Excitons in van der Waals materials: from monolayer to bulk hexagonal boron nitride

Jaakko Koskelo,1 Giorgia Fugallo,2,3 Mikko Hakala,1 Matteo Gatti,2,3,4 Francesco Sottile,2,3 and Pierluigi Cudazzo2,3

1Department of Physics, P.O. Box 64, FI-00014 University of Helsinki, Helsinki, Finland
2Laboratoire des Solides Irradiés, École Polytechnique, CNRS, CEA, Université Paris-Saclay, F-91128 Palaiseau, France
3European Theoretical Spectroscopy Facility (ETSF)
4Synchrotron SOLEIL, L’Orme des Merisiers, Saint-Aubin, BP 48, F-91192 Gif-sur-Yvette, France

(Dated: April 5, 2018)

We present a general picture of the exciton properties of layered materials in terms of the excitations of their single-layer building blocks. To this end, we derive a model excitonic Hamiltonian by drawing an analogy with molecular crystals, which are other prototypical van der Waals materials. We employ this simplified model to analyse in detail the excitation spectrum of hexagonal boron nitride (hBN) that we have obtained from the ab initio solution of the many-body Bethe-Salpeter equation as a function of momentum. In this way we identify the character of the lowest-energy excitons in hBN, discuss the effects of the interlayer hopping and the electron-hole exchange interaction on the exciton dispersion, and illustrate the relation between exciton and plasmon excitations in layered materials.

PACS numbers: 71.35.-y,78.67.-n,78.20.Bh

I. INTRODUCTION

In many nanostructured materials, while strong covalent bonding provides the stability of the sub-nanometric elementary units, the whole assembly is held together by weak van der Waals interactions. The individual building blocks hence maintain most of their intrinsic characteristics also when arranged together to form a crystalline solid. In principle, novel materials properties can be thus tailored by controlling those of the elementary units. This bottom-up strategy in the synthesis of new materials is dominated by strong electron-hole (e-h) interactions for improved nanoelectronics and optoelectronics applications with customized functionalities.

In order to design materials with desired features for improved nanoelectronics and optoelectronics applications, the optical properties of layered materials need to be understood in detail. Due to the reduced effective screening, the optical response of 2D materials is dominated by strong electron-hole (e-h) interactions giving rise to bound e-h pairs, i.e. excitons. Nowadays, the state-of-the-art method to describe excitonic effects in condensed matter is the solution of the Bethe-Salpeter equation (BSE) within the framework of time-dependent density-functional theory. Those studies already addressed general questions like the effects of crystal local fields due to spatial inhomogeneities in the charge-density variation of the Hartree potential, the role of the interlayer coupling due to the long-range Coulomb interaction between charge oscillations on different layers, and the possibility to adopt a local-response approxima-
tion to formally relate 2D and 3D response functions. More recently, a “quantum-electrostatic heterostructure” model was similarly derived to describe the dielectric properties of complex multilayers starting from those of the single-layer building blocks, also taking into account the long-range coupling between layers due to the Coulomb interaction. However, in both cases hybridisation effects were neglected: in the present work they will be analysed in detail in terms of interlayer hopping mechanisms.

II. THEORETICAL FRAMEWORK AND COMPUTATIONAL DETAILS

The BSE is a formally exact Dyson-like equation relating the electron-hole correlation function $L$ to its independent-particle version $L_{0}$. Within the GWA to the self-energy, the BSE reads:

$$L(1,2,3,4) = L_{0}(1,2,3,4) + L_{0}(1,2,5,6) \times \varepsilon(5,7)\delta(5,6)\delta(7,8) - W(5,6)\delta(5,7)\delta(6,8)\{L(7,8,3,4),\}$$

where we have used the shorthand notation (1) for position, time and spin ($r_{f}, t_{f}, \sigma_{f}$) and repeated indices are integrated over. In $v$ is the bare Coulomb interaction and $W$ its statistically screened version calculated at the RPA level. The former enters the kernel of the BSE as an e-h exchange repulsive interaction and is responsible for crystal local-field effects. The latter is a direct attractive e-h interaction that is at the origin of excitonic effects, including the formation of bound excitons. For triplet excitons the e-h exchange interaction $v$ is absent.

The diagonal of the correlation function $L$ yields the density-density response function $\chi(1,2) = L(1,1,2,2)$. In a crystal, by taking the Fourier transform of $\chi$ to frequency and reciprocal-lattice space, one directly obtains the loss function $-\text{Im} \varepsilon_{M}^{-1}$ as:

$$-\text{Im} \varepsilon_{M}^{-1}(q, \omega) = \frac{4\pi}{q^{2}} \text{Im} \chi(q, q, \omega).$$

Here $\varepsilon_{M}$ is the macroscopic dielectric function and $q$ is a wave vector such that $q = q_{F} + G_{0}$, where $q_{F}$ belongs to first Brillouin zone (1BZ) and $G_{0}$ is a reciprocal-lattice vector. The loss function, which can be measured by inelastic x-ray scattering (IXS) and electron energy loss spectroscopy (EELS), describes the longitudinal linear response of the system to an external potential. It gives hence access to collective excitations such as plasmons, and (screened) electron-hole excitations.

Optical absorption spectra are related to the vanishing-$q$ limit of $\text{Im} \varepsilon_{M}(q, \omega)$, which can be obtained from the Fourier transform of the modified response function $\tilde{\chi}(1,2) = \tilde{L}(1,1,2,2)$:

$$\text{Im} \varepsilon_{M}(q, \omega) = -\frac{4\pi}{q^{2}} \text{Im} \tilde{\chi}(q, q, \omega),$$

where $\tilde{L}$ satisfies the BSE with the modified Coulomb interaction $\tilde{v}$ at the place of $v$. In the reciprocal space, $\tilde{v}$ is defined to be equal to $v$ except for the $G_{0}$ component for which it is set to 0. Therefore, the difference between optical absorption and loss function at $q = 0$ is given by the long-range $G_{0} = 0$ component of the Coulomb interaction which is absent for $\text{Im} \varepsilon_{M}$ in the BSE.

The loss function can be also explicitly written in terms of the imaginary and real parts of the macroscopic dielectric function:

$$-\text{Im} \varepsilon_{M}^{-1}(q, \omega) = \frac{\text{Im} \varepsilon_{M}(q, \omega)}{|\text{Re} \varepsilon_{M}(q, \omega)|^{2} + |\text{Im} \varepsilon_{M}(q, \omega)|^{2}}.$$  

Plasmon excitations are peaks in $-\text{Im} \varepsilon_{M}^{-1}$ corresponding to the frequencies where $\text{Re} \varepsilon_{M}$ is zero and $\text{Im} \varepsilon_{M}$ (which provides the damping of the plasmon) is not too large.

In order to describe correlated e-h pairs explicitly, the BSE (with $\tilde{v}$ at the place of $v$) can be cast in the form of a two-particle Schrödinger equation with an excitonic Hamiltonian:

$$\hat{H}_{ex}(q) = \sum_{ck} E_{ck}^1 a_{ck}^{\dagger} a_{ck} + \sum_{rk} E_{rk}^1 b_{rk}^{\dagger} b_{rk} + \sum_{vck,k'} \left[ 2\delta_{m} \bar{v}_{vck,k'}^2(q) - W_{vck,k'}^2(q) \right] a_{vck}^{\dagger} b_{rk}^{\dagger} a_{vck'} b_{rk}.$$  

Here $k$, belonging to the 1BZ, and $v(c)$ denote a valence (conduction) Bloch state of energy $E_{ck}$ ($E_{rk}$) calculated within the GWA; $a_{v}^{\dagger}$ ($a_{c}$) and $b_{v}^{\dagger}$ ($b_{c}$) are creation (annihilation) operators for electrons and holes, respectively; $\delta_{m}$ is 1 for the singlet and 0 for the triplet channel.

The first line of (5) is an independent particle Hamiltonian $\hat{H}_{ip}$ (corresponding to $L_{0}$ in the Dyson equation), while the second line contains the interaction terms stemming from the kernel of (1). The matrix elements of $\tilde{v}$ and $W$ are calculated in the basis of Bloch states $\psi_{vck}$.

$$\tilde{v}_{v'c'k'}^{vck}(q) = \int \! d\mathbf{r}d\mathbf{r}' \psi_{vck}^{*}(\mathbf{r}) \psi_{v'c'k'}(\mathbf{r}') \tilde{v}(\mathbf{r}, \mathbf{r}')$$

$$W_{v'c'k'}^{vck}(q) = \int \! d\mathbf{r}d\mathbf{r}' \psi_{vck}^{*}(\mathbf{r}) \psi_{v'c'k'}(\mathbf{r}) W(\mathbf{r}, \mathbf{r}')$$

In Eq. (5) we have adopted the Tamm-Dancoff approximation (TDA), which amounts to neglecting antiresonant $c \rightarrow v$ transitions and their coupling with resonant $v \rightarrow c$ transitions (extension to the general case can be seen in (22)).

The macroscopic dielectric function

$$\epsilon_{M}(q, \omega) = 1 - \frac{8\pi}{q^{2}} \sum_{\lambda} \frac{\left| \sum_{vck} A_{vck}^{\lambda}(q) \bar{v}_{vck}(q) \right|^{2}}{\omega - E_{\lambda}(q) + i\eta},$$

where $\bar{A}_{vck}(q)$ was similarly derived to describe the dielectric function.

$$\bar{A}_{vck}(q) = \frac{\text{Im} \varepsilon_{M}(q, \omega)}{|\text{Re} \varepsilon_{M}(q, \omega)|^{2} + |\text{Im} \varepsilon_{M}(q, \omega)|^{2}}.$$
with the oscillator strengths $\hat{\rho}_{\text{exc}}(\mathbf{q})$ defined as:

$$\hat{\rho}_{\text{exc}}(\mathbf{q}) = \langle \mathbf{k} e^{-i\mathbf{q} \cdot \mathbf{r}} | \mathbf{c} \rangle ,$$

(9)

and the exciton wavefunction

$$\Psi^\lambda(\mathbf{q}) = \sum_{\alpha \beta} \tilde{A}_\alpha^\lambda a^\dagger_\alpha \tilde{A}_\beta^\lambda a^\dagger_\beta |0\rangle ,$$

(10)

where $\mathbf{q}$ is the total momentum of the two-particle state, can be thus written in terms of the eigenvectors $A^\lambda(\mathbf{q})$ and the eigenvalues $E^\lambda(\mathbf{q})$ of the excitonic hamiltonian $H_{\text{exc}}(\mathbf{q})$:

$$\hat{H}_{\text{exc}}(\mathbf{q}) A^\lambda(\mathbf{q}) = E^\lambda(\mathbf{q}) A^\lambda(\mathbf{q}) .$$

(11)

The excitonic eigenvalues $E^\lambda(\mathbf{q})$ of $\hat{H}_{\text{exc}}$ are hence the poles of the $\hat{L}$ and $\epsilon_M$ functions in the frequency domain. They give rise to peaks in the spectrum of $\text{Im} \epsilon_M(\mathbf{q}, \omega)$ whose intensity is given by the numerator of Eq. (8). If it is zero, the corresponding excitonic state is said to be dark.

The inverse macroscopic dielectric function $\epsilon_M^{-1}(\mathbf{q}, \omega)$ can be analogously obtained from the eigenvectors and eigenvalues of the excitonic hamiltonian $H_{\text{exc}}$ that, in addition to $H_{\text{exc}}(\mathbf{q})$, also includes the long-range component of the Coulomb interaction:

$$\epsilon_M^{-1}(\mathbf{q}, \omega) = 1 + \frac{8\pi}{q^2} \sum_\lambda \frac{\sum_{\alpha \beta} A^\lambda_{\alpha \beta}(\mathbf{q}) \hat{\rho}_{\text{exc}}(\mathbf{q})}{\omega - E^\lambda(\mathbf{q}) + i\eta} .$$

(12)

Therefore also the loss function $-\text{Im} \epsilon_M^{-1}(\mathbf{q}, \omega)$ can be decomposed in terms of the eigenvalues $E^\lambda(\mathbf{q})$ and the eigenvectors $A^\lambda(\mathbf{q})$ of $H_{\text{exc}}$.

In our first-principles calculations we obtain the single-particle states $\psi_{\text{exc}}$ using Kohn-Sham (KS) density-functional theory within the local-density approximation (LDA). We use Troullier-Martins pseudopotentials, and expand the KS wavefunctions in a plane-wave basis set with a cutoff of 30 Hartree. The lattice parameters for the bulk are optimized using the LDA. We also consider hBN systems with variable interlayer distances $d$ for which the in-plane lattice vectors are kept constant to the bulk value. On the basis of the results of GW calculations for hBN bulk, we apply a scissor operator of 1.96 eV to correct for the LDA underestimation of the single-particle band gap. For larger interlayer distances $d$ the GW correction increases, for example it becomes 2.47 eV for $d = 1.5d_0$, where $d_0$ is the interlayer separation of the bulk. For the GW-BSE computational details of the hBN monolayer we refer to Ref. 33. In all the other cases, we sample the Brillouin zone using a $48 \times 48 \times 4$ Γ-centered grid. For the BSE calculations at finite $\mathbf{q}$ we follow the same procedure as described in Ref. 34. To simplify the analysis of the results in Sec. LVII here we use a minimal e-h transition basis set comprising 2 valence and 2 conduction bands and solve the BSE within TDA. As a consequence of the Kramers-Kronig relations, $\text{Re} \epsilon_M$ converges more slowly than $\text{Im} \epsilon_M$ with the number of higher-energy e-h transitions and (especially at small $\mathbf{q}$) is affected by the coupling with antiresonant transitions neglected in the TDA. While in the present case the main interest is to establish a direct connection between the electronic excitations characterising $\text{Im} \epsilon_M(\mathbf{q}, \omega)$ and the loss function $-\text{Im} \epsilon_M^{-1}(\mathbf{q}, \omega)$, for the comparison of the calculated loss-function spectra with experiment we refer to Ref. 34. In the construction of the BSE hamiltonian, we expand the single-particle states and static dielectric function with plane-wave cutoffs up to 387 and 133 eV, respectively. We perform the KS and static screening calculations using ABINIT, and BSE calculations with EXC. All the spectra presented in the following sections are calculated for in-plane momentum transfer $\mathbf{q}$ along the ΓM direction.

### III. RESULTS

The two panels of Fig. II display the real and imaginary parts of macroscopic dielectric function $\epsilon_M$ and the loss function $-\text{Im} \epsilon_M^{-1}$ of the bulk crystal of hBN calculated by solving the BSE for two different in-plane momenta $\mathbf{q}$. At vanishing $\mathbf{q}$ (top panel of Fig. II) the prominent peak at 5.67 eV in the absorption spectrum $\text{Im} \epsilon_M$ is a tightly bound exciton, located well within the direct band gap (which in GW amounts to 6.47 eV and is marked by the vertical arrow in the top panel of Fig. II). In the plot we have labeled the main peak as “A” (the explanation of the identification of the various excitations will be the subject of the detailed analysis in Sec. LV). Other structures are visible in the spectrum at higher energies, but for simplicity here and in the following we will focus on the lowest-energy excitations. As explained in previous works, the main absorption peak derives from $\pi - \pi^*$ transitions between top-valence and bottom-conduction bands that are visible for in-plane light polarization. Through the Kramers-Kronig relation, this “A” peak of $\text{Im} \epsilon_M$ induces a strong oscillation in $\text{Re} \epsilon_M$ which crosses the zero axis with a positive slope at 5.99 eV. $\text{Im} \epsilon_M$ being small at this energy, this zero of $\text{Re} \epsilon_M$ gives rise to a plasmon resonance in the loss function $-\text{Im} \epsilon_M^{-1}$, which shows a peak at the same energy (see Eq. (47)). It is here worth noticing that in hBN also this plasmon excitation lies within the direct gap, since the collective charge excitation of the $\pi$ electrons is strongly affected by the e-h attraction. As discussed in details in Refs. 33 and 34, for increasing $\mathbf{q}$ this $\pi$ plasmon disperses to higher energies and at larger $\mathbf{q}$ it enters the continuum of particle-hole excitations.

As a matter of example, the bottom panel of Fig. II shows the spectra obtained for the second smallest finite $\mathbf{q}$ that we have considered in our calculations (for the other momentum transfers, not shown here, similar considerations can be made). Globally the spectra at finite $\mathbf{q}$ remain qualitatively similar to the $\mathbf{q} = 0$ case shown in the top panel of Fig. II. Still we can recognize that in
that becomes active at a new many-body electron-hole excitation. This new peak (which does not take place in correspondence with a zero of $\epsilon_M$, hence it is not a plasmon) matches a new very weak peak in $\epsilon_M$, so it has to be ascribed to a new many-body electron-hole excitation that becomes active at $q \neq 0$.

As discussed in Sec. II the spectra for $\epsilon_M$ and $-\epsilon_M\lambda^q$ can be also analysed in detail by making use of the eigenvalues and eigenvectors of the excitonic hamiltonian that enter Eq. (8) and Eq. (12), respectively. Fig. 2(a) shows the 18 lowest energies $\epsilon_q^\lambda$ as a function of $q$ for the singlet excitons that are obtained from the diagonalization of excitonic hamiltonian $H_{\text{ex}}$ [4]. They are hence the poles of $\epsilon_M$ [5] and of the modified two-particle correlation function $\tilde{L}$. The color scale represents their intensity $|\lambda^q|\rho$ at the numerator of Eq. (5) where the e-h exchange interaction $\tilde{L}$ is the only contribution. Red squares are for states that have a visible peak in Im $\epsilon_M$, while blue squares are dark exciton states with no intensity in the spectrum. The other two panels of Fig. 2 use the same representation. Fig. 2(b) displays the exciton eigenvalues $\epsilon_q^\lambda(q)$ obtained from the diagonalization of $H_{\text{ex}}$ that includes the long-range Coulomb interaction: they enter the loss function spectra $-\text{Im} \epsilon_M^\lambda(q,\omega)$ (12). Finally, in Fig. 2(c) the triplet exciton eigenvalues are also reported for comparison (they cannot be directly measured by loss or absorption spectroscopies). They are calculated from the excitonic hamiltonian $H_{\text{ex}}$ [5] where the e-h exchange interaction $\tilde{L}$ is absent. With respect to the singlet excitons the triplet eigenvalues are globally lower [compare Fig. 2(a) and 2(c)], as the e-h exchange interaction is repulsive and hence yields singlet states that have higher energies than the corresponding triplets.

The first and third $q$ points in Figs. 2(a), 2(b) allow us to understand better the absorption and loss spectra plotted in the two panels of Fig. 1. For example, in Fig. 2(a) we discover that in the optical limit $q \rightarrow 0$ the first visible exciton “A+” is degenerate with a dark state “B+” (labelled “B+” here) and that below them there are other two degenerate dark excitons “A−” and “B−” that do not contribute to the $q \rightarrow 0$ absorption spectrum in the top panel of Fig. 1 (this point was already subject of discussion in Refs. 28, 30, and 31). We can also see that finite $q$ one of the two lowest dark excitons becomes visible, giving rise to the low-energy peak “A−” in the absorption spectrum of Fig. 1, bottom panel. Finally, the weak peak “X” at 5.88 eV is due to another exciton state that is dark at $q = 0$ and switches on at $q \neq 0$. For all wavevectors $q$, at higher energies the exciton states become very dense, forming a continuum of excitations. In Sec. V we will focus on the 4 lowest-energy discrete states that are well within the fundamental gap.

We can now repeat the same analysis for the loss functions $-\text{Im} \epsilon_M^\lambda(q,\omega)$ in Fig. 1 using the poles of $L$ represented in Fig. 2(b). We thus discover that at $q = 0$ the plasmon excitation at 5.99 eV is not the lowest-energy eigenvalue. It is actually located already in the energy region where e-h excitations are rather dense. So it is not easy to track its dispersion after the first few $q$ points. At the bottom of the eigenvalue spectrum there are instead 3 dark states (2 of them are degenerate at $q = 0$) that are well separated from the other excitations. They have a dispersion as a function of $q$ that is similar to that of the lowest poles of $L$ in Fig. 2(a). It is hence tempting to make a connection between them. In Sec. V we will explain rigorously why this is indeed the case (so they are labeled “A−” and “B+” here) and why the plasmon ex-

\[\begin{align*}
\text{FIG. 1.} & \quad \text{The real and imaginary parts of the macroscopic dielectric function Re} \epsilon_M & \text{and Im} \epsilon_M & \text{and the loss function} \\
& \quad \text{Re} \epsilon_M & \text{and Im} \epsilon_M & \text{calculated at two different wave vectors q along} \\
& \quad \text{the in-plane FM direction. For improved visibility the loss} & \text{functions have been rescaled. The vertical arrows mark the} \\
& \quad \text{smallest independent-particle GW transition energy (which} & \text{for q = 0 corresponds to the direct band gap).}
\end{align*}\]
FIG. 2. Exciton eigenvalue spectrum $E^\lambda(q)$ in bulk hBN for in-plane $q$ along $\Gamma M$ for (a) the electron-hole correlation function $L$ corresponding to $\text{Im} \epsilon_M(q, \omega)$ featuring singlet excitons, (b) the electron-hole correlation function $L$ corresponding to the loss function $-\text{Im} \epsilon_M(q, \omega)$, displaying plasmons and e-h excitations, and (c) for triplet excitons. The $\Gamma M$ length is $1.45 \text{ Å}^{-1}$. For the explanation of the peak labels see the main text. In the loss-function plot, panel (b), the solid black line is a guide for the eye in order to better track the plasmon dispersion (corresponding to the $A^+$ feature).

FIG. 3. Same as Fig. 2 for increased interlayer distance $d = 1.5d_0$ (where $d_0$ is the experimental interlayer separation of hBN).

citation instead has a “$A^+$” character. Finally, at $q = 0$ at 5.88 eV we recognize the same “$X$” excitation that is present also in the spectrum of $\bar{L}$ in Fig. 2(a) and is responsible for the weak structures in the absorption and loss spectra in the bottom panel of Fig. 1.

IV. DISCUSSION

A. The exciton hamiltonian in layered crystals

In order to interpret the numerical results of the previous section, here we generalize the approach that some of us introduced in Ref. 23 to explain the excitonic properties of molecular crystals. We thus rewrite the excitonic hamiltonian $\hat{H}_{ex}$ (which in Eq. (5) is expressed in terms
of Bloch wave functions delocalised over the crystal in the basis of wave functions localized on the elementary units of the system. While in molecular crystals the elementary units are the single molecules, in the present case they are the single layers of BN (stacked along the $z$ axis). We assume that the one-particle wave functions $\psi(r)$ localized on different layers do not overlap and can be factorized in an in-plane $\phi(\rho)$ and out-of-plane $\chi(z)$ components, with $r = (\rho, z)$. Specifically, for given in-plane wave vector $k$ and out-of-plane $k_z$, the single-particle wave function $\psi_{k,k_z}(r)$ is expanded in the basis of single-layer wave functions as: $\sum_{k_i} c_{ki,k_z} R_{ki}(\rho) \chi_i(z - R)$. Here $R$ is the lattice vector along $z$ and the index $i$ denotes the layers inside the unit cell. We also consider the possibility that the various layers stacked along $z$ are rotated one with respect to another by an angle $\beta$ (in hBN $\beta = 60^\circ$), and therefore also the 2D Brillouin zones are rotated by an angle $\beta$. Hence, choosing a reference layer $i = 1$, we define $k^{(i)} = k$ for $i = 1$ and $k^{(i)} = \beta^{-1}k$ for $i \neq 1$, $\beta k$ being the wave vector obtained rotating $k$ by an angle $\beta$ (see App. A for more details). For simplicity we further consider for each layer a two-bands system, with only one valence $v$ and one conduction $c$ bands. Under these assumptions, the whole excitonic Hamiltonian $H_{ex}$ of the crystal Eq. \ref{eq:Hex} takes the simple form of the sum of three terms $H_{ex} = H_{ip} + \hat{K}_{CT} + \hat{K}_{FR}$:

\begin{align}
\hat{H}_{ip} &= \sum_{k_1,k_2} \sum_{RS} \sum_{ij} E_c^{R_i S_j}(k_1, k_2) a_{c,k_1 R_i}^\dagger a_{c,k_2 S_j} - \sum_{k_1,k_2} \sum_{RS} \sum_{ij} E_v^{R_i S_j}(k_1, k_2) b_{c,k_1 R_i}^\dagger b_{c,k_2 S_j}, \\
\hat{K}_{FR} &= \sum_{k_1,k_2} \sum_{RS} [\delta_{R_i S_j} (v k_1 c k_2 v k_3 c k_4) - \delta_{R_i S_j} W_{R_i R_i}^{S_i S_j}(v k_1 c k_2 v k_3 c k_4)] a_{c,k_2 R_i}^\dagger b_{c,k_1 S_j} a_{c,k_1 R_i} b_{c,k_2 S_j} \\
\hat{K}_{CT} &= - \sum_{k_1,k_2} \sum_{R_i S_j} \big(1 - \delta_{R_i S_j}\big) \sum_{i} W_{S_i S_j}^{S_i S_j}(v k_1 c k_2 v k_3 c k_4) \phi_{v,k_2}^{\dagger} \phi_{v,k_1} (\rho) \phi_{c,k_3} (\rho) \chi_{v}^{\dagger} (z - R) \phi_{v,k_1}^\dagger (\rho) \chi_{v} (z - S)
\end{align}

with

\begin{align}
W_{S_i S_j}^{S_i S_j}(v k_1 c k_2 v k_3 c k_4) &= \int d\rho d\rho' \phi_{v,k_1}^\dagger (\rho) \chi_{v}^{\dagger} (z - R) \phi_{v,k_1}^\dagger (\rho) \chi_{v}^{\dagger} (z - S) W(\rho, \rho') \phi_{v,k_3} (\rho) \chi_{v} (z - R) \phi_{v,k_3} (\rho) \chi_{v} (z - S) \\
\delta_{R_i S_j} W_{R_i R_i}^{S_i S_j}(v k_1 c k_2 v k_3 c k_4) &= \int d\rho d\rho' \phi_{v,k_1}^\dagger (\rho) \chi_{v}^{\dagger} (z - R) \phi_{v,k_1}^\dagger (\rho) \chi_{v}^{\dagger} (z - S) \delta(\rho, \rho') \phi_{v,k_3} (\rho) \chi_{v} (z - R) \phi_{v,k_3} (\rho) \chi_{v} (z - S).
\end{align}

In the Bloch picture $\hat{H}_{ip}$ contains independent e-h transitions between single-particle bands. Equivalently, here $H_{ip}$ describes scattering processes from layer to layer, independently for electrons and holes, being

\begin{equation}
E_{v(c)}^{RS}(k_1, k_2) = E_{v(c)}^{i}(k_1) \delta_{R_i S_j} k_1 - k_2 + \delta_{v(c)}^{R_i S_j}(k_1, k_2),
\end{equation}

where $E_{v(c)}^{i}(k_1)$ is the single-layer band dispersion and $\delta_{v(c)}^{R_i S_j}(k_1, k_2)$ are interlayer hopping matrix elements (see App. A) that give rise to the finite $k_z$ dispersion of the bands in the crystal (see Fig. S in App. B).

In Eq. \ref{eq:Hex} the second and third terms $\hat{K}_{FR}$ and $\hat{K}_{CT}$ describe the interaction between an electron and a hole that are localized on the same layer or on different layers, respectively. In order to keep a closer contact with the exciton physics of molecular crystals, here we name the intralayer configuration as a “Frenkel” (FR) exciton and the interlayer configuration as a “charge-transfer” (CT) exciton. In other words, in the present context we call FR an exciton that is fully localized on a single layer, independently of being localised or not within the layer. Therefore this definition applies equivalently for excitons with different in-plane localisation characters, as for example in hBN (where the exciton is tightly bound also within the single layer) or in MoS$_2$ (where it is weakly bound). We note that the e-h exchange interaction $\tilde{v}$ is different from zero only for e-h pairs localized on the same layer, therefore it is absent for CT excitons in Eq. \ref{eq:Hex}.

The FR and CT interaction terms in Eq. \ref{eq:Hex} are coupled by the interlayer hopping terms in $\hat{H}_{ip}$. Without the interlayer hopping $\delta_{v(c)}^{R_i S_j}(k_1, k_2)$ the excitonic Hamiltonian \ref{eq:Hex} factorizes into two independent blocks: a CT Hamiltonian $H_{CT} = H_{ip} + \hat{K}_{CT}$ describing an interacting e-h pair localised on different layers and a FR Hamiltonian $H_{FR} = H_{ip} + \hat{K}_{FR}$ describing an interacting e-h pair on the same layer (in both cases we set $\delta_{v(c)}^{R_i S_j}(k_1, k_2) = 0$ in $H_{ip}$).

The CT exciton wavefunctions\cite{56}:

\begin{equation}
|\Psi_{ex}^{CT}(q)\rangle = \sum_{\lambda} \sum_{ij \tau} c_{ij}^{T} |\Psi_{ij,\tau}^{\lambda}(q)\rangle
\end{equation}
Frenkel Hamiltonian $\hat{H}_{FR}$ instead contains also an interlayer coupling that needs additional consideration.

By further splitting the e-h exchange interaction $\tilde{v}$ into a long-range contribution $\tilde{v}_0$ (corresponding to the $G_{||} = 0$ component in reciprocal space) and a short-range contribution $\tilde{v}$ such that $\tilde{v} = \tilde{v}_0 + \tilde{v}$, the FR Hamiltonian $\hat{H}_{FR}$ can be separated into an intralayer term $\hat{H}_L$ and an interlayer coupling $\hat{V}$, with:

\[
\hat{V} = \sum_{k_1,k_2,k_3,k_4} \sum_{R\|} \tilde{v}_{R\|}^{S_j S_j} (v_{k_1 c k_2 v k_3 c k_4}) a_{c k_2 R\|}^\dagger b_{k_1 R\|}^\dagger b_{k_3 R\|} a_{c k_4 R\|}
\]

where

\[
\tilde{v}_{R\|}^{S_j S_j} (v_{k_1 c k_2 v k_3 c k_4}) = \sum_{q_{||},G_{||} \neq 0} \frac{4\pi}{|q_{||} + G_{||}|^2} \tilde{\rho}^{S_j S_j}_{c k_2 k_1} (q_{||}) \tilde{\rho}^{S_j S_j}_{c k_4 k_3} (q_{||})
\]

Equivality, Eq. (21) can be written in terms of the partial Fourier transform of the Coulomb potential:

\[
v(q_{||} + G_{||}, z, z') = \frac{2\pi}{|q_{||} + G_{||}|} e^{-i|q_{||} + G_{||}|z - z'}
\]

as:

\[
\tilde{v}_{R\|}^{S_j S_j} (v_{k_1 c k_2 v k_3 c k_4}) = \sum_{q_{||},G_{||} \neq 0} \frac{2\pi}{|q_{||} + G_{||}|} \tilde{\rho}^{S_j S_j}_{c k_2 k_1} (q_{||}) \tilde{\rho}^{S_j S_j}_{c k_4 k_3} (q_{||})
\]

From Eq. (24) we can conclude that the off-diagonal elements $S \neq R$ and $i \neq j$ of $\tilde{v}_{R\|}^{S_j S_j}$ are actually zero, for the presence of the exponential terms $e^{-i|q_{||} + G_{||}|z - z'}$ with $G_{||} \neq 0$. For its short-range nature, the $\tilde{v}$ interaction therefore does not couple different layers.

With respect to the Bloch picture, such a transformation and decomposition of the excitonic hamiltonian illustrates much more clearly the physics of excitons in layered materials that we want to uncover. Here the eigenstates of $\hat{H}_L$ represent the excitations of an elementary unit of our van der Waals material, namely a single BN layer embedded in the bulk crystal. They are formally analogous to the excitations of a single molecule in a molecular solid. Thus, by analogy with molecular crystals, a FR exciton in the present case can be seen as an elementary excitation of a single layer, which can scatter from one layer to another due to the interlayer coupling $\hat{V}$. From a mathematical point of view, this means that we expand the FR exciton wave functions (which are the eigenfunctions of $\hat{H}_{FR}$) on the basis of the eigenstates of
\[ \hat{H}_L: \]
\[ |\Psi_{ex}^{FR}(\mathbf{q})\rangle = \frac{1}{\sqrt{N}} \sum_{\lambda, \mathbf{R}_i} \tilde{c}_\lambda^*(\mathbf{q}) \Psi_{\hat{R},ii}(\mathbf{q}) \] (26)

where
\[ |\Psi_{\hat{R},ii}(\mathbf{q})\rangle = \sum_{\mathbf{k}} A_{\hat{R},ii}^{\lambda,\mathbf{k}}(\mathbf{q}) \tilde{b}^\dagger_{\hat{R},(\mathbf{i})}\mathbf{k} \tilde{b}^{\dagger}_{\mathbf{k}(\mathbf{i})} \Psi_{\mathbf{k}(\mathbf{i})}(0). \] (27)

and where we have used the fact that for in-plane \( q \) \( e^{iq \mathbf{R}} = 1 \). The matrix elements of \( \hat{V} \) are:
\[ \langle \Psi_{\hat{R},ii}(\mathbf{q}) | \hat{V} | \Psi_{\hat{R},jj}(\mathbf{q}) \rangle = \sum_{G_z \neq 0, G_z = 0} \frac{4\pi}{|q + G|^2} \]
\[ \times |S_i^\lambda(q)|^2 \int dz \chi_v^*(z) e^{iG_zz} \chi_v(z) \bigg| e^{-iG_zd_{ij}}, \] (28)

where \( S_i^\lambda(q) \) is the oscillator strength of the exciton \( \lambda \) of the layer \( i \):
\[ S_i^\lambda(q) = \sum_{\mathbf{k}} A_{\mathbf{v},\mathbf{k}}^{\lambda,\mathbf{i}}(q) \tilde{\rho}^{\dagger}_{\mathbf{v},\mathbf{k}}(q^{\mathbf{i}}), \] (29)

and where we have used the fact that \( \chi_{i}^+(z) = \chi_{i}^-(z - d_{ij}) \equiv \chi_{i}(z - d_{ij}) \) for both \( n = v, c \), with \( d_{ij} \) the distance between the layers \( i \) and \( j \). From Eq. (28) we realise that \( \hat{V} \) operates only on visible excitons and cannot couple visible and dark excitons for which \( S_i^\lambda(q) = 0 \).

**B. The exciton hamiltonian in hBN**

If we consider a crystal with two inequivalent layers per unit cell, as it is the case for hBN, for each quantum number \( \lambda \) that defines an excitation of the single layer one has four excitons in the bulk [we take into account only first nearest-neighbor CT excitons and assume \( \tau = 0 \) in Eq. (17)]. The FR and CT excitons that diagonalize the excitonic hamiltonian \( \hat{H}_{ex} \) in absence of interlayer hopping are then the symmetric and antisymmetric combinations with respect to the exchange of the e-h pair between two inequivalent layers:
\[ \langle CT_{\pm}^\lambda \rangle = \frac{1}{\sqrt{N}} \sum_{\mathbf{R}} \frac{1}{\sqrt{2}} \left[ |\Psi_{\hat{R},112}^\lambda \rangle \pm |\Psi_{\hat{R},211}^\lambda \rangle \right] \] (30)
\[ |FR_{\pm}^\lambda \rangle = \frac{1}{\sqrt{N}} \sum_{\mathbf{R}} \frac{1}{\sqrt{2}} \left[ |\Psi_{\hat{R},111}^\lambda \rangle \pm |\Psi_{\hat{R},222}^\lambda \rangle \right]. \] (31)

The \( \langle CT_{\pm}^\lambda \rangle \) states are degenerate, while the energy separation between the \( |FR_{\pm}^\lambda \rangle \) states in the context of molecular crystals is usually called Davydov splitting.\(^{27}\)

In the case of hBN, the two lowest excitons of the BN monolayer, which are degenerate at \( q = 0 \), are a visible exciton \( A \) and a dark exciton \( B \) [see Fig. 4(a)]. They originate from transitions from the top-valence to the bottom-conduction bands with \( q \) vectors located around the \( K \) or \( K' \) points of the Brillouin zone, respectively.\(^{15,58}\) These two intralayer \( A \) and \( B \) excitons hence produce eight excitons (four FR and four CT excitons) in the bulk crystal. Since the \( B \) exciton is dark for all \( q \) along \( \Gamma M \), in the bulk the \( A \) and \( B \) excitons are not mixed by \( \hat{V} \) [see Eq. (28)] and preserve their identity. The four \( |CT_{\pm}^{A,B} \rangle \) excitons are located at higher energies since they have smaller binding energies, as a result of the e-h attraction being smaller for interlayer e-h pairs than for intralayer e-h pairs. In the following we focus on the four \( |FR_{\pm}^{A,B} \rangle \) excitons that are the lowest-energy excitations in the bulk.

The \( A \) exciton of the single layer of energy \( E_A^A(q) = E_{11}^A(q) = E_{22}^A(q) \) gives rise to the two \( |FR_{\pm}^A \rangle \) excitons:
\[ E_{\pm}^A(q) = E_A^A(q) \pm \bar{I}_A(q) \pm I_A(q) \] (32)

where \( \bar{I}_A(q) \pm I_A(q) \) are the contribution to the exchange e-h interaction \( \hat{V} \) for the symmetric (+) and antisymmetric (−) states, respectively. They are the excitation-transfer interactions that are responsible for the interlayer propagation of the FR exciton in the crystal.\(^{23}\) \( \bar{I}^A \) is related to the scattering process of an e-h pair between two inequivalent layers and, analogously, \( I^A \) between equivalent layers in different unit cells. Explicitly
they read:

\[
\bar{I}^A(q) = \sum_{G_x \neq 0, G_y = 0} \frac{4\pi}{|q + G|^2} |S^A(q)|^2 \times \left| \int dz \chi^*_v(z)e^{iG_z z} \chi_v(z) \right|^2
\]

\[
\bar{J}^A(q) = \sum_{G_x \neq 0, G_y = 0} \frac{4\pi}{|q + G|^2} |S^A(q)|^2 \times \left| \int dz \chi^*_v(z)e^{iG_z z} \chi_v(z) \right|^2 e^{-iG_d d}.
\]

We note that \(\bar{I}^A(q)\) and \(\bar{J}^A(q)\) are both zero at \(q = 0\), since the oscillator strength \(S^A(q)\) in the dipole limit \(q \to 0\) is proportional to \(q \cdot \mu^A\). Therefore \(\bar{I}\) and \(\bar{J}\) in layered systems do not yield any Davydov splitting between symmetric and antisymmetric excitons at \(q = 0\), in contrast to the molecular crystal case. Instead at finite \(q\) and \(q = 0\), the hopping induces a finite Davydov splitting between symmetric and antisymmetric states. Indeed this effect of the interlayer hopping that at \(q = 0\) becomes visible at \(d = 1.5d_0\), where \(d_0\) is the experimental interlayer distance of hBN, are displayed in Fig. 3(a) and (c).

The symmetric \(|FR^A\rangle\) exciton is hence visible, while the antisymmetric \(|FR^B\rangle\) exciton is dark, since the two integrals in Eq. (33) exactly cancel in this case. For the B exciton the matrix element of \(\hat{V}\) is zero [since \(S^B(q) = 0\) in Eq. (28)]. Therefore the two \(|FR^B\rangle\) excitons remain degenerate in the bulk:

\[
E^B_{\pm}(q) = E^B(q).
\]

Moreover, as \(S^B(q) = 0\), they are both dark [see Eq. (33)].

In summary, by neglecting the interlayer hopping terms in the exciton hamiltonian \(H^A\), we would expect that the two lowest A and B excitons of the BN single layer give rise to 3 FR dark excitons and 1 FR visible exciton in the bulk (together with CT excitons at high energies).

The effect of the hopping is, in general, to couple FR and CT excitons. This coupling produces states with mixed character, FR+CT and CT+FR respectively, and modifies their energies. In hBN, as demonstrated in the App. A at \(q = 0\) \((CT^+_{2d}|FR^A_{2d}) = 0\): excitons with different parities do not couple, giving rise to \(|(FR + CT)^+_{2d}\) states with well defined parity. Moreover, since \((CT^+_{2d}|FR^A_{2d}) \neq (CT^+_{2d}|FR^B_{2d})\) (see App. A), at \(q = 0\) the hopping induces a finite Davydov splitting between symmetric and antisymmetric excitons. Instead at finite \(q\) the various excitons formally lose their parity character as FR and CT states with different parities are generally allowed to mix together.

C. Exciton dispersion: electron-hole exchange and interlayer hopping

On the basis of the previous analysis, we can now examine in detail the properties of the four lowest-energy singlet excitons in hBN [see Fig. 2(a)]. In particular, we can understand the effect of e-h exchange by comparing singlet and triplet excitons [see Fig. 2(a) and (c)], because in the latter there is no e-h exchange. Moreover, we can suppress also the interlayer hopping by artificially increasing the interlayer distance \(d\). The singlet and triplet exciton band structures obtained with \(d = 1.5d_0\), where \(d_0\) is the experimental interlayer distance of hBN, are displayed in Fig. 3(a) and (c). With this increased separation between BN layers, the interlayer hopping is reduced so much that the \(k_z\) dispersion of the top-valence and bottom-conduction single-particle bands becomes negligible (see App. B).

At \(q = 0\) the four lowest singlet excitons are grouped in two pairs [see Fig. 2(a)]. Since in the single layer the A and B excitons are degenerate at \(q = 0\) [see Fig. 2(a)], and the e-h exchange terms \(I(q = 0)\) and \(J(q = 0)\) are zero for all of them [see Eqs. (33)-(34)], the energy splitting between the two pairs must derive from the interlayer hopping (which we reasonably assume to be the same for A and B excitons). At \(q = 0\) the hopping conserves the parity character, removing the degeneracy between symmetric and antisymmetric states. Indeed this energy splitting is present also for the triplet excitons [see Fig. 2(c)], whereas it becomes zero for an increased interlayer distance [see Fig. 3(a) and (c)]. Therefore we can conclude that at \(q = 0\) the two excitons of the lowest pair, which are both dark, are the antisymmetric \(|(FR+CT)^A\rangle\) and \(|(FR+CT)^B\rangle\) states, while the two excitons of the other pair are the symmetric \(|(FR+CT)^+\rangle\) (which is visible) and \(|(FR+CT)^-\rangle\) (which is dark). For simplicity, in Figs. 3(a) and (c) we have labeled “A\(\pm\)” and “B\(\pm\)” respectively the states \(|(FR+CT)^A\rangle\) and \(|(FR+CT)^B\rangle\).

Having established the character of the excitons at \(q = 0\), we can now track their dispersion as a function of \(q\). The fact that one of the excitons of the lowest pair that is dark at \(q = 0\) becomes visible at \(q \neq 0\) for both the singlet and triplet cases [see Fig. 2(a) and (c)] is another effect of the interlayer hopping that at \(q \neq 0\) mixes FR and CT states with different parities. This means that the parity is no more a good quantum number and the eigenstates of the excitonic hamiltonian are combinations of \(|(FR + CT)^\pm\rangle\) and \(|(FR + CT)^\mp\rangle\) states. In this way the dark exciton \(|(FR + CT)^\pm\rangle\) is switched on by the effective coupling with the visible exciton \(|(FR + CT)^\mp\rangle\).

Formally all the excitons lose their defined parity, but here for simplicity we still call them “A\(\pm\)” (the two visible states) and “B\(\pm\)” (the two dark states).

In order to infer the effect of the interlayer hopping on the exciton dispersion, we compare the behavior of the triplet excitons in the bulk [see Fig. 2(c)], for the increased interlayer distance [see Fig. 3(c)] and in the monolayer [see Fig. 2(b)]. In the monolayer the A and B
triplet excitons are almost degenerate: there is a tiny separation due to the direct e-h attraction \( W_{\text{short}} \). The same holds for \( d = 1.5 d_0 \), where there is no effect of the interlayer hopping. In the bulk, instead, the hopping acts differently for the various excitons, giving rise to a finite dispersion that removes the degeneracies. The energy level ordering remains the same for all \( q \). From the bottom to the top one has the following states: “B\(^-\)”, “A\(^-\)”, “B\(^+\)”, and “A\(^+\)”. The difference between the dispersions of the singlet and the triplet excitons in Fig. 2(a) and (c) illustrates the role of the e-h exchange interaction in the bulk as a function of \( q \). While the “B\(^\pm\)” excitons keep the same dispersion in the two channels (as the e-h exchange \( \tilde{I}(q) = \tilde{J}(q) = 0 \) for B excitons), for the “A\(^\pm\)” excitons we observe that the effect of the e-h exchange interaction becomes larger for small \( q \) than for large \( q \), where the dispersion of singlet and triplet excitons tend to be the same, being determined by the single-particle band dispersion only.

At increased interlayer distance \( d = 1.5 d_0 \), in contrast to the bulk, both for the singlet and the triplet channels also at finite \( q \) there remain one visible and three dark excitons, as in the limit \( q = 0 \) [see Fig. 2(a) and (c)]. This confirms that by suppressing the interlayer hopping the antisymmetric \([FR^A]\) exciton cannot couple with excitons of different parity and continues to be dark. All the excitons keep the same parity as at \( q = 0 \). The two dark \([FR + CT]_B^A \) excitons remain degenerate, since the e-h interaction \( \tilde{V} \) has no effect on them. They are located at lower energies than the \([FR+CT]^A_+\) excitons as \( \tilde{V} \) is repulsive. In particular, the dark \([FR + CT]^A_\pm\) shows a larger dispersion for the visible \([FR + CT]^A_\pm\), implying that the effect of \( \tilde{I} - \tilde{J} \) is larger than \( \tilde{I} + \tilde{J} \). In general, the energy-level ordering is, from the bottom to the top: \([FR + CT]_B^A \) (degenerate), \([FR + CT]_+^A\) and \([FR + CT]_-^A\).

By increasing \( d \), the screening of the e-h attraction \( W \) is reduced and, as a consequence, the binding energies of all the excitons increase (however their absolute positions remain almost constant\(^\text{39}\)). In order to directly compare, for increasing interlayer distances \( d \), the dispersion of the visible “A\(^+\)” exciton as a function of \( q = 2 \pi / \lambda \), in Fig. 4 we have hence aligned, for the different separations \( d \), the exciton energies to their \( q = 0 \) value. By increasing the interlayer distance, the dispersion becomes more steep at small \( q \) and tends to be the same at large \( q \). As a result of the competition between the e-h exchange interaction and the single-particle band dispersion, in the exciton dispersions we can always distinguish two regimes: (i) at large \( q \) (i.e. for \( \lambda \ll d \)) the sum over \( G_\|=0 \) in Eq. 33 can be approximated with an integral. So \( \tilde{I}^A \) and \( \tilde{J}^A \) become:

\[
\tilde{I}^A(q) \approx \frac{2 \pi}{q} \beta(q) |S^A(q)|^2 
\]

\[
\tilde{J}^A(q) \approx \tilde{I}^A(q) e^{-q d} 
\]

Under these conditions, as shown in Ref. 33, \( \tilde{I}^A \) reaches a constant value at large \( q \). Moreover, since \( \lambda \ll d \), the exponential factor in Eq. 38 goes to zero and \( \tilde{J}^A \) becomes negligible. As a consequence, in this regime the dispersion of the symmetric and antisymmetric excitons becomes the same and, at large \( q \), is set by the hopping only. (ii) at small \( q \) (i.e. for \( \lambda \gg d \)) the sums over \( G_\|=0 \), Eqs. 33-34 is independent of \( q \) and \( S^A(q) = q \cdot \mu^A \); \( \tilde{I} \) and \( \tilde{J} \) are quadratic in \( q \). Therefore in this regime the exciton dispersion is also determined by the e-h exchange \( \tilde{V} \), in addition to the hopping contribution that is always present. At small \( q \) the e-h exchange interaction becomes more and more important as \( d \) increases, until in the 2D limit it becomes the dominant contribution [compare the dispersion of singlet in Fig. 2(a) and triplet in Fig 2(b)]. Indeed, in the 2D limit, when Eqs. 47 and 48 are exact for every \( q \), the e-h exchange contribution becomes linear in \( q \), as explained in detail in Ref. 33.

D. Plasmon dispersion: long-range Coulomb interaction

In order to describe the plasmon properties, in the excitonic Hamiltonian \( \hat{H}_\text{exc} \) one has to replace the short-range \( \tilde{v} \) with the full Coulomb interaction \( v \). This implies that in the long-range \( G_\|=0 \) contribution to the e-h exchange \( \tilde{v} \) also the \( G_\|=0 \) component has to be included. The excitation transfer interactions [with \( \tilde{I} \) and

![FIG. 5. Dispersion of the visible “A\(^+\)” exciton for different interlayer distances \( d \) (\( d_0 \) is the experimental value). For each case the exciton energies are defined with respect to the corresponding \( q = 0 \) value.](image)
The repulsive long-range interaction is felt only by the symmetric state \((FR+CT)^{A+}\) that is the plasmon excitation in \(L\). As a consequence, its energy at \(q = 0\) is upshifted with respect to corresponding \("A+\"\) pole of \(L\) by \(\sim (8\pi/q^2)S(q = 0) = 8\pi|\mathbf{q} \cdot \mathbf{\mu}|^2\). At finite \(q\) the plasmon energy displays a quadratic dependence on \(q\). Without interlayer hopping (i.e. for interlayer spacing \(d > 1.5d_0\)), the plasmon dispersion is hence similar to that of the triplet exciton energy. This is a consequence of the cancellation at finite \(q\) occurring to a large extent between the first and second terms in Eqs. (40)-(41). While the first terms account for the difference between plasmon and singlet exciton (see Fig. 6), the second terms are responsible for the difference between singlet and triplet excitons (see Fig. 7). As a matter of fact, by comparing Fig. 6 and Fig. 7 we notice that for each interlayer separation they have an opposite behavior as a function of \(q\).

At large \(q\) (i.e. for \(\lambda \ll d\)) the long-range contribution becomes negligible. As shown in Fig. 6 for increasing \(d\) the plasmon energy approaches the visible-exciton energy for smaller and smaller \(q\); the loss function \(-\text{Im} \epsilon_M^{-1}\) becomes equal to \(\text{Im} \epsilon_M\) when \(qd \gg 1\). In the 2D limit (i.e. \(d \to \infty\), as for any completely isolated system\(^44,45\), \(-\text{Im} \epsilon_M^{-1}\) and \(\text{Im} \epsilon_M\) mathematically coincide for all \(q\).

\section{V. SUMMARY}

From the solution of the \textit{ab initio} Bethe-Salpeter equation (BSE) as a function of momentum \(q\), we have obtained the eigenvalue spectrum of the excitonic Hamiltonian for the electronic excitations of hexagonal boron nitride and we have established the connection with measured optical absorption and energy loss spectra. We have discussed the properties of both visible and dark excitons on the basis of a simplified model that we have derived from the full \textit{ab initio} BSE and by analogy with the case of molecular solids. This model has allowed us to provide an efficient description of the excitations in the bulk crystal starting from the knowledge of the excitons in the single layer. In this way we have obtained a general picture of the exciton physics in layered materials. Our analysis uncovers the interplay between the electronic band dispersion and the electron-hole exchange interaction in setting the exciton properties in this important class of materials. Holding a general validity, it can be similarly applied to other van der Waals systems.

\section{ACKNOWLEDGMENTS}

This research was supported by the MATRENA Doctoral Programme and Academy of Finland (Contract No. 1260204), by an Énergies Durables Research Grant from the École Polytechnique, the École Polytechnique Foundation, and the EDF Foundation, by a Marie Curie FP7 Integration Grant within the 7th European Union.
is given by the wave functions basis for both excitonic and single-particle Hamiltonians complete basis set for the representation of the bulk wave with energy \( E \). Choosing the wave vector \( \mathbf{k} \) where \( n \) related by:

\[
\phi_{nk,k_2}(z - R) \text{ (i here denotes the layer in the unit cell and } \mathbf{R} \text{ the lattice vector along } z).
\]

For a system characterized by two layers per unit cell, \( \phi_{nk}^1(\mathbf{R})\chi_n^k(z - R) \) is a set of \( 2N \) degenerate states \( N \) is the number of unit cells) corresponding to the eigenvalues \( E_n^1(\mathbf{R}) \) of \( \hat{H}_L(\mathbf{R}, z) \) and represent a complete basis set for the bulk wave function. Moreover, in the case of hBN with the AB stacking, the two layers in the unit cell are rotated one with respect to the other by an angle \( \beta = 60^\circ \) and therefore also the corresponding 2D first Brillouin zones are rotated by the angle \( \beta \). For a given wave vector \( \mathbf{k} \) the in-plane components of the electronic wave functions associated to two inequivalent layers are related by:

\[
\phi^2_{nk}(\mathbf{R}) = \phi^1_{n\beta k}(\mathbf{R}) \tag{A1}
\]

where \( \beta \mathbf{k} \) is the wave vector obtained rotating \( \mathbf{k} \) by an angle \( \beta \). Similarly for the corresponding eigenvalues one has:

\[
E^2_n(\mathbf{R}) = E^1_n(\beta \mathbf{k}). \tag{A2}
\]

Choosing the wave vector \( \mathbf{k} \) in the first Brillouin zone of the reference layer \( i = 1 \), the single-layer basis set is split in two subsets of \( N \) wave functions \( \phi_{nk}^1(\mathbf{R})\chi_n^k(z - R) \) and with energy \( E^1_n(\mathbf{R}) \) and \( \phi_{n\beta^{-1}k}^1(\mathbf{R})\chi_n^k(z - R) \) with energy \( E^2_n(\beta^{-1}\mathbf{k}) \). The ensemble of the two subsets represents a complete basis set for the representation of the bulk wave functions. In a more compact notation, the single-layer basis for both excitonic and single-particle Hamiltonians is given by the wave functions \( \phi_{nk,i}(\mathbf{R})\chi_n^k(z - R) \) with \( \mathbf{k}^{(i)} = \mathbf{k} \) for \( i = 1 \) and \( \mathbf{k}^{(i)} = \beta^{-1}\mathbf{k} \) for \( i = 2 \).

The single-particle Hamiltonian (written in second quantisation) hence takes the form:

\[
\hat{H} = \sum_{n\mathbf{k} \mathbf{k}'} \sum_{ij} \sum_{RS} F_{n\mathbf{k} Sj}(\mathbf{k}, \mathbf{k}') a_{n\mathbf{k} RS}^\dagger a_{n\mathbf{k} Sj} \tag{A3}
\]

where \( F_{n\mathbf{k} Sj}(\mathbf{k}, \mathbf{k}') \) are the matrix elements of \( \hat{H}_L(\mathbf{R}, z) + \delta U(\mathbf{z}) \) and are given by the expression:

\[
F_{n\mathbf{k} Sj}(\mathbf{k}, \mathbf{k}') = E^1_n(\mathbf{k})\delta_{RS} \delta_{n\mathbf{k}'} + i^{nk}\mathbf{k}' \tag{A4}
\]

with \( i^{nk}\mathbf{k}' \) denoting the effective interlayer hopping:

\[
i^{nk}\mathbf{k}' = \int d\rho \phi^2_{n\mathbf{k}}(\rho) \phi^1_{n\mathbf{k}'}(\rho) \int dz \chi_n^1(z - \mathbf{R}) \delta U(z) \chi_n^2(z - \mathbf{S}). \tag{A5}
\]

Defining

\[
t_{\mathbf{R} \mathbf{S} j}^{nk} = \int dz \chi_n^1(z - \mathbf{R}) \delta U(z) \chi_n^2(z - \mathbf{S}), \tag{A6}
\]

we have:

\[
i^{nk}\mathbf{k}' = t_{\mathbf{R} \mathbf{S} j}^{nk} \delta_{RS} \delta_{n\mathbf{k}'} \text{ for } i = j \text{ and } t_{\mathbf{R} \mathbf{S} j}^{nk}\mathbf{k}' = t_{\mathbf{R} \mathbf{S} j}^{n\beta^{-1}k} \text{ for } i \neq j.
\]

We note that in the present case the hopping \( t_{\mathbf{R} \mathbf{S} j}^{nk}\mathbf{k}' \) is not diagonal in \( \mathbf{k} \) and in this way the single-particle energies \( E^1_{n\mathbf{k} Sj}(\mathbf{k}, \mathbf{k}') \) acquire a dependence on both \( \mathbf{k} \) and \( \mathbf{k}' \).

We consider a two-band system \((n = c, v)\) and we take into account only the interlayer hopping between first nearest-neighbour layers \((i \neq j)\). In this case the hopping operators acting on electrons and holes are given by the following expressions:

\[
\hat{T}^c = \sum_{\mathbf{R}, \mathbf{S}, k} t^{c} [a_{c\mathbf{k}\mathbf{R}1} a_{c\beta-1\mathbf{k}\mathbf{R}-\mathbf{R} - \mathbf{\tau}} + a_{c\beta-1\mathbf{k}\mathbf{R}2}^\dagger a_{c\mathbf{k}\mathbf{R}1}] \tag{A7}
\]

\[
\hat{T}^v = \sum_{\mathbf{R}, \mathbf{S}, k} t^{v} [b_{v\mathbf{k}\mathbf{R}1} b_{v\beta-1\mathbf{k}\mathbf{R} - \mathbf{R} - \mathbf{\tau}} + b_{v\beta-1\mathbf{k}\mathbf{R}2}^\dagger b_{v\mathbf{k}\mathbf{R}1}] \tag{A8}
\]

where \( \mathbf{\tau} \) is the smallest lattice vector \((0, 0, 1)\) and \( t^{c(v)} = t^{c(v)}_{\mathbf{R}\mathbf{R} - \mathbf{\tau}} \). The effect of the hopping is to induce a dispersion along the \( z \) axis in reciprocal space and a splitting of the single-layer bands without modifying their in-plane dispersion. This is a consequence of the decoupling approximation between in-plane and out-of-plane coordinates. It is justified by the fact that in hBN the excitons originate from a limited area in the Brillouin zone, so that we can assume that the \( k_z \) dispersion in the single-particle band structure is constant for all the relevant \( \mathbf{k} \) points.

First of all, we neglect the hopping terms in such a way that the charge-transfer and Frenkel excitons are decoupled [see Eq. (13)]. We analyse here the interlayer charge-transfer exciton state, where the electron and the hole are localized on different layers. The charge-transfer wave function for the exciton state \( \lambda \) is

\[
\Psi^\lambda_{ij,\mathbf{R} + \mathbf{\tau}}(\mathbf{q}) = \frac{1}{\sqrt{N}} \sum_{\mathbf{R}} \Psi^\lambda_{ij,\mathbf{R} + \mathbf{\tau}}(\mathbf{q}) \tag{A9}
\]

with

\[
\Psi^\lambda_{ij,\mathbf{R} + \mathbf{\tau}}(\mathbf{q}) = \sum_{\mathbf{k}} A^\lambda_{\mathbf{c},\mathbf{k}}(\mathbf{q}) a_{c\mathbf{k}i\mathbf{R}1}^\dagger a_{c\mathbf{k}j\mathbf{R}1} b_{v\mathbf{k}i\mathbf{R}1}^\dagger b_{v\mathbf{k}j\mathbf{R}1} \tag{A10}
\]

where \( \mathbf{k}^{(i)} = \mathbf{k} \) for \( i = 1 \) and \( \mathbf{k}^{(i)} = \beta^{-1}\mathbf{k} \) for \( i = 2 \) (the same applies for the wave vector \( \mathbf{q} \)), while the coefficients \( A^\lambda_{\mathbf{c},\mathbf{k}}(\mathbf{q}) \) satisfy the excitonic eigenvalue equation:
Comparing Eq. (A14) and Eq. (A13) we see that, being $i$ identifies the index of the layer where the electron of the CT e-h pair is located, while $j$ the layer of the corresponding hole; $\tau$ defines the lattice-vector separation along z of the two unit cells to which the layers $i$ and $j$ belong. In the following we will focus on the first nearest-neighbour CT states for which $i \neq j$ and $\tau = (0, 0, 0)$ [for the other first nearest-neighbour CT state $\tau$ would be $(0, 0, -1)$]. In this case we have two possible configurations for the e-h pair: $i=1$ and $j=2$ or $i=2$ and $j=1$. They are described respectively by the equations:

\[
\begin{align*}
[E_c(k) - E_v(\beta^{-1}k + \beta^{-1}q)]A^\lambda_{vc,k}^{\lambda,12}(q) - &\sum_{k'} W^{R_i+\tau_j R_i}(v k + q^{(j)} c k^{(j)} + q^{(j)} c k^{(j)}) A^\lambda_{vc,k'}^{\lambda,12}(q) = E_{ij}^\lambda(q) A^\lambda_{vc,k}^{\lambda,12}(q) \\
[E_c(\beta^{-1}k) - E_v(k + q)]A^\lambda_{vc,k}^{\lambda,21}(q) - &\sum_{k'} W(v k + q c^{(i)} v k' + q c^{(i)} k') A^\lambda_{vc,k'}^{\lambda,21}(q) = E_{ij}^\lambda(q) A^\lambda_{vc,k}^{\lambda,21}(q)
\end{align*}
\]  

(A12)  

(A13)

where we have dropped the indices $i,j$ since the functional form of both the single-particle energies $E_v$ and $E_c$ and the interlayer effective electron-hole interaction $W$ is invariant under the exchange of the layer index. By applying the rotation $\beta$ to the $k$ space, Eq. (A12) becomes:

\[
\begin{align*}
[E_c(\beta k) - E_v(k + q)]A^\lambda_{vc,\beta k}^{\lambda,12}(\beta q) - &\sum_{k'} W(v k + q c^{(i)} v k' + q c^{(i)} k') A^\lambda_{vc,\beta k'}^{\lambda,12}(\beta q) = E_{ij}^\lambda(\beta q) A^\lambda_{vc,\beta k}^{\lambda,12}(\beta q).
\end{align*}
\]

(A14)

Comparing Eq. (A12) and Eq. (A13), we see that, being $E_v(\beta k) = E_v(k)$ (the layer is invariant under rotation of $\pm60^\circ$), the Hamiltonian in Eq. (A12) is the same as in Eq. (A13). This results in the following property for the energies $E^\lambda$ and coefficients $A^\lambda$ of the CT excitonic state:

\[
\begin{align*}
E_{ij}^\lambda(q) &= E_{ij}^\lambda(\beta q) \\
A^\lambda_{vc,k}^{\lambda,12}(q) &= A^\lambda_{vc,\beta k}^{\lambda,12}(\beta q)
\end{align*}
\]

(A15)  

(A16)

We now analyse the intralayer Frenkel exciton. In this case the excitonic state is

\[
\Psi(q) = \frac{1}{\sqrt{N}} \sum_{\lambda,R_i} c_{\lambda}^\lambda(q) \Psi_{R_i,(q)},
\]

(A17)

where

\[
\Psi_{R_i,(q)} = \sum_{k} A^\lambda_{vc,k}^{\lambda,i}(q) a^\dagger_{ek^{(i)}} R_i b^\dagger_{ek^{(i)} + q^{(i)} R_i}(0).
\]

(A18)

The electron and the hole of the excitonic pair in this case both belong to the same layer $i$. The coefficients $A^\lambda_{vc,k}^{\lambda,i}(q)$ satisfy the following excitonic eigenvalue equation:

\[
\begin{align*}
[E_c(k) - E_v(k + q)]A^\lambda_{vc,k}^{\lambda,1}(q) + &\sum_{k'} \left[6(v k + q^{(i)} c k^{(i)} + q^{(i)} c k^{(i)}) - W(v k + q^{(i)} c k^{(i)} + q^{(i)} c k^{(i)})\right] A^\lambda_{vc,k'}^{\lambda,1}(q) = E_{1}^\lambda(q) A^\lambda_{vc,k}^{\lambda,1}(q).
\end{align*}
\]

(A19)

Writing explicitly the eigenvalue equations for the $i = 1$ and $i = 2$ configurations, we have respectively:

\[
\begin{align*}
[E_c(k) - E_v(k + q)]A^\lambda_{vc,k}^{\lambda,1}(q) + &\sum_{k'} \left[6(v k + q^{(i)} c k^{(i)} + q^{(i)} c k^{(i)}) - W(v k + q^{(i)} c k^{(i)} + q^{(i)} c k^{(i)})\right] A^\lambda_{vc,k'}^{\lambda,1}(q) = E_{1}^\lambda(q) A^\lambda_{vc,k}^{\lambda,1}(q)
\end{align*}
\]

(A20)


\[
\begin{align*}
[ E_v(\beta^{-1}k) - E_v(\beta^{-1}k + \beta^{-1}q)] A^\lambda_{\gamma\epsilon,\delta k}(q) + \sum_{k'} [\vec{v}(v\beta^{-1}k + \beta^{-1}q) E_v(q) \beta^{-1}k + \beta^{-1}q) - \beta^{-1}q)] W(v\beta^{-1}k + \beta^{-1}q) E_v(q) - \beta^{-1}q) k' - \beta^{-1}q) k'] - A^\lambda_{\gamma\epsilon,\delta k}(q) = E^\lambda_2(q) A^\lambda_{\gamma\epsilon,\delta k}(q)
\end{align*}
\]

following what we have done for the CT exciton, we apply a rotation \( \beta \) to the whole \( k \) space in Eq. (A21):

\[
[ E_v(k) - E_v(k + q)] A^\lambda_{\gamma\epsilon,\delta k}(\beta q) + \sum_{k'} [\vec{v}(v k + q) v q k' + q k') - W(v k + q) v q k' q - \beta^{-1} q) k' - \beta^{-1} q) k' A^\lambda_{\gamma\epsilon,\delta k}(\beta q) = E^\lambda_2(q) A^\lambda_{\gamma\epsilon,\delta k}(\beta q).
\]

(A22)

Comparing Eq. (A22) with Eq. (A20) we find that in analogy with the CT state, for the FR exciton the following properties hold:

\[
\begin{align*}
E^\lambda_2(\beta q) &= E^\lambda_1(q) \quad \text{(A23)} \\
A^\lambda_{\gamma\epsilon,\delta k}(\beta q) &= A^\lambda_{\gamma\epsilon,\delta k}(q). \quad \text{(A24)}
\end{align*}
\]

Finally, we discuss the effect of the hopping, which enters the excitonic hamiltonian through the operator \( \hat{T} = \hat{T}_c - \hat{T}_v \), coupling intralayer Frenkel and interlayer charge-transfer states [see Eq. (13)]. In particular this coupling is given by the matrix elements of the hopping operator \( \hat{T} \) between \( |CT^\lambda\rangle \) and \( |FR^\lambda\rangle \) states. We have the following possibilities:

\[
\langle CT^\lambda_+ | \hat{T} | FR^\lambda_+ \rangle = \frac{t^c}{2} \left[ \sum_k A^\lambda_{\gamma\epsilon,\delta k}(q) A^\lambda_{\gamma\epsilon,\delta k}(q) + \sum_k A^\lambda_{\gamma\epsilon,\delta k}(q) A^\lambda_{\gamma\epsilon,\delta k}(q) \right]
\]

(A25)

\[
\langle CT^\lambda_+ | \hat{T} | FR^\lambda_+ \rangle = \frac{t^c}{2} \left[ \sum_k A^\lambda_{\gamma\epsilon,\delta k}(q) A^\lambda_{\gamma\epsilon,\delta k}(q) - \sum_k A^\lambda_{\gamma\epsilon,\delta k}(q) A^\lambda_{\gamma\epsilon,\delta k}(q) \right]
\]

(A26)

\[
\langle CT^\lambda_+ | \hat{T} | FR^\lambda_+ \rangle = \frac{t^c}{2} \left[ \sum_k A^\lambda_{\gamma\epsilon,\delta k}(q) A^\lambda_{\gamma\epsilon,\delta k}(q) - \sum_k A^\lambda_{\gamma\epsilon,\delta k}(q) A^\lambda_{\gamma\epsilon,\delta k}(q) \right]
\]

(A27)

\[
\langle CT^\lambda_+ | \hat{T} | FR^\lambda_+ \rangle = \frac{t^c}{2} \left[ \sum_k A^\lambda_{\gamma\epsilon,\delta k}(q) A^\lambda_{\gamma\epsilon,\delta k}(q) + \sum_k A^\lambda_{\gamma\epsilon,\delta k}(q) A^\lambda_{\gamma\epsilon,\delta k}(q) \right]
\]

(A28)

Using the properties of the excitonic coefficients \( A^\lambda \) from Eq. (A16) for the CT state and from Eq. (A24) for the FR state, the previous relations become:

\[
\langle CT^\lambda_+ | \hat{T} | FR^\lambda_+ \rangle = \frac{t^c}{2} \left[ \sum_k A^\lambda_{\gamma\epsilon,\delta k}(q) A^\lambda_{\gamma\epsilon,\delta k}(q) + \sum_k A^\lambda_{\gamma\epsilon,\delta k}(q) A^\lambda_{\gamma\epsilon,\delta k}(q) \right]
\]

(A29)
\[
\langle CT^\lambda|\hat{T}|FR^\lambda_+\rangle = \frac{t^c}{2} \sum_k A^\lambda_{vc,k}(q)A^{\lambda,12*}_{vc,k}(q) - \sum_k A^\lambda_{vc,k}(\beta q)A^{\lambda,21*}_{vc,k}(\beta q) \\
+ \frac{t^v}{2} \sum_k A^\lambda_{vc,k}(q)A^{\lambda,12*}_{vc,k}(q) - \sum_k A^\lambda_{vc,k}(\beta q)A^{\lambda,12*}_{vc,k}(\beta q) \\
\]
\[
\langle CT^\lambda|\hat{T}|FR^\lambda_-\rangle = \frac{t^c}{2} \sum_k A^\lambda_{vc,k}(q)A^{\lambda,12*}_{vc,k}(q) - \sum_k A^\lambda_{vc,k}(\beta q)A^{\lambda,21*}_{vc,k}(\beta q) \\
- \frac{t^v}{2} \sum_k A^\lambda_{vc,k}(q)A^{\lambda,12*}_{vc,k}(q) - \sum_k A^\lambda_{vc,k}(\beta q)A^{\lambda,12*}_{vc,k}(\beta q) \\
\]
\[
\langle CT^\lambda|\hat{T}|FR^\lambda_\beta\rangle = \frac{t^c}{2} \sum_k A^\lambda_{vc,k}(q)A^{\lambda,12*}_{vc,k}(q) + \sum_k A^\lambda_{vc,k}(\beta q)A^{\lambda,21*}_{vc,k}(\beta q) \\
+ \frac{t^v}{2} \sum_k A^\lambda_{vc,k}(q)A^{\lambda,12*}_{vc,k}(q) + \sum_k A^\lambda_{vc,k}(\beta q)A^{\lambda,12*}_{vc,k}(\beta q) .
\]

We can thus conclude that at \(q = 0\) the first and second term for each row of Eq. (A30) and Eq. (A31) cancel each other so that \(\langle CT^\lambda|\hat{T}|FR^\lambda_+\rangle = \langle CT^\lambda|\hat{T}|FR^\lambda_-\rangle = 0\). This means that at \(q = 0\) the hopping couples only Frenkel and charge-transfer states of the same parity. As a consequence, also in presence of the hopping the parity of the excitonic states remains a good quantum number. Instead, at finite \(q\) there is no more exact cancellation and a mixing between symmetric and antisymmetric states occurs. The parity is no more a good quantum number.

**Appendix B: Single-particle band structure**

Fig. 8 shows the single-particle band structures calculated within the GWA for bulk hBN (interlayer distance \(d_0\)) and for increased interlayer distance \(d = 1.5d_0\).

---

1. M. S. Dresselhaus, G. Dresselhaus, and P. C. Eklund, Science of Fullerenes and Carbon Nanotubes: Their Properties and Applications (Academic Press, 1996).
2. M. Dresselhaus, R. Smalley, G. Dresselhaus, and P. Avouris, Carbon Nanotubes: Synthesis, Structure, Properties, and Applications, Topics in Applied Physics (Springer Berlin Heidelberg, 2003).
3. P. Moriarty, Reports on Progress in Physics 64, 297 (2001).
4. V. I. Klimov, ed., Semiconductor and Metal Nanocrystals: Synthesis and Electronic and Optical Properties (CRC Press, 2003).
5. K. S. Novoselov, D. Jiang, F. Schedin, T. J. Booth, V. V. Khotkevich, S. V. Morozov, and A. K. Geim, Proceedings of the National Academy of Sciences of the United States of America 102, 10451 (2005).
6. S. Z. Butler, S. M. Hollen, L. Cao, Y. Cui, J. A. Gupta, H. R. Gutierrez, T. F. Heinz, S. S. Hong, J. Huang, A. F. Ismach, E. Johnston-Halperin, M. Kuno, V. V. Flashnitsa, R. D. Robinson, R. S. Ruoff, S. Salahuddin, J. Shan, L. Shi, M. G. Spencer, M. Terrones, W. Windl, and J. E. Goldberger, ACS Nano 7, 2898 (2013).
7. G. R. Bhimanapati, Z. Lin, V. Meunier, Y. Jung, J. Cha, S. Das, D. Xiao, Y. Son, M. S. Strano, V. R. Cooper, L. Liang, S. G. Louie, E. Ringe, W. Zhou, S. S. Kim, R. R. Naik, B. G. Sumpter, H. Terrones, F. Xia, Y. Wang, J. Zhu, D. Akinwande, N. Alem, J. A. Schuller, R. E. Schaak, M. Terrones, and J. A. Robinson, ACS Nano 9, 11509 (2015).
8. A. K. Geim and I. V. Grigorieva, Nature 499, 419 (2013).
9. K. S. Novoselov, A. Mishchenko, A. Carvalho, and A. H. Castro Neto, Science 353 (2016), 10.1126/science.aac9439.
10. Q. H. Wang, K. Kalantar-Zadeh, A. Kis, J. N. Coleman, and M. S. Strano, Nat Nano 7, 699 (2012).
11. F. Xia, H. Wang, D. Xiao, M. Dubey, and A. Ramanathan, Nat Photon 8, 899 (2014).
12. P. Cudazzo, I. V. Tokatly, and A. Rubio, Phys. Rev. B 84, 085406 (2011).
13. W. Hanke and L. J. Sham, Phys. Rev. Lett. 43, 387 (1979).
14. Semiconductor and Metal Nanocrystals: Synthesis and Electronic and Optical Properties (CRC Press, 2003).
15. L. Hedlin, Phys. Rev. 139, A796 (1965).
16. R. M. Martin, L. Reining, and D. M. Ceperley, Interacting Electrons: Theory and Computational Approaches (Cambridge University Press, 2016).
17. G. Onida, L. Reining, R. W. Godby, R. Del Sole, and W. Andreoni, Phys. Rev. Lett. 75, 818 (1995).
18. S. Albrecht, L. Reining, R. Del Sole, and G. Onida, Phys. Rev. Lett. 80, 4510 (1998).
19. L. X. Benedict, E. L. Shirley, and R. B. Bohn, Phys. Rev. Lett. 80, 4514 (1998).
FIG. 8. GW electronic band structure for hBN with 1.0 $d_0$ and for 1.5 $d_0$.
e-prints (2016), arXiv:1605.09581 [cond-mat.mtrl-sci].

59 H. Yu, Y. Wang, Q. Tong, X. Xu, and W. Yao, Phys. Rev. Lett. 115, 187002 (2015).

60 P. Rivera, K. L. Seyler, H. Yu, J. R. Schaibley, J. Yan, D. G. Mandrus, W. Yao, and X. Xu,