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Universality of the optical absorptance quantization in two-dimensional group-IV, III-V, II-VI and IV-VI semiconductors

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The optical absorptance of a single graphene layer over a wide range of wavelengths is known to be remarkably constant at the universal value $\pi\alpha$ where α is the fine structure constant. Using atomistic tight-binding calculations, we show that the absorptance spectra of nanometer-thin layers (quantum wells) of group-IV, III-V, II-VI or IV-VI semiconductors are characterized by marked plateaus at integer values of $\pi\alpha$, in the absence of excitonic effects. In the case of InAs, the results obtained are in excellent agreement with the currently available experimental data. By revisiting experimental data on semiconductor superlattices, we show that $\pi\alpha$ is also a metric of their absorption when normalized to a single period. We conclude that the $\pi\alpha$ quantization is universal in semiconductor quantum wells provided that excitonic effects are weak, and is therefore not specific to the zero-gap graphene case. The physical origin of this universality and its limits are discussed using analytical models that capture the main underlying physics of the lowest optical transitions in III-V and II-VI semiconductor quantum wells. These models show that the absorptance is ruled by $\pi\alpha$ independent of the material characteristics because of the presence of a dominant term in the Hamiltonian, linear in the wave vector \mathbf{k} ($\sim \mathbf{V} \cdot \mathbf{k}$), which couples the conduction band to the valence bands. However, the prefactor in front of $\pi\alpha$ is not unity as in graphene due to the different nature of the electronic states. In particular, the spin-orbit coupling plays an important role in bringing the absorptance plateaus closer to integer values of $\pi\alpha$. The case of IV-VI semiconductor (PbSe) quantum wells characterized by a rocksalt lattice and multi-valley physics is very similar to that of graphene, with the specification that a “massful gap” is formed around the Dirac points.

I. INTRODUCTION

The fine structure constant α , introduced by Arnold Sommerfeld [1], characterizes the strength of the interactions between charged particles and the electromagnetic field [2]. In condensed matter physics, this constant quantifies the response to electromagnetic excitation of a large number of very different systems, such as materials with topological phases [3] including the quantum Hall effect [4], graphene [5–10], black phosphorus multilayers [11] or plasmonic networks [12]. In particular, if an undoped graphene monolayer is irradiated, the amount of absorbed light in a wide range of wavelengths is given by the universal constant $\pi\alpha \approx 2.3\%$. Remarkably, this same value appears in the quantization of optical absorption through InAs nanometer thick layers (hereinafter referred to as quantum wells) [13]. In this case, after correction of local-field effects, the absorptance spectrum is characterized by a succession of several plateaus whose respective heights are very close to $\pi\alpha$. Surprisingly, however, the case of quantum wells of semiconductors other than InAs has not been studied so far.

This quantization is not universal. For example, ultra-fine monolayers obtained by exfoliation of van der Waals

materials, such as transition metal dichalcogenides, often show spectra of great richness, with bound exciton absorptances much higher than $\pi\alpha$, especially under the action of strong excitonic effects [14–19]. These effects are also very strong in ultrathin ($\lesssim 2$ nm) layers of semiconductors synthesized by colloidal chemistry, often called nanoplatelets [20, 21].

These observations have inspired theoretical attempts to identify the origin of the $\pi\alpha$ quantization and its intrinsic limits [22–28]. In the case of graphene, deviations from $\pi\alpha$ appear when electrons lose their Dirac fermion character as a non-zero mass, for example at high energy relative to the Dirac point [26–28]. The same effect appears when a gap is opened at the Dirac point, for example under the action of spin-orbit coupling (SOC), as predicted in the case of silicene, germanene and stanene [25], and as shown in Sect. S8 of the Supplemental Material [29] (see, also, Refs. [30–33] therein).

All previous theoretical studies [22–28] were based on two-band models from which it was concluded that the $\pi\alpha$ quantization of the absorptance results from a numerical compensation between the joint density-of-states (JDOS) and the oscillator strength. However, this cancellation of terms is only perfect in the case of pristine graphene, for which the Fermi velocity determines both the band dispersion and the momentum matrix elements over a large energy range [5–10, 28]. In the case of quantum wells of zinc-blende semiconductors which are char-

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TABLE I. Summary of the results for the different types of materials considered in this work. CB and VB stand for conduction and valence bands, respectively. The number of bands per valley must be multiplied by two if spin is taken into account. In the case of Ge, we consider the direct bands at Γ . The integer $n > 0$ indicates that the absorptance present several plateaus, multiple of $\pi\alpha$ or $2\pi\alpha$.

Material	Lattice	Valleys	Number of bands per valley	Band dispersion	SOC	Absorptance plateaus	Unit per valley
Graphene	Honeycomb	K ($\times 2$)	CB:1 VB:1	Linear	No	$\pi\alpha$	$\pi\alpha/2$
III-V or II-VI semiconductor	Zinc-blende	$\Gamma(\times 1)$	CB:1 VB:3	CB: Parabolic at the edge, non-parabolic beyond	Yes, sets VB SOC splitting	$\sim n\pi\alpha$	$\sim \pi\alpha$
Ge	Diamond	$\Gamma(\times 1)$		VB: Complex, anisotropy, warping			
IV-VI semiconductor	Rocksalt	L ($\times 4$)	CB:1 VB:1	Parabolic at edges, almost linear beyond	Yes, opens the gap	$\sim 2n\pi\alpha$	$\sim \pi\alpha/2$

acterized by heavy, light and split-off hole bands with marked anisotropies that couple under the effect of the confinement, a two-band model is not justified [34]. Previous theoretical studies [13, 23, 24, 26] assumed a constant JDOS, typical of two-dimensional (2D) materials with parabolic bands. In reality, however, InAs is characterized by marked non-parabolic effects due to a very strong coupling between valence and conduction bands [35–37]. As a result, the fundamental physical reasons for the $\pi\alpha$ quantization of the absorptance in InAs quantum wells remains largely to be revealed, and so does the universality of this characteristics. In this paper, we show that absorptance quantization can be found in 2D materials characterized by very different band structures, such as graphene, group-IV, III-V and II-VI semiconductors, and even in IV-VI semiconductors with a rocksalt lattice.

In the following, we present detailed atomistic tight-binding calculations of electronic structures and absorptance spectra of semiconductor quantum wells. As summarized in Table I, we predict that, despite band structures of great variety and complexity, quantum wells of many semiconductors are characterized by simple absorptance spectra with marked plateaus, directly interpretable in terms of the universal constant $\pi\alpha$. We show that InAs is not an singular example, but is on the contrary representative of other conventional semiconductors with weak excitonic effects, which is already the case for InAs layers thicker than 3 nm [13]. These predictions are reinforced by published absorption measurements in semiconductor superlattices whose results we revisit [23].

We show that this universality of the absorptance quantization has a common fundamental origin that we are able to reveal through simple analytical models. The nature of the electronic states involved in the optical transitions plays essentially only on the prefactor in front of $\pi\alpha$, which differs according to the categories of materials. This universal behavior thus provides simple rules for the design of photodetectors based on semiconductor quantum wells. It creates an unexpected bridge with the world of graphene, not by coincidence but due to very fundamental reasons. Finally, we remark that reports

show that the fine structure constant also rules spontaneous emission and gain, a topic beyond the scope of the present work [23].

II. METHODOLOGY

A. Expression for the absorptance

The absorptance \mathcal{A} is the measure of the absorbed light flux, such that $\mathcal{R} + \mathcal{A} + \mathcal{T} = 1$ where \mathcal{T} is the transmittance and \mathcal{R} is the reflectance. In the case of a single quantum well deposited on a transparent substrate, the absorptance at photon energy $\hbar\omega$ can be written as

$$\mathcal{A}(\hbar\omega) = F^2 \mathcal{A}_0(\hbar\omega) \quad (1)$$

where F is equal to $2/(n_s + 1)$ in which n_s is the refractive index of the substrate (assumed to be very thick). The prefactor F^2 , which is a local-field factor, describes the fact that the quantum well experiences interfering incident and reflected electric fields. In the case of a sample irradiated perpendicularly, a quantum well being a quasi-2D system, the external electric field (plus its reflected component) is unscreened (for wavelength \gg thickness), the polarization charges being repelled at the edges.

The rate for an optical transition between a valence subband v and a conduction subband c at wavevector \mathbf{k} is determined by the oscillator strength $f_{c,v}(\mathbf{k}) = (2/m_0) |\langle c, \mathbf{k} | \mathbf{p} \cdot \mathbf{e} | v, \mathbf{k} \rangle|^2 / [E_c(\mathbf{k}) - E_v(\mathbf{k})]$ where \mathbf{e} is the light polarization vector, \mathbf{p} is the momentum operator, m_0 is the free electron mass, $E_c(\mathbf{k})$ [$E_v(\mathbf{k})$] is the energy in the conduction [valence] subband. In absence of excitonic effects, the bare absorptance of a quantum well is given by [38, 39]

$$\mathcal{A}_0(\hbar\omega) = \frac{\pi \hbar e^2}{2m_0 c \epsilon_0 S} \sum_{c,v,\mathbf{k}} f_{c,v}(\mathbf{k}) \delta(\hbar\omega - E_c(\mathbf{k}) + E_v(\mathbf{k})) \quad (2)$$

where S is the area of the sample on which we apply periodic boundary conditions that define the components of \mathbf{k} . For numerical implementation, the number of \mathbf{k} vectors is increased until the results are converged, the Dirac delta function $\delta(x)$ in Eq. (2) being replaced by a Gaussian function $\exp[-x^2/(2\eta^2)]/(\eta\sqrt{2\pi})$ where η characterizes all sources of broadening. Unless otherwise specified, η is fixed at 10 meV.

B. Tight-binding calculations of the absorptance

The electronic structure of (001) quantum wells is calculated using an atomistic tight-binding method. Each atom of the compound is described by a set of 20 atomic orbitals, $sp^3d^5s^*$ for each spin degree of freedom where s^* represents a second s orbital. The dangling bonds at quantum well surfaces are saturated with pseudo-hydrogen atoms described by a single s orbital [38], except for PbSe for which it is not necessary [40]. The Hamiltonian matrix elements are restricted to the nearest-neighbor interactions and onsite terms. Spin-orbit coupling is included. We used the tight-binding model of Ref. 41 for InAs, Ref. 40 for PbSe, Ref. 42 for (zinc-blende) CdSe, Ref. 43 for GaAs, Ref. 44 for Ge, Ref. 45 for HgTe. We have verified that very similar results have been obtained with other sets of parameters for PbSe [46] and for Ge [47].

Following a well-established recipe for the description in tight-binding of the optical properties of semiconductors, [38, 48] the momentum matrix elements are determined by those of the Hamiltonian $\mathbf{H}(\mathbf{k})$,

$$\langle c, \mathbf{k} | \mathbf{p} \cdot \mathbf{e} | v, \mathbf{k} \rangle \rightarrow \frac{m_0}{\hbar} \langle c, \mathbf{k} | \nabla H(\mathbf{k}) \cdot \mathbf{e} | v, \mathbf{k} \rangle \quad (3)$$

in which the gradient of $\mathbf{H}(\mathbf{k})$ is relative to \mathbf{k} .

C. Condition for the existence of plateaus in the absorptance

It is interesting to consider the case where the system is isotropic, i.e., where all quantities depend only on k , the modulus of \mathbf{k} . As shown in Appendix A, and as already found by Huang *et al.* [28], the bare absorptance can be written from Eq. (A4) as

$$\mathcal{A}_0(\hbar\omega) = 2\pi\alpha \sum_{c,v} \frac{|\langle c, \mathbf{k} | \nabla H(\mathbf{k}) | v, \mathbf{k} \rangle|^2}{\partial |E_v(k) - E_c(k)|^2 / \partial (k^2)} \quad (4)$$

with k such that $E_c(k) - E_v(k) = \hbar\omega$. ∇H , the gradient of the Hamiltonian with respect to k , comes from the expression of the momentum [Eq. (3)]. The numerator and denominator of Eq. (4) have thus the same unit, i.e., (energy.length)². Within one constant, the inverse of the

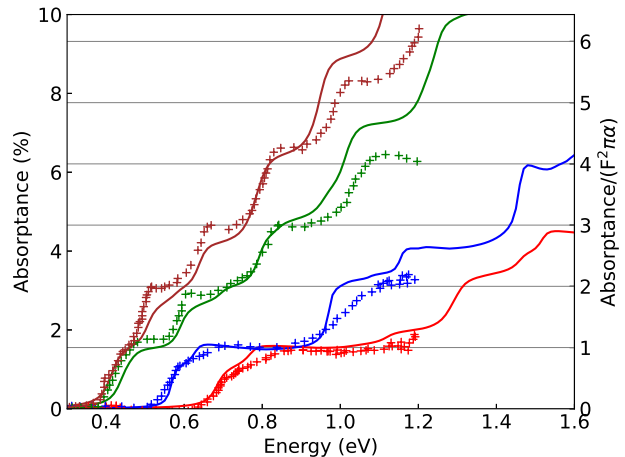


FIG. 1. Crosses: Measured absorptance [13] for InAs quantum wells of different thickness, 6.0 nm (red), 9.0 nm (blue), 14 nm (green), 19 nm (brown). Solid lines: absorptance $\mathcal{A}(\hbar\omega)$ calculated for quantum wells with slightly smaller thickness, 4.0 nm (red), 6.0 nm (blue), 12.5 nm (green), 17.5 nm (brown). Broadening: $\eta = 20$ meV.

numerator is equal to the joint density of states divided by $\hbar\omega$ (Appendix A).

Equation (4) shows that, for the absorptance to be constant, there must be compensation between the optical matrix element $|\langle c, \mathbf{k} | \nabla H(\mathbf{k}) | v, \mathbf{k} \rangle|^2$ and the denominator. As we show in the following, this is likely to occur when a non-diagonal term of the Hamiltonian, linear in k ($\sim \mathbf{V} \cdot \mathbf{k}$ where \mathbf{V} is a constant vector), coupling conduction and valence subbands, dominates. In this case, both the numerator and denominator are proportional to $|\mathbf{V}|^2$ and $\mathcal{A}_0(\hbar\omega)$ is a constant determined by the symmetry of the electronic states.

III. RESULTS AND DISCUSSION

A. InAs

Figure 1 presents the absorptance for InAs quantum wells. The results of the calculations are compared with the experimental data of Ref. 13. We have adjusted the thickness of the quantum wells in order to obtain the correct optical threshold (gap) compared to experiments. A reduction in thickness of the order of 2 nm is needed (Fig. 1), which can be understood by band-bending effects which are well-known in the case of InAs surfaces, due to the pinning of the Fermi level by surface states [49].

The agreement between theory and experiments in Fig. 1 is quite remarkable, showing that excitonic effects which are not taken into account in our calculations are weak in these InAs quantum wells. The absorptance is characterized by very clear steps. Their position, their amplitude, and even their shape when they exhibit fine

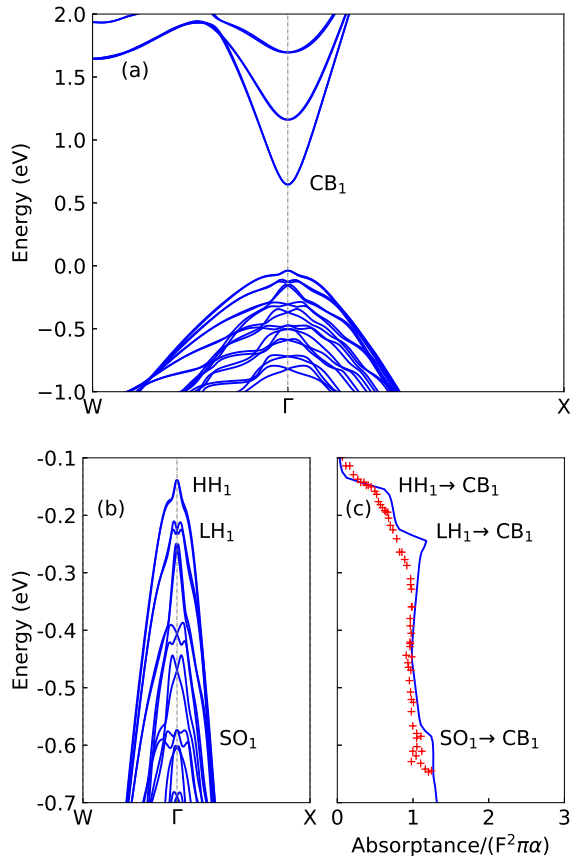


FIG. 2. (a) Lowest conduction subbands and highest valence subbands of a 4 nm thick InAs quantum well. (b) Zoom on the highest valence subbands and, facing each other, (c) plot of the bare absorbance calculated with a smaller broadening ($\eta = 6$ meV) than in Fig. 1 in such a way that each absorbance plateau can be associated to a transition from a valence subband (HH_1 , LH_1 , SO_1) to the lowest conduction subband (CB_1). Red crosses: experiments [13] for a 6 nm thick quantum well. High symmetry points of the Brillouin zone: $W = \pi/a[1, 1, 0]$, $X = 2\pi/a[1, 0, 0]$, a being the bulk lattice parameter.

structures are well described by the calculations. The main plateaus in the spectra correspond to a $n\pi\alpha$ quantization of the bare absorbance $\mathcal{A}_0(\hbar\omega)$.

As shown in Fig. 2a for a 4 nm quantum well, the band structure is composed of subbands due to the vertical quantum confinement effect. The valence subbands (Fig. 2b) exhibit a complex dispersion, due to the contribution of the Heavy-Hole (HH), Light-Hole (LH) and Split-Off (SO) bands [34]. In spite of this complexity, the absorbance shown in Fig. 1 has relatively simple behavior, with main plateaus of $\mathcal{A}_0(\hbar\omega)$ at $n\pi\alpha$ with $n \in \mathcal{N}$ (for additional results for a thickness of 12.5 nm, see Fig. S1 of Supplemental Material [29]).

Figure 2c presents the absorbance calculated using a smaller broadening to reveal the details. This shows that the first plateau is in fact composed of three steps

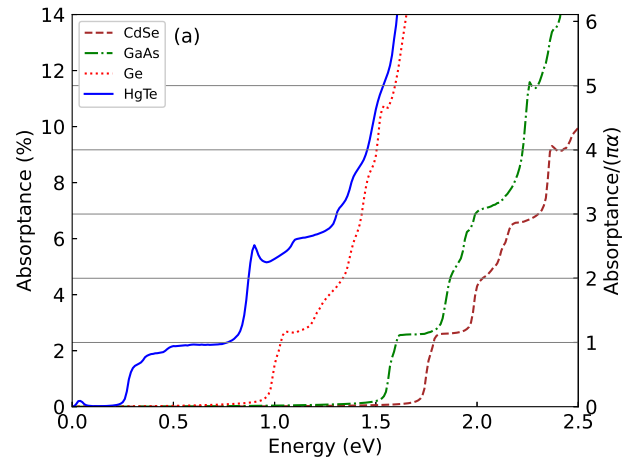


FIG. 3. Bare absorbance $\mathcal{A}_0(\hbar\omega)$ calculated for 6 nm thick quantum wells of CdSe (brown dashed line), Ge (red dotted line), GaAs (green dotted-dashed line), and for a 3.9 nm thick quantum well of HgTe (blue solid line).

corresponding to the transitions from valence subbands HH_1 , LH_1 and SO_1 to the lowest conduction subband CB_1 (Fig. 2a and Fig. 2b). Here the index 1 represents the first quantum mode induced by the vertical confinement in the quantum well. Similarly, a detailed analysis shows that the plateau at $n\pi\alpha$ can be associated with transitions $(\text{HH}+\text{LH}+\text{SO})_n \rightarrow \text{CB}_n$ (Supplemental Material [29], Fig. S1 in Sect. S1).

It is important to point out that the description of the valence subbands in the form HH_n , LH_n and SO_n is only a simplified representation. Indeed, it is not easy to follow the HH, LH, SO character of the subbands as a function of k because of their non-parabolicity, their warping, and their k -dependent mixing under quantum confinement, which lead to complex dispersion including band crossings and anti-crossings (Fig. 2b). However, as shown in Appendix B and in the following sections, this simple representation is justified by extracting a 2D $\mathbf{k} \cdot \mathbf{p}$ model from the atomistic tight-binding calculations. In other words, this very common description [34] of the bands in terms of HH, LH and SO is well justified as long as we only want to describe the main optical transitions which tend to hide the subtle features of the band structure.

B. Other semiconductors

InAs is a semiconductor characterized by a small band gap. It is therefore interesting to test the quantization of absorption in the case of semiconductors with a higher band gap. Fig. 3 shows that the absorbance spectra of GaAs and CdSe quantum wells are also characterized by several well-marked steps close to multiples of $\pi\alpha$. The deviations from $\pi\alpha$ are here larger than for InAs but remain relatively modest.

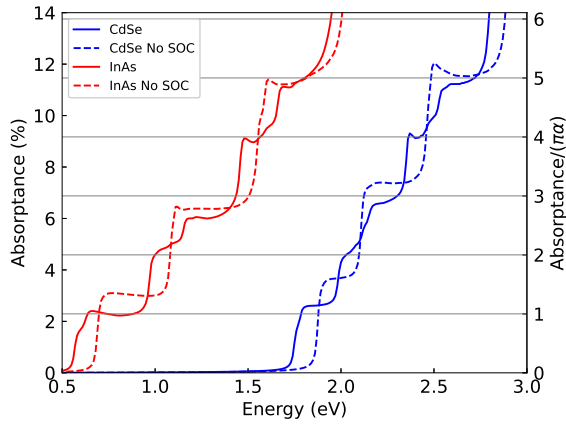


FIG. 4. Bare absorptance $\mathcal{A}_0(\hbar\omega)$ calculated in tight-binding for 6-nm-thick layers of InAs (red curves) or CdSe (blue curves), with (solid lines) or without (dashed lines) SOC.

Remarkably, the first step of $\mathcal{A}_0(\hbar\omega)$ at $\approx \pi\alpha$ is also clearly visible in the case of a Ge quantum well, yet an indirect semiconductor. In fact, in bulk Ge, the minimum of the conduction band at Γ is just above the minima at L points, so the first step corresponds to the vertical transitions in the vicinity of Γ . At higher energy, the steps are less visible, due to the complex conduction band structure of Ge in this region [50].

Another particularly interesting system is HgTe, a semi-metal, which can be seen as an inverted band semiconductor due to the strong SOC. Under the effect of a weak quantum confinement, a band gap opens, leading to the formation of a topological insulator as long as band remains inverted [51–53]. When the thickness of the layers is reduced, the band gap closes and then opens again. In this case, the band order of a conventional semiconductor is restored. Remarkably, a band gap of the order of one electron-volt can even be obtained in ultrathin layers, colloidal nano-platelets [54]. We have therefore considered a 3.9 nm thick (001) HgTe quantum well, characterized by a band gap of 0.19 eV, thin enough to be in the non-inverted regime, thick enough to minimize the excitonic effects. Figure 3 shows that the absorptance spectrum is again characterized by a very marked plateau at $\pi\alpha$. However, this plateau is preceded by several intermediate steps, which are relatively long, but whose contributions converge towards $\pi\alpha$. These steps, as well as the sharp peaks visible at higher energy, are due to the complex energy dispersion of the valence subbands with heavy-hole and light-hole components (Supplemental Material [29], Fig. S2).

C. Role of spin-orbit coupling

Figure 4 illustrates the effect of the SOC. In the absence of SOC, $\mathcal{A}_0(\hbar\omega)$ in InAs and CdSe quantum wells is still quantized, but the height of the first plateau is

closer to $1.5\pi\alpha$ than $\pi\alpha$. This is also true for the following steps which are clearly larger than $\pi\alpha$. When the SOC is restored, the quantization in $\pi\alpha$ becomes remarkable.

In the following, in order to understand these behaviors, we will work in two steps, first, by proposing an analytical model of $\mathcal{A}_0(\hbar\omega)$ in the absence of SOC, second, by understanding the effect of SOC on the absorptance step height.

IV. FUNDAMENTALS OF THE ABSORPTANCE PLATEAUS

A. The prototypical model of graphene

To illustrate the compensation mechanism in Eq. (4), it is interesting to return to the case of graphene to understand what is happening in semiconductor quantum wells. In each valley K or K' of graphene, the Hamiltonian matrix near the Dirac point (zero of energy) can be written, in the basis of the Bloch functions made of p_z orbitals on the two sublattices of the honeycomb, as follows [55]

$$H = \begin{bmatrix} 0 & V(k_x + ik_y) \\ V^*(k_x - ik_y) & 0 \end{bmatrix} \quad (5)$$

where $V = \hbar v_F$ is related to the Fermi velocity v_F .

By defining bottom (Ψ_-) and top (Ψ_+) band states, the numerator and the denominator of Eq. (4) are given by

$$|\langle \Psi_+ | \nabla H(\mathbf{k}) | \Psi_- \rangle|^2 = |V|^2 \quad (6)$$

$$\partial |E_+(k) - E_-(k)|^2 / \partial k^2 = 4|V|^2 \quad (7)$$

There is a total compensation between these two terms in Eq. (4), so that $|V|^2$ disappears [28]. The bare absorptance per valley is therefore $\mathcal{A}_0(\hbar\omega) = \pi\alpha/2$, independent of V and any material characteristics [5–10].

This compensation is no longer effective in the case of massive Dirac bands (Sect. S8 in Supplemental Material [29] and Refs. [25–28]).

B. A prototypical analytical model for semiconductor quantum wells

In this section, we present a three-band model based on the Kane Hamiltonian [56] that provides the simplest description of the absorptance in III-V or II-VI semiconductor quantum wells, in absence of SOC. $\mathcal{A}_0(\hbar\omega)$ is calculated analytically as the sum of two components, the first of which is equal to $\pi\alpha$, the second depends only on the quantum well gap energy and tends to zero at high

energy. This model thus sheds light on the fundamental origins of the quantization of the absorptance and its relative invariance from one semiconductor to another.

The model Hamiltonian H is defined in a basis of three vectors, corresponding to the Bloch states at $\mathbf{k} = 0$, one of s character ($|s\rangle$) for the conduction band, two of p character ($|x\rangle, |y\rangle$) for valence bands, translating the 2D character of the system. Keeping only the principal coupling terms at the first-order in k ($\propto P$), we write

$$H = \begin{bmatrix} E_g & iPk_x & iPk_y \\ -iPk_x & 0 & 0 \\ -iPk_y & 0 & 0 \end{bmatrix} \quad (8)$$

where E_g is the energy gap of the quantum well. Introducing two vectors,

$$\begin{aligned} |p_{\parallel}\rangle &= \frac{k_x|x\rangle + k_y|y\rangle}{k}, \\ |p_{\perp}\rangle &= \frac{k_y|x\rangle - k_x|y\rangle}{k}, \end{aligned} \quad (9)$$

the Hamiltonian in the basis $\{|s\rangle, |p_{\parallel}\rangle, |p_{\perp}\rangle\}$ becomes

$$H = \begin{bmatrix} E_g & iPk & 0 \\ -iPk & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix}. \quad (10)$$

The solutions are $|\Psi_{hh}\rangle = |p_{\perp}\rangle$ of energy $E_{hh} = 0$ (flat band), and the states $|\Psi_e\rangle$ and $|\Psi_{lh}\rangle$ of energy

$$E_{lh}^e(k) = \left(\frac{E_g}{2}\right) \pm \sqrt{\left(\frac{E_g}{2}\right)^2 + P^2k^2}, \quad (11)$$

respectively. $E_e(k)$ is the energy of the conduction subband. $E_{hh}(k)$ and $E_{lh}(k)$ are the energies of HH and LH subbands, respectively. Interestingly, the energy dispersions in this model are basically the same as for massive Dirac fermions.

The absorptance coming from transitions $|\Psi_{hh}\rangle \rightarrow |\Psi_e\rangle$ and $|\Psi_{lh}\rangle \rightarrow |\Psi_e\rangle$ is written as $\mathcal{A}_0(\hbar\omega) = \mathcal{A}_{hh-e}(\hbar\omega) + \mathcal{A}_{lh-e}(\hbar\omega)$ and is calculated analytically using Eq. (4), replacing $(|v, \mathbf{k}\rangle, |c, \mathbf{k}\rangle)$ by $(|\Psi_{hh}\rangle, |\Psi_e\rangle)$ or $(|\Psi_{lh}\rangle, |\Psi_e\rangle)$, respectively. Details on the calculations are given in Supplemental Material [29], Sect. S4.

1. Calculation of the absorptance for the heavy-hole electron transition

For the transition $\Psi_{hh} \rightarrow \Psi_e$ at the energy $\hbar\omega = E_e(k) - E_{hh}(k)$, the optical matrix element is given by (Supplemental Material [29], Sect. S4.1)

$$|\langle \Psi_e | \nabla H(\mathbf{k}) | \Psi_{hh} \rangle|^2 = \frac{P^2 \hbar\omega}{2\hbar\omega - E_g} \quad (12)$$

and the denominator of Eq. (4) by

$$\frac{\partial |E_e(k) - E_{hh}(k)|^2}{\partial k^2} = \frac{2P^2 \hbar\omega}{2\hbar\omega - E_g}. \quad (13)$$

We deduce the absorptance

$$\mathcal{A}_{hh-e}(\hbar\omega) = \pi\alpha \theta(\hbar\omega - E_g) \quad (14)$$

where $\theta(x)$ is the unit step function.

Remarkably, as in the case of graphene, there is a complete compensation between the different terms so that the absorptance is a constant, independent of P , E_g , and $\hbar\omega$. This compensation takes place while the JDOS is not constant (Supplemental Material [29], Sect. S4.1), unlike the ideal case of 2D parabolic valence and conduction bands.

2. Calculation of the absorptance for the light-hole electron transition

For the transition $\Psi_{lh} \rightarrow \Psi_e$ at the energy $\hbar\omega = E_e(k) - E_{lh}(k)$, the optical matrix element is given by (Supplemental Material [29], Sect. S4.2)

$$|\langle \Psi_e | \nabla H | \Psi_{lh} \rangle|^2 = \left(\frac{E_g P}{\hbar\omega}\right)^2 \quad (15)$$

and the denominator of Eq. (4) by

$$\partial |E_e(k) - E_{lh}(k)|^2 / \partial k^2 = 4P^2. \quad (16)$$

The absorptance is thus given by

$$\mathcal{A}_{lh-e}(\hbar\omega) = \frac{\pi\alpha}{2} \left(\frac{E_g}{\hbar\omega}\right)^2 \theta(\hbar\omega - E_g). \quad (17)$$

Also remarkably, this contribution does not depend on P , is equal to $\pi\alpha/2$ when $\hbar\omega = E_g$, and tends to 0 when $\hbar\omega \gg E_g$.

NB: This is at variance with graphene where the transition between the two band gives a plateau at $\pi\alpha/2$. Note that, as in the case of graphene, the JDOS is proportional to $\hbar\omega$ [Eq. (S32) in Supplemental Material [29]]. The difference is therefore due to different local 2D symmetries, i.e., to the different nature of the Bloch states.

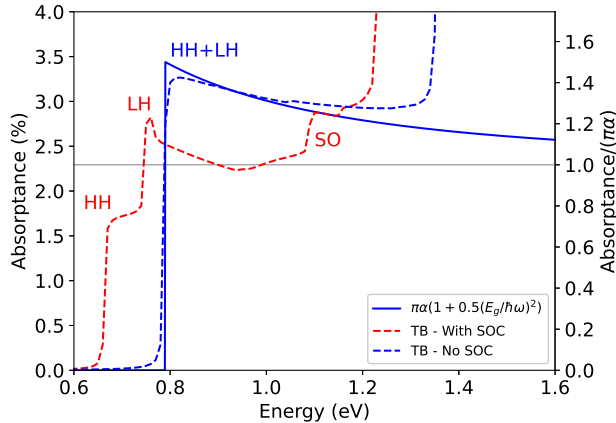


FIG. 5. Bare absorbance $\mathcal{A}_0(\hbar\omega)$ for 4.2-nm-thick layer of InAs calculated in tight-binding (TB, dashed curves) with (red) or without (blue) SOC. The blue solid line shows $\mathcal{A}_0(\hbar\omega)$ given by the simple analytical model, Eq. (18) with $E_g = 0.79$ eV. For the sake of comparison, the tight-binding calculations were performed with a small broadening η of 2 meV.

3. Comparison to tight-binding calculations

The total absorbance in this simple model is therefore given by

$$\mathcal{A}_0(\hbar\omega) = \pi\alpha \left[1 + \frac{1}{2} \left(\frac{E_g}{\hbar\omega} \right)^2 \right] \theta(\hbar\omega - E_g). \quad (18)$$

Figure 5 shows that, despite its great simplicity, the analytical model reproduces remarkably well the results obtained by the tight-binding method, in the absence of SOC. This model describes the essential characteristics of the optical transitions from valence subbands to the first conduction subband. It gives a simple explanation to the quantization of the absorbance, and to its universal character since the Eq. (18) only involves the energy gap as a characteristic quantity of the semiconductor quantum well. As in the case of graphene, $\pi\alpha$ is a metric of the absorbance.

4. Justification of the simple analytical model

We have shown in the previous sections that the valence band structure of III-V or II-VI semiconductor quantum wells is in general of great complexity. In this context, it is not obvious that a three-band model, including one flat band and two fully symmetric bands is representative of real systems.

The general justification of a 2D $\mathbf{k} \cdot \mathbf{p}$ model, starting from a bulk Kane Hamiltonian, is presented in Supplemental Material [29], Sect. S3. We also present and justify in Appendix B a more elaborate (3×3) 2D $\mathbf{k} \cdot \mathbf{p}$ model

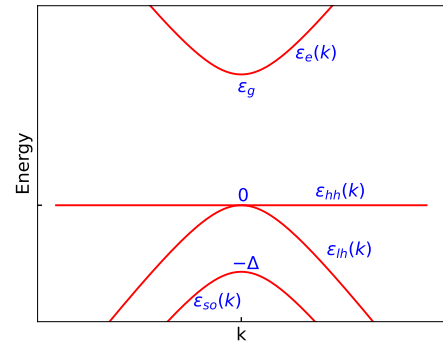


FIG. 6. Band structure in the analytical model including SOC. The energies are noted by the Greek letter ε instead of E to distinguish from the case without SOC.

whose parameters are directly deduced from tight binding calculations, following the methodology described in Supplemental Material [29], Sect. S5. This model includes all matrix elements up to the second order in components of \mathbf{k} . We deduce from Appendix B that the simple analytical model described above is justified because the terms of the form Pk_x and Pk_y of the 2D $\mathbf{k} \cdot \mathbf{p}$ Hamiltonian dominate in the expression of the different components of the absorbance. The simple analytical model is justified even more when the energy gap of the semiconductor is small. The deviations from the absorption values predicted by Eq. (18) remain nevertheless small even for large gap materials.

C. Influence of the SOC on the absorbance

We have seen in Fig. 4 and Fig. 5 that the SOC contributes to bring the plateaus from about $1.5\pi\alpha$ (at E_g) to about $\pi\alpha$. Here we discuss the reasons, on the one hand with the help of a simplest-as-possible analytical model describing the main underlying effects, on the other hand by analyzing the results of the tight binding calculations

1. A prototypical analytical model including SOC

We start from the Kane Hamiltonian [56] including SOC in the basis of Bloch states $|s \uparrow\rangle$, $|x \uparrow\rangle$, $|y \uparrow\rangle$, $|z \uparrow\rangle$, $|s \downarrow\rangle$, $|x \downarrow\rangle$, $|y \downarrow\rangle$, $|z \downarrow\rangle$, where Oz remains perpendicular to the quantum well. We write the Hamiltonian as two identical 4×4 blocks [34]

$$H = \begin{bmatrix} \varepsilon_g & i\sqrt{\frac{2}{3}}Pk & -\frac{i}{\sqrt{3}}Pk & 0 \\ -i\sqrt{\frac{2}{3}}Pk & 0 & 0 & 0 \\ \frac{i}{\sqrt{3}}Pk & 0 & -\Delta & 0 \\ 0 & 0 & 0 & 0 \end{bmatrix} \quad (19)$$

for one block in the basis $|s \uparrow\rangle, |\frac{3}{2}, \frac{1}{2}\rangle, |\frac{1}{2}, \frac{1}{2}\rangle, |\frac{3}{2}, \frac{3}{2}\rangle$, for the other one in the basis $|s \downarrow\rangle, |\frac{3}{2}, -\frac{1}{2}\rangle, |\frac{1}{2}, -\frac{1}{2}\rangle, |\frac{3}{2}, -\frac{3}{2}\rangle$, eigenvectors of \mathbf{J} and J_Z with Z oriented along \mathbf{k} . ε_g is the gap in presence of SOC, and Δ is the splitting between $J = 3/2$ and $J = 1/2$ states due to the SOC. The expression of the $J = 3/2$ and $J = 1/2$ states in terms of $|x \uparrow\rangle \dots |z \downarrow\rangle$ is given in Sect. S7 of the Supplemental Material [29].

In the same spirit as before, following Sect. S3 of the Supplemental Material [29], we treat the 2D problem with $k_z = 0$ ($k \equiv k_{\parallel}$). The corresponding band structure is shown schematically in Fig. 6. The heavy-hole band is flat [$\varepsilon_{hh}(k) = 0$] and is degenerate at $k = 0$ with the light-hole band [$\varepsilon_{lh}(k)$]. The electron band $\varepsilon_e(k)$ starts at ε_g at $k = 0$, the top of the split-off band $\varepsilon_{so}(k)$ is at $-\Delta$.

Therefore in this model, the absorptance spectrum exhibits two steps, the first one at $\hbar\omega = \varepsilon_g$ that include two contributions (heavy and light holes \rightarrow electrons), the second one at $\hbar\omega = \varepsilon_g + \Delta$ (split-off \rightarrow electrons). As described in Sect. S7 of the Supplemental Material [29], the height of these steps can be calculated analytically at $k_{\parallel} = 0$, the energy dispersions of the different bands being obtained by treating Pk terms within second order perturbation theory (becomes invalid when $\Delta \rightarrow 0$). The height of the first and second steps is given by

$$\mathcal{A}_{0,1}(\varepsilon_g) = \pi\alpha \left[\frac{3}{4 \left(1 + \frac{\varepsilon_g}{2(\varepsilon_g + \Delta)}\right)} + \frac{1}{2 \left(1 + \frac{\varepsilon_g}{4(\varepsilon_g + \Delta)}\right)} \right] \quad (20)$$

$$\delta\mathcal{A}_{0,2}(\varepsilon_g + \Delta) = \pi\alpha \left[\frac{1}{2 \left(1 + \frac{\varepsilon_g + \Delta}{\varepsilon_g}\right)} \right] \quad (21)$$

where $\delta\mathcal{A}_{0,2}$ in the second equation means that it must be added to the other contributions at $\hbar\omega = \varepsilon_g + \Delta$ to get the total absorptance.

Using the data corresponding to the InAs quantum well of 4.2 nm (Fig. 5), i.e., $\varepsilon_g \approx 0.7$ eV and $\Delta \approx 0.4$ eV, the height of the first step [Eq. (20)] is about $\pi\alpha$, and the height of the second step [Eq. (21)] is about $0.2\pi\alpha$, in excellent agreement with the tight binding results of Fig. 5 and the experimental data of Ref. 13. Remarkably, Eq. (20) and Eq. (21) only depend on $x = \varepsilon_g/\Delta$, and $\mathcal{A}_0(\varepsilon_g)$ is found to be very close to $\pi\alpha$ in a wide range of values of x . Figure S5 of the Supplemental Material [29] shows a deviation of Eq. (20) from $\pi\alpha$ below 10% for x between 0.5 and 5. This explains the universality of the $\pi\alpha$ quantization of the absorptance in presence of SOC.

2. Comparison to tight-binding calculations

Figure 5 allows to understand the effect of the SOC on the absorptance spectrum calculated in tight-binding. The first step found in absence of SOC transforms in

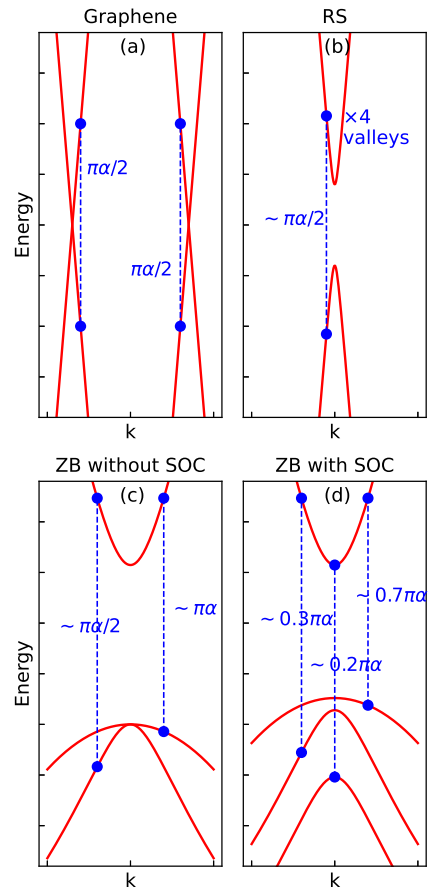


FIG. 7. Schematic representation of electronic bands and optical transitions in graphene (a), in a rocksalt (RS) semiconductor quantum well (b), in a zinc-blende (ZB) semiconductor quantum well without (c) or with (d) SOC. The intensity at the optical thresholds of the absorptance steps is indicated (the values are indicative, they may slightly vary from one material to another). The corresponding data are also summarized in Table I.

three steps. A short plateau at about $0.7\pi\alpha$ is followed, after a small bump, by a long plateau close to $\pi\alpha$. This corresponds to transitions from HH and LH bands, respectively, which were degenerate at $k_{\parallel} = 0$ by construction in the analytical model. Then, at higher energy (~ 1.1 eV), there is a third small step (split-off) such that the absorptance finally reaches the value obtained without SOC. Apart from the fact that it does not give a splitting between LH and HH bands, the analytical model accounts for this behavior very well.

Note that the respective weights of the steps from HH and LH depend significantly on the nature of the semiconductor, the spin-orbit coupling and the vertical confinement.

3. Summary of our understanding

Figure 7 summarizes our understanding of the origin of absorptance quantization in III-V or II-VI semiconductor quantum wells and in graphene. This quantization comes from the compensation between numerator and denominator in Eq. (4). In the case of graphene, this compensation is exact which leads to an absorptance $\pi\alpha/2$ for each of the two K valleys.

In the case of a semiconductor quantum well, the situation seems at first sight very different because there are optical transitions between valence bands mainly derived from p orbitals and a conduction band of strong s character. However, under strong vertical confinement, in absence of SOC, the valence band can be described by two bands coupled to the conduction band by terms like Pk_x and Pk_y . Transitions between the LH subband and the conduction subband lead to an absorptance of the order of $\pi\alpha/2$ at the threshold, decreasing to zero at higher energy. Transitions between the HH subband and the conduction subband give a constant absorptance at $\pi\alpha$. In absence of SOC, the total absorptance is thus characterized by a single step at $\sim 3\pi\alpha/2$ which decreases to $\pi\alpha$ at higher energy (Fig. 5). This reasoning remains correct as long as P^2/E_g is large compared to the other terms of the $\mathbf{k}\cdot\mathbf{p}$ Hamiltonian (Appendix B), which is the situation in many conventional semiconductors [34]. The height of the different contributions of the absorptance can vary from one compound to another, but the previous description remains nevertheless qualitatively correct for all the studied zinc-blende materials, especially those with a small bandgap.

When the SOC is switched on, the absorptance remains quantized because of the compensation effect due to the dominance of the coupling terms Pk_i . However, the bands are split and a certain part of the oscillator strength is transferred to the SO band which is lower in energy. This leads to an absorptance $\approx 0.7\pi\alpha$ for the transitions from HH subband, $\approx 0.3\pi\alpha$ for those from LH subband, and $\approx 0.2\pi\alpha$ for those from SO subband [Fig. 7(d)]. The total contribution from transitions from HH and LH subbands is thus close to $\pi\alpha$, which explains the wide plateau in Fig. 5. Of course, the $\pi\alpha$ quantization of the absorptance is not exact, numbers slightly depend on the nature of the semiconductor, the plateaus are not totally flat because the compensation is not perfect, and therefore do not form a standard for the fine structure constant [24]. However, our calculations show that the quantization is universal, provided that excitonic effects can be neglected.

This quantification of the absorptance is visible as long as the energy gap between the $\pi\alpha$ steps remains smaller than the broadening factors. This gap becomes smaller and smaller as the thickness of the quantum well decreases. In III-V and II-VI materials, this gap is essentially determined by the quantum confinement in the conduction band and thus varies approximately as the inverse of the well thickness square.

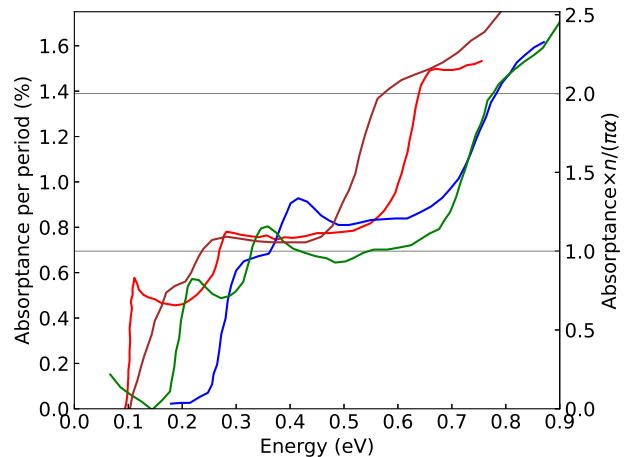


FIG. 8. Absorbance per period in HgTe/Hg_xCd_{1-x}Te superlattices deduced from the measurement of their absorption coefficient (period = well thickness L_w plus barrier thickness L_b). The vertical axis on the right gives the absorptance multiplied by the sample refractive index n normalized by $\pi\alpha$. Red line: data of Ref. 57 for $x = 0.05$, $L_w = 4.15$ nm, $L_b = 8.95$ nm. Brown line: data of Ref. 58 for $x = 0.15$, $L_w = 5.8$ nm, $L_b = 4.2$ nm. Green line: data of Ref. 59 for $x = 0.37$, $L_w = 3.47$ nm, $L_b = 7.60$ nm. Blue line: data of Ref. 60 for $x = 0.32$, $L_w = 3.4$ nm, $L_b = 7.70$ nm.

V. REVISITING EXPERIMENTAL DATA ON SUPERLATTICES

The physical effects explaining the quantization of absorption should be operative also in semiconductor superlattices, provided that the barriers are high and the couplings between quantum wells are weak (multilayers). As already proposed in Ref. 23, this invites us to revisit experimental results obtained on these systems. Typically, the experiments consist of measuring the transmission through relatively thick layers composed of a large number (25-50) of periods, each formed by a quantum well and a barrier. The absorption coefficient $a(\hbar\omega)$ of the medium, deduced from these measurements, is given quite generally by

$$a(\hbar\omega) = \frac{\omega}{cn} \varepsilon''(\hbar\omega) \quad (22)$$

where ε'' is the imaginary part of the dielectric constant and n is the refractive index of the medium. Consequently, the absorptance of a single period can be defined as

$$\mathcal{A}(\hbar\omega) = a(\hbar\omega)L_p \quad (23)$$

where $L_p = L_w + L_b$ is the length of the period, L_w (L_b) being the thickness of the quantum well (barrier). However, we can deduce from Eq. (22) and the theory that we discussed above, that the quantity of interest, the one

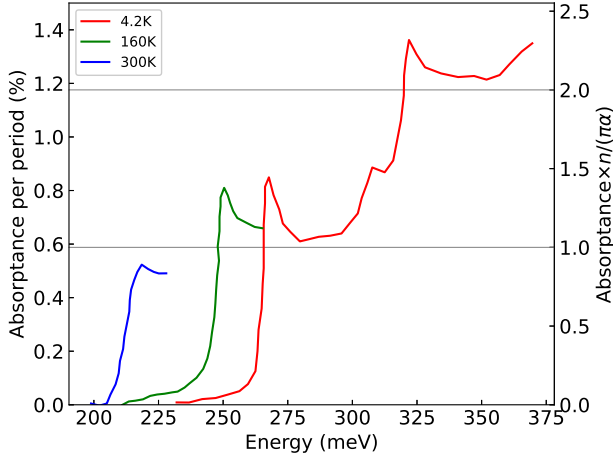


FIG. 9. Absorbance per period in InSb/Al_{0.09}In_{0.91}Sb superlattices deduced from the measurement [61] of their absorption coefficient (well thickness = 22.5 nm, barrier thickness = 50 nm) at different temperatures, 4.2K (red line), 160K (green line), or 300K (blue line). The right vertical axis gives the absorbance multiplied by the sample refractive index ($n = 3.9$) normalized by $\pi\alpha$. Spectra at 160K and 300K were measured in a limited energy range [61].

to be quantized in units of $\pi\alpha$, should be $n\mathcal{A}(\hbar\omega)$ [23]. In the following, we revisit experimental results from the literature, deliberately presenting them by increasing width of their band gap.

Figure 8 presents the absorbance per period which was measured in HgTe/Hg_xCd_{1-x}Te superlattices. The spectra are characterized by a wide plateau for which $n\mathcal{A}(\hbar\omega)$ is approximately equal to $\pi\alpha$. Just at the optical threshold, there is also a smaller step at approximately $3\pi\alpha/4$, which can be attributed to transitions to the heavy-hole subband. Interestingly, the absorbance spectrum that we predicted for a 3.9-nm-thick quantum well of HgTe (Fig. 3) is close to that obtained for HgTe/Hg_{0.37}Cd_{0.63}Te superlattices with $L_w = 3.47$ nm and $L_b = 7.60$ nm. However, the experimental spectra exhibit clear bumps at the step edges, which are absent in calculated ones and can be attributed to excitonic effects.

The quantity $n\mathcal{A}(\hbar\omega)$ measured in InSb/Al_{0.09}In_{0.91}Sb superlattices at 4.2K is also characterized by clear steps at $\pi\alpha$ and $2\pi\alpha$ (Fig. 9), plus marked peaks which were attributed to excitonic effects [61]. These peaks tend to decrease at higher temperature.

The $\pi\alpha$ quantization is also clearly visible in the case of a Ga_{0.47}In_{0.53}As/Al_{0.48}In_{0.52}As superlattice (Fig. 10), although the excitonic peaks become more prominent.

Figure 11 presents results for GaAs/AlAs superlattices, deduced from measurements made at 2K [62]. The average value of $n\mathcal{A}(\hbar\omega)$ is clearly above $\pi\alpha$. Similar values are obtained in GaAs/AlGaAs (not shown) [63]. These values higher than $\pi\alpha$ must be seen as a consequence of strong excitonic effects. However, $\pi\alpha$ remains a good metric of the absorbance.

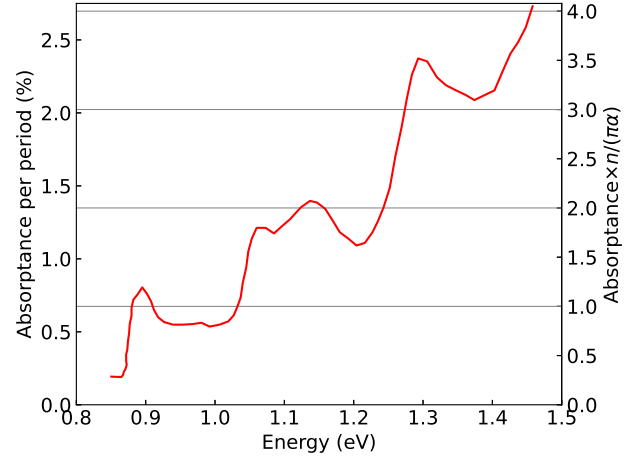


FIG. 10. Absorbance per period in a Ga_{0.47}In_{0.53}As/Al_{0.48}In_{0.52}As superlattice deduced from the measurement [34] of its absorption coefficient (well thickness = 8.5 nm, barrier thickness = 8.5 nm) at 77K (red line). The right vertical axis gives the absorbance multiplied by the sample refractive index ($n = 3.4$) normalized by $\pi\alpha$.

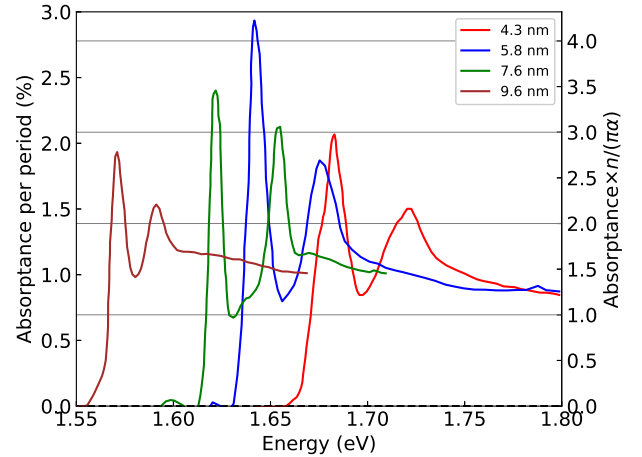


FIG. 11. Absorbance per period in GaAs/AlAs superlattices deduced from the measurement [62] at 2K of their absorption coefficient (period = well thickness L_w plus barrier thickness L_b). The right vertical axis gives the absorbance multiplied by the sample refractive index n normalized by $\pi\alpha$. Red line: $L_w = 4.3$ nm, $L_b = 6.2$ nm. Blue line: $L_w = 5.8$ nm, $L_b = 7.1$ nm. Green line: $L_w = 7.6$ nm, $L_b = 3.3$ nm. Brown line: $L_w = 9.6$ nm, $L_b = 9.1$ nm.

It is important to realize that deviations from $\pi\alpha$ are expected to be stronger in materials characterized by a wide band gap. On the one hand, P^2/E_g becomes less dominant compared to the other components of the $\mathbf{k} \cdot \mathbf{p}$ Hamiltonian [the A , L and M terms in Eq. (B6) and Eq. (B9) of Appendix B]. Larger deviations are clearly visible in Fig. 3 and Fig. 4. On the other hand, the dielectric constant is lower, the effective masses are usually higher (especially in the conduction and light-hole bands

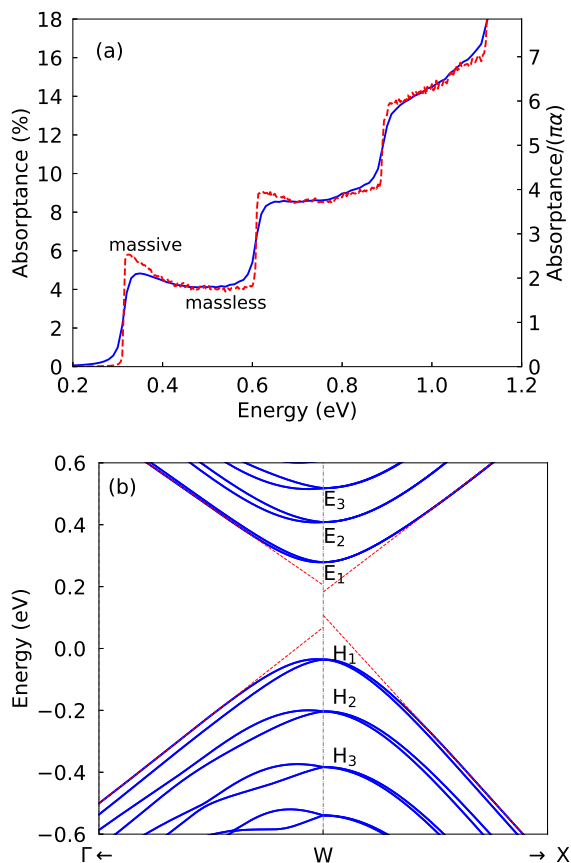


FIG. 12. (a) Bare absorbance $A_0(\hbar\omega)$ calculated for a 6 nm thick PbSe quantum wells for two values of the broadening, $\eta = 10$ meV (blue solid line) or $\eta = 1$ meV (red dashed line). (b) Zoom on the electron (E_i) and hole (H_i) subbands in the vicinity of the gap. The red dotted lines show a linear variation of the subbands. The full band structure is presented in Fig. S8, Sect. S9 of Supplemental Material [29].

where $1/m_{e,lh} \propto P^2/E_g$), which tends to increase the excitonic effects.

VI. QUANTUM WELLS OF PbSe

By comparison with InAs and other III-V or II-VI compounds, the case of PbSe (and other IV-VI semiconductors with rocksalt lattice) is interesting since the bulk material is characterized in conduction and valence bands by four nonequivalent valleys at the L point of the Brillouin zone [40]. In absence of experimental data, we have performed calculations of the absorbance for a 6 nm thick (001) PbSe quantum well. The results, presented in Fig. 12a, show that $A_0(\hbar\omega)$ is also characterized by clear plateaus, but of height $2\pi\alpha$ instead of $\pi\alpha$.

Figure 12b shows the band structure of the quantum well. Due to the quantum confinement, the energy spectrum is characterized by subbands which are (almost) twofold degenerate, fourfold with spin degeneracy (there

is small splitting due to inter-valley coupling [40]). In addition, there are two nonequivalent W ($\pi/a[1, 1, 0]$) valleys in the Brillouin zone of the 2D layer. This behavior can be understood by the projection of the four nonequivalent L points ($\pi/a[1, 1, 1]$) of the bulk on the 2D Brillouin zone. The different absorption steps correspond to the transitions allowed only between subbands sharing the same quantum number associated with vertical confinement, i.e., $H_i \rightarrow E_i$ (Fig. 12b).

Since there are four valleys, absorbance plateaus at $2\pi\alpha$ mean $\pi\alpha/2$ contributions from each valley. This behavior is attributed to quasi-linear dispersions of the conduction and valence subbands as one moves away from the band gap (Fig. 12b), for reasons which are discussed in Ref. 40. We then find the situation of graphene where the absorbance per valley is $\pi\alpha/2$. According to the $\mathbf{k}\cdot\mathbf{p}$ theory, in each valley, subbands H_i and E_i can be described by a 2×2 Hamiltonian in which the non-diagonal coupling, of the form $\mathbf{T} \cdot \mathbf{k}$ where \mathbf{T} is a 2D vector, becomes dominant when moving away from the edges of the subbands. Our tight-binding calculations thus show that the absorbance quantization in PbSe quantum wells has a physical origin much closer to the case of graphene than to the case of III-V or II-VI semiconductors [Fig. 7(b)].

However, the subbands are parabolic in the vicinity of their edges (Fig. 12b). Consequently, if the broadening η is reduced, the absorbance spectrum develops peaks just above each optical threshold (Fig. 12a). We then recover the situation of tight-binding models on honeycomb lattices in which a band gap is opened under the effect of SOC or an asymmetry between the two sublattices of the honeycomb (see Sect. S8 in the Supplemental Material [29]). The peaks followed by the plateaus reflect the transformation from massive particles to massless particles [26–28].

VII. CONCLUSION

In conclusion, we have shown that the quantization of the absorbance in semiconductor quantum wells in units of $\pi\alpha$ is universal. It is predicted for a large number of compounds, provided that the excitonic effects are small. This behavior results from several factors, the main one being the strong coupling of the form Pk_x and Pk_y , between conduction and valence subbands. Using a minimum model incorporating only these dominant terms in the Hamiltonian, which nevertheless describes the main underlying physics, the absorbance is found to be directly related to $\pi\alpha$, independent of P , largely independent of the physical quantities characterizing the semiconductor, as in graphene. The SOC also helps bring the absorbance steps closer to integer values of $\pi\alpha$, redistributing the oscillator strengths. Experimental results obtained on InAs layers, and on superlattices of other materials, support these conclusions. The situation for IV-VI semiconductors such as PbSe is extremely close to that of graphene in the presence of a band gap. We

believe that this work will provide a simplified view of optical absorption phenomena in 2D or quasi-2D materials for which excitonic effects are not too strong.

Appendix A: Alternative expression for the absorptance

1. General case

Using Eq. (3), Eq. (2) can be rewritten in the following form

$$\mathcal{A}_0(\hbar\omega) = \frac{\alpha}{\hbar\omega} \sum_{c,v} \int \frac{|\langle c, \mathbf{k} | \nabla H(\mathbf{k}) | v, \mathbf{k} \rangle|^2}{\delta(\hbar\omega - E_c(\mathbf{k}) + E_v(\mathbf{k})) d^2k} \quad (\text{A1})$$

where we are assuming that, in a cubic semiconductor, the absorptance does not depend on the in-plane orientation of the polarization vector, we can sum the contributions for the two in-plane polarizations and divide by 2 ($|\langle \nabla H(\mathbf{k}) \cdot \mathbf{x} \rangle|^2 + |\langle \nabla H(\mathbf{k}) \cdot \mathbf{y} \rangle|^2 = |\langle \nabla H(\mathbf{k}) \rangle|^2$).

For functions f and g defined over a n -dimensional space, we have:

$$\int_{(n)} f(\mathbf{x}) \delta(g(\mathbf{x})) d\mathbf{x} = \int_{(n-1)} \frac{f(\mathbf{x})}{|\nabla g|} d\sigma(\mathbf{x}) \quad (\text{A2})$$

where the second integral is on the $(n-1)$ -dimensional surface defined by $g(\mathbf{x}) = 0$. Equation A1 can be rewritten as

$$\mathcal{A}_0(\hbar\omega) = \frac{\alpha}{\hbar\omega} \sum_{c,v} \int_{\mathcal{L}_{vc}} \frac{|\langle c, \mathbf{k} | \nabla H(\mathbf{k}) | v, \mathbf{k} \rangle|^2}{|\nabla(E_v(\mathbf{k}) - E_c(\mathbf{k}))|} dl \quad (\text{A3})$$

where \mathcal{L}_{vc} is the path in \mathbf{k} -space defined by $\hbar\omega - E_c(\mathbf{k}) + E_v(\mathbf{k}) = 0$.

2. Isotropic case

In the case where all terms only depend on $k = |\mathbf{k}|$, Eq. A3 becomes

$$\mathcal{A}_0(\hbar\omega) = \frac{2\pi\alpha k}{\hbar\omega} \sum_{c,v} \frac{|\langle c, \mathbf{k} | \nabla H(\mathbf{k}) | v, \mathbf{k} \rangle|^2}{\partial|E_v(k) - E_c(k)|/\partial k} \quad (\text{A4})$$

with k such that $E_c(k) - E_v(k) = \hbar\omega$. This expression can be rewritten in another form given in Eq. (4).

The JDOS per unit area can be calculated in a similar way. For the pair of bands v and c , the JDOS is

$$J_{c,v}(\hbar\omega) = \frac{1}{S} \sum_{\mathbf{k}} \delta(\hbar\omega - E_c(\mathbf{k}) + E_v(\mathbf{k})) \quad (\text{A5})$$

which, in the isotropic case, transforms into

$$J_{c,v}(\hbar\omega) = \frac{1}{\pi} \frac{k}{\partial|E_c(k) - E_v(k)|/\partial k}. \quad (\text{A6})$$

The bare absorptance can be written as

$$\mathcal{A}_0(\hbar\omega) = \frac{2\pi^2\alpha}{\hbar\omega} \sum_{c,v} J_{c,v}(\hbar\omega) |\langle c, \mathbf{k} | \nabla H(\mathbf{k}) | v, \mathbf{k} \rangle|^2 \quad (\text{A7})$$

with k such that $E_c(k) - E_v(k) = \hbar\omega$. $|\langle c, \mathbf{k} | \nabla H(\mathbf{k}) | v, \mathbf{k} \rangle|^2 / (\hbar\omega)$ is related to the oscillator strength.

Appendix B: 2D $\mathbf{k} \cdot \mathbf{p}$ model deduced from tight-binding

Section S3 of the Supplemental Material [29] gives the rationale for describing, in quantum wells, the bands in the vicinity of the band gap with a 2D $\mathbf{k} \cdot \mathbf{p}$ model. Here we describe how we deduce a $\mathbf{k} \cdot \mathbf{p}$ Hamiltonian directly from tight binding calculations, without adjusting any parameters. We also present calculations of the absorptance in this model, to go beyond the approximations made in the analytic model of Sect. IV B.

1. Derivation of the 2D model

We consider a InAs quantum well of thickness 4.2 nm, chosen because it is characterized by two symmetric surfaces. The system is thus characterized by a perfect in-plane square symmetry, which simplifies the derivation of a 2D $\mathbf{k} \cdot \mathbf{p}$ model. Figure 13 shows that the band structure calculated without SOC is again characterized by many subbands, with complex dispersion in the valence band. However, the absorptance spectrum exhibits a simple behavior (Fig. 5). The analysis of the optical matrix elements shows that the first plateau corresponds to transitions from the two highest valence bands (heavy hole and light hole), degenerate at $k_{\parallel} = 0$, to the lowest conduction subband (Fig. S6 of the Supplemental Material [29]).

Next, we deduce a 3×3 $\mathbf{k} \cdot \mathbf{p}$ matrix using the calculation of the first-order and second-order derivatives of the tight-binding Hamiltonian matrix with respect to k_x and k_y , restricted to the lowest conduction subband and the two highest valence subbands, but renormalizing the $\mathbf{k} \cdot \mathbf{p}$ parameters to include the effects of the remote bands (see Sect. S5 of the Supplemental Material [29] for the methodology). We obtain

$$\tilde{H} = \begin{bmatrix} E_c + Ak^2 & iPk_x & iPk_y \\ -iPk_x & E_v + Lk_x^2 + Mk_y^2 & Nk_x k_y \\ -iPk_y & Nk_x k_y & E_v + Lk_y^2 + Mk_x^2 \end{bmatrix} \quad (\text{B1})$$

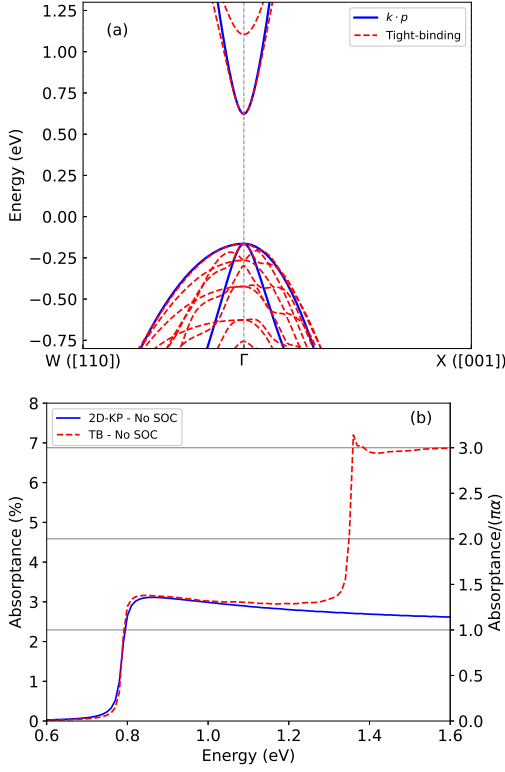


FIG. 13. Comparison between tight-binding (red dashed lines) and 2D $\mathbf{k} \cdot \mathbf{p}$ (blue solid lines) calculations for 4.2-nm-thick InAs quantum well. SOC is not included. (a) Band structure. (b) Bare absorbance $\mathcal{A}_0(\hbar\omega)$.

in the basis of the three vectors $|s\rangle$, $|x\rangle$ and $|y\rangle$. The parameters directly derived from the tight-binding Hamiltonian are given by (in atomic units): $A = 0.00242$, $P = 0.52684$, $E_c = 0.02288$, $E_v = -0.00604$, $L = 2.78493$, $M = -0.81699$, $N = 2.71777$. E_c and E_v differ from the bulk values, since they include the effect of the confinement along z ($E_v = 0$ in the bulk).

Figure 13 presents the comparison between tight-binding and 2D $\mathbf{k} \cdot \mathbf{p}$ band structures, and the comparison between absorbance spectra. In spite of its apparent simplicity, since it only describes two valence subbands of the quantum well, the 2D $\mathbf{k} \cdot \mathbf{p}$ model gives a perfect description of the first step of $\mathcal{A}_0(\hbar\omega)$.

2. Simplified 2D $\mathbf{k} \cdot \mathbf{p}$ isotropic model

In order to obtain analytical expressions for the absorbance, it is useful to derive a simpler 2D $\mathbf{k} \cdot \mathbf{p}$ isotropic model. We use the two vectors $|p_{\parallel}\rangle$ and $|p_{\perp}\rangle$ previously defined [Eq. (9)].

The Hamiltonian matrix element $\langle p_{\parallel} | H | p_{\perp} \rangle$ vanishes when $N = L - M$. In the bulk, this takes place when the Luttinger parameters γ_2^L and γ_3^L are equal, which corresponds to the spherical symmetry [64]. In this condition, the Hamiltonian written in $|s\rangle$, $|p_{\parallel}\rangle$ and $|p_{\perp}\rangle$ is simply

given by:

$$H = \begin{bmatrix} E_s(k) & iPk & 0 \\ -iPk & E_{\parallel}(k) & 0 \\ 0 & 0 & E_{\perp}(k) \end{bmatrix} \quad (\text{B2})$$

with

$$E_s(k) = E_c + Ak^2 \quad E_{\parallel}(k) = E_v + Lk^2 \quad E_{\perp}(k) = E_v + Mk^2 \quad (\text{B3})$$

The solutions are $|\Psi_{hh}\rangle = |p_{\perp}\rangle$, $|\Psi_e\rangle$, $|\Psi_{lh}\rangle$ of energy $E_{hh}(k)$ and

$$E_{lh}^e(k) = \left(\frac{E_s(k) + E_{\parallel}(k)}{2} \right) \pm \sqrt{\left(\frac{E_s(k) - E_{\parallel}(k)}{2} \right)^2 + P^2k^2}, \quad (\text{B4})$$

respectively. $E_e(k)$ is the energy of the conduction subband. $E_{hh}(k)$ and $E_{lh}(k)$ are the energies of the heavy and light hole subbands, respectively.

3. Calculation of the absorbance

The absorbance $\mathcal{A}_0(\hbar\omega) = \mathcal{A}_{hh-e}(\hbar\omega) + \mathcal{A}_{lh-e}(\hbar\omega)$ is calculated analytically using Eq. A4 of Appendix A. In this isotropic approximation, we use the previous parameters for the 2D $\mathbf{k} \cdot \mathbf{p}$ model except that we set $N = L - M$. After some algebra (Sect. S6 of the Supplemental Material [29]), the different terms can be calculated to the second order in k as

$$\mathcal{A}_{hh-e}(\hbar\omega) = \pi\alpha(C_0 + C_2k^2) + O(k^3) \quad (\text{B5})$$

$$C_0 = \frac{P^2}{(AE_g - E_gM + P^2)} \quad (\text{B6})$$

$$C_2 = -\frac{U}{E_g(A - M + U)^2} \quad (\text{B7})$$

$$\times (A^2 + 2AL - 4AM + AU - 2LM + 4LU + 3M^2 - 5MU)$$

with $U = P^2/E_g$ and k is solution of $E_e(k) - E_{hh}(k) = \hbar\omega$. Similarly, we obtain

$$\mathcal{A}_{lh-e}(\hbar\omega) = \pi\alpha(D_0 + D_2k^2) + O(k^3) \quad (\text{B8})$$

$$D_0 = \frac{P^2}{AE_g - E_gL + 2P^2} \quad (\text{B9})$$

$$D_2 = -\frac{U(5A^2 - 10AL + 12AU + 5L^2 - 12LU + 8U^2)}{E_g(A - L + 2U)^2} \quad (\text{B10})$$

in which k is solution of $E_e(k) - E_{lh}(k) = \hbar\omega$.

With the parameters given above for the 4.2-nm-thick InAs quantum well, $C_0 = 0.92$, which is close to 1 because

$P^2/E_g \gg A - M$. Numerical evaluations show that the variation with k and with $\hbar\omega$ is weak (Fig. S4 of the Supplemental Material [29]), i.e., $C_2k^2 \ll C_0$. It means that $\mathcal{A}_{hh-e}(\hbar\omega)$ is very close to $\pi\alpha$, but smaller.

With the same parameters, $D_0 = 0.58$ is close to $1/2$, because the term $2P^2$ is dominant in the denominator of Eq. B9. $\mathcal{A}_{lh-e}(\hbar\omega)$ is equal to $0.58\pi\alpha$ at the energy gap but it decreases at higher energy as D_2k^2 becomes non negligible compared to D_0 .

We conclude that the 2D $\mathbf{k} \cdot \mathbf{p}$ model and its simplified version under isotropic approximation explain very well the absorptance spectrum calculated in tight-binding in absence of SOC. This also completely justifies the simple analytical model of Sect. IV B in which $A, L, M, N = 0$.

An additional discussion of compensation mechanisms leading to absorptance quantization is presented in Sect. S6.4 of the Supplemental Material [29].

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Supplemental Material

Universality of the optical absorptance quantization in two-dimensional group-IV, III-V, II-VI and IV-VI semiconductors

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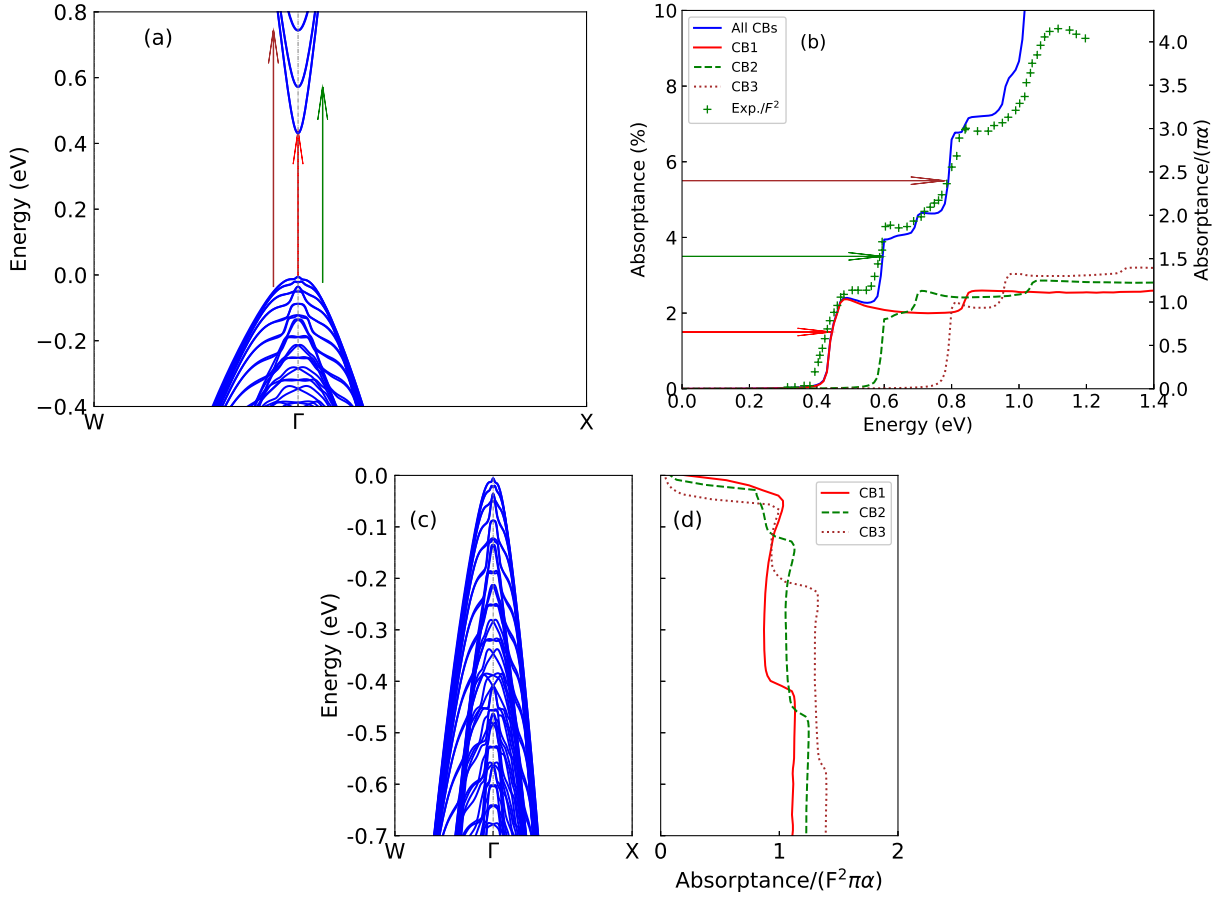


Figure S1: (a) Lowest conduction subbands and highest valence subbands of a 12.5 nm thick InAs quantum well. (b) Bare absorbance calculated including all conduction subbands (blue solid line), only the first conduction subband (red solid line, CB1), only the second one (green dashed line, CB2) or only the third one (brown dotted line, CB3). Green crosses: experimental data divided by F^2 for a quantum well thickness of 14 nm [1]. (c) Zoom on the highest valence subbands and, facing each other, (d) plot of the bare absorbance presented in such a way that each absorbance plateau can be associated to a transition from valence subbands to one of the conduction subbands, the first conduction subband (CB1, red solid line), the second one (CB2, green dashed line), and the third one (CB3, brown dotted line). $\eta = 6$ meV.

S1 Results for a 12.5-nm-thick quantum well of InAs

Figure S1 present additional results of tight-binding calculations for a thick InAs quantum well (12.5 nm) for which the absorbance was compared to experiments in Fig. 1 of the main document. The different steps at $\approx n\pi\alpha$ shown in Fig. S1b can be attributed to transitions $(\text{HH}+\text{LH}+\text{SO})_i \rightarrow \text{CB}_i$, where i denotes the quantum number related to the vertical confinement effect (HH = heavy-hole, LH = light hole, SO = split-off, CB = conduction). These results show that the absorbance contributions from the different sets of transitions (i) all behave approximately the same and therefore can be described by the same model.

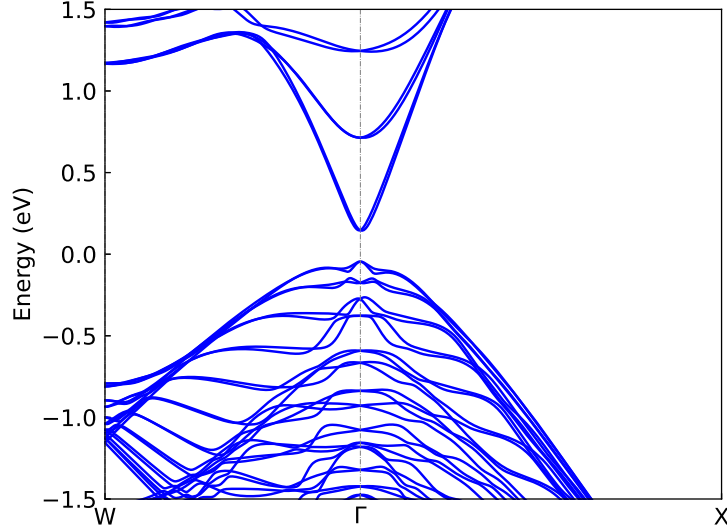


Figure S2: Lowest conduction subbands and highest valence subbands of the HgTe quantum well (thickness of 3.9 nm).

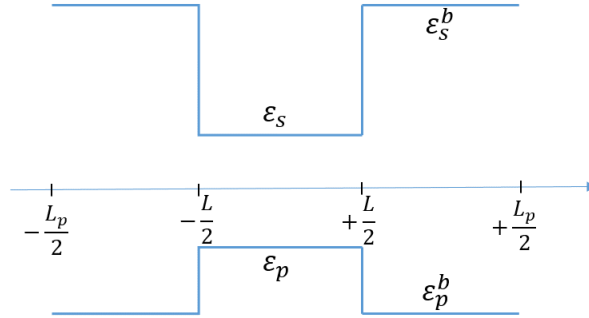


Figure S3: Band diagram for a superlattice along the z direction. We define $\Delta_s = \epsilon_s^b - \epsilon_s > 0$ and $\Delta_p = \epsilon_p^b - \epsilon_p < 0$.

S2 Band structure of a quantum well of HgTe

The band structure of a HgTe quantum well is presented in Fig. S2. The corresponding absorptance spectrum is shown in Fig. 3 of the main document.

S3 Justification of a 2D $\mathbf{k} \cdot \mathbf{p}$ model from a 3D one

In this section, we provide a theoretical basis for the use of a 2D $\mathbf{k} \cdot \mathbf{p}$ Kane model. We consider a periodic system (superlattice) of width L_p (Fig. S3) where the quantum well is positioned at the center. We employ the usual 4-band $\mathbf{k} \cdot \mathbf{p}$ hamiltonian without Spin-Orbit Coupling (SOC) [2]. For simplicity, we work in the spherical approximation ($L - M = N$) but extension to the more general case is straightforward. We expand the envelope function along z into plane waves $\exp(ik_{z,n}z)/\sqrt{L_p}$ where $k_{z,n} = 2n\pi/L_p$ due to the periodicity.

We thus write the hamiltonian into the basis set $u_l \exp(i\mathbf{k}_{\parallel} \cdot \mathbf{r}_{\parallel}) \exp(ik_{z,n}z)/\sqrt{L_p}$ where u_l are the periodic Bloch functions at $k = 0$ [$\mathbf{k}_{\parallel} = (k_x, k_y)$]. In this basis set, we assume the Kane matrix to be the same all along z except for the discontinuities in conduction $[\varepsilon_s(z)]$ and valence $[\varepsilon_p(z)]$ band edges (Fig. S3). For a given value of n , the 4×4 Kane matrix becomes

$$\tilde{H}_{n,n}^0 = \begin{bmatrix} \bar{\varepsilon}_s & Vk_x & Vk_y & Vk_{z,n} \\ V^\dagger k_x & \bar{\varepsilon}_p + Nk_x^2 + Mk_n^2 & Nk_x k_y & Nk_x k_{z,n} \\ V^\dagger k_y & Nk_x k_y & \bar{\varepsilon}_p + Nk_y^2 + Mk_n^2 & Nk_y k_{z,n} \\ V^\dagger k_{z,n} & Nk_x k_{z,n} & Nk_y k_{z,n} & \bar{\varepsilon}_p + Nk_{z,n}^2 + Mk_n^2 \end{bmatrix} \quad (\text{S1})$$

where $k_n^2 = k_x^2 + k_y^2 + k_{z,n}^2$. $\bar{\varepsilon}_s$ and $\bar{\varepsilon}_p$ are the average values of $\varepsilon_s(z)$ and $\varepsilon_p(z)$ over a period.

For a couple n, m , the only matrix element is due to the functions $\varepsilon_s(z)$ and $\varepsilon_p(z)$. We can write

$$(\varepsilon_s)_{n,m} = \frac{1}{L_p} \left\{ \int_{-\frac{L_p}{2}}^{\frac{L_p}{2}} \varepsilon_s^b e^{i(k_{z,m} - k_{z,n})z} dz + \int_{-\frac{L_p}{2}}^{\frac{L_p}{2}} (\varepsilon_s - \varepsilon_s^b) e^{i(k_{z,m} - k_{z,n})z} dz \right\} \quad (\text{S2})$$

$$= \bar{\varepsilon}_s \delta_{n,m} - (1 - \delta_{n,m}) \gamma_{s,n,m} \quad (\text{S3})$$

with

$$\gamma_{s,n,m} = \Delta_s \frac{\sin[\pi(n-m)L/L_p]}{\pi(n-m)} \quad (\text{S4})$$

This is also valid for p , i.e., $(\varepsilon_p)_{n,m} = \bar{\varepsilon}_p \delta_{n,m} - (1 - \delta_{n,m}) \gamma_{p,n,m}$. The 4×4 coupling matrix between states $n \neq m$ is thus

$$\tilde{H}_{n,m}^0 = \begin{bmatrix} \gamma_{s,n,m} & 0 & 0 & 0 \\ 0 & \gamma_{p,n,m} & 0 & 0 \\ 0 & 0 & \gamma_{p,n,m} & 0 \\ 0 & 0 & 0 & \gamma_{p,n,m} \end{bmatrix}. \quad (\text{S5})$$

In the spirit of self-energy calculations, we write the effective Hamiltonian in the n, n subspace as $\tilde{H}_{n,n}^0 + \delta\tilde{H}_{n,n}$ with

$$\delta\tilde{H}_{n,n} = \sum_{m \neq n} \tilde{H}_{n,m}^0 \tilde{g}_{m,m} \tilde{H}_{m,n}^0 \quad (\text{S6})$$

where $\tilde{g}_{m,m}$ is the Green's functions $(\varepsilon - \tilde{H}_{m,m}^0)^{-1}$.

S3.1 Eigenstates of $\tilde{H}_{m,m}^0$

Starting from $\tilde{H}_{m,m}^0$, this leads to the following equations

$$(\varepsilon - \bar{\varepsilon}_s)|s, m\rangle = V k_m |p_{\parallel}, m\rangle \quad (\text{S7})$$

$$(\varepsilon - \bar{\varepsilon}_p)|x, m\rangle = V^\dagger k_x |s, m\rangle + (Nk_x^2 + Mk_n^2)|x, m\rangle + Nk_x(k_y|y, m\rangle + k_{z,m}|z, m\rangle)$$

$$(\varepsilon - \bar{\varepsilon}_p)|y, m\rangle = V^\dagger k_y |s, m\rangle + (Nk_y^2 + Mk_n^2)|y, m\rangle + Nk_y(k_x|x, m\rangle + k_{z,m}|z, m\rangle)$$

$$(\varepsilon - \bar{\varepsilon}_p)|z, m\rangle = V^\dagger k_{z,m} |s, m\rangle + (Nk_{z,m}^2 + Mk_n^2)|z, m\rangle + Nk_{z,m}(k_y|y, m\rangle + k_x|x, m\rangle) \quad (\text{S8})$$

with

$$|p_{\parallel}, m\rangle = \frac{k_x|x, m\rangle + k_y|y, m\rangle + k_{z,m}|z, m\rangle}{k_m} \quad (\text{S9})$$

From this, we can deduce an equation for $|p_{\parallel}, m\rangle$, taking $L = M + N$,

$$(\varepsilon - \bar{\varepsilon}_p - Lk_m^2)|p_{\parallel}, m\rangle = V^\dagger k_m |p_{\parallel}, m\rangle \quad (\text{S10})$$

This, with Eq. (S7), leads to two dispersive bands, for electrons and light holes:

$$\varepsilon_m^\pm = \frac{\bar{\varepsilon}_s + \bar{\varepsilon}_p + Lk_m^2}{2} \pm \sqrt{\Delta_m^2 + V^2 k_m^2} \quad (\text{S11})$$

with $\Delta_m = (\bar{\varepsilon}_s - \bar{\varepsilon}_p - Lk_m^2)/2$. The eigenstates are

$$\begin{aligned} |+, m\rangle &= c_{s,m}^+ |s, m\rangle + c_{p_{\parallel},m}^+ |p_{\parallel}, m\rangle \\ |-, m\rangle &= c_{p_{\parallel},m}^{+*} |s, m\rangle - c_{s,m}^+ |p_{\parallel}, m\rangle \end{aligned} \quad (\text{S12})$$

with

$$\begin{aligned} c_{s,m}^+ &= \frac{D_m}{\sqrt{D_m^2 + 1}} \\ c_{p_{\parallel},m}^+ &= \frac{1}{\sqrt{D_m^2 + 1}} \frac{V^\dagger}{|V|} \\ D_m &= \frac{\Delta_m + \sqrt{\Delta_m^2 + V^2 k_m^2}}{|V| k_m} \end{aligned} \quad (\text{S13})$$

To this one can add the two pure p states, heavy holes, perpendicular to the previous two

$$\begin{aligned} |p_{\perp 1}, m\rangle &= \frac{k_y|x, m\rangle - k_x|y, m\rangle}{k_{\parallel}} \\ |p_{\perp 2}, m\rangle &= \frac{k_{z,m}}{k_{\parallel}} \left\{ \frac{k_x|x, m\rangle + k_y|y, m\rangle}{k_m} - \frac{k_{\parallel}^2}{k_m k_{z,m}} |z, m\rangle \right\} \end{aligned} \quad (\text{S14})$$

with energy

$$\varepsilon_{\perp 1,m} = \varepsilon_{\perp 2,m} = \varepsilon_{\perp,m} = \bar{\varepsilon}_p + M k_m^2 \quad (\text{S15})$$

S3.2 Calculation of the $\delta\tilde{H}_{n,n}$ matrix

From Eq. (S5) and Eq. (S6) we get

$$\delta\tilde{H}_{n,n} = \sum_{m \neq n} \begin{bmatrix} \gamma_{s,n,m}^2 \mathcal{G}_{sm,sm} & \gamma_{s,n,m} \gamma_{p,n,m} \mathcal{G}_{sm,xm} & \gamma_{s,n,m} \gamma_{p,n,m} \mathcal{G}_{sm,ym} & \gamma_{s,n,m} \gamma_{p,n,m} \mathcal{G}_{sm,zm} \\ \gamma_{s,n,m} \gamma_{p,n,m} \mathcal{G}_{xm,sm} & \gamma_{p,n,m}^2 \mathcal{G}_{xm,xm} & \gamma_{p,n,m}^2 \mathcal{G}_{xm,ym} & \gamma_{p,n,m}^2 \mathcal{G}_{xm,zm} \\ \gamma_{s,n,m} \gamma_{p,n,m} \mathcal{G}_{ym,sm} & \gamma_{p,n,m}^2 \mathcal{G}_{ym,xm} & \gamma_{p,n,m}^2 \mathcal{G}_{ym,ym} & \gamma_{p,n,m}^2 \mathcal{G}_{ym,zm} \\ \gamma_{s,n,m} \gamma_{p,n,m} \mathcal{G}_{zm,sm} & \gamma_{p,n,m}^2 \mathcal{G}_{zm,xm} & \gamma_{p,n,m}^2 \mathcal{G}_{zm,ym} & \gamma_{p,n,m}^2 \mathcal{G}_{zm,zm} \end{bmatrix} \quad (\text{S16})$$

in which the terms $g_{im,jm}$ are matrix elements of $\tilde{g}_{m,m}$.

The matrix $\delta\tilde{H}_{n,n}$ contains terms like k_x or $k_x k_y$ except for $g_{sm,sm}$. To show this in detail, let us examine the form of the $\tilde{g}_{m,m}$ matrix. We start from the general formula

$$g_{\alpha m, \beta m} = \sum_{\phi_m} \frac{\langle \alpha | \phi_m \rangle \langle \phi_m | \beta \rangle}{\epsilon - \epsilon_{\phi_m}} \quad (\text{S17})$$

where $|\phi_m\rangle$ are the eigenstates given by Eq. (S12) and Eq. (S14) of energies given by Eq. (S11) and Eq. (S15). We detail these for each different type of matrix element:

$$g_{sm,sm} = \frac{|c_{s,m}^+|^2}{\epsilon - \epsilon_m^+} + \frac{|c_{s,m}^-|^2}{\epsilon - \epsilon_m^-} \quad (\text{S18})$$

$$g_{sm,xm} = \left(\frac{c_{s,m}^+ c_{p\parallel,m}^{+*}}{\epsilon - \epsilon_m^+} + \frac{c_{s,m}^- c_{p\parallel,m}^{-*}}{\epsilon - \epsilon_m^-} \right) \frac{k_x}{k_m} = \eta_{sm,pm} \frac{k_x}{k_m} \quad \text{idem for } k_x \text{ and } k_y \quad (\text{S19})$$

$$g_{xm,ym} = \left(\frac{|c_{p\parallel,m}^+|^2}{\epsilon - \epsilon_m^+} + \frac{|c_{p\parallel,m}^-|^2}{\epsilon - \epsilon_m^-} - \frac{1}{\epsilon - \epsilon_{\perp,m}} \right) \frac{k_x k_y}{k_m^2} = \eta_{xm,ym} \frac{k_x k_y}{k_m^2} \quad (\text{S20})$$

$$g_{xm,xm} = \left(\frac{|c_{p\parallel,m}^+|^2}{\epsilon - \epsilon_m^+} + \frac{|c_{p\parallel,m}^-|^2}{\epsilon - \epsilon_m^-} \right) \frac{k_x^2}{k_m^2} + \frac{k_y^2 + k_{z,m}^2}{k_m^2 (\epsilon - \epsilon_{\perp,m})} = \eta_{xm,xm}^{(1)} \frac{k_x^2}{k_m^2} + \eta_{xm,xm}^{(2)} \frac{k_y^2 + k_{z,m}^2}{k_m^2} \quad (\text{S21})$$

idem for permutations of x, y, z with $k_z = k_{z,m}$.

Conclusion: This directly shows that $g_{\alpha m, \beta m}$ and thus $\delta\tilde{H}_{n,n}$ has the same form as the Kane matrix and will thus provide renormalized parameters.

S3.3 Application to the lowest transitions

We start from $\tilde{H}_{0,0}^0$ given by Eq. (S1) for which $k_{z,0} = 0$ which splits in a 3×3 2D matrix plus one heavy hole p state. The renormalized matrix to consider is thus $\tilde{H}_{0,0} = \tilde{H}_{0,0}^0 + \delta\tilde{H}_{0,0}$ in which $\delta\tilde{H}_{0,0}$ is given by the sum in Eq. (S16). In that summation, all terms linear in $k_{z,n}$ will vanish since $k_{z,-n} = -k_{z,-n}$. This corresponds to the lowest symmetrical state in the potential well. This gives

$$\tilde{H}_{n,n}^0 = \left[\begin{array}{cccc} \bar{\epsilon}_s & V k_x & V k_y & 0 \\ V^\dagger k_x & \bar{\epsilon}_p + L k_x^2 + M k_y^2 & N k_x k_y & 0 \\ V^\dagger k_y & N k_x k_y & \bar{\epsilon}_p + L k_y^2 + M k_x^2 & 0 \\ 0 & 0 & 0 & \bar{\epsilon}_p + M k_\parallel^2 \end{array} \right] + \sum_{m>0} \quad (\text{S22})$$

$$\left[\begin{array}{ccc} 2\gamma_{s,0,m}^2 g_{sm,sm} & 2\gamma_{s,0,m} \gamma_{p,0,m} \eta_{sm,pm} \frac{k_x}{k_m} & 2\gamma_{s,0,m} \gamma_{p,0,m} \eta_{sm,pm} \frac{k_y}{k_m} & 0 \\ 2\gamma_{s,0,m} \gamma_{p,0,m} \eta_{sm,pm}^* \frac{k_x}{k_m} & 2\gamma_{p,0,m}^2 \left(\eta_{xm,xm}^{(1)} \frac{k_x^2}{k_m^2} + \eta_{xm,xm}^{(2)} \frac{k_y^2 + k_{z,m}^2}{k_m^2} \right) & 2\gamma_{p,0,m}^2 \eta_{xm,ym} \frac{k_x k_y}{k_m^2} & 0 \\ 2\gamma_{s,0,m} \gamma_{p,0,m} \eta_{sm,pm}^* \frac{k_y}{k_m} & 2\gamma_{p,0,m}^2 \eta_{xm,ym}^* \frac{k_x k_y}{k_m^2} & 2\gamma_{p,0,m}^2 \left(\eta_{xm,xm}^{(1)} \frac{k_y^2}{k_m^2} + \eta_{xm,xm}^{(2)} \frac{k_x^2 + k_{z,m}^2}{k_m^2} \right) & 0 \\ 0 & 0 & 0 & 2\gamma_{p,0,m}^2 g_{zm,zm} \end{array} \right]$$

For optical transitions with polarization vector perpendicular to Oz only the 3×3 submatrix will be involved which justifies the use of a 2D 3×3 Kane matrix with renormalized parameters.

S4 Details on the simple analytical model without SOC

We consider the simplest 2D model described in Section IV.B of the main document. The Hamiltonian is

$$H = \begin{bmatrix} E_g & iPk_x & iPk_y \\ -iPk_x & 0 & 0 \\ -iPk_y & 0 & 0 \end{bmatrix} \quad (\text{S23})$$

where E_g is the gap of the quantum well.

S4.1 Calculation of the absorptance for the first transition $\Psi_{hh} \rightarrow \Psi_e$

Transition energy:

$$E_e(k) - E_{hh}(k) = \frac{E_g}{2} + \frac{\sqrt{E_g^2 + 4P^2k^2}}{2} \quad (\text{S24})$$

Optical matrix element:

$$|\langle \Psi_e | \nabla H(\mathbf{k}) | \Psi_{hh} \rangle|^2 = \frac{4P^4k^2}{4P^2k^2 + \left(E_g - \sqrt{E_g^2 + 4P^2k^2}\right)^2} = \frac{P^2\hbar\omega}{2\hbar\omega - E_g} \quad (\text{S25})$$

Derivative of energy:

$$\frac{1}{k} \frac{\partial(E_e(k) - E_{hh}(k))}{\partial k} = \frac{2P^2}{\sqrt{E_g^2 + 4P^2k^2}} \quad (\text{S26})$$

The JDOS including spin degeneracy is (inverse of previous equation / π):

$$J_{hh-e}(\hbar\omega) = \frac{2\hbar\omega - E_g}{4\pi P^2} \quad (\text{S27})$$

The denominator of Eq. (4) of the main document is given by

$$\frac{\partial |E_e(k) - E_{hh}(k)|^2}{\partial k^2} = \frac{P^2\hbar\omega}{\sqrt{(E_g/2)^2 + P^2k^2}} = \frac{2P^2\hbar\omega}{2\hbar\omega - E_g} \quad (\text{S28})$$

S4.2 Calculation of the absorptance for the second transition $\Psi_{lh} \rightarrow \Psi_e$

Transition energy:

$$\hbar\omega = E_e(k) - E_{lh}(k) = 2\sqrt{\frac{E_g^2}{4} + P^2k^2} \quad (\text{S29})$$

Optical matrix element:

$$|\langle \Psi_e | \nabla H | \Psi_{lh} \rangle|^2 = \frac{E_g^2 P^2}{E_g^2 + 4P^2k^2} = \left(\frac{E_g P}{\hbar\omega}\right)^2 \quad (\text{S30})$$

Derivative of energy:

$$\frac{1}{k} \frac{\partial(E_e(k) - E_{lh}(k))}{\partial k} = \frac{4P^2}{\sqrt{E_g^2 + 4P^2k^2}} \quad (\text{S31})$$

The JDOS including spin degeneracy is (inverse of previous equation / π):

$$J_{lh-e}(\hbar\omega) = \frac{\hbar\omega}{4\pi P^2} \quad (\text{S32})$$

S5 Derivation of a 2D $\mathbf{k} \cdot \mathbf{p}$ model from tight-binding calculations

In this section, we present the method used to derive the $\mathbf{k} \cdot \mathbf{p}$ matrix directly from tight-binding calculations for a quantum well. First, the tight-binding Hamiltonian H for the quantum well is written in the basis of Bloch functions as function of k_x and k_y . Second, H is diagonalized at $\mathbf{k} = 0$. The eigenvectors form a complete basis of the 2D system. Third, we write H in this basis up to the second order in \mathbf{k} ,

$$\begin{aligned} H &= H^{(0)} + H^{(1)} + H^{(2)} \\ H^{(1)} &= \sum_{i=x,y} \left(\frac{dH}{dk_i} \right) k_i \\ H^{(2)} &= \frac{1}{2} \sum_{i,j=x,y} \left(\frac{d^2H}{dk_i dk_j} \right) k_i k_j \end{aligned} \quad (\text{S33})$$

in which the derivatives are calculated numerically at $\mathbf{k} = 0$.

Fourth, we define the subset of the basis in which the $\mathbf{k} \cdot \mathbf{p}$ model will be defined (in our case, the lowest conduction state and the two highest valence states). We represent the states of the subspace by upper case letters, e.g., I of energy E_I , and the other states of the basis by lower case ones, e.g., n of energy E_n .

Fifth, we use the Luttinger-Kohn theory of perturbation [3]. If we write H as $\mathcal{H} + V$ where \mathcal{H} contains just the terms restricted to the subspace and V the others, the renormalized Hamiltonian \tilde{H} in the subspace is given to the second order in V by

$$\tilde{H}_{IJ} = \mathcal{H}_{IJ} + \frac{1}{2} \sum_n V_{In} V_{nJ} \left(\frac{1}{E_{In}} + \frac{1}{E_{Jn}} \right) \quad (\text{S34})$$

If we use for H its expression given in Eq. (S33), the $\mathbf{k} \cdot \mathbf{p}$ Hamiltonian restricted to second order terms in \mathbf{k} is given by

$$\tilde{H}_{IJ} = H_{IJ}^{(0)} + H_{IJ}^{(1)} + H_{IJ}^{(2)} + \frac{1}{2} \sum_n H_{In}^{(1)} H_{nJ}^{(1)} \left(\frac{1}{E_{In}} + \frac{1}{E_{Jn}} \right) \quad (\text{S35})$$

Therefore, the computation of the matrices of the first and second derivatives of H allows to completely define the $\mathbf{k} \cdot \mathbf{p}$ Hamiltonian, without adjustable parameters. The last term of Eq. (S35) describes the effect of remote bands that lead to a renormalization of the $\mathbf{k} \cdot \mathbf{p}$ parameters.

S6 2D $\mathbf{k} \cdot \mathbf{p}$ model derived from tight binding

In this section, we provide details on the calculations presented in Appendix B of the main document.

S6.1 Hamiltonian and eigenstates

We give details on 2D $\mathbf{k} \cdot \mathbf{p}$ model and its transformation in the isotropic approximation. We have introduced two vectors:

$$|p_{\parallel}\rangle = \frac{k_x|x\rangle + k_y|y\rangle}{k} \quad (\text{S36})$$

$$|p_{\perp}\rangle = \frac{k_y|x\rangle - k_x|y\rangle}{k} \quad (\text{S37})$$

$$(\text{S38})$$

The matrix element (see definition of \tilde{H} in Eq. (B1) of the main document)

$$\langle p_{\parallel} | \tilde{H} | p_{\perp} \rangle = \frac{k_x k_y}{k^2} (k_x^2 - k_y^2) (L - M - N) \quad (\text{S39})$$

vanishes when $N = L - M$. In this condition corresponding to the isotropic approximation, the Hamiltonian written in $|s\rangle$, $|p_{\parallel}\rangle$ and $|p_{\perp}\rangle$ is simply given by:

$$H = \begin{bmatrix} E_s(k) & iPk & 0 \\ -iPk & E_{\parallel}(k) & 0 \\ 0 & 0 & E_{\perp}(k) \end{bmatrix} \quad (\text{S40})$$

with

$$E_s(k) = E_c + Ak^2 \quad E_{\parallel}(k) = E_v + Lk^2 \quad E_{\perp}(k) = E_v + Mk^2 \quad (\text{S41})$$

The solutions are

$$|\Psi_e\rangle = \gamma|s\rangle - i\beta|p_{\parallel}\rangle \quad (\text{S42})$$

$$|\Psi_{lh}\rangle = -i\beta|s\rangle + \gamma|p_{\parallel}\rangle \quad (\text{S43})$$

$$|\Psi_{hh}\rangle = |p_{\perp}\rangle \quad (\text{S44})$$

with

$$\gamma = \frac{Pk}{\sqrt{(E_s(k) - E_e(k))^2 + P^2k^2}} \quad (\text{S45})$$

$$\beta = \frac{E_e(k) - E_s(k)}{\sqrt{(E_s(k) - E_e(k))^2 + P^2k^2}} \quad (\text{S46})$$

and

$$E_{lh}^e(k) = \left(\frac{E_s(k) + E_{\parallel}(k)}{2} \right) \pm \sqrt{\left(\frac{E_s(k) - E_{\parallel}(k)}{2} \right)^2 + P^2k^2} \quad (\text{S47})$$

S6.2 Effective masses

From the previous equations, we can derive the effective masses for electron (m_e), heavy hole (m_{hh}) and light hole (m_{lh}) bands. We have:

$$\begin{aligned}\frac{\hbar^2}{2m_e} &= A + U \\ \frac{\hbar^2}{2m_{lh}} &= -L + U \\ \frac{\hbar^2}{2m_{hh}} &= -M\end{aligned}\tag{S48}$$

with $U = P^2/E_g$.

S6.3 Calculation of the absorptance

The absorption comes from transitions $|\Psi_{lh}\rangle \rightarrow |\Psi_e\rangle$ and $|\Psi_{hh}\rangle \rightarrow |\Psi_e\rangle$. The absorptance is calculated analytically using Eq. (A4) of the Appendix A of the main document.

S6.3.1 Transitions $|\Psi_{hh}\rangle \rightarrow |\Psi_e\rangle$

We consider k for which $E_e(k) - E_{hh}(k) = \hbar\omega$ and we calculate the different contributions to the expression of the absorptance. After some algebra, we obtain:

$$\begin{aligned}|\langle\Psi_e|\nabla H(\mathbf{k})|\Psi_{hh}\rangle|^2 &= (\gamma P + \beta N k)^2 \\ &= \frac{k^2 \left(2P^2 + (L - M) \left(Ak^2 + E_g - Lk^2 - \sqrt{4P^2k^2 + (Ak^2 + E_g - Lk^2)^2} \right) \right)^2}{4P^2k^2 + \left(Ak^2 + E_g - Lk^2 - \sqrt{4P^2k^2 + (Ak^2 + E_g - Lk^2)^2} \right)^2}\end{aligned}\tag{S49}$$

$$\frac{1}{k} \frac{\partial(E_e(k) - E_{hh}(k))}{\partial k} = A + L - 2M + \frac{(A - L) \left(\frac{E_s(k) - E_{||}(k)}{2} \right) + P^2k^2}{\sqrt{\left(\frac{E_s(k) - E_{||}(k)}{2} \right)^2 + P^2k^2}}\tag{S50}$$

$$\mathcal{A}_{hh-e}(\hbar\omega) = \frac{2\pi\alpha k}{E_e(k) - E_{hh}(k)} \frac{|\langle\Psi_e|\nabla H(\mathbf{k})|\Psi_{hh}\rangle|^2}{|\partial(E_e(k) - E_{hh}(k))/\partial k|}\tag{S51}$$

For simplification, it is interesting to perform a second order expansion of the different terms with respect to k :

$$|\langle\Psi_e|\nabla H(\mathbf{k})|\Psi_{hh}\rangle|^2 = -\frac{P^4k^2}{E_g^2} + \frac{2MP^2k^2}{E_g} - \frac{2LP^2k^2}{E_g} + P^2 + O(k^3)\tag{S52}$$

$$\frac{1}{k} \frac{\partial(E_e(k) - E_{hh}(k))}{\partial k} = -\frac{4P^4k^2}{E_g^3} + \frac{4LP^2k^2}{E_g^2} + \frac{2P^2}{E_g} - 2M - \frac{4AP^2k^2}{E_g^2} + 2A + O(k^3)\tag{S53}$$

$$E_e(k) - E_{hh}(k) = \frac{P^2 k^2}{E_g} - Mk^2 + E_g + Ak^2 + O(k^3) \quad (\text{S54})$$

We deduce the absorbance to the second order in k :

$$\mathcal{A}_{hh-e}(\hbar\omega) = \pi\alpha(C_0 + C_2 k^2) + O(k^3) \quad (\text{S55})$$

$$C_0 = \frac{U}{(A - M + U)} \quad (\text{S56})$$

$$C_2 = -\frac{U(A^2 + 2AL - 4AM + AU - 2LM + 4LU + 3M^2 - 5MU)}{E_g(A - M + U)^2} \quad (\text{S57})$$

S6.3.2 Transitions $|\Psi_{lh}\rangle \rightarrow |\Psi_e\rangle$

The contribution of $|\Psi_{lh}\rangle \rightarrow |\Psi_e\rangle$ transitions to the absorbance can be calculated similarly. Using the different contributions

$$\hbar\omega = E_e(k) - E_{lh}(k) = 2\sqrt{P^2 k^2 + \frac{(Ak^2 + E_g - Lk^2)^2}{4}} \quad (\text{S58})$$

$$|\langle\Psi_e|\nabla H|\Psi_{lh}\rangle|^2 \quad (\text{S59})$$

$$\begin{aligned} &= (P(2\gamma^2 - 1) - 2\gamma\beta k[A - L])^2 \\ &= \frac{P^2(A^2 k^4 - 2AE_g k^2 - 2ALK^4 + E_g^2 + 2E_g Lk^2 + L^2 k^4)}{A^2 k^4 + 2AE_g k^2 - 2ALK^4 + E_g^2 - 2E_g Lk^2 + L^2 k^4 + 4P^2 k^2} \\ &= \frac{E_g^2 P^2 + 4E_g P^2 k^2(-A + L) - 4P^4 k^2}{E_g^2} + O(k^3) \end{aligned}$$

$$\frac{1}{k} \frac{\partial(E_e(k) - E_{lh}(k))}{\partial k} = \frac{2(2P^2 + (A - L)(Ak^2 + E_g - Lk^2))}{\sqrt{4P^2 k^2 + (Ak^2 + E_g - Lk^2)^2}} \quad (\text{S60})$$

we obtain $\mathcal{A}_{lh-e}(\hbar\omega)$ to the second order in k :

$$\mathcal{A}_{lh-e}(\hbar\omega) = \pi\alpha \times (D_0 + D_2 k^2) \quad (\text{S61})$$

$$D_0 = \frac{U}{A - L + 2U} \quad (\text{S62})$$

$$D_2 = -\frac{U(5A^2 - 10AL + 12AU + 5L^2 - 12LU + 8U^2)}{E_g(A - L + 2U)^2} \quad (\text{S63})$$

S6.3.3 Comparison with tight-binding calculations

Figure S4 shows that the total absorbance $\mathcal{A}_0(\hbar\omega)$ calculated under the isotropic approximation is in excellent agreement with $\mathcal{A}_0(\hbar\omega)$ calculated numerically using the 2D $\mathbf{k} \cdot \mathbf{p}$ model originally deduced from the tight-binding Hamiltonian, which itself is in excellent agreement with tight-binding calculations. It demonstrates the validity of the isotropic approximation.

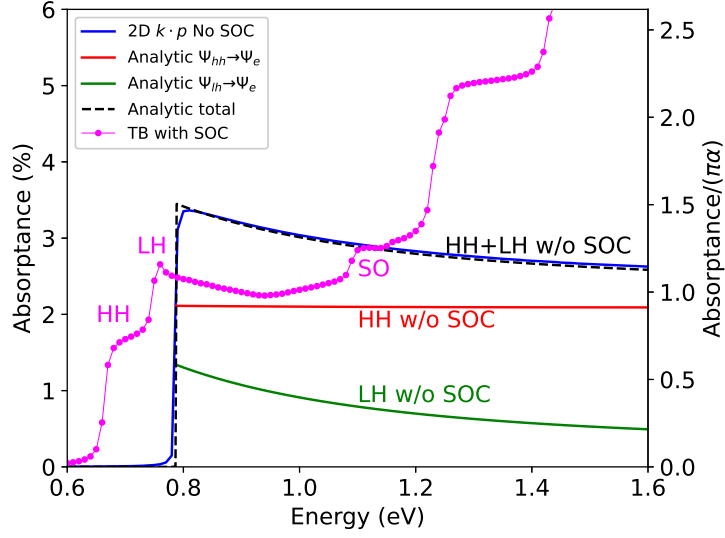


Figure S4: Comparison between different calculations of the absorptance spectrum for a 4.2-nm-thick InAs quantum well. Blue solid line: 2D $\mathbf{k} \cdot \mathbf{p}$ model (no SOC) directly deduced from the tight-binding Hamiltonian, calculated using a fully numerical approach ($\eta = 1$ meV). Black dashed line: Same but with the analytical 2D $\mathbf{k} \cdot \mathbf{p}$ model under isotropic approximation. The contributions coming from $|\Psi_{hh}\rangle \rightarrow |\Psi_e\rangle$ (HH \rightarrow CB) and $|\Psi_{lh}\rangle \rightarrow |\Psi_e\rangle$ (LH \rightarrow CB) transitions are shown with red and green solid lines, respectively. Magenta dotted solid line: Tight-binding calculations with SOC.

S6.4 Importance of the term P^2/E_g on the $\pi\alpha$ quantization

Equations (S56) and (S62) show that $C_0 \approx 1$ and $D_0 \approx 1/2$ when $P^2/E_g \gg A, |L|, |M|$, which leads to the $\pi\alpha$ quantization of the absorptance after inclusion of the spin-orbit coupling. However, Eq. (S48) shows that we could have the same effective masses with a much lower value of $U = P^2/E_g$, compensating via A and L . In that case, the effective masses would be mainly determined by the couplings with the bands outside the subspace. Interestingly, C_0 and D_0 can be rewritten as

$$C_0 = \frac{U}{\frac{\hbar^2}{2} \left(\frac{1}{m_e} + \frac{1}{m_{hh}} \right)} \quad (\text{S64})$$

$$D_0 = \frac{U}{\frac{\hbar^2}{2} \left(\frac{1}{m_e} + \frac{1}{m_{lh}} \right)} \quad (\text{S65})$$

A compensation between numerator and denominator is done when $U \gg A, |L|, |M|$, i.e., when $\hbar^2/(2m_e)$ and $\hbar^2/(2m_{lh})$ are approximately equal to U [Eq. (S48)]. But, if we decrease U while keeping the effective masses constant, C_0 and D_0 become very small compared to 1 and 1/2, the absorptance is then no longer quantized in units of $\pi\alpha$, but can take any (smaller) value depending on U .

Therefore, we deduce that the compensation in $\mathcal{A}_0(\hbar\omega)$ between the joint density of states and the optical matrix elements comes mainly from the combination of three effects. First, the dispersion of the electron and light-hole bands is mainly governed by couplings of type Pk between them. Second, P also determines the optical matrix elements between valence and conduction bands. Third, the valence band whose dispersion

does not depend on the Pk coupling is characterized by a heavy effective mass ($M \ll U, m_e \ll m_{hh}$).

Since U is equal to P^2/E_g , and P is known to be rather constant in different types of III-V and II-VI semiconductors, we expect that the absorptance quantization will be less accurate in semiconductors with a large gap E_g . This is what we found with tight-binding calculations (see main document).

S7 Analytical model with SOC

S7.1 Details on the analytical calculations

As discussed in Sect. IV.C of the main document, the Kane Hamiltonian is written in the basis of the eigenstates of \mathbf{J}^2 and J_Z where Z is defined along the \mathbf{k} vector, i.e., $|Z \uparrow\rangle \propto k_x|x \uparrow\rangle + k_y|y \uparrow\rangle + k_z|z \uparrow\rangle$ (idem for \downarrow).

In the case of a quantum well, we reduce it to a 2D problem with $k_z = 0$ ($k = k_{\parallel}$), i.e.,

$$|Z \uparrow\rangle = \frac{k_x|x \uparrow\rangle + k_y|y \uparrow\rangle}{k_{\parallel}}. \quad (\text{S66})$$

We define the two orthogonal states

$$|X \uparrow\rangle = \frac{k_y|x \uparrow\rangle - k_x|y \uparrow\rangle}{k_{\parallel}}, \quad (\text{S67})$$

$$|Y \uparrow\rangle = |z \uparrow\rangle, \quad (\text{S68})$$

with similar expressions for spin down states.

The eigenstates of \mathbf{J}^2 and J_Z are given by [4]

$$|\frac{3}{2}, \frac{1}{2}\rangle = -\sqrt{\frac{2}{3}}|Z \uparrow\rangle + \frac{1}{\sqrt{6}}|X + iY, \downarrow\rangle \quad (\text{S69})$$

$$|\frac{1}{2}, \frac{1}{2}\rangle = \frac{1}{\sqrt{3}}|Z \uparrow\rangle + \frac{1}{\sqrt{3}}|X + iY, \downarrow\rangle \quad (\text{S70})$$

$$|\frac{3}{2}, -\frac{1}{2}\rangle = -\sqrt{\frac{2}{3}}|Z \downarrow\rangle - \frac{1}{\sqrt{6}}|X - iY, \uparrow\rangle \quad (\text{S71})$$

$$|\frac{1}{2}, -\frac{1}{2}\rangle = \frac{1}{\sqrt{3}}|Z \downarrow\rangle - \frac{1}{\sqrt{3}}|X - iY, \uparrow\rangle \quad (\text{S72})$$

$$|\frac{3}{2}, \frac{3}{2}\rangle = \frac{1}{\sqrt{2}}|X + iY, \uparrow\rangle \quad (\text{S73})$$

$$|\frac{3}{2}, -\frac{3}{2}\rangle = \frac{1}{\sqrt{2}}|X - iY, \downarrow\rangle. \quad (\text{S74})$$

The Hamiltonian is given by two blocks like Eq. (19) of the main document.

By treating Pk within second order perturbation theory, the energy dispersion with respect to $k = 0$ is given by

$$\varepsilon_e(k) = \varepsilon_g + \frac{2}{3} \frac{P^2 k^2}{\varepsilon_g} + \frac{1}{3} \frac{P^2 k^2}{\varepsilon_g + \Delta} \quad (\text{S75})$$

$$\varepsilon_{hh}(k) = 0 \quad (\text{S76})$$

$$\varepsilon_{lh}(k) = -\frac{2}{3} \frac{P^2 k^2}{\varepsilon_g} \quad (\text{S77})$$

$$\varepsilon_{so}(k) = -\Delta - \frac{1}{3} \frac{P^2 k^2}{\varepsilon_g + \Delta}. \quad (\text{S78})$$

This gives at $k = 0$

$$\frac{\partial(\varepsilon_e(k) - \varepsilon_{lh}(k))^2}{\partial k^2} = \frac{8P^2}{3} \left(1 + \frac{\varepsilon_g}{4(\varepsilon_g + \Delta)} \right), \quad (\text{S79})$$

$$\frac{\partial(\varepsilon_e(k) - \varepsilon_{so}(k))^2}{\partial k^2} = \frac{4P^2}{3} \left(1 + \frac{\varepsilon_g + \Delta}{\varepsilon_g} \right), \quad (\text{S80})$$

$$\frac{\partial(\varepsilon_e(k) - \varepsilon_{hh}(k))^2}{\partial k^2} = \frac{4P^2}{3} \left(1 + \frac{\varepsilon_g}{2(\varepsilon_g + \Delta)} \right). \quad (\text{S81})$$

The optical matrix elements at $k = 0$ are:

$$\left| \langle s \uparrow | \nabla H(\mathbf{k}) | \frac{3}{2}, \frac{1}{2} \rangle \right|^2 = \frac{2P^2}{3}, \quad (\text{S82})$$

$$\left| \langle s \uparrow | \nabla H(\mathbf{k}) | \frac{1}{2}, \frac{1}{2} \rangle \right|^2 = \frac{P^2}{3}, \quad (\text{S83})$$

$$\left| \langle s \uparrow | \nabla H(\mathbf{k}) | \frac{3}{2}, \frac{3}{2} \rangle \right|^2 = \frac{P^2}{2}. \quad (\text{S84})$$

From this, we could calculate the absorptance using Eq. (4) of the main document.

S7.2 Step heights versus semiconductor characteristics

Figure S5 presents the step height calculated using Eq. (20) [$\mathcal{A}_0(\varepsilon_g)$] and Eq. (21) [$\delta\mathcal{A}_0(\varepsilon_g + \Delta)$] of the main document, versus $x = \varepsilon_g/\Delta$.

The development in series with respect to $x = 1$ of Eq. (20) of the main document gives

$$\pi\alpha \left\{ 1.225 - 0.341x + 0.241x^2 - 0.098x^3 + 0.017x^4 + O\left((x-1)^5; x \rightarrow 1\right) \right\}. \quad (\text{S85})$$

The term in braces is close to unity. $\mathcal{A}_0(\varepsilon_g)$ remains close to $\pi\alpha$, within $\pm 10\%$ for x varying from 0.5 to 5.

S7.3 Effect of the SOC from tight binding calculations

Figure S6 presents the variation of the valence subband energies and optical matrix elements at $k = 0$ when we vary the strength of the SOC. First, the SOC splits the two highest degenerate energy subbands. Second,

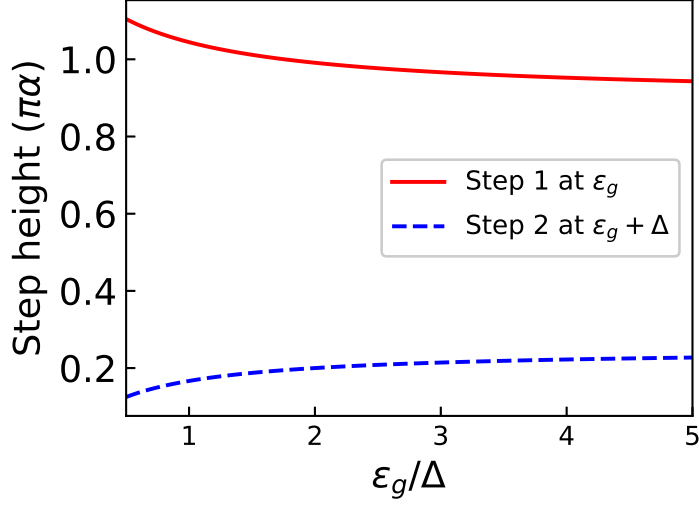


Figure S5: Height of the steps in unit of $\pi\alpha$ calculated using the analytical model including SOC versus ϵ_g/Δ (b). Red solid line: first step, $\mathcal{A}_0(\epsilon_g)$. Blue dashed line: second step, $\delta\mathcal{A}_0(\epsilon_g + \Delta)$.

it couples the lowest of these two subbands to a third subband located at lower energy (~ -0.3 eV), which leads to a redistribution of the optical oscillator strengths, their sum remaining approximately constant. At increasing SOC, this leads to the conventional picture of three valence subbands derived from heavy-hole, light-hole, and split-off bands. [4]

S8 Absorptance spectra in tight-binding models on a honeycomb lattice

S8.1 Model Hamiltonians

As already demonstrated in Ref. [5], the case of Hamiltonians on the honeycomb lattice is particularly interesting since it allows to study the effect on the absorptance spectrum of the transition from massless particles to massive ones. In this appendix, we illustrate different physical effects that induce the opening of a band gap at the Dirac point of a graphene band structure. These effects are, the asymmetry between sublattices A and B of the honeycomb, the intrinsic Spin-Orbit Coupling (SOC), and the Rashba SOC. We start from the tight-binding model that describes the π bands of graphene [6], with one orbital on each lattice site (two including spin). The Hamiltonian is written as $H = H_0 + H_{\text{NN}} + H_{\text{ISO}} + H_{\text{RSO}}$. Here, H_0 incorporates the on-site energies E_A and E_B on sublattice A and B, respectively. The term

$$H_{\text{NN}} = t \sum_{\langle i,j \rangle} \sum_{\alpha} c_{i,\alpha}^{\dagger} c_{j,\alpha}, \quad (\text{S86})$$

includes hopping terms, where $\langle i, j \rangle$ denotes nearest-neighbor (NN) sites, α denotes spin, and $t = -2.8$ eV is the hopping parameter typically used for graphene [7].

The intrinsic SOC term H_{ISO} couples the electron orbital momentum \mathbf{L} and spin \mathbf{S} . There is no on-site term because the orbital momentum is frozen. However, as shown by Kane and Mele [8], the intrinsic SOC can introduce a next-nearest-neighbour (NNN) hopping term, which is written as

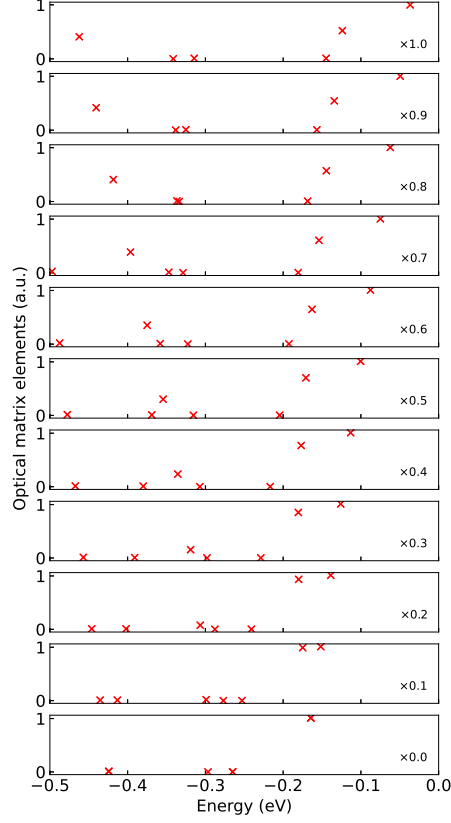


Figure S6: Evolution of the energy levels in a 4.2-nm-thick InAs layer, at $k = 0$ in the valence band when the SOC is scaled by a factor varying from 0.0 to 1.0. The vertical axis indicates the optical matrix element $|\langle c, \mathbf{0} | \nabla H(\mathbf{k}) | v, \mathbf{0} \rangle|^2$ for the transition to the lowest conduction subband (same scale for all figures).

$$H_{\text{ISO}} = i\lambda \sum_{\langle\langle i,j \rangle\rangle} \sum_{\alpha} c_{i,\alpha}^{\dagger} S_{\alpha\alpha}^z v_{ij} c_{j,\alpha}. \quad (\text{S87})$$

Here, the summation is over NNNs, and $v_{ij} = \pm 1$, with the sign depending on the outer product of the two NN vectors that connect sites i and j . The Rashba SOC term is written as a NN-hopping term

$$H_{\text{RSO}} = i\gamma \sum_{\langle i,j \rangle} \sum_{\alpha,\beta} c_{i,\alpha}^{\dagger} [\hat{z} \cdot (\mathbf{S} \times \mathbf{r}_{ij})]_{\alpha\beta} c_{j,\beta} \quad (\text{S88})$$

λ and γ are coupling constants that we adjust to open a visible gap. In any case, we set the Fermi level at the zero of energy, i.e., at the Dirac point of graphene.

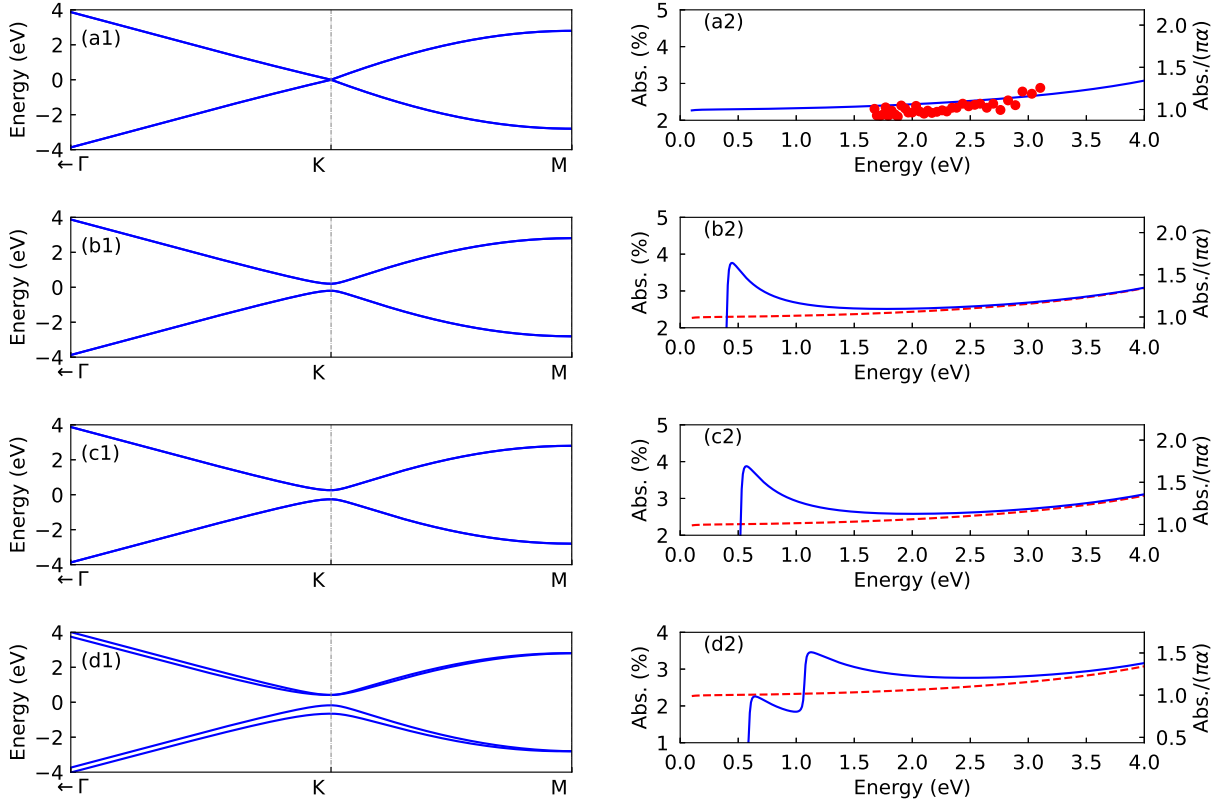


Figure S7: Band structures (x1) and absorbance spectra (x2) calculated in the following configurations. (a1,a2) Model of graphene, no lattice asymmetry ($E_A = E_B = 0$), and no SOC. The red points represent the experimental data of Ref. [9]. (b1,b2) Same but with a sublattice asymmetry, $E_A = 0.2$ eV and $E_B = -0.2$ eV. (c1,c2) No asymmetry ($E_A = E_B = 0$) but intrinsic SOC ($\lambda = 0.1$ eV). (d1,d2) No asymmetry but intrinsic SOC ($\lambda = 0.16$ eV) and Rashba SOC ($\gamma = 0.16$ eV). The red dashed lines in (b2,c2,d2) reproduce the absorbance spectrum (a2) of graphene, for comparison.

S8.2 Results

The results of Fig. S7(a1,a2) concerning graphene have already been the subject of much work[10, 11, 12, 9, 13, 14, 15]. The absorbance is equal to $\pi\alpha$ over a wide energy range, as long as the bands involved in the optical transitions keep a linear dispersion. The other figures show that the opening of a band gap, whether due to the asymmetry of the honeycomb sublattices [Fig. S7(b1,b2)] or to the intrinsic SOC [Fig. S7(c1,c2)], induces just above the optical threshold the appearance of a peak reflecting an absorbance considerably greater than $\pi\alpha$. This is attributed to the non-zero mass that appears at the edges of the bands, in the vicinity of the band gap [5]. When both charge carriers transform from massless to massive quasiparticles, the absorbance per valley is evolving as $\pi\alpha/2 \rightarrow \pi\alpha$. This transformation, predicted in the case of 2D group-IV honeycomb crystals [16], is partial in the case where the massive character of the bands disappears very quickly as soon as we deviate in energy from the band edges. The same effect occurs in the case of PbSe quantum wells (Sect. VI of the main document).

When the spin degeneracy is lifted under the effect of Rashba SOC coupling, the peak reflecting the massive character splits into two peaks [Fig. S7(d1,d2)]. In any case, by moving towards the highest energies,

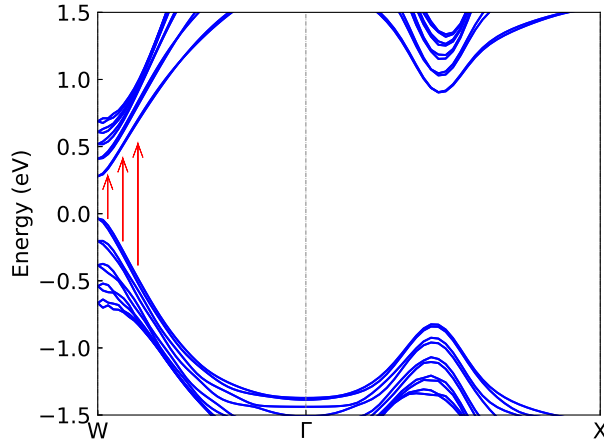


Figure S8: Lowest conduction subbands and highest valence subbands in a 6 nm quantum well of PbSe. The steps in the absorptance (Fig. 12 of the main document) are associated with the transitions indicated by red arrows.

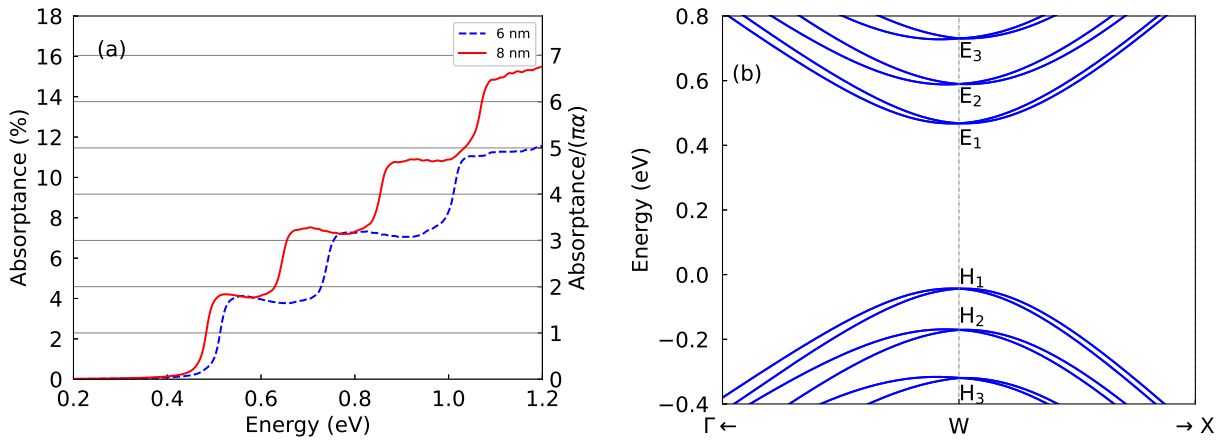


Figure S9: (a) Bare absorptance spectra calculated for (001) quantum wells of PbS (red solid line: thickness = 8 nm; blue dashed line: 6 nm). (b) Lowest conduction subbands and highest valence subbands in a 8 nm quantum well of PbS. The steps in the absorptance are associated with the transitions $H_i \rightarrow E_i$.

the absorptance reaches the graphene reference value.

S9 Additional results for quantum wells of rocksalt IV-VI semiconductors

S9.1 Band structure of the PbSe quantum well

The band structure of the quantum well of PbSe discussed in the main document is presented in Fig. S8.

S9.2 Absorptance spectrum of a PbS quantum well

The calculated band structure and absorptance spectra of PbS quantum wells are presented in Fig. S9. The first absorptance plateau is close to $2\pi\alpha$, like in PbSe (main document). The height of the next plateaus is smaller than $2\pi\alpha$, which could be due to the strongly anisotropic dispersion of higher subbands (Fig. S9b).

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