

Dynamical linear response of TDDFT with LDA+*U* functional: Strongly hybridized Frenkel excitons in NiO

Chi-Cheng Lee (李啟正),^{1,2} H. C. Hsueh (薛宏中),² and Wei Ku (顧威)^{1,3}

¹*Condensed Matter Physics and Materials Science Department, Brookhaven National Laboratory, Upton, New York 11973, USA*

²*Department of Physics, Tamkang University, Tamsui, Taiwan 25137, Republic of China*

³*Physics Department, State University of New York, Stony Brook, New York 11790, USA*

(Received 5 August 2010; published 30 August 2010)

Within the framework of time-dependent density-functional theory (TDDFT), we derive the dynamical linear response of local-density approximation plus *U* functional and benchmark it on NiO, a prototypical Mott insulator. Formulated using real-space Wannier functions, our computationally inexpensive framework gives detailed insights into the formation of tightly bound Frenkel excitons with reasonable accuracy. Specifically, a strong hybridization of multiple excitons is found to significantly modify the exciton properties. Furthermore, our study exposes a significant generic limitation of adiabatic approximation in TDDFT with hybrid functionals and in existing Bethe-Salpeter-equation approaches, advocating the necessity of strongly energy-dependent kernels in future development.

DOI: [10.1103/PhysRevB.82.081106](https://doi.org/10.1103/PhysRevB.82.081106)

PACS number(s): 71.15.Qe, 71.27.+a, 71.35.-y, 78.20.Bh

Computing many-body excitations of weakly to strongly interacting materials continues to be a major challenge for first-principles studies.¹ Following the great achievement of density-functional theory (DFT) in ground-state calculations using the local-density approximation (LDA),^{2,3} time-dependent DFT (TDDFT) (Refs. 4 and 5) has promised to be an affordable and accurate theoretical framework to study the dynamical linear response of weakly interacting materials ranging from finite⁶⁻⁸ to extended systems.^{1,9-11} While TD-DFT is formally exact in describing the time-dependent density, in practice the lack of knowledge of the proper analytical form of the action functional highlights the immaturity of its full potential in studying dynamics of real materials. For example, there is not yet an affordable first-principles local functional that includes proper particle-hole interactions at the two-particle level to allow the exciton formation, leaving the computationally expensive perturbation theory^{1,12} (assuming small enough interactions) or the highly parametrized cluster model¹³ the only option. Due to this well-known limitation, the applicability of TDDFT (with existing approximation) to strongly correlated materials remains basically unexplored.

Recently, a significant improvement in the ground-state calculations of strongly interacting Mott insulators is made via the new LDA+*U* functional,^{14,15} in which strong intra-atomic Coulomb interactions are introduced at the screened Hartree-Fock level. Similar to the hybrid functionals,¹⁶ the inclusion of nonlocal exchange allows opening of the Mott gap and gives also reasonable quasiparticle excitation energies. (Here we do not limit TDDFT or DFT to the Kohn-Sham framework, and we consider LDA+*U* and other hybrid functionals as implicit functionals of density within DFT.) This physically motivated approach has led to important understanding of orbital- and charge-ordered systems.¹⁷⁻²⁰ On the other hand, to date the study of charge excitation within LDA+*U* scheme has been very limited and its applicability to strongly interacting Mott insulators remains unclear. It is thus interesting and timely to explore the dynamical linear response of LDA+*U* functional within the framework of TD-DFT.

In this Rapid Communication, we examine the strength and weakness of LDA+*U* in describing charge excitations of strongly interacting Mott insulators, by developing diagrammatically the dynamical linear response of LDA+*U* functional within TDDFT framework (TDLDA+*U*). The resulting formula in tackling local excitations connects TDLDA+*U* (and other hybrid functionals) to a Bethe-Salpeter equation (BSE) (Ref. 21) with intra-atomic Hartree-Fock kernels. The framework is then implemented on the basis of symmetry-respecting Wannier functions (WFs),^{19,22} which not only dramatically reduce the computation expense but also facilitate a comprehensive real-space picture of local excitons. The integrated methodologies are applied to the study of tightly bound *d-d* Frenkel excitons in NiO, a representative Mott insulator. Our diagrammatic approach allows a step-by-step elucidation of the effects of different components of the interaction kernel. Specifically, multiple tightly bound (by ~ 6 eV) excitons form inside the Mott gap and are found to hybridize strongly to give reasonable excitation energies and highly anisotropic *q*-dependent spectral weights observed recently.²³ Furthermore, our results illustrate a serious general limitation of adiabatic approximation widely employed in most state-of-the-art TD-hybrid methods²⁴ and BSE approaches,²⁵⁻²⁷ and advocate the necessity of energy dependence in future design of approximations within these theoretical frameworks.

Following the standard procedure,²⁸ the linear response can be derived from the equation of motion (Dyson equation) of Kohn-Sham particle shown diagrammatically in Fig. 1.²⁹ In addition to LDA potentials, site-local (*s*-local) screened Hartree-Fock interactions and the double-counting terms among local orbitals are introduced in LDA+*U*. Taking the derivative of LDA+*U* Green's function with respect to the external potential, the response function χ ,

$$\chi(\mathbf{x}_1 t_1; \mathbf{x}_2 t_2) = \sum_{p_1 h_1 p_2 h_2} M_{p_1 h_1}^{*x_1} L_{p_1 h_2; h_1 p_2}(t_1 t_2; t_1 t_2) M_{p_2 h_2}^{x_2}, \quad (1)$$

is formulated by creating particle-hole (*p-h*) pairs at position \mathbf{x}_2 and time t_2 , with probability amplitude $M_{p_2 h_2}^{x_2}$

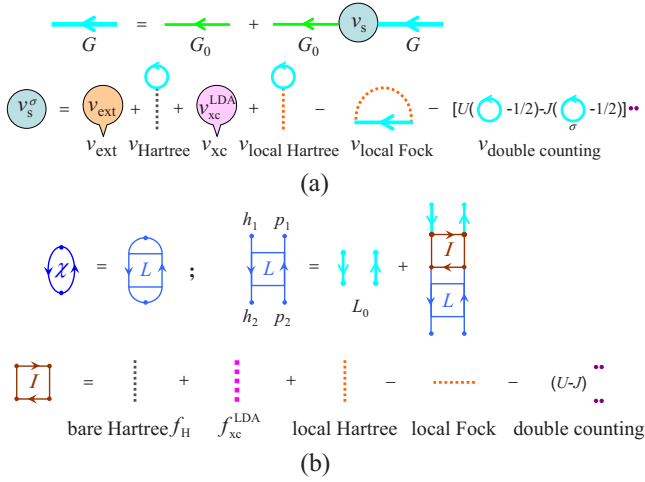


FIG. 1. (Color online) (a) Dyson equation with LDA+ U potential v_s . (b) Dynamical linear-response function χ , correlation function $L(p_1 h_2; h_1 p_2)$, and the corresponding BSE with LDA+ U kernel.

$\equiv \phi_{p_2}^*(\mathbf{x}_2) \phi_{h_2}(\mathbf{x}_2)$, followed by the propagation of the p - h pairs expressed in terms of the p - h correlation function L , and finally the annihilation of p - h pairs at \mathbf{x}_1 and a later time t_1 with probability amplitude $M_{p_1 h_1}^{**x_1}$ as shown in Fig. 1(b). The equation of motion of L (BSE) involves irreducible kernel I , including bare Hartree (f_{H}), exchange correlation f_{xc} , s-local screened Hartree-Fock, and double-counting terms. The s-local Fock interaction directly provides the attraction between p - h pairs of orbitals that U is applied to. For the study of local d - d excitations, the original f_{LDA} and the double-counting contributions are meant to counter each other approximately and should be neglected, leaving only the s-local screened Hartree-Fock contribution. (Besides, the long-range screening of f_{H} is inefficient at short distance.) This diagrammatic representation of the equation of motion makes it apparent that one is allowed to visualize the physical effects of interaction kernels by turning them on one after another, as presented below.

For the study of local excitations, it is most convenient and efficient to employ a real-space s-local basis.³⁰ Furthermore, it is advantageous to have orbitals that diagonalize the one-particle density matrix such that they enter only as either pure particle or hole orbitals. To this end, we implemented the above TDLDA+ U framework using the energy-resolved symmetry-respecting Wannier functions^{19,22} that are constructed without mixing the occupied and unoccupied bands.²⁹ In this basis, the complicated six-dimensional exciton wave function, $\psi_i(\mathbf{x}\mathbf{x}')$, can be decomposed into small number of “bare” exciton wave functions as direct products of one particle (ϕ_p) and one hole (ϕ_h) orbitals, $\psi_i(\mathbf{x}\mathbf{x}') = \sum_{ph} c_{ph}^i \phi_p(\mathbf{x}) \phi_h^*(\mathbf{x}')$. This gives an easy visualization of exciton wave functions and a direct computation of various experimental “form factor.” Moreover, the strong binding suppresses significantly the kinetics of the exciton such that only very few short-range neighbors are necessary in solving the BSE, significantly reducing the computational expense. In fact, even a single-site calculation is found quite accurate for NiO.²⁹

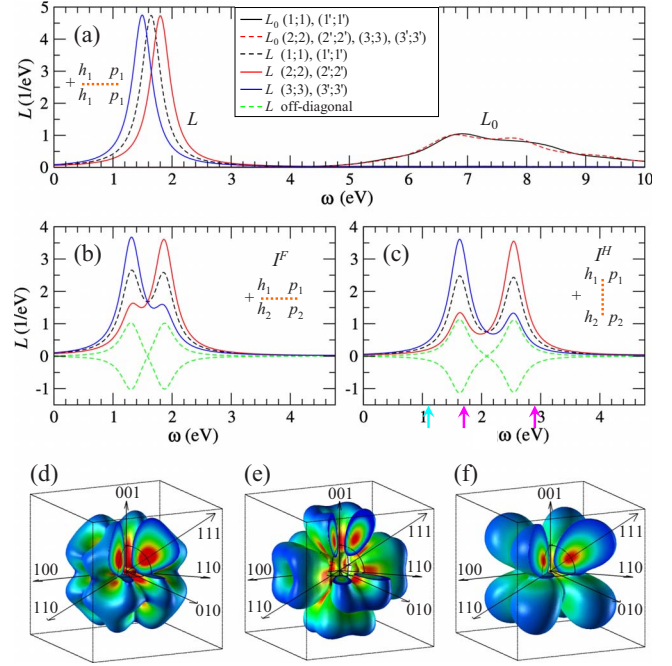


FIG. 2. (Color online) Imaginary parts of p - h correlation function L_{ij} denoted as $(i;j)$ by solving BSE for (a) L_0 : unbound p - h pair and L : bare exciton, (b) L with s-local Fock, and (c) L with s-local Hartree-Fock. Experimental excitation energies are indicated by the arrows. (d), (e), and (f) illustrate the spectral weight of L_0 , low-energy exciton in L , and high-energy exciton in L , respectively, via three-dimensional isovalue contours in \mathbf{q} space. The black solid circles indicate $q = 7 \text{ \AA}^{-1}$. Spectral weight in (e)/(f) resembles very well the experimental ones corresponding to the blue(leftmost)/red arrows.

The LDA+ U calculation for type-II antiferromagnetic (AFM) NiO ($U=8 \text{ eV}$, $J=0.95 \text{ eV}$) is performed by an all-electron full-potential method using linearized augmented plane-wave basis.³¹ It gives the correct ground state of NiO with high-spin configuration, leaving only the spin-minority channel active for s-local d - d charge excitations. The rhombohedral symmetry, dictated by the AFM order along $[111]$ direction, splits the d -Wannier orbitals into two unoccupied e_g' , two occupied e_g' , and one occupied a_g orbitals. In the rest of the Rapid Communication, we denote $1 \rightarrow (e_{g1}, a_g)$, $1' \rightarrow (e_{g2}, a_g)$, $2 \rightarrow (e_{g1}, e_{g1}')$, $2' \rightarrow (e_{g2}, e_{g2}')$, $3 \rightarrow (e_{g2}, e_{g1}')$, and $3' \rightarrow (e_{g1}, e_{g2}')$ for different p - h pairs (p, h) forming bare excitons. Within this notation, $\chi(\mathbf{q}, \omega)$ is thus $\sum_{ij} M_i^{*q} L_{ij}(\mathbf{q}, \omega) M_j^q$. Here $M_i^q \equiv \int e^{i\mathbf{q}\cdot\mathbf{x}} M_i^x d\mathbf{x}$ gives the probability amplitudes in momentum space corresponding to the anisotropic form factor of inelastic x-ray and electron scattering.^{23,32}

We now demonstrate in Fig. 2 the physical effects of the above interaction kernels on forming the low-energy Frenkel excitons in NiO, by switching them on one after another in the corresponding BSE presented in Fig. 1(b). The BSE is solved as a linear equation via the standard sparse iterative method.²⁹ (A broadening of $\eta=0.2 \text{ eV}$ is introduced in calculating L in accordance to the experimental resolution.²³) For clarity, only contributions that lead to the bound Frenkel excitons are shown. As a reference, the unbound p - h excita-

tions, L_0 , containing no interaction kernel, is shown in Fig. 2(a). L_0 gives basically the creations of p - h pairs across sites (intersite d - d excitations) that contribute at small momentum but are eventually overwhelmed by the on-site excitations at larger momentum. Not surprisingly, the excitation energies correspond to the energy difference of LDA+ U eigenvalues encapsulating a large Coulomb interaction (of order U) due to the addition of a particle and a hole at different atoms. In addition, a broad spectrum (5–10 eV) is obtained, reflecting the high mobility of such unbound p - h pairs and their strong coupling to the oxygen p orbitals.

For intrasite excitations, however, the interaction kernel can dramatically modify the excitations. As shown in Fig. 2(a), upon switching on the Fock kernel within the *same* pairs, which encapsulates the strong binding of p - h pairs, the excitations collapse into three well-defined doubly degenerate bare excitons at 1–2 eV via binding energies ~ 6 eV. This seemingly large binding energy is a necessary consequence of the large s -local Coulomb repulsion present in NiO and other strongly correlated materials. Preserving the particle number at each site, the s -local charge excitations should not be subject to the Hubbard- U energy scale of adding or removing one particle. On the other hand, since one-particle Green's function used to build L_0 includes the energy scale of U , such effects must be countered by a strong attraction of the same scale in any proper theory of p - h pairs, as demonstrated here. Benefiting from the lack of decay process deep inside the Mott gap, these bare excitons have very long lifetime (negligible line width beyond the experimental resolution). Due to the well-defined point-group symmetry of these Wannier orbitals (and thus M_i^q), the resulting bare excitons possess well-defined angular structure in their spectral weight. As shown in Fig. 2(d), a prominent anisotropy of χ is found with the strongest weight along [111] directions and vanishing weight along [001] directions, in agreement with the recent experiment.²³ In addition, the dipole-forbidden nature is revealed clearly by a hollow center at $q \rightarrow 0$.

Next, we activate the scattering process among different p - h pairs in the remaining part of the Fock kernel, I^F . These additional couplings between bare excitons introduce an interesting effect of strong hybridization and split the exciton energies into two sets. As illustrated in Fig. 2(b), this strong hybridization results in new large off-diagonal elements of L in the bare exciton basis and the double-peak structure in the diagonal elements. From the conventional eigenvector point of view, this is equivalent to having exciton eigenfunctions of each energy containing superposition of the above bare excitons. Indeed, the \mathbf{q} dependence of excitons for the low-energy and the high-energy peaks now shows significant differences, as shown in Figs. 2(e) and 2(f). Along [111] directions, the former has a vanishing weight (new nodal directions), while the latter shows strong enhancement, reflecting the antibonding and bonding nature of the superposition, respectively. Such strong hybridization of multiexcitons is absent in weakly correlated systems but should be expected from most strongly correlated systems with open shells.

Finally, upon addition of the s -local Hartree kernel, I^H , the overall exciton spectrum is further modified. As shown in Fig. 2(c), an overall blueshift of the spectrum is observed,

originated from the screening (weakening) of the exciton binding energy via the diagonal elements of I^H . In addition, without altering the \mathbf{q} -dependent spectral weight significantly, the off-diagonal elements of I^H further split the exciton energies. Both features are expected from the Kramers-Kronig relation in the typical dynamical screening process.

So far, the above analysis has been performed with Wannier functions associated with only one Ni site, which makes the proposed framework extremely inexpensive and allows a very clean real-space physical picture. More extensive calculations²⁹ including non- s -local (beyond one Ni site) effects resulted in a negligible energy reduction (<0.1 eV) in the overall spectrum, and, in perfect agreement with the experiment,²³ a negligible dispersion. Indeed, the Frenkel excitons in NiO are so tightly bound and so heavy to propagate that the physics can be captured quite completely even with only a single site. This further advocates strongly our choice of employing Wannier basis for strongly correlated materials in general.^{19,20,22,23}

Overall, compared with the recently measured local d - d excitations,^{23,32} the resulting theoretical spectrum of above TDLDA+ U performs unexpectedly well for such a simple approximation. In great contrast to the qualitative failure of TDLDA (Ref. 33) or the random-phase-approximation response of LDA+ U that give not even a hint of excitons, TDLDA+ U generates successfully the tightly bound Frenkel excitons deep inside the Mott gap with good energy scales at 1–3 eV. Considering the enormous binding energies of ~ 6 eV and the complexity of the multiexciton coupling, this degree of agreement is quite impressive, not to mention the good \mathbf{q} -dependent spectral weight that reflects the high quality of the exciton wave functions. Since the analytical structure of LDA+ U is the same as the hybrid functionals and its equation of motion contains similar kernel to that of existing GW-BSE (Refs. 25 and 26) approximation, the same degree of success should be expected from these approaches as well, so should above general physical picture of the formation of Frenkel excitons.

On the other hand, our study also reveals a generic limitation of TDLDA+ U and all these state-of-the-art approximations. While TDLDA+ U puts the excitons in the right energy range, it is unable to reproduce the fine structures of the experimental excitons satisfactorily. Indeed, while three charge excitation energies in the Mott gap were found experimentally (c.f. Fig. 2), TDLDA+ U only manages to produce two distinct energies. It turns out that the inability to split resulting peaks into the fine structures is dictated by the analytical structure of the approximations. This can be elucidated by reformulating rigorously the BSE into an effective Hamiltonian via the creation operator of bare excitons $|i\rangle$, b_i^\dagger ,

$$H_{eff} = \sum_i \epsilon_i b_i^\dagger b_i - \sum_{i \neq j} I_{ij}^F b_i^\dagger b_j + \sum_{ij} I_{ij}^H b_i^\dagger b_j, \quad (2)$$

where ϵ_i denotes the energy of the bare excitons in Fig. 2(a). In this representation, all the above physical effects are made transparent: the hybridization due to the off-diagonal elements of I^F and I^H , and the blueshift of the excitonic energy via the diagonal elements of I^H . This representation also proves that without terms that encapsulate interboson inter-

actions, e.g., $I_{ijj'j'}b_i^\dagger b_j^\dagger b_j b_{i'}$, this class of approximations is only capable of producing excitons in number equal to that of the bare excitons ($3 \times 2 = 6$ for NiO), subject to degeneracy under the point-group symmetry. Thus, this generic limitation can only be removed via higher-order kernel in BSE or new generations of hybrid-functional kernels that explicitly include time dependence.

This general conclusion is actually more stringent than those in the present TDDFT literature. Without ω -dependent I , our results demonstrate exciton formation with correct scale of large binding energy. Clearly, in this crudest level of approximation, the known necessity of nonadiabatic (memory-dependent) kernel f_{xc} of TDDFT in the Kohn-Sham framework^{34–36} would originate solely from the spatial reduction in nonlocal I_{xc} into f_{xc} .¹² On the other hand, to allow any fine (multiplets) structure in strongly interacting systems, a nonadiabatic kernel is absolutely necessary even without the spatial reduction. Obviously, this is one key aspect that almost all the existing approximate functionals lack and presents an essential and necessary step toward a proper description of local excitations in strongly interacting systems, within all the existing theoretical frameworks.

In summary, a TDLDA+ U method is derived within TD-DFT framework and implemented on the basis of Wannier functions. As a benchmark and an illustration, the formation of tightly bound excitons in NiO are analyzed step by step. A intriguing strong hybridization of multiple excitons is found that gives reasonable excitation energies with a large ~ 6 eV binding energy and the observed highly anisotropic spectral weight. Our computationally inexpensive approach not only provides detailed insights into the formation of Frenkel excitons but also demonstrates an intrinsic nonadiabaticity of the interaction kernel in allowing fine structures in the excitation spectra of strongly interacting systems. The lack of such essential feature in nowadays hybrid functional and BSE methods advocates the inclusion of explicit time dependence in future design of approximate functionals.

This work is supported by the U.S. Department of Energy (DOE), Office of Basic Energy Science, under Contract No. DE-AC02-98CH10886, and DOE CMSN. H. C. Hsueh thanks National Science Council (Grant No. NSC 99-2112-M-032-007) and NCTS of ROC for support, and also NCHC of ROC for CPU time.

-
- ¹G. Onida *et al.*, *Rev. Mod. Phys.* **74**, 601 (2002).
²P. Hohenberg and W. Kohn, *Phys. Rev.* **136**, B864 (1964).
³W. Kohn and L. J. Sham, *Phys. Rev.* **140**, A1133 (1965).
⁴E. Runge and E. K. U. Gross, *Phys. Rev. Lett.* **52**, 997 (1984).
⁵R. van Leeuwen, *Phys. Rev. Lett.* **82**, 3863 (1999).
⁶M. Petersilka, U. J. Gossmann, and E. K. U. Gross, *Phys. Rev. Lett.* **76**, 1212 (1996).
⁷I. Vasiliev, S. Ögüt, and J. R. Chelikowsky, *Phys. Rev. Lett.* **82**, 1919 (1999).
⁸K. Yabana and G. F. Bertsch, *Int. J. Quantum Chem.* **75**, 55 (1999).
⁹N. E. Maddocks *et al.*, *Europhys. Lett.* **27**, 681 (1994).
¹⁰W. Ku and A. G. Eguiluz, *Phys. Rev. Lett.* **82**, 2350 (1999).
¹¹W. Ku, W. E. Pickett, R. T. Scalettar, and A. G. Eguiluz, *Phys. Rev. Lett.* **88**, 057001 (2002).
¹²I. V. Tokatly and O. Pankratov, *Phys. Rev. Lett.* **86**, 2078 (2001).
¹³M. W. Haverkort, A. Tanaka, L. H. Tjeng, and G. A. Sawatzky, *Phys. Rev. Lett.* **99**, 257401 (2007).
¹⁴V. I. Anisimov, J. Zaanen, and O. K. Andersen, *Phys. Rev. B* **44**, 943 (1991).
¹⁵A. B. Shick, A. I. Liechtenstein, and W. E. Pickett, *Phys. Rev. B* **60**, 10763 (1999).
¹⁶F. Tran, P. Blaha, K. Schwarz, and P. Novák, *Phys. Rev. B* **74**, 155108 (2006).
¹⁷I. Leonov, A. N. Yaresko, V. N. Antonov, M. A. Korotin, and V. I. Anisimov, *Phys. Rev. Lett.* **93**, 146404 (2004).
¹⁸H.-T. Jeng, G. Y. Guo, and D. J. Huang, *Phys. Rev. Lett.* **93**, 156403 (2004).
¹⁹W.-G. Yin, D. Volja, and W. Ku, *Phys. Rev. Lett.* **96**, 116405 (2006).
²⁰D. Volja *et al.*, *EPL* **89**, 27008 (2010).
²¹E. E. Salpeter and H. A. Bethe, *Phys. Rev.* **84**, 1232 (1951).
²²W. Ku, H. Rosner, W. E. Pickett, and R. T. Scalettar, *Phys. Rev. Lett.* **89**, 167204 (2002).
²³B. C. Larson, W. Ku, J. Z. Tischler, C.-C. Lee, O. D. Restrepo, A. G. Eguiluz, P. Zschack, and K. D. Finkelstein, *Phys. Rev. Lett.* **99**, 026401 (2007).
²⁴J. Paier, M. Marsman, and G. Kresse, *Phys. Rev. B* **78**, 121201(R) (2008).
²⁵G. Onida, L. Reining, R. W. Godby, R. Del Sole, and W. Andreoni, *Phys. Rev. Lett.* **75**, 818 (1995).
²⁶M. Rohlfing and S. G. Louie, *Phys. Rev. Lett.* **81**, 2312 (1998).
²⁷L. X. Benedict, E. L. Shirley, and R. B. Bohn, *Phys. Rev. Lett.* **80**, 4514 (1998).
²⁸G. Baym and L. P. Kadanoff, *Phys. Rev.* **124**, 287 (1961).
²⁹See supplementary material at <http://link.aps.org/supplemental/10.1103/PhysRevB.82.081106> for the details of TDLDA+ U calculations and non-local effects on the Frenkel excitons in NiO.
³⁰J. A. Soininen, A. L. Ankudinov, and J. J. Rehr, *Phys. Rev. B* **72**, 045136 (2005).
³¹K. Schwarz *et al.*, *Comput. Phys. Commun.* **147**, 71 (2002).
³²F. Müller and S. Hufner, *Phys. Rev. B* **78**, 085438 (2008).
³³V. Turkowski, A. Leonardo, and C. A. Ullrich, *Phys. Rev. B* **79**, 233201 (2009).
³⁴Y.-H. Kim and A. Görling, *Phys. Rev. B* **66**, 035114 (2002).
³⁵M. Hellgren and U. von Barth, *Phys. Rev. B* **78**, 115107 (2008).
³⁶N. T. Maitra, K. Burke, and C. Woodward, *Phys. Rev. Lett.* **89**, 023002 (2002).