delta_k$) systematically improves all the effective masses with respect to the experimental values. Closer agreement with experiment could be achieved by including the small effect of polaronic mass enhancement to the *ab initio* calculations [76]. We also note that the electron effective mass has been confirmed by quantum magnetotransport measurements [77,78], but a corresponding high-accuracy measurement has not been achieved yet for holes in GaN.

IV. CARRIER MOBILITY IN UNSTRAINED GaN

A. Linearized Boltzmann transport equation

The carrier drift mobility μ describes the change of steady-state carrier current $J_\alpha = e(n_e \mu_{e,\alpha\beta} + n_h \mu_{h,\alpha\beta}) E_\beta$ due to an applied external electric field $\mathbf{E}$, where Greek indices denote Cartesian coordinates, n_e and n_h the electron and hole density, respectively. The mobility can be computed using the linearized Boltzmann transport equation (BTE) [79–87], which for electrons reads as

$$\mu_{e,\alpha\beta} = \frac{-1}{n_e \Omega} \sum_{n \in \text{CB}} \int \frac{d\mathbf{k}}{\Omega_{\text{BZ}}} v_{n\mathbf{k},\alpha} \partial_{E_\beta} f_{n\mathbf{k}}. \quad (9)$$

Here, $v_{n\mathbf{k},\alpha} = \hbar^{-1} \partial \varepsilon_{n\mathbf{k}} / \partial k_\alpha$ is the group velocity of the band state of energy $\varepsilon_{n\mathbf{k}}$, band index n , and wave vector $\mathbf{k}$. CB stands for conduction bands, $\partial_{E_\beta} f_{n\mathbf{k}}$ is the perturbation to the Fermi-Dirac distribution induced by the applied electric field $\mathbf{E}$; Ω and Ω_{BZ} are the volumes of the crystalline unit cell and first Brillouin zone, respectively. We note that the additional term in the velocity arising from the Berry curvature contribution vanishes in bulk GaN due to time-reversal symmetry and does not contribute to the BTE mobility [88]. The perturbation to the equilibrium carrier distribution is obtained by solving the following self-consistent equation:

$$\begin{aligned} \partial_{E_\beta} f_{n\mathbf{k}} = & e \frac{\partial f_{n\mathbf{k}}^0}{\partial \varepsilon_{n\mathbf{k}}} v_{n\mathbf{k},\beta} \tau_{n\mathbf{k}} + \frac{2\pi \tau_{n\mathbf{k}}}{\hbar} \sum_{m\nu} \int \frac{d\mathbf{q}}{\Omega_{\text{BZ}}} |g_{m\nu}(\mathbf{k}, \mathbf{q})|^2 \\ & \times [(n_{q\nu} + 1 - f_{n\mathbf{k}+\mathbf{q}}^0) \delta(\Delta \varepsilon_{\mathbf{k},\mathbf{q}}^{nm} + \hbar \omega_{q\nu}) \\ & + (n_{q\nu} + f_{n\mathbf{k}}^0) \delta(\Delta \varepsilon_{\mathbf{k},\mathbf{q}}^{nm} - \hbar \omega_{q\nu})] \partial_{E_\beta} f_{m\mathbf{k}+\mathbf{q}}, \end{aligned} \quad (10)$$

where $\Delta \varepsilon_{\mathbf{k},\mathbf{q}}^{nm} = \varepsilon_{n\mathbf{k}} - \varepsilon_{m\mathbf{k}+\mathbf{q}}$, $f_{n\mathbf{k}}^0$ is the equilibrium distribution function, and $n_{q\nu}$ is the Bose-Einstein occupation. The matrix elements $g_{m\nu}(\mathbf{k}, \mathbf{q})$ are the probability amplitudes for scattering from an initial electronic state $n\mathbf{k}$ to a final state $m\mathbf{k} + \mathbf{q}$ via a phonon of branch index ν , crystal momentum $\mathbf{q}$, and frequency $\omega_{q\nu}$:

$$g_{m\nu}(\mathbf{k}, \mathbf{q}) = \left[\frac{\hbar}{2M_\kappa \omega_{q\nu}} \right]^{1/2} \langle \psi_{m\mathbf{k}+\mathbf{q}} | \partial_{q\nu} V | \psi_{n\mathbf{k}} \rangle, \quad (11)$$

where M_κ is the mass of the atom κ and $\partial_{q\nu} V$ is the derivative of the self-consistent potential associated with a phonon of wave vector $\mathbf{q}$. $\psi_{n\mathbf{k}}$ is the electronic wave function for band n and wave vector $\mathbf{k}$.

The quantity $\tau_{n\mathbf{k}}$ in Eq. (10) is the relaxation time, and is given by [89,90]

$$\begin{aligned} \frac{1}{\tau_{n\mathbf{k}}} = & \frac{2\pi}{\hbar} \sum_{m\nu\sigma} \int \frac{d\mathbf{q}}{\Omega_{\text{BZ}}} |g_{m\nu}(\mathbf{k}, \mathbf{q})|^2 \\ & \times [(n_{q\nu} + 1 - f_{m\mathbf{k}+\mathbf{q}}^0) \delta(\Delta \varepsilon_{\mathbf{k},\mathbf{q}}^{nm} - \hbar \omega_{q\nu}) \\ & + (n_{q\nu} + f_{m\mathbf{k}+\mathbf{q}}^0) \delta(\Delta \varepsilon_{\mathbf{k},\mathbf{q}}^{nm} + \hbar \omega_{q\nu})]. \end{aligned} \quad (12)$$

In our calculations we first compute Eq. (12), then solve Eq. (10) iteratively to obtain $\partial_{E_\beta} f_{n\mathbf{k}}$, and we use the result inside Eq. (9).

A common approximation for calculating mobilities is to neglect the second term on the right-hand side of Eq. (10). In this case, the relaxation time is explicitly given by Eq. (12), and the equations are solved non-self-consistently.

In Ref. [85] we called this simplification the self-energy relaxation time approximation (SERTA), and the corresponding mobility is given explicitly by

$$\mu_{e,\alpha\beta}^{\text{SERTA}} = \frac{e}{n_e \Omega} \sum_{n \in \text{CB}} \int \frac{d\mathbf{k}}{\Omega_{\text{BZ}}} \frac{\partial f_{n\mathbf{k}}^0}{\partial \varepsilon_{n\mathbf{k}}} v_{n\mathbf{k},\alpha} v_{n\mathbf{k},\beta} \tau_{n\mathbf{k}}. \quad (13)$$

The main advantage of the SERTA is that the grids of $\mathbf{k}$ points and $\mathbf{q}$ points do not need to be commensurate. In practice, this allows for a denser sampling of the momentum regions that contribute the most to the mobility, i.e., regions close to the band edges. These grids typically converge faster than homogeneous or random grids for the same number of points. We now detail our computational setup used to evaluate Eq. (9).

B. Electron-phonon matrix elements and Brillouin-zone integrals

The key challenge in numerically evaluating Eq. (9) is related to the fact that the mobility converges very slowly with the number of $\mathbf{k}$ and $\mathbf{q}$ points included [85]. This translates into having to compute the electron-phonon matrix elements $g_{m\nu}(\mathbf{k}, \mathbf{q})$ from Eq. (11) for millions of momentum points.

In the case of SERTA calculations, we interpolate the electron-phonon matrix elements using Wannier functions [91], from a coarse $6 \times 6 \times 6$ Brillouin-zone grid to a dense grid with 73 000 $\mathbf{k}$ points and 205 000 $\mathbf{q}$ points. The $\mathbf{q}$ points follow a Cauchy distribution of width $0.02 \text{ \AA}^{-1}$ centered at Γ , and are weighted according to their Voronoi volume [92]. All the mobility calculations in this work are based on density-functional theory including spin-orbit coupling (SOC) for the Kohn-Sham states, density-functional perturbation theory for phonons and electron-phonon matrix elements, and many-body perturbation theory for *GW* quasiparticle corrections, as implemented in the software packages QUANTUM ESPRESSO [39], YAMBO [63], WANNIER90 [93], and EPW [91,94]. The convergence tests for the SERTA drift mobility are presented on Figs. 4(a) and 4(b). We observe a fast convergence with sampling size. We obtained a room-temperature electron and hole mobility of 452 and $18 \text{ cm}^2/\text{Vs}$, respectively.

In the case of the complete self-consistent solution of Eqs. (9)–(12), we use a homogeneous grid with $100 \times 100 \times 100$ $\mathbf{k}$ points and $\mathbf{q}$ points. In this latter case we rely on the crystal symmetry operations on the $\mathbf{k}$ -point grid to reduce the number of electron-phonon matrix elements to be explicitly computed. We emphasize that, while in some systems the SERTA is accurate enough for predictive calculations of carrier mobilities [82,85], this is not true in general [86]. In the present case of wurtzite GaN, we find that the self-consistent solution of the BTE yields enhancement factors of about 2 of the electron and hole mobility upon the values obtained within the SERTA, as shown on Figs. 4(c) and 4(d). This leads to electron and hole room-temperature drift mobilities of 830 and $42 \text{ cm}^2/\text{Vs}$, respectively. Therefore, it is very important to always benchmark SERTA results versus the complete solution of the BTE. The corresponding Hall mobility of 1034 and $52 \text{ cm}^2/\text{Vs}$ for electron and hole, respectively; will be discussed in Sec. IV E.

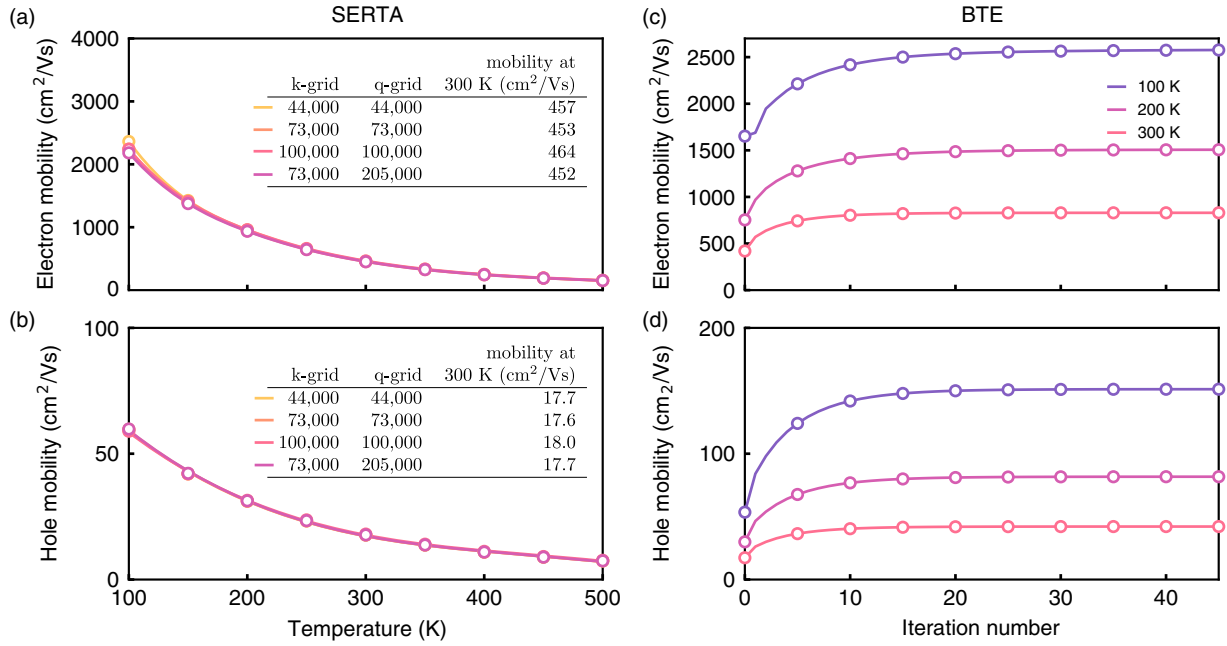


FIG. 4. Convergence tests for the electron and hole mobilities of wurtzite GaN. (a), (b) Electron and hole mobility vs temperature, for different sizes of the Brillouin-zone grids of electrons (**k**) and phonons (**q**). The calculations are performed within the SERTA using LDA and Cauchy grids. (c), (d) Electron and hole mobilities of GaN using uniform 100×100×100 grids, calculated iteratively from the self-consistent BTE, as a function of the number of iterations.

We found that the self-consistent calculations converge rapidly, in about 20 iterations, and without the need for linear mixing. The convergence could be accelerated by methods such as conjugate gradients [82], but since the iterations are fast in comparison to the calculation of scattering rates, we find it unnecessary to improve the iterative solver.

C. Band velocity

In Ref. [85] the band velocities $v_{n\mathbf{k},\alpha}$ appearing in Eqs. (9) and (10) for silicon were computed by neglecting the **k** derivatives of the ionic pseudopotentials. We named this approach the “local velocity approximation”:

$$v_{n\mathbf{k},\alpha} = \mathbf{k}\delta_{mn} + \sum_{\mathbf{G}} c_{n\mathbf{k}}(\mathbf{G})^* c_{m\mathbf{k}}(\mathbf{G}) \mathbf{G}, \quad (14)$$

where $c_{n\mathbf{k}}$ are the plane-wave coefficients. The “exact” band velocity can be computed as [95,96]

$$v_{n\mathbf{k},\alpha} = \frac{1}{\hbar} H_{n\mathbf{k},\alpha} - \frac{i}{\hbar} (\varepsilon_{m\mathbf{k}} - \varepsilon_{n\mathbf{k}}) A_{m\mathbf{k},\alpha}, \quad (15)$$

where $H_{n\mathbf{k},\alpha}$ and $A_{m\mathbf{k},\alpha} = i\langle u_{n\mathbf{k}} | \partial_{\alpha} u_{m\mathbf{k}} \rangle$ are the **k** derivatives of the Hamiltonian and position operator in the direction α , interpolated on the fine momentum grids, and $u_{n\mathbf{k}}$ is the periodic part of the wave function. In both cases, the velocity $v_{n\mathbf{k},\alpha}$ is obtained by taking the diagonal elements of $v_{n\mathbf{k},\alpha}$ from Eqs. (14) or (15).

We find that the local velocity approximation is inadequate in GaN, and it can lead to an overestimation of the mobilities by up to 50%. More specifically, the room-temperature electron and hole mobility increases by 46% and 30%, respectively, when using Eq. (15) instead of Eq. (14). In the case of silicon and the LDA, the electron and hole mobility increase

by 15% and 16%, respectively, when using Eq. (15) instead of Eq. (14). As shown in Fig. 5, this effect is not sensitive to the choice of the pseudopotential, but it depends strongly on the system under consideration. Throughout this paper, we use the velocities given by Eq. (15).

D. Hall factor

In many experiments it is common to measure the Hall mobility μ_H instead of the drift mobility μ of Eq. (9). In order to perform meaningful comparisons, we calculate the Hall factor r_H and obtain the Hall mobilities $\mu_H = r_H \mu$. Following Refs. [97] (p. 118) and [98] [Eq. (3.12)], we calculate the temperature-dependent Hall factor as the ratio

$$r_H = \langle \tau^2 \rangle / \langle \tau \rangle^2, \quad (16)$$

where

$$\langle \tau^n \rangle = \frac{\int_0^\infty \tau^n(x) x^{3/2} e^{-x} dx}{\int_0^\infty x^{3/2} e^{-x} dx} \quad (17)$$

is an energy-averaged carrier scattering rate, and $x = \varepsilon/k_B T$. The energy-dependent scattering rates are obtained through

$$\tau(\varepsilon) = \sum_n \int \frac{d\mathbf{k}}{\Omega_{\text{BZ}}} \delta(\varepsilon - \varepsilon_{n\mathbf{k}}) \tau_{n\mathbf{k}}, \quad (18)$$

where the Dirac deltas are evaluated using Gaussian of width 1 meV. The calculated Hall factors for electrons and holes as a function of temperature are reported in Fig. 6. The values range from 1.07 to 1.37 across the whole temperature range. These data are for unstrained GaN. We checked that the Hall factor is not sensitive to strain for the other cases considered in this work.

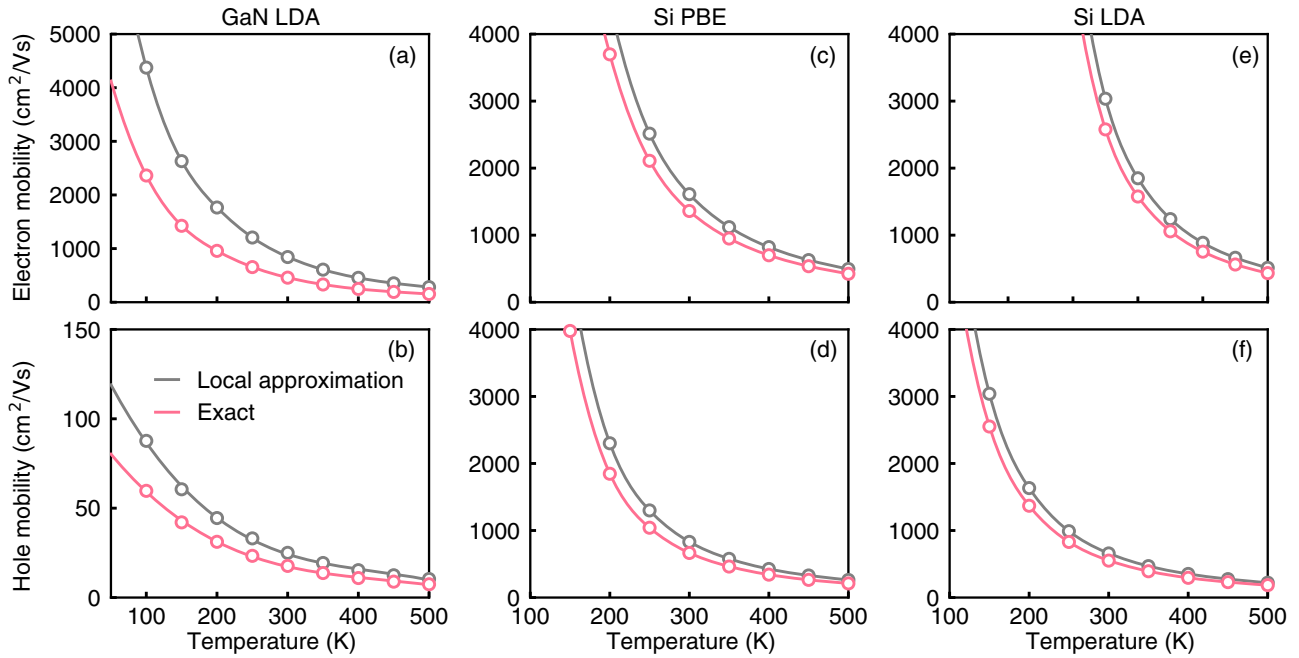


FIG. 5. Comparison between calculations of mobility using the “local approximation” to the band velocity and the “exact” velocity, which takes into account the contribution from the nonlocal part of the ionic pseudopotentials. (a), (b) Electron and hole mobility in GaN versus temperature, both in the local approximation (gray) and using exact velocities (red). Same comparison, this time for silicon, using a PBE pseudopotential (c), (d) or LDA pseudopotential (e), (f).

E. Carrier mobility in unstrained GaN

Using Eq. (9), we computed the drift and Hall mobilities for electrons and holes in intrinsic GaN as a function of temperature. In Table II, we compare our results with available experimental data. We find that the Hall mobility is about 15% higher than the drift mobility at room temperature, as expected [101]. Our predicted electron and hole Hall mobilities at 300 K are 1034 and 52 cm²/Vs, respectively. They are in good agreement with the measured values 1265 cm²/Vs [23] and 31 cm²/Vs [22], respectively. The Hall mobility at 100 K is computed to be 3941 cm²/Vs for electrons and 230 cm²/Vs for holes. Since at room temperature the Cauchy grid yields mobilities which are converged within 1%, and

uniform grids yield mobilities converged within 10%, we use the ratio between the BTE and SERTA mobilities on uniform grids to estimate the BTE mobilities on dense Cauchy grids.

TABLE II. Electron and hole mobilities of wurtzite GaN, calculated using the *ab initio* Boltzmann formalism in the self-energy relaxation time approximation (SERTA) and iterative form (BTE), compared with experiment. We show both the drift mobilities computed via Eqs. (9)–(12) and the Hall mobilities obtained by applying the Hall factor shown in Fig. 6 (bold).

Temperature (K)	Electron mobility (cm ² /Vs)				
	Drift mobility		Hall mobility		
	SERTA	BTE	BTE+ scaling	Experiments	
100	2363	3686	3941	3332 [23]	2202 [99]
200	958	1916	2157	2420 [23]	1700 [99]
				540 [24]	540 [100]
300	457	905	1034	1265 [23]	840 [99]
				330 [24]	370 [100]
400	247	480	541	400 [99]	160 [24]
				245 [100]	
500	154	299	326	250 [99]	100 [24]
				150 [100]	
Temperature (K)	Hole mobility (cm ² /Vs)				
	SERTA	BTE	BTE+ scaling	Experiments	
100	60	168	230		
200	31	85	105	83 [22]	
300	18	44	52	31 [22]	
400	11	25	28	14 [22]	
500	7	15	16		

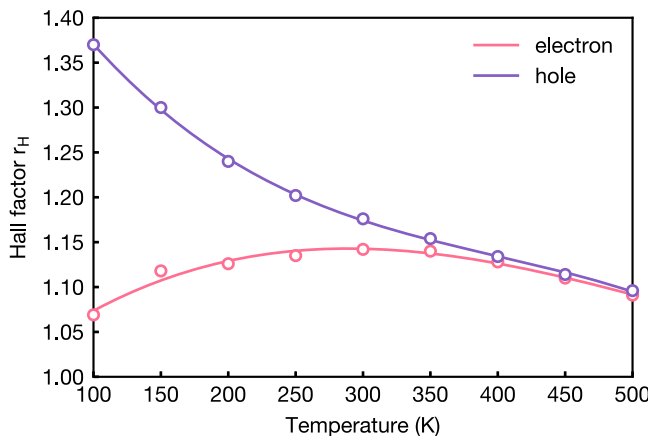


FIG. 6. Calculated temperature-dependent Hall factor of unstrained wurtzite GaN, for electrons and holes.

Direct BTE calculations are not possible on such grids due to the commensurability requirement in Eq. (10). The results reported throughout the paper correspond to this ratio; the conclusions of the paper remain unchanged if we use the BTE results for homogeneous grids.

In Table II, we see that the electron mobility is $1034/52 \approx 20$ times higher than the hole mobility. Experimentally, this ratio is even larger, $1265/31 \approx 41$. To understand the origin of the large difference between the electron and hole mobilities in GaN, we refer to Eqs. (9)–(12). In the simplified case of parabolic bands, the mobility in Eq. (9) scales as $e\tau/m^*$ following Drude's law, with m^* and τ being the average effective mass and relaxation rate, respectively. As discussed in Sec. III, Table I, and Fig. 3, the ratio between the conductivity effective masses $3/(m_{\parallel}^{-1} + 2m_{\perp}^{-1})$ of electrons and holes is 2.4/2.9 for the hh/lh case, respectively. These values are significantly lower than the observed ratio of electron to hole mobilities, therefore, the difference between electron and hole effective masses alone cannot fully account for the order-of-magnitude difference in carrier mobilities.

To determine the origin of the residual difference between electron and hole mobilities, we analyzed in Ref. [29] the angular averages of the carrier relaxation rate $1/\tau$. Although every electronic state has its own lifetime τ_{nk} in our calculation, we have shown previously that the most representative carrier energy [66] comes out from an energy $k_B T = 25$ meV away from the band edges, and these are the values that we used for our analysis in Ref. [29].

By examining the scattering rates and their spectral decomposition $\partial\tau^{-1}/\partial\omega$ in terms of phonon energy, we found that the dominant scattering channel is from long-wavelength acoustic phonons around a phonon energy of 2 meV (77% and 84% of the scattering rates for electrons and holes, respectively). The remaining contribution is from polar Fröhlich longitudinal-optical (LO) phonons near Γ , and located around a phonon energy of 91 meV both for electron and holes. The largest source of acoustic scattering in the case of holes is from acoustic-deformation-potential (ADP) scattering.

In the case of ADP scattering, the scattering rate scales with the electronic density of states, and hence with the effective masses, as $1/\tau \sim (m^*)^{3/2}$ [101]. Using the angular averages, we computed the electron lifetimes to be in the range of 17 fs, while the hole lifetimes of around 4 fs are much shorter. Their ratio is similar to the ratio between the conductivity effective masses $(m_{\text{hh}}^*)^{3/2}/(m_{\text{e}}^*)^{3/2} = 3.7$ and $(m_{\text{hh}}^*)^{3/2}/(m_{\text{e}}^*)^{3/2} = 4.9$. This highlights the fact that the high density of lh and hh states plays a central role in reducing the hole mobility.

The combination of higher effective masses and larger density of states accounts for most of the mobility difference between electron and hole in GaN. The remaining difference can be attributed to other effects that are also responsible for reducing hole mobility. For example, the strong non-parabolicity of the hh in-plane band will increase the effective masses for states with momentum slightly away from the zone center [49,102]; the fact that GaN has multiple scattering channels for the holes (two spin-split sets of bands) will also increase the overall scattering [50,74]; and longitudinal-optical phonons contribute about 20% of additional scattering.

TABLE III. Comparison between calculated elastic constants C_{ij} , bulk B , Young E , shear G modulus, and Poisson's ratio ν (LDA calculations without spin-orbit coupling) with prior theoretical and experimental work.

	C_{11} (GPa)	C_{12} (GPa)	C_{13} (GPa)	C_{33} (GPa)	C_{44} (GPa)	C_{66} (GPa)
This work						
LDA	435	108	65	474	115	163
Previous work						
LDA [108]	346	148	105	405	76	99
LDA [109]	367	135	103	405	95	116
LDA [110]			104	414		
LDA [111]	374	127	81	442	99	124
Exp. [112]	390	145	106	398	105	123
Exp. [113]	365	135	114	381	109	115
Exp. [114]	370	145	110	390	90	112
Exp. [115]	373	141	80	387	94	118
		B	E	G	ν	
This work		(GPa)	(GPa)	(GPa)		
LDA		202	358	148	0.205	
Previous work						
LDA [102]		207				
LDA [109]		202				
LDA [110]		207	373		0.202	
LDA [111]		196	303	122	0.240	
Exp. [31]		188				
Exp. [112]		210	356	120	0.198	
Exp. [113]		205	293	116	0.261	
Exp. [114]		207	276	108	0.278	
Exp. [115]		192	286	114	0.252	
Exp. [104]			295		0.250	
Exp. [105]			295	116	0.250	

All these effects are fully accounted for in our *ab initio* BTE formalism.

Now that we have a better understanding of the various mechanisms behind the low hole mobility, we can proceed to computational design of higher-mobility *p*-type GaN. Since the low mobilities stem primarily from the presence of two adjacent bands with heavy masses, we investigate whether we can employ strain to change the energetics and ordering of the valence band maximum states.

V. CARRIER MOBILITY OF STRAINED GaN

A. Elastic properties

We studied the elastic properties of wurtzite GaN using the THERMO_PW code [40]. The stiffness matrix C_{ij} was obtained by third-order polynomial fitting under 12 deformations geometries of small strain intervals of 0.001 to remain in the linear regime. For each strain, the ions were relaxed to their equilibrium positions. The stiffness matrix of Laue class D_{6h} for wurtzite crystals has five independent elastic constants C_{11} , C_{12} , C_{13} , C_{33} , and C_{44} . The other coefficients follow the symmetry relationships $C_{23} = C_{13}$, $C_{55} = C_{44}$, and $C_{66} = (C_{11} - C_{12})/2$. The computed stiffness constants are given in Table III and are compared to prior theoretical and

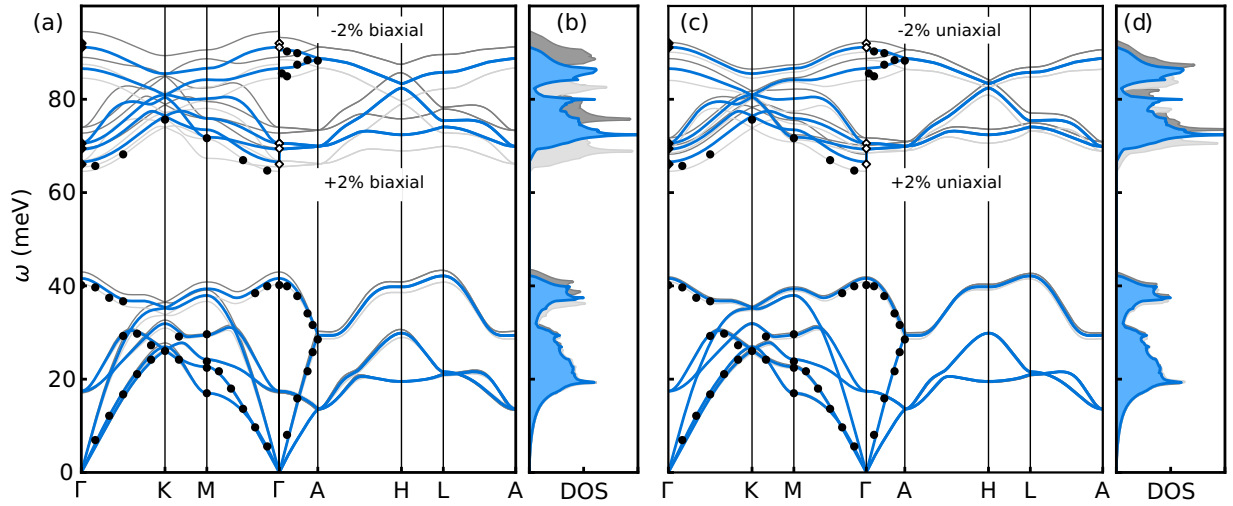


FIG. 7. Phonon dispersion relations (a), (c) and phonon density of states (b), (d) of wurtzite GaN at the relaxed (blue lines) or strained (gray lines) atomic positions. The experimental data are from Ref. [116] (filled disks, inelastic x-ray scattering) and from Ref. [117] (empty diamonds, Raman).

experimental values. In the Voigt approximation, the bulk and shear moduli are given by [103]

$$9B_V = C_{11} + C_{22} + C_{33} + 2(C_{12} + C_{13} + C_{23}), \quad (19)$$

$$15G_V = C_{11} + C_{22} + C_{33} - (C_{12} + C_{13} + C_{23}) + 3(C_{44} + C_{55} + C_{66}), \quad (20)$$

while in the Reuss approximation, the bulk and shear modulus are defined as [103]

$$B_R^{-1} = S_{11} + S_{22} + S_{33} + 2(S_{12} + S_{13} + S_{23}), \quad (21)$$

$$15G_R^{-1} = 4(S_{11} + S_{22} + S_{33}) - 4(S_{12} + S_{13} + S_{23}) + 3(S_{44} + S_{55} + S_{66}), \quad (22)$$

where $S_{ij} = C_{ij}^{-1}$ is the elastic compliance matrix. The Voigt approximation provides an upper bound for the bulk and shear moduli, while the Reuss approximation gives a lower bound. We can therefore define the arithmetic mean, referred to as the Voigt-Reuss-Hill approximation [103], as $B = (B_V + B_R)/2$ and $G = (G_V + G_R)/2$. We then express effective Young E modulus and Poisson ratio ν as

$$E = 9BG/(3B + G), \quad (23)$$

$$\nu = (3B - 2G)/(6B + 2G). \quad (24)$$

Using Eqs. (19)–(24), we obtained a bulk modulus of 202 GPa, a Young modulus of 358 GPa, a shear modulus of 148 GPa, and a Poisson's ratio of 0.205. As expected with the LDA, our calculations overestimate the experimental values of 188 GPa [31], 295 GPa [104], and 116 GPa [105] for the bulk, Young, and shear modulus, respectively. In contrast, the calculated Poisson's ratio sits in-between the experimental values 0.183 [106] and 0.23 [107].

B. Phonon dispersion relations

We compute the phonon dispersions and phonon density of states using density functional perturbation theory [118,119] as implemented in QUANTUM ESPRESSO [39], for unstrained GaN as well as under $\pm 1\%$ and $\pm 2\%$ biaxial and uniaxial strain. The phonon dispersions of the unstrained GaN are reported in Fig. 7 in blue. The theoretical curves follow closely the experimental data from inelastic x-ray scattering [116] and the Raman measurements [117]. The calculated unstrained in-plane ($\epsilon_{\infty}^{\perp}$) and out-of-plane ($\epsilon_{\infty}^{\parallel}$) high-frequency dielectric constants are 5.60 and 5.77, respectively. These values slightly overestimate the experimental values of $\epsilon_{\infty}^{\perp} = 5.14$ [120], 5.25 [121], 5.29 [122], or 5.35 [123] as well as $\epsilon_{\infty}^{\parallel} = 5.31$ [120], 5.41 [121]. The corresponding in-plane and out-of-plane Born effective charges are 2.59 and 2.73, respectively; these values are in agreement with earlier theoretical values of 2.60 and 2.74, respectively [110].

We now investigate how the phonon dispersions change under strain. As shown in Fig. 7(a), the zone-center highest E_1 LO phonon hardens from 91.2 to 94.5 meV under 2% biaxial compressive strain, and softens to 87.5 meV under 2% tensile biaxial strain. The same behavior occurs under uniaxial strain, although changes are more modest, as seen in Fig. 7(b): the highest phonon mode hardens to 92.2 meV under 2% compression, and softens to 90.1 meV under 2% traction. This behavior can easily be understood by the fact that a fixed uniaxial strain imposed to a crystal has a smaller effect than a corresponding fixed biaxial strain. Indeed, a $\pm 2\%$ uniaxial strain modifies the overall volume from 98.8% to 101.2% while for biaxial strain, the change of volumes goes from 97.0% to 103.0%.

The high-frequency dielectric constants $\epsilon_{\infty}^{\perp}$ and $\epsilon_{\infty}^{\parallel}$ are almost insensitive to uniaxial and biaxial strain, respectively; while $\epsilon_{\infty}^{\parallel}$ increases from 5.62 (−2% strain) to 5.96 (+2% strain) under uniaxial strain, and $\epsilon_{\infty}^{\perp}$ increases from 5.44 (−2% strain) to 5.80 (+2% strain) under biaxial strain. Finally, the Ga and N Born effective charges have opposite

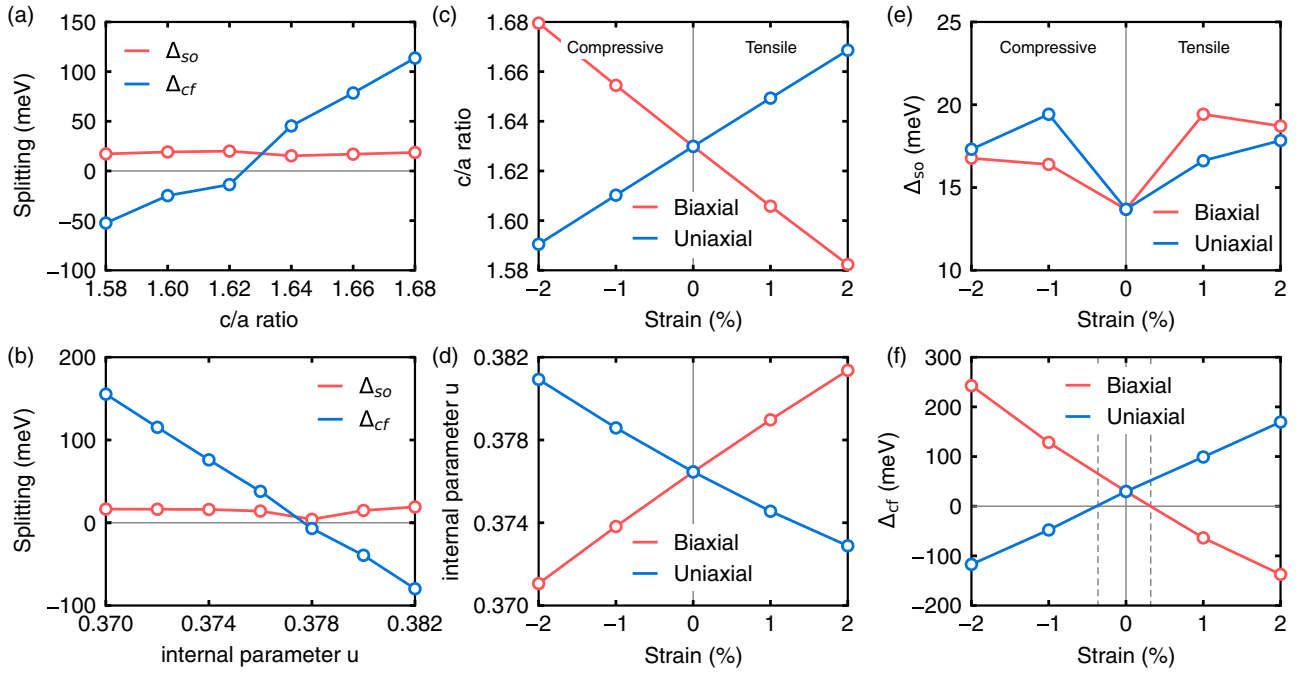


FIG. 8. (a), (b) Sensitivity of the spin-orbit splitting Δ_{so} and the crystal-field splitting Δ_{cf} to the c/a ratio and the internal u parameter in wurtzite GaN. In (a) we fix the internal parameter to $u = 0.3765$, in (b) we fix the aspect ratio to $c/a = 1.6299$. (c), (d) Dependence of c/a and u on biaxial and uniaxial strain, respectively. (e), (f) Dependence of Δ_{so} and Δ_{cf} on the biaxial and uniaxial strain, respectively. The region outside of the dashed lines in (f) leads to a reversal of Δ_{cf} .

values and the largest change is observed for in-plane biaxial strain, going from 2.56 (−2% strain) to 2.62 (+2% strain).

C. Band structures of strained GaN

We now turn to the changes of electronic properties under uniaxial and biaxial strain. As seen in Eq. (7), the lh and hh bands are separated by the spin-orbit splitting Δ_{so} , which we found to be relatively insensitive to the c/a ratio and internal parameter u [see Figs. 8(a) and 8(b)]. However, the separation between the lh/hh and the sh bands is controlled by the crystal-field splitting, Δ_{cf} which is given by Eq. (8). In contrast to Δ_{so} , the crystal-field splitting is known to be sensitive to the internal parameter u of the wurtzite structure, or equivalently to a change of the c/a ratio [102,124]. As seen in Figs. 8(a) and 8(b), Δ_{cf} increases from −52 meV to +114 meV following the increase of the c/a ratio from 1.58 to 1.68. In contrast, the crystal-field splitting decreases linearly from 155 to −80 meV as the internal parameter u increases from 0.370 to 0.382. In Figs. 8(c) and 8(d), we can see that there is a linear correlation between the applied strain and the c/a or internal parameter u . In the case of biaxial strain, the c/a ratio decreases with strain while u increases and the situation is reversed for uniaxial strain.

We therefore see how by combining Figs. 8(a) and 8(b) and 8(c) and 8(d) we can modify Δ_{so} and Δ_{cf} via strain. In particular, we see in Fig. 8(e) that the Δ_{so} has a minimum for unstrained GaN, and slightly increases with strain; while in Fig. 8(f) we see a drastic linear reduction of Δ_{cf} with increasing biaxial strain, going from +243 meV to −137 meV. A smaller linear increase of Δ_{cf} is observed with increasing

uniaxial strain, from −117 to +169 meV for −2% and +2% strain, respectively. We emphasize that the reason for a smaller splitting in Figs. 8(a) and 8(b) is that these results are obtained at fixed u or c/a ratio, while in Figs. 8(e) and 8(f) both c/a and u are changing with strain, contributing to increase the splitting even more. We see that in both types of applied strain, a reversal of the crystal-field splitting is observed. Such reversal of the crystal-field splitting happens for biaxial tensile strain (along $[2\bar{1}10]$ and $[\bar{1}210]$) and for uniaxial compressive strain (along $[0001]$). Using linear interpolation of our results from Fig. 8(f), we estimate that the crystal-field splitting reversal will happen at +0.46% biaxial tensile strain and −0.62% uniaxial compressive strain. Under these conditions, the split-off hole band is lifted above the light-hole and heavy-hole bands, as shown in Fig. 9(a) for the case of uniaxial compression.

In contrast, in the case of applied uniaxial tensile strain shown in Fig. 9(b), we can see that the hh band is almost unaffected with respect to the unstrained cases, while the sh and lh bands get significantly pushed down in energy. The largest change of band energy with strain is indicated with black arrows in Fig. 9. A similar but reversed effect is observed with biaxial strain, as shown in Ref. [29]. In that case the sh band goes above the hh and lh bands under biaxial tensile strain. This effect alters the ordering of the valence band top, as well as the character of the wave functions. As discussed in Ref. [29], the hole wave function at the valence band maximum has a dominant $N-p_{x,y}$ character. In both uniaxial and biaxial strain, the hole wave function keeps this character if no band reversal occurs, but as soon as the sh band goes above the hh and lh bands, it abruptly changes character to a dominant $N-p_z$ state.

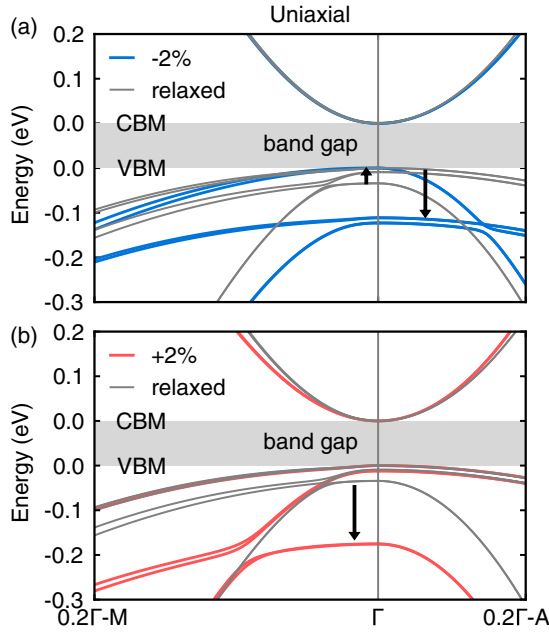


FIG. 9. (a), (b) Change in the GW quasiparticle band structure of GaN upon uniaxial compression and dilation, respectively. The energy levels have been aligned to the band edges.

As seen in Sec. III, the conductivity effective mass of the sh band is $m_{sh}^* = 0.45 m_e$ at the zone center, but away from Γ it quickly decreases to $m_{sh}^* = 0.22 m_e$ due to strong nonparabolicity. This effective mass is much smaller than the masses of the light- and heavy-hole bands. For this reason, we expect the hole mobility of GaN to sharply increase upon reversal of the sign of Δ_{cf} . In the next section, we validate this assumption by performing first-principles calculations of mobility on strained GaN.

D. Mobility of strained GaN

We performed transport calculations for uniaxially and biaxially strained GaN. We computed the drift mobilities of GaN for several strain levels and temperatures using the SERTA approximation with Cauchy grids, and using the more accurate BTE with uniform grids. The uniform grids consist of $100 \times 100 \times 100$ $\mathbf{k}$ and $\mathbf{q}$ points while the random Cauchy grids consist of 45 000 points. The calculated mobilities at different temperatures within the SERTA or BTE are presented in Table IV. First, we computed the mobilities on the homogeneous grid within the SERTA and the iterative solution. We observed an increase in hole mobility ranging between a factor 1.98 to 2.8 across the entire strain and temperature ranges when using the BTE compared to the SERTA. For electrons the ratio is slightly more modest, ranging from 1.17 to 1.98.

As expected, the room-temperature hole mobility significantly increases from $42 \text{ cm}^2/\text{Vs}$ to $113 (117) \text{ cm}^2/\text{Vs}$ upon 2% biaxial tensile (uniaxial compressive) strain. Conversely, the hole mobility remains almost constant when the crystal experiences no reversal of the crystal-field splitting, i.e., under compressive biaxial or tensile uniaxial strain.

As discussed in Sec. IV E, Cauchy grids converge faster than homogeneous grids, but do not allow for BTE calculations due to incommensurability of the momentum grids. However, we noted that the ratio between the BTE to SERTA results is converging faster than the value themselves. Therefore, we used the ratio between the BTE and SERTA mobilities on uniform grids to estimate the BTE mobilities on Cauchy grids. The results are shown in Table IV in bold, and represent our most accurate estimates.

In contrast to the hole mobility, the electron mobility remains close to the value for unstrained GaN in the case of biaxial strain, but decreases slightly in the case of uniaxial strain due to a small increase in the electron effective mass. For this reason, the use of biaxial strain might be more attractive than uniaxial strain, as it leads to a doubling of

TABLE IV. Calculated drift mobilities of GaN for several strain levels and temperatures within the SERTA approximation and the more accurate BTE using uniform grids, SERTA results using Cauchy grids, and BTE results estimated from these data (BTE in boldface). We use the ratio between the BTE (b) and SERTA (a) mobilities on uniform grids to estimate the BTE mobilities on Cauchy grids $e = (b/a)c$.

Strain	Temperature (K)	Holes					Electrons				
		Homogenous grid			Cauchy grid		Homogenous grid			Cauchy grid	
		Mobility (cm ² /Vs)			Mobility (cm ² /Vs)		Mobility (cm ² /Vs)			Mobility (cm ² /Vs)	
		SERTA <i>a</i>	BTE <i>b</i>	Ratio <i>c = b/a</i>	SERTA <i>d</i>	BTE <i>e = cd</i>	SERTA <i>f</i>	BTE <i>g</i>	Ratio <i>h = g/f</i>	SERTA <i>i</i>	BTE <i>j = hi</i>
2% Biaxial	100	289	571	1.98	231	457	1995	3482	1.75	2466	4315
	300	54	113	2.09	46	96	438	847	1.93	459	886
1% Biaxial	100	226	478	2.11	177	373	1945	2836	1.46	2754	4021
	300	38	84	2.21	32	71	453	835	1.84	506	931
Unstrained	100	54	151	2.80	60	168	1652	2584	1.56	2363	3686
	300	17	42	2.47	18	44	420	830	1.98	457	905
−1% Uniaxial	100	176	379	2.15	139	299	1567	2243	1.43	2178	3114
	300	30	67	2.23	25	56	404	730	1.81	439	795
−2% Uniaxial	100	249	519	2.08	206	428	2014	2350	1.17	1976	2312
	300	53	117	2.21	46	102	431	724	1.68	408	685

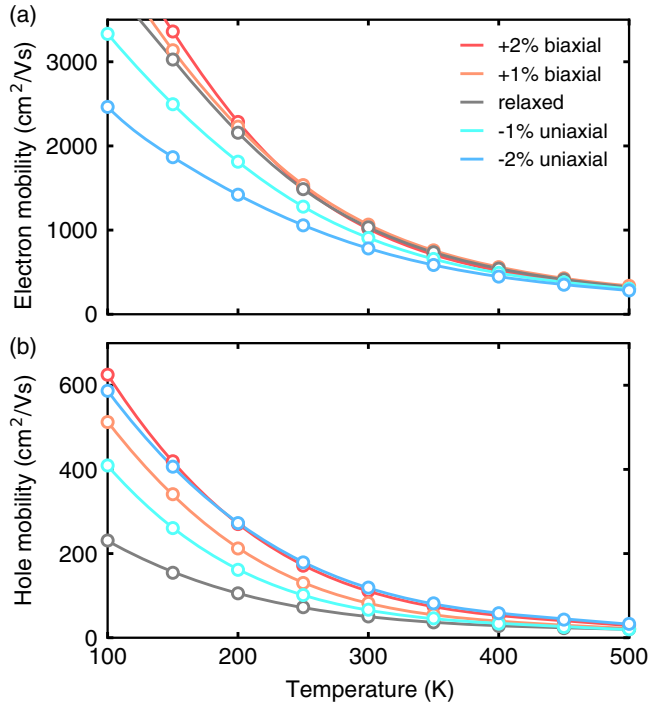


FIG. 10. Predicted temperature-dependent Hall (a) electron and (b) hole mobility in wurtzite GaN as a function of biaxial and uniaxial strain.

the hole mobility with no change to the electron mobility. However, given that the electron mobility is already high and the reduction under strain is in the order of 20%, one could easily imagine a successful device based on uniaxially strained GaN.

Figure 10 shows the electron and hole Hall mobility of GaN computed for biaxial tensile and uniaxial compressive strains of 1% and of 2%. At room temperature the hole Hall mobility increases from 50 cm^2/Vs for the relaxed GaN to 111 and 119 cm^2/Vs for +2% biaxial and -2% uniaxial strain, respectively. This represents a 230% increase in hole mobility. In contrast, the electron mobility is much less affected by strain. The results confirm our expectation that, as soon as we change the sign of Δ_{cf} , we have an enhancement in the hole mobility. We emphasize that these results are not sensitive to the details of the calculations and rests on the *change of ordering* between the split-off band and the light- and heavy-hole bands under applied strain. We confirmed this finding by performing additional calculations of the variation of Δ_{cf} with strain using the PBE functional, obtaining similar results.

VI. FEASIBILITY OF STRAINED GAN

We now investigate the feasibility of realizing high hole mobility GaN experimentally. First, we have computed in Fig. 2 the GaN phase diagram and showed that the wurzite structure remains the lowest-enthalpy phase in a large pressure and temperature range. Second, we noticed that biaxial strain of up to 4% has already been realized experimentally by epitaxial growth on substrates such as AlN or 6H-SiC [110,125,126]. However, in these experiments the large film

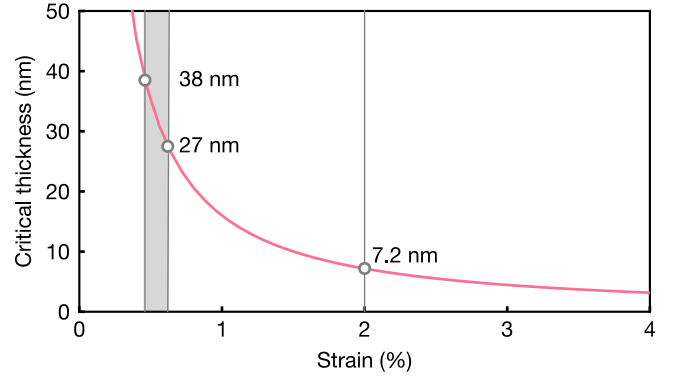


FIG. 11. Critical layer thickness of GaN as a function of strain, estimated using Eq. (25). The gray area represents the minimal strain required for crystal-field splitting inversion under uniaxial or biaxial strain.

thickness induces misfit dislocations [127] to release the strain in the sample. The dislocations increase defect scattering [128], yielding low hole mobility. This may be the reason why high hole mobility GaN has not been observed to date. Therefore, to realize high-hole-mobility GaN we have to devise a plan for preventing dislocation nucleation.

When growing an epilayer on a substrate with a different lattice parameter, dislocations might occur in the epitaxial layer. The most common plastic relaxation mechanism is through the formation of misfit dislocations, to accommodate the strain induced by the substrate [129]. The relaxation of misfit strain via plastic flow occurs for an epitaxial layer with a thickness larger than a critical thickness h_c . Numerous models have been developed over the years to compute the critical thickness. Energy balance models have been developed [130] where the energy of adding a misfit dislocation is balanced with the energy gained by the system from its addition. Another popular model developed by Matthews and Blakeslee [131] is based on the force equilibrium method, in which the forces required to move misfit dislocations are balanced against the elastic stress field due to dislocation interactions. Such a model was later refined by Fischer [132] using an image-force method where the critical thickness h_c for a given strain ϵ is obtained by solving the following nonlinear equation [132]:

$$h_c = \frac{b \cos \lambda}{2\epsilon} \left[1 + \ln \left(\frac{h_c}{b} \right) \left(\frac{1 - \nu/4}{4\pi(1 + \nu) \cos^2 \lambda} \right) \right]. \quad (25)$$

Here, $b = 6.026$ bohrs is the magnitude of the Burgers vector, $\nu = 0.183$ is the experimental Poisson ratio [106], and $\cos \lambda = 0.5$ is the angle between the dislocation Burgers vector and its line direction. Cracks will typically appear for a film thickness above h_c [133,134]. As shown in Fig. 11, we see that at 2% strain, films with thickness of up to 7 nm should not exhibit cracks or misfit dislocations.

It is also possible that the same effect could be achieved using smaller strain levels. Indeed, as soon as reversal of the crystal-field splitting is achieved, the hole mobility should significantly increase. As discussed in Sec. V C, the sh band goes above the lh and hh bands for strain levels above 0.46% in the case of biaxial tensile strain, and above 0.62% for

uniaxial compressive strain. These values correspond to critical film thicknesses of 38 and 27 nm, respectively, as shown in Fig. 11. These values are in agreement with observed critical thicknesses in GaN and AlN, which were found to range between 3 and 30 monolayers depending on the growth temperature [135]. We also note that such type layer thicknesses have recently become accessible for GaN [136,137], making our proposal realistic. In addition, as shown in Table I, our theoretical approach slightly overestimates the crystal-field splitting with respect to experiment and some theoretical studies. As a result, the critical strain required to reverse the crystal-field splitting might be even lower than our estimate. We emphasize that the engineering of mobility via strain is a common strategy in semiconductors such as Si, Ge, and III-V compounds [138–140], but it has become possible only recently in the case of GaN [22,49,50,102,110,124,141].

Finally, an alternative to induce strain via lattice mismatch would be to modify the crystal-field splitting by directly changing the internal parameter u [see Fig. 8(b)]. Given that the internal parameter can be tuned by the A_1 transverse-optical phonon at Γ , it should be possible to reverse the crystal-field splitting by coherently exciting this optical phonon with femtosecond infrared pulses [142,143]. This means that we might be able to control the hole mobility in GaN with light instead of strain.

VII. CONCLUSION

In summary, we have computed the phase diagram of GaN and shown that the wurzite phase is the thermodynamical stable phase for a very wide range of pressure and temperatures. We have analyzed in detail the electronic band structure using many-body corrections and spin-orbit coupling, and showed that the crystal-field splitting heavily depends on the internal parameter of the wurzite structure, and could be tuned via strain engineering. We predicted the room-temperature electron and hole Hall mobilities in unstrained GaN to be 1034 and 52 cm²/Vs, respectively. We showed that the hole mobility can be increased by modifying the ordering of the valence

band top such that split-off holes rise above the light holes and heavy holes. This can be achieved using either biaxial tensile strain or uniaxial compressive strain. We analyzed the effect of strain in GaN including the elastic constants, the high-frequency dielectric constants, Born-effective charges, and phonons. We predict over 200% increase in the hole mobility under strain with respect to the unstrained crystal, reaching values of 120 cm²/Vs under 2% biaxial tensile or uniaxial compressive strain. In contrast, the electron mobility remains mostly unaffected. We propose to realize such band inversion by reversing the the crystal-field splitting via strain engineering or via optical phonon pumping. To avoid cracks or misfit dislocations, we propose the use of ultrathin GaN films (7–40 nm) grown for example by molecular-beam epitaxy on substrates of larger lattice constant than GaN. We hope that this work will stimulate further experimental research in high-hole-mobility GaN, and will accelerate progress toward GaN-based CMOS technology and nitride-based high-power electronics.

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