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Proximity Eliashberg theory of electrostatic field-effect doping in superconducting filmsG. A. Ummarino,^{1,2,*} E. Piatti,¹ D. Daghero,¹ R. S. Gonnelli,¹ Irina Yu. Sklyadneva,³ E. V. Chulkov,^{3,4,5,6} and R. Heid⁷¹*Dipartimento di Scienza Applicata e Tecnologia, Politecnico di Torino, Corso Duca degli Abruzzi 24, 10129 Torino, Italy*²*National Research Nuclear University MEPhI (Moscow Engineering Physics Institute), Kashira Hwy 31, Moskva 115409, Russia*³*Donostia International Physics Center (DIPC), 20018 San Sebastian/Donostia, Basque Country, Spain*⁴*St. Petersburg State University, 199034, St. Petersburg, Russian Federation*⁵*Departamento de Física de Materiales, Facultad de Ciencias Químicas, UPV/EHU, Apdo. 1072, 20080 San Sebastian/Donostia, Basque Country, Spain*⁶*Centro de Física de Materiales CFM - Materials Physics Center MPC, Centro Mixto CSIC-UPV/EHU, 20018 San Sebastian/Donostia, Basque Country, Spain*⁷*Karlsruher Institut für Technologie, Institut für Festkörperphysik, D-76021 Karlsruhe, Germany*

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We calculate the effect of a static electric field on the critical temperature of a *s*-wave one band superconductor in the framework of proximity effect Eliashberg theory. In the weak electrostatic field limit the theory has no free parameters while, in general, the only free parameter is the thickness of the surface layer where the electric field acts. We conclude that the best situation for increasing the critical temperature is to have a very thin film of a superconducting material with a strong increase of electron-phonon (boson) constant upon charging.

DOI: [10.1103/PhysRevB.96.064509](https://doi.org/10.1103/PhysRevB.96.064509)**I. INTRODUCTION**

In the last decade, electric double layer (EDL) gating has come to the forefront of solid state physics due to its capability to tune the surface carrier density of a wide range of different materials well beyond the limits imposed by solid-gate field-effect devices. The order-of-magnitude enhancement in the gate electric field allows this technique to reach doping levels comparable to those of standard chemical doping. This is possible due to the extremely large specific capacitance of the EDL that builds up at the interface between the electrolyte and the material under study [1–5].

EDL gating was first exploited to control the surface electronic properties of relatively low-carrier density systems, where the electric field effect is more readily observable. Field-induced superconductivity was first demonstrated in strontium titanate [6] and zirconium nitride chloride [7,8], and subsequently on other insulating systems such as perovskites [9] and layered transition-metal dichalcogenides [10–16]. Significant effort was also invested in the control of the superconducting properties of cuprates [17–25], although in this class of materials the mechanism behind the carrier density modulation is still debated [25].

More recently however, several experimental studies have been devoted to the exploration of field effect in superconductors [26] with a large ($\gtrsim 1 \times 10^{22} \text{ cm}^{-3}$) native carrier density. The interplay between two different ground states, namely superconductivity and charge density waves, was explored in titanium and niobium diselenides [27–29]. The thickness and gate voltage dependence of a high-temperature superconducting phase were studied in iron selenide, both in thin-film [30] and thin flake [31,32] forms. The effect of the ultrahigh interface electric fields achievable via EDL gating were also probed in standard BCS superconductors, namely niobium [33] and niobium nitride [34,35].

With the exception of the work of Ref. [29] on niobium diselenide, all of these studies have been performed on relatively thick samples ($\gtrsim 10 \text{ nm}$) with a thickness larger than the electrostatic screening length. These systems are thus expected to develop a strong dependence of their electronic properties along the *z* direction (*z* being perpendicular to the sample surface). As a first approximation, this dependence can be conceptualized by schematizing the system as the parallel of a surface layer (where the carrier density is modified by the electric field) and an unperturbed bulk. The two electronic systems can be expected to couple via superconducting proximity effect, resulting in a complicated response to the applied electric field that goes well beyond a simple modification of the superconducting properties of the surface layer alone [35] and is strongly dependent on both the electrostatic screening length and the total thickness of the film.

So far, the only quantitative assessment of this phenomenon has been reported in the framework of the strong-coupling limit of the BCS theory of superconductivity [35]. A proper theoretical treatment for field effect on more complex materials, which can be described only by means of the more complete Eliashberg theory [36,37], is lacking. Given the rising interest in the control of the properties of superconductors by means of surface electric fields, the development of such a theoretical approach would be very convenient not only to quantitatively describe the results of future experiments, but also to determine beforehand the experimental conditions (e.g., device thickness) most suitable for an optimal control of the superconducting order via electric fields.

In this work, we use the Eliashberg theory of proximity effect to describe a junction composed by the perturbed surface layer ($T_c = T_{c,s}$), where the carrier density is modulated (with a doping level per unitary cell *x*), and the underlying unperturbed bulk ($T_c = T_{c,b}$). Here *s* and *b* indicate “surface” and “bulk”, respectively (see Fig. 1). Under the application of an electric field, $T_{c,s} \neq T_{c,b}$ and the material behaves like a junction between a superconductor and a normal metal in the temperature range bounded by $T_{c,s}$ and $T_{c,b}$. If the application

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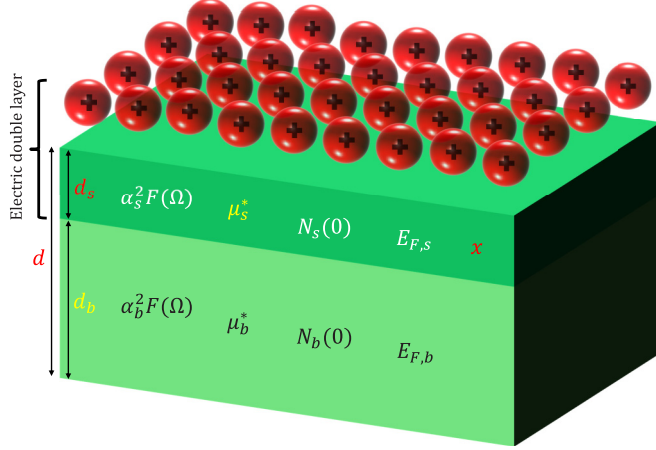


FIG. 1. Scheme of an EDL-gated superconducting thin film. The layer of adsorbed ions and the surface layer where the carrier density is perturbed (dark green region) compose the EDL. The unperturbed bulk of the film is indicated in light green color. For both layers, we indicate the relevant parameters of the proximity Eliashberg equations (see text for details). Parameters in red, black, and white indicate the free parameters of the model, data obtained from the literature, and the output of the DFT calculations, respectively. Parameters in yellow are obtained from these by simple calculations.

of the electric field increases (decreases) $T_{c,s}$, then the surface layer will be the superconductor (normal metal) and the bulk will be the normal metal (superconductor). We perform the calculation for Pb since all the input parameters of the theory are well known in the literature for this strong-coupling superconductor [36].

II. MODEL: PROXIMITY ELIASHBERG EQUATIONS

In general, a proximity effect at a superconductor/normal metal junction is observed as the opening of a finite superconducting gap in the normal metal together with its reduction in a thin region of the superconductor close to the junction. In our model we use the one band s -wave Eliashberg equations [36,37] with proximity effect to calculate the critical temperature of the system. In this case we have to solve four coupled equations for the gaps $\Delta_{s,b}(i\omega_n)$ and the renormalization functions $Z_{s,b}(i\omega_n)$, where ω_n are the Matsubara frequencies. The imaginary-axis equations with proximity effect [38–42] are

$$\omega_n Z_b(i\omega_n) = \omega_n + \pi T \sum_m \Lambda_b^Z(i\omega_n, i\omega_m) N_b^Z(i\omega_m) + \Gamma_b N_s^Z(i\omega_n), \quad (1)$$

$$Z_b(i\omega_n) \Delta_b(i\omega_n) = \pi T \sum_m [\Lambda_b^\Delta(i\omega_n, i\omega_m) - \mu_b^*(\omega_c)] \times \Theta(\omega_c - |\omega_m|) N_b^\Delta(i\omega_m) + \Gamma_b N_s^\Delta(i\omega_n), \quad (2)$$

$$\omega_n Z_s(i\omega_n) = \omega_n + \pi T \sum_m \Lambda_s^Z(i\omega_n, i\omega_m) N_s^Z(i\omega_m) + \Gamma_s N_b^Z(i\omega_n), \quad (3)$$

$$Z_s(i\omega_n) \Delta_s(i\omega_n) = \pi T \sum_m [\Lambda_s^\Delta(i\omega_n, i\omega_m) - \mu_s^*(\omega_c)] \times \Theta(\omega_c - |\omega_m|) N_s^\Delta(i\omega_m) + \Gamma_s N_b^\Delta(i\omega_n), \quad (4)$$

where $\mu_{s(b)}^*$ are the Coulomb pseudopotentials in the surface and in the bulk, respectively, Θ is the Heaviside function, and ω_c is a cutoff energy at least three times larger than the maximum phonon energy. Thus we have

$$\Lambda_{s(b)}^\Delta(i\omega_n, i\omega_m) = 2 \int_0^{+\infty} d\Omega \Omega \alpha_{s(b)}^2 F(\Omega) / [(\omega_n - \omega_m)^2 + \Omega^2], \quad (5)$$

$$\Gamma_{s(b)} = \pi |t|^2 A d_{b(s)} N_{b(s)}(0), \quad (6)$$

and thus $\frac{\Gamma_s}{\Gamma_b} = \frac{d_b N_b(0)}{d_s N_s(0)}$,

$$N_{s(b)}^\Delta(i\omega_m) = \Delta_{s(b)}(i\omega_m) / \sqrt{\omega_m^2 + \Delta_{s(b)}^2(i\omega_m)}, \quad (7)$$

and

$$N_{s(b)}^Z(i\omega_m) = \omega_m / \sqrt{\omega_m^2 + \Delta_{s(b)}^2(i\omega_m)}, \quad (8)$$

where $\alpha_{s(b)}^2 F(\Omega)$ are the electron-phonon spectral functions, A is the junction cross-sectional area, d_s and d_b are the surface and bulk layer thicknesses respectively, such that $d_s + d_b = d$ (where d is the total film thickness), and $N_{s(b)}(0)$ are the densities of states at the Fermi level $E_{F,s(b)}$ for the surface and bulk material. Since we only consider electrostatic perturbations to the system, we assume the transmission matrix $|t|^2 = 1$ as we expect the interface between the surface and bulk layers to be nearly ideal. This assumption is supported by our experimental findings on niobium nitride [35], where the experimental doping dependence of T_c turned out to be compatible with a high interface transparency.

The electron-phonon coupling constants are defined as

$$\lambda_{s(b)} = 2 \int_0^{+\infty} d\Omega \frac{\alpha_{s(b)}^2 F(\Omega)}{\Omega} \quad (9)$$

and the representative energies as

$$\ln(\omega_{ln,s(b)}) = \frac{2}{\lambda_{s(b)}} \int_0^{+\infty} d\Omega \ln \Omega \frac{\alpha_{s(b)}^2 F(\Omega)}{\Omega}. \quad (10)$$

The solution of these equations requires 11 input parameters: the two electron-phonon spectral functions $\alpha_{s(b)}^2 F(\Omega)$, the two Coulomb pseudopotentials $\mu_{s(b)}^*$, the values of the normal density of states at the Fermi level $N_{s(b)}(0)$, the shift of the Fermi energy $\Delta E_F = E_{F,s} - E_{F,b}$ that enters in the calculation of the surface Coulomb pseudopotential (as shown later), the thickness of the surface layer d_s , the total film thickness d , and the junction cross-sectional area A . The values of d and A are experimental data. The exact value of d_s , in particular in the case of very strong electric fields at the surface of a thin film, is in general difficult to be determined *a priori* [35]. Thus we leave it as a free parameter of the model, and we perform our calculations for different reasonable estimations of its value, as we discuss in detail in Sec. IV.

Typically, the bulk values of $\alpha_b^2 F(\Omega)$, μ_b^* , $N_b(0)$, and $E_{F,b}$ are known and can be found in the literature. Thus, we assume

that we need to determine only their surface values. In the next section we will use density functional theory (DFT) to calculate $\alpha_s^2 F(\Omega)$, ΔE_F , and $N_s(0)$.

The value of the Coulomb pseudopotential in the surface layer μ_s^* can be obtained in the following way: In the Thomas-Fermi theory, the dielectric function is [43] $\varepsilon(q) = 1 + \frac{k_{\text{TF}}^2}{q^2}$ and the bare Coulomb pseudopotential μ is the angular average of the ratio between the screened electrostatic potential $V(q)$ and $\varepsilon(q)$:

$$\mu = \frac{1}{4\pi^2 \hbar v_F} \int_0^{2k_F} \frac{V(q)}{\varepsilon(q)} q dq. \quad (11)$$

Since [44] $V(q) = \frac{4\pi e^2}{q^2}$ it turns out that

$$\mu = \frac{k_{\text{TF}}^2}{8k_F^2} \ln\left(1 + \frac{4k_F^2}{k_{\text{TF}}^2}\right). \quad (12)$$

Hence we write

$$\mu_b = \frac{a_b^2}{2} \ln\left(1 + \frac{1}{a_b^2}\right), \quad (13)$$

with $a_b = 2k_{\text{TF},b}/k_{F,b}$. Since a_b can be calculated by numerically solving Eq. (13), and by remembering that the square of Thomas-Fermi wave number $k_{\text{TF},b(s)}$ is proportional to $N_{b(s)}(0)$, we have

$$a_s^2 = a_b^2 \left(\frac{N_s(0)}{N_b(0)} \right) / \left(\frac{1 + \Delta E_F}{E_{F,b}} \right) \quad (14)$$

and thus

$$\mu_s = \frac{a_s^2}{2} \ln\left(1 + \frac{1}{a_s^2}\right). \quad (15)$$

The new Coulomb pseudopotential [44] in the surface layer is thus

$$\mu_s^*(\omega_c) = \frac{\mu_s}{1 + \mu_s \ln[(E_{F,b} + \Delta E_F)/\omega_c]}. \quad (16)$$

We note that, usually, the effect of electrostatic doping on μ^* is very small and can be neglected. We can quantify the effect on T_c of this small modulation of μ^* by computing it in the case of maximum doping $x = 0.40$ and very thin film ($d = 5$ nm), i.e., when the effect is the largest. As discussed in the next section, the unperturbed Coulomb pseudopotential is $\mu^*(x = 0) = 0.14164$, while for the maximum doping Eqs. (12)–(16) give $\mu^*(x = 0.4) = 0.14048$. If we use $d_s = d_{\text{TF}}$ we find, respectively, $T_c = 7.3770$ K for the bulk (unperturbed) value of the Coulomb pseudopotential and $T_c = 7.3768$ K for the surface value of the Coulomb pseudopotential. Thus, if we consider the Coulomb pseudopotential to be doping independent we underestimate the critical temperature of a $\Delta T_c|_{\Delta\mu^*} = -0.0002$ K ($\Delta T_c|_{\Delta\mu^*}/T_c = 0.0027\%$).

However, a possible critical situation can appear when the applied electric field is very strong and the Thomas-Fermi approximation does not hold anymore. In such a case, μ^* becomes ill-defined as the Thomas-Fermi dielectric function is no longer strictly valid for very large electric fields. Nevertheless, the true dielectric function $\varepsilon(q)$ should still be a function of the ratio k_{TF}/k_F [45], which—in the free-electron model—is independent of the normal density of states at the

Fermi level. Thus, Eq. (11) should still be able to describe the behavior of the system as a first approximation.

III. CALCULATION OF $\alpha_s^2 F(\Omega)$, ΔE_F , AND $N_s(0)$

DFT calculations are performed within the mixed-basis pseudopotential method (MBPP) [46]. For Pb a norm-conserving relativistic pseudopotential including $5d$ semicore states and partial core corrections is constructed following the prescription of Vanderbilt [47]. This provides both scalar-relativistic and spin-orbit components of the pseudopotential. Spin-orbit coupling (SOC) is then taken into account within each DFT self-consistency cycle (for more details on the SOC implementation see [48]). The MBPP approach utilizes a combination of local functions and plane waves for the basis set expansion of the valence states, which reduces the size of the basis set significantly. One d type local function is added at each Pb atomic site to efficiently treat the deep $5d$ potential. Sufficient convergence is then achieved with a cutoff energy of 20 Ry for the plane waves. The exchange correlation is treated in the local density approximation (LDA) [49]. Brillouin zone (BZ) integrations are performed on regular k -point meshes in conjunction with a Gaussian broadening of 0.2 eV. For phonons, $16 \times 16 \times 16$ meshes are used, while for the calculations of density of states and electron-phonon coupling (EPC) even denser $32 \times 32 \times 32$ meshes are employed.

Phonon properties are calculated with the density-functional perturbation theory [50,51] as implemented in the MBPP approach [52], which also provides direct access to the electron-phonon coupling matrix elements. The procedure to extract the Eliashberg function is outlined in Ref. [48]. Dynamical matrices and corresponding EPC matrix elements are obtained on a $16 \times 16 \times 16$ phonon mesh. These quantities are then interpolated using standard Fourier techniques to the whole BZ, and the Eliashberg functions are calculated by integration over the BZ using the tetrahedron method on a $40 \times 40 \times 40$ mesh. SOC is consistently taken into account in all calculations including lattice dynamical and EPC properties. It is well known from previous work that SOC is necessary for a correct quantitative description of both the phonon anomalies and EPC of undoped bulk Pb [48].

Charge induction is simulated by adding an appropriate number of electrons during the DFT self-consistency cycle, compensated by a homogeneous background charge to retain overall charge neutrality. Electronic structure properties and the Eliashberg function are calculated for face-centered cubic (fcc) Pb with the lattice constant $a = 4.89$ Å as obtained by optimization for the undoped case. For doping levels considered here, we found that to a good approximation charge induction does not change the band structure but merely results in a shift of E_F . In a previous study, the variation of the EPC was studied as function of the averaging energy [53]. The present approach goes beyond this analysis by taking into account explicitly the effect of charge induction on the screening properties, which modifies both the phonon spectrum and the EPC matrix elements.

Finally we point out that, since this DFT approach simulates the effect of the electric field by adding extra charge carriers to the system together with a uniform compensating countercharge (Jellium model [54]), it is unable to describe

inhomogeneous distributions caused by the screening of the electric field itself. A more complete approach has been developed in Ref. [55], and requires the self-consistent solution of the Poisson equation; however, this method is currently unable to compute the phonon spectrum of the gated material, making it unsuitable for the application to the proximity Eliashberg formalism.

IV. RESULTS AND DISCUSSION

We start our calculations by fixing the input parameters for bulk Pb according to the established literature. We set $T_{c,b}$ to its experimental value [36] $T_{c,b} = 7.22$ K. The undoped $\alpha_b^2 F(\Omega)$ gives a corresponding electron-phonon coupling $\lambda_b = 1.5596$. Assuming a cutoff energy $\omega_c = 60$ meV and a maximum energy $\omega_{\max} = 70$ meV in the Eliashberg equations, we are thus able to determine the bulk Coulomb pseudopotential to be $\mu_b^* = 0.14164$ to obtain the exact experimental critical temperature $T_{c,b}$.

In Fig. 2(a) we show the calculated electron-phonon spectral functions $\alpha^2 F(\Omega)$ resulting at the increase of the doping level x . Specifically, we plot the curves corresponding to $x = 0.000, 0.075, 0.150, 0.300, 0.400$ e^- /unit cell. We calculate the spectral functions up to $x = 0.4$ e^- /cell because for larger values of doping an instability emerges in the calculation processes. We can see the phonon softening evidenced by a reduction of ω_{in} with increasing doping level. The increase of the carrier density gives rise to two competing effects: the value of ω_{in} (i.e., the representative phonon energy) decreases while the value of electron-phonon coupling constant λ increases [see Fig. 2(b)]. In the Eliashberg theory, the critical temperature is an increasing function of both ω_{in} and λ . This can be easily visualized by the well-known, approximated Allen-Dynes formula [56]:

$$T_c = \frac{\omega_{\text{in}}}{1.2} \exp\left(\frac{1.04(1 + \lambda)}{\lambda - \mu^*(1 + 0.62\lambda)}\right), \quad (17)$$

which features a linearly increasing dependence on ω_{in} . Hence, this could result in either a net enhancement or suppression of T_c , depending on which of the two contributions is dominant. Consequently the ideal situation for obtaining the largest possible critical temperature in an electric field doped material is to have a strong increase of λ and ω_{in} concurrently. In the case of Pb the contribution from the increase of λ is dominant over that from the reduction of ω_{in} , giving rise to a net increase of the superconducting critical temperature [as we report in Fig. 2(c)]. In addition, in Table I we summarize all the input parameters of the proximity Eliashberg equations as obtained from the DFT calculations.

Having determined the response of the superconducting properties of a homogeneous lead film to a modulation of its carrier density, we can now consider the behavior of the more realistic junction between the perturbed surface layer and the unperturbed bulk. In order to do so, however, it is now mandatory to select a value for the thickness of the perturbed surface layer. Close to T_c , the superfluid density is small [57] and the screening is dominated by unpaired electrons. Thus, a very rough approximation would be to set d_s equal to the Thomas-Fermi screening length d_{TF} , which for lead can be estimated to be 0.15 nm [58]. However, we have recently

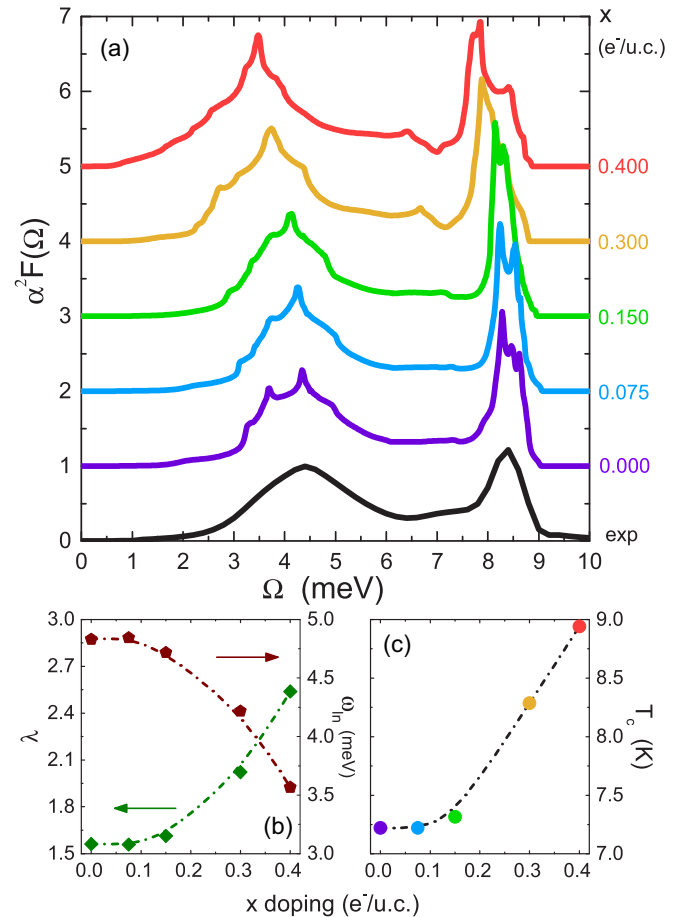


FIG. 2. (a) Calculated Pb electron-phonon spectral function for five different values of charge doping (electrons/unitary cell): 0.00 (violet solid line), 0.075 (blue solid line), 0.15 (green solid line), 0.30 (orange solid line), and 0.40 (red solid line). We also show the experimental electron-phonon spectral function determined via tunneling measurements [60] (black solid line). All curves are shifted by a constant offset equal to one. (b) Calculated values of electron-phonon coupling constants λ (green diamonds) and representative energies ω_{in} (brown pentagons) versus charge doping. (c) Calculated critical temperature versus charge doping for a system without proximity effect. Symbol colors correspond to those used in (a). All dash-dotted lines act as guides to the eye.

shown [35] that this assumption might not be satisfactory in the presence of the very large electric fields that build up in the electric double layer. Indeed, our experimental findings on niobium nitride indicated that the screening length increases for very large doping values [35]. However, it is reasonable to assume the exact entity of this increase to be specific to each material. Thus, while the qualitative behavior can be expected to be general, the exact values of d_s determined for niobium nitride cannot be directly applied to lead.

In order not to lose the generality of our approach, we calculate the behavior of our system for three different choices of the behavior of d_s . We start by expressing $d_s = d_{\text{TF}}[1 + m\Theta(x - x_0)]$, where m is a dimensionless parameter indicating how much d_s expands beyond the Thomas-Fermi value for large doping levels, and x_0 is the specific doping value at which this increase in d_s takes place. By selecting $x_0 = 0.2$, we

TABLE I. Input parameters calculated by DFT and T_c calculated by Eliashberg theory without proximity effect.

x (e^- /cell)	λ	ω_{in} (meV)	$N(0)$ states/(eV spin)	ΔE_F (meV)	μ^*	T_c (K)
0.000	1.5612	4.8431	0.25866	0.00	0.14164	7.2200
0.075	1.5582	4.8432	0.25754	108.42	0.14136	7.2197
0.150	1.6137	4.7176	0.25611	218.77	0.14116	7.3165
0.300	2.0237	4.2175	0.26770	435.07	0.14074	8.2862
0.400	2.5392	3.5668	0.27833	571.62	0.14048	8.9406

allow the upper half of our doping values to bring the system beyond the Thomas-Fermi approximation. We then perform proximity-coupled Eliashberg calculations for $m = 0, 1, 4$ and five different film thicknesses $d = 5, 10, 20, 30, 40$ nm, always assuming the junction area to be $A = 10^{-7}$ m². Note that the case $m = 0$ of course corresponds to the case where the

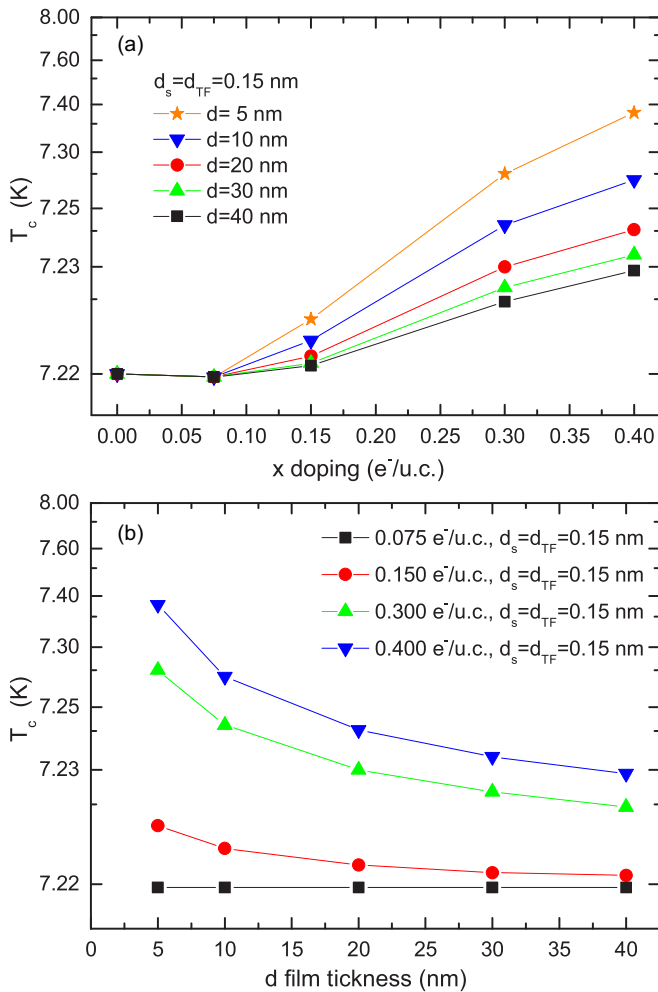


FIG. 3. (a) Calculated critical temperature versus charge doping for five different values of film thickness $d = 5$ nm (orange stars), $d = 10$ nm (blue down triangles), $d = 20$ nm (red circles), $d = 30$ nm (green up triangles), and $d = 40$ nm (black squares) with surface layer thickness $d_s = d_{\text{TF}}$. (b) Calculated critical temperature versus film thickness for four different charge doping (electrons/unitary cell): 0.075 (black squares), 0.150 (red circles), 0.300 (green up triangles), and 0.400 (blue down triangles) with $d_s = d_{\text{TF}}$. The two graphs are in semilogarithmic scale [$\log(T_c - 7.218)$].

material satisfies the Thomas-Fermi model for any value of doping: in this case, the model has no free parameters.

In Fig. 3 we plot the evolution of T_c upon increasing electron doping and assuming that the Thomas-Fermi model always holds ($m = 0$ and $d_s = d_{\text{TF}}$), for different values of film thickness. The calculations show that the qualitative increase in T_c with increasing doping level that we observed in the homogeneous case is retained also in proximized films of any thickness [see Fig. 3(a)]. However, the presence of a coupling between surface and bulk induced by the proximity effect gives rise to a key difference with respect to the homogeneous

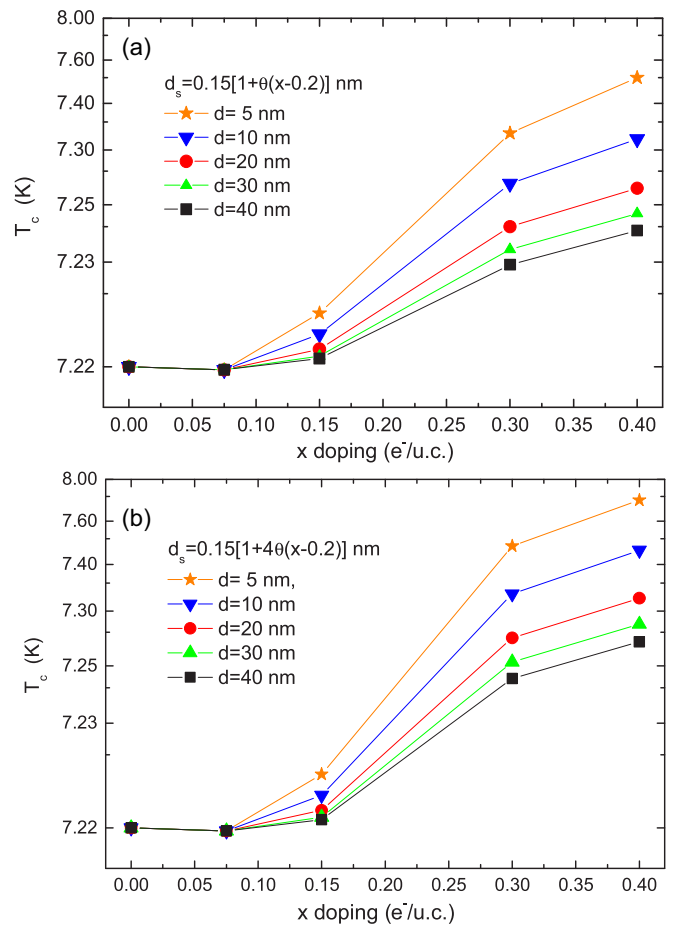


FIG. 4. Calculated critical temperature versus charge doping for five different values of film thickness $d = 5$ nm (orange stars), $d = 10$ nm (blue down triangles), $d = 20$ nm (red circles), $d = 30$ nm (green up triangles), and $d = 40$ nm (black squares) with surface layer thickness $d_s = d_{\text{TF}}[1 + m\Theta(x - 0.2)]$ [(a) $m = 1$ and (b) $m = 4$]. The two graphs are in semilogarithmic scale [$\log(T_c - 7.218)$].

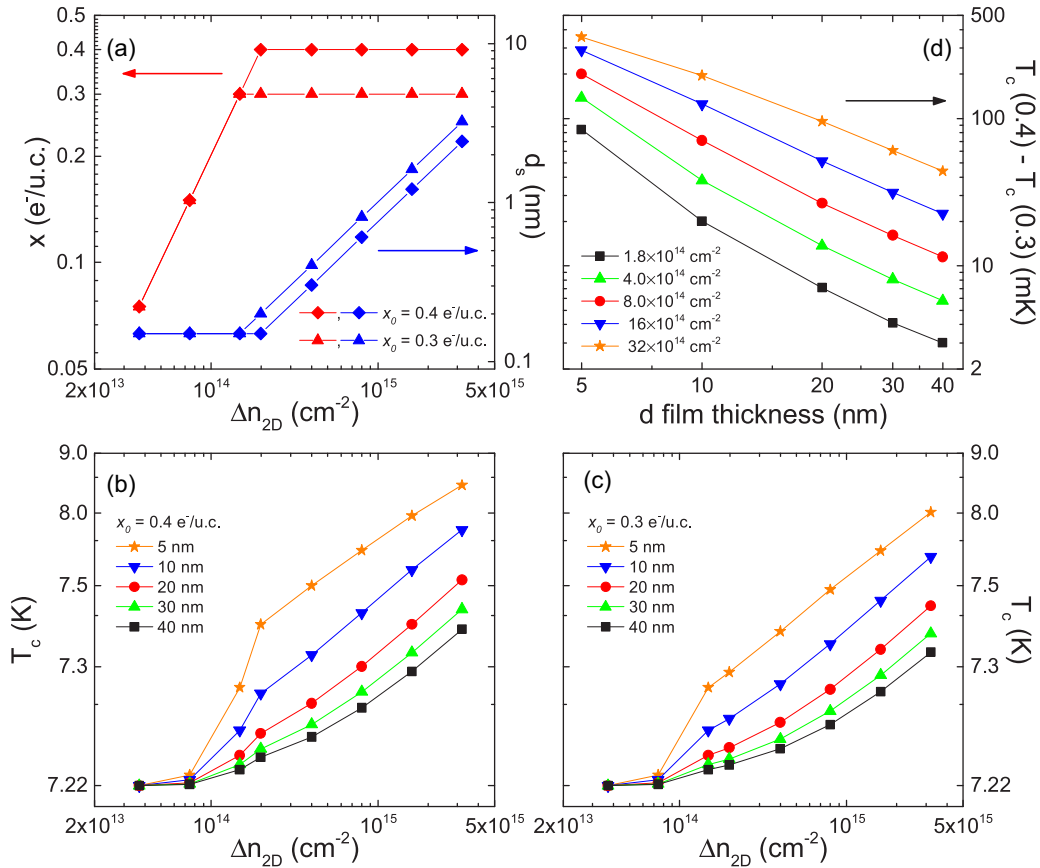


FIG. 5. (a) Dependence of the doping *per unit volume* x (red up triangles and diamonds) and surface layer thickness d_s (blue up triangles and diamonds) on the induced carrier density *per unit surface* Δn_{2D} , for two different values of the maximum doping level $x_0 = 0.3$ and $x_0 = 0.4$ e⁻/unit cell. (b) and (c) T_c versus induced carrier density *per unit surface* Δn_{2D} for five different film thicknesses [$d = 5$ nm (orange stars), $d = 10$ nm (blue down triangles), $d = 20$ nm (red circles), $d = 30$ nm (green up triangles), and $d = 40$ nm (black squares)] in the cases $x_0 = 0.4$ and 0.3 e⁻/unit cell, respectively. (b) and (c) In semilogarithmic scale [$\log(T_c - 7.2)$]. (d) $T_c(x = 0.4) - T_c(x = 0.3)$ versus induced carrier density *per unit surface* Δn_{2D} for the five different film thicknesses.

case, namely, a strong dependence of T_c on film thickness in the doped films. Indeed, the magnitude of the T_c shift with respect to the homogeneous case is heavily suppressed already in films as thin as 5 nm. This behavior is best seen in Fig. 3(b), where we plot the same data as a function of the total film thickness for all doping levels. As we can see, the increase of critical temperature drops dramatically with increasing film thickness. We have not calculated the critical temperature for monolayer films since the approximations of the model would no longer apply in this case: in particular the unperturbed electron-phonon spectral function would be different from the bulklike one we employed in our calculations [59].

We now consider the effect of the different degrees of confinement for the induced charge carriers at the surface of the films. We do so by allowing the perturbed surface layer to spread further in the depth of the film for large electron doping, i.e., by increasing the m parameter in the definition of d_s . In Fig. 4 we plot the evolution of T_c with increasing electron doping and for different film thicknesses, in the two cases $m = 1$ (d_s is allowed to expand up to $2d_{TF} = 0.3$ nm) and $m = 4$ (d_s is allowed to expand up to $5d_{TF} = 0.75$ nm). We can first observe how a different value of d_s does not change the qualitative behavior of the curves. The evolution

of T_c with increasing electron doping is still comparable to both the homogeneous case and the proximized films in the Thomas-Fermi limit. The suppression of the T_c increase with increasing film thickness is also similar to the latter case. However, the magnitude of the T_c shift *for the same values of film thickness and doping level per unit cell* is clearly the more enhanced the larger the value of d_s . This is to be expected, as larger values of d_s increase the fraction of the film that is perturbed by the application of the electric field and reduce the T_c shift dampening operated by the proximity effect. In principle, for values of m large enough (or film thickness d small enough) one could reach the limit value $d_s \simeq d$ and recover the homogeneous case where the T_c shift is maximum.

All the calculations we performed so far assume that one could directly control the induced carrier density *per unit volume* x in the surface layer, without an explicit upper limit. However, this is not an experimentally achievable goal in a field-effect device architecture. In this class of devices, the polarization of the gate electrode allows one to tune the electric field at the interface and thus the induced carrier density *per unit surface* Δn_{2D} required to screen it. Within our model, $\Delta n_{2D} = \int_0^{d_s} \Delta n_{3D} dz$ is distributed within a layer of thickness d_s . In general, the determination of the exact

depth profile of this distribution requires the self-consistent solution of the Poisson equation [55]; however, as a first approximation we can consider this distribution to be constant, obtaining an effective doping level per unit volume simply as $x = \Delta n_{2D}/d_s$. This procedure allows one to employ the same DFT-Eliashberg formalism we developed before in order to simulate a field effect experiment on a superconducting thin film.

In addition, according to our recent experimental findings on niobium nitride [35], the surface layer thickness d_s is a monotonically increasing function of the induced carrier density *per unit surface* Δn_{2D} , while the doping level *per unit volume* x saturates for large Δn_{2D} . In other words, the previously defined threshold value x_0 for the breakdown of the Thomas-Fermi approximation now takes on also the meaning of a *maximum* achievable doping level. Figure 5(a) shows the resulting dependence of x and d_s on Δn_{2D} , for two different values of the maximum doping level $x_0 = 0.3$ and $x_0 = 0.4$ e^- /unit cell. When Δn_{2D} is small enough so that $x < x_0$, the Thomas-Fermi screening holds, $d_s = d_{TF}$ is constant, and x linearly increases with Δn_{2D} . As soon as Δn_{2D} becomes large enough that $x = x_0$ is constant [$\Delta n_{2D}(x_0)$], the Thomas-Fermi screening is no longer valid and $d_s(\Delta n_{2D}) = \Delta n_{2D}x_0$ increases linearly with Δn_{2D} .

In Figs. 5(b) and 5(c) we plot the resulting modulation of T_c for five different film thicknesses in the cases $x_0 = 0.4$ and 0.3 e^- /unit cell, respectively. In both cases we can readily distinguish between two regimes of Δn_{2D} . When $\Delta n_{2D} \lesssim \Delta n_{2D}(x_0)$, Thomas-Fermi screening holds and we reproduce the behavior we observed in Fig. 3(a). In this regime, the induced carrier density directly modulates x and thus the electron-phonon spectral function $\alpha^2F(\Omega)$. The T_c modulation is thus a result of a direct modification of the material properties at the surface, with proximity effect simply operating a “smoothing” which is more effective the thinner the film. On the other hand, when $\Delta n_{2D} > \Delta n_{2D}(x_0)$, the surface properties [$\alpha^2F(\Omega)$] are no longer modified by the extra charge carriers, and the further modulation of T_c originates entirely from the proximity effect as determined by the increase in d_s .

We can also compare the T_c shifts for different maximum doping levels x_0 . Figure 5(d) shows the difference between the T_c corresponding to $x_0 = 0.4$ and 0.3 e^- /unit cell as a function of the total film thickness, for different values of Δn_{2D} . We can clearly see how T_c is always larger for the films with larger x_0 , for any value of film thickness, even if the associated values of d_s are always smaller. This indicates that the maximum achievable value of x_0 is dominant with respect to the increase of d_s to determine the final value of T_c , also in the doping regime $\Delta n_{2D} > \Delta n_{2D}(x_0)$.

Of course, in a real sample we do not expect the transition between the two regimes to be so clear-cut, as the saturation of x to x_0 would occur over a finite range of Δn_{2D} . In this intermediate region, the modulation of $\alpha^2F(\Omega)$ and d_s would both contribute in a comparable way to the final value of T_c in the film. We stress, however, that in both regimes the proximity effect is fundamental in determining the T_c of the gated film. We also note that the proximized Eliashberg equations are able to account for a nonuniform scaling of the T_c shift for different values of film thickness,

unlike the models that use approximated analytical equations for T_c .

V. CONCLUSIONS

In this work we have developed a general method for the theoretical simulation of field-effect doping in superconducting thin films of arbitrary thickness, and we have benchmarked it on Pb as a standard strong-coupling superconductor. Our method relies on *ab initio* DFT calculations to compute how the increasing doping level x per unit volume modifies the structural and electronic properties of the material [shift of Fermi level ΔE_F , density of states $N(0)$, and electron-phonon spectral function $\alpha^2F(\Omega)$]. The Coulomb pseudopotential μ^* is determined by simple calculations from some of these parameters. The properties of the pristine thin film (critical temperature T_c , device area A , and total film thickness d) can be obtained either from the literature or experimentally from standard transport measurements. For doping values where the Thomas-Fermi theory of screening is satisfied, the perturbed surface layer thickness is constant ($d_s = d_{TF}$) and the theory has no free parameters.

Once all the input parameters are known, our method allows us to compute the transition temperature T_c for arbitrary values of film thickness d and electron doping in the surface layer x by solving the proximity-coupled Eliashberg equations in the surface layer and unperturbed bulk. On the other hand, if no reliable estimations of the surface layer thickness d_s are available, our method allows one to determine $d_s(x)$ by reproducing the experimentally measured $T_c(x)$. This allows us to probe deviations from the standard Thomas-Fermi theory of screening in the presence of very large interface electric fields.

We also show how, even in the case where the Thomas-Fermi approximation breaks down and the doping level x can no longer be increased, the transition temperature T_c of a thin film can still be indirectly modulated by the electric field by changing the surface layer thickness d_s . For what concerns artificial enhancements of T_c in superconducting thin films, we conclude that very thin films ($d \lesssim d_s$, in order to minimize the smoothing operated by the proximity effect) of a superconductive material characterized by a strong increase of the electron-phonon (boson) coupling upon changing its carrier density are required to optimize the effectiveness of the field-effect-device architecture.

Finally, our calculations indicate that sizable T_c enhancements of the order of ~ 0.5 K should be achievable in thin films of a standard strong-coupling superconductor such as Pb, for easily realizable thicknesses of ~ 10 nm and doping levels routinely induced via EDL gating in metallic systems [5,35]. These features may open the possibility for superconducting switchable devices and electrostatically reconfigurable superconducting circuits above liquid helium temperature.

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- [1] T. Fujimoto and K. Awaga, *Phys. Chem. Chem. Phys.* **15**, 8983 (2013).
- [2] K. Ueno, H. Shimotani, H. T. Yuan, J. T. Ye, M. Kawasaki, and Y. Iwasa, *J. Phys. Soc. Jpn.* **83**, 032001 (2014).
- [3] A. M. Goldman, *Annu. Rev. Mater. Res.* **44**, 45 (2014).
- [4] Y. Saito, T. Nojima, and Y. Iwasa, *Supercond. Sci. Technol.* **29**, 093001 (2016).
- [5] D. Daghero, F. Paolucci, A. Sola, M. Tortello, G. A. Ummarino, M. Agosto, R. S. Gonnelli, J. R. Nair, and C. Gerbaldi, *Phys. Rev. Lett.* **108**, 066807 (2012).
- [6] K. Ueno, S. Nakamura, H. Shimotani, A. Ohtomo, N. Kimura, T. Nojima, H. Aoki, Y. Iwasa, and M. Kawasaki, *Nat. Mater.* **7**, 855 (2008).
- [7] J. T. Ye, S. Inoue, K. Kobayashi, Y. Kasahara, H. T. Yuan, H. Shimotani, and Y. Iwasa, *Nat. Mater.* **9**, 125 (2010).
- [8] Y. Saito, Y. Kasahara, J. T. Ye, Y. Iwasa, and T. Nojima, *Science* **350**, 409 (2015).
- [9] K. Ueno, S. Nakamura, H. Shimotani, H. T. Yuan, N. Kimura, T. Nojima, H. Aoki, Y. Iwasa, and M. Kawasaki, *Nat. Nanotech.* **6**, 408 (2011).
- [10] J. T. Ye, Y. J. Zhang, R. Akashi, M. S. Bahramy, R. Arita, and Y. Iwasa, *Science* **338**, 1193 (2012).
- [11] S. Jo, D. Costanzo, H. Berger, and A. F. Morpurgo, *Nano Lett.* **15**, 1197 (2015).
- [12] J. M. Lu, O. Zheliuk, I. Leermakers, N. F. Q. Yuan, U. Zeitler, K. T. Law, and J. T. Ye, *Science* **350**, 1353 (2015).
- [13] W. Shi, J. T. Ye, Y. Zhang, R. Suzuki, M. Yoshida, J. Miyazaki, N. Inoue, Y. Saito, and Y. Iwasa, *Sci. Rep.* **5**, 12534 (2015).
- [14] Y. Yu, F. Yang, X. F. Lu, Y. J. Yan, Y.-H. Cho, L. Ma, X. Niu, S. Kim, Y.-W. Son, D. Feng, S. Li, S.-W. Cheong, X. H. Chen, and Y. Zhang, *Nat. Nanotechnol.* **10**, 270 (2015).
- [15] D. Costanzo, S. Jo, H. Berger, and A. F. Morpurgo, *Nat. Nanotechnol.* **11**, 339 (2016).
- [16] Y. Saito, Y. Nakamura, M. S. Bahramy, Y. Kohama, J. T. Ye, Y. Kasahara, Y. Nakagawa, M. Onga, M. Tokunaga, T. Nojima, Y. Yanase, and Y. Iwasa, *Nat. Phys.* **12**, 144 (2016).
- [17] A. T. Bollinger, G. Dubuis, J. Yoon, D. Pavuna, J. Misewich, and I. Božović, *Nature (London)* **472**, 458 (2011).
- [18] X. Leng, J. Garcia-Barriocanal, S. Bose, Y. Lee, and A. M. Goldman, *Phys. Rev. Lett.* **107**, 027001 (2011).
- [19] X. Leng, J. Garcia-Barriocanal, B. Yang, Y. Lee, J. Kinney, and A. M. Goldman, *Phys. Rev. Lett.* **108**, 067004 (2012).
- [20] S. Maruyama, J. Shin, X. Zhang, R. Suchoski, S. Yasui, K. Jin, R. L. Greene, and I. Takeuchi, *Appl. Phys. Lett.* **107**, 142602 (2015).
- [21] K. Jin, W. Hu, B. Zhu, J. Yuan, Y. Sun, T. Xiang, M. S. Fuhrer, I. Takeuchi, and R. L. Greene, *Sci. Rep.* **6**, 26642 (2016).
- [22] A. Fête, L. Rossi, A. Augieri, and C. Senatore, *Appl. Phys. Lett.* **109**, 192601 (2016).
- [23] L. Burlachkov, I. B. Khalfin, and B. Ya. Shapiro, *Phys. Rev. B* **48**, 1156 (1993).
- [24] M. Ghinovker, V. B. Sandomirsky, and B. Ya. Shapiro, *Phys. Rev. B* **51**, 8404 (1995).
- [25] J. Walter, H. Wang, B. Luo, C. D. Frisbie, and C. Leighton, *ACS Nano* **10**, 7799 (2016).
- [26] P. Lipavsky, J. Kolacek, K. Morawetz, E. H. Brandt, and T.-J. Yang, *Bernoulli Potential in Superconductors How the Electrostatic Field Helps to Understand Superconductivity*, Lecture Notes in Physics, Vol. 733 (Springer, Berlin, 2008).
- [27] L. J. Li, E. C. T. O'Farrell, K. P. Loh, G. Eda, B. Özyilmaz, and A. H. Castro Neto, *Nature (London)* **529**, 185 (2016).
- [28] M. Yoshida, J. T. Ye, T. Nishizaki, N. Kobayashi, and Y. Iwasa, *Appl. Phys. Lett.* **108**, 202602 (2016).
- [29] X. X. Xi, H. Berger, L. Forró, J. Shan, and K. F. Mak, *Phys. Rev. Lett.* **117**, 106801 (2016).
- [30] J. Shiogai, Y. Ito, T. Mitsuhashi, T. Nojima, and A. Tsukazaki, *Nat. Phys.* **12**, 42 (2016).
- [31] B. Lei, J. H. Cui, Z. J. Xiang, C. Shang, N. Z. Wang, G. J. Ye, X. G. Luo, T. Wu, Z. Sun, and X. H. Chen, *Phys. Rev. Lett.* **116**, 077002 (2016).
- [32] K. Hanzawa, H. Sato, H. Hiramatsu, T. Kamiya, and H. Hosono, *Proc. Natl. Acad. Sci. USA* **113**, 3986 (2016).
- [33] J. Choi, R. Pradheesh, H. Kim, H. Im, Y. Chong, and D. H. Chae, *Appl. Phys. Lett.* **105**, 012601 (2014).
- [34] E. Piatti, A. Sola, D. Daghero, G. A. Ummarino, F. Laviano, J. R. Nair, C. Gerbaldi, R. Cristiano, A. Casaburi, and R. S. Gonnelli, *J. Supercond. Novel Magn.* **29**, 587 (2016).
- [35] E. Piatti, D. Daghero, G. A. Ummarino, F. Laviano, J. R. Nair, R. Cristiano, A. Casaburi, C. Portesi, A. Sola, and R. S. Gonnelli, *Phys. Rev. B* **95**, 140501 (2017).
- [36] J. P. Carbotte, *Rev. Mod. Phys.* **62**, 1027 (1990).
- [37] G. A. Ummarino, in *Emergent Phenomena in Correlated Matter*, edited by E. Pavarini, E. Koch, and U. Schollwöck (Forschungszentrum Jülich GmbH and Institute for Advanced Simulations, Jülich, Germany, 2013), pp. 13.1–13.36.
- [38] W. L. McMillan, *Phys. Rev.* **175**, 537 (1968).
- [39] E. Schachinger and J. P. Carbotte, *J. Low Temp. Phys.* **54**, 129 (1984).
- [40] H. G. Zarate and J. P. Carbotte, *Phys. Rev. B* **35**, 3256 (1987).
- [41] H. G. Zarate and J. P. Carbotte, *Physica B+C* **135**, 203 (1985).
- [42] V. Z. Kresin, H. Morawitz, and S. A. Wolf, *Mechanisms of Conventional and High Tc Superconductivity* (Oxford University Press, Oxford, 1999).
- [43] G. A. Ummarino and R. S. Gonnelli, *Phys. Rev. B* **66**, 104514 (2002).
- [44] P. Morel and P. W. Anderson, *Phys. Rev.* **125**, 1263 (1962).
- [45] G. Grosso and G. Pastori Parravicini, *Solid State Physics* (Academic, New York, 2014).
- [46] B. Meyer, C. Elsässer, and M. Fähnle, FORTRAN90 Program for Mixed-Basis Pseudopotential Calculations for Crystals, Max-Planck-Institut für Metallforschung, Stuttgart (unpublished).
- [47] D. Vanderbilt, *Phys. Rev. B* **32**, 8412 (1985).
- [48] R. Heid, K.-P. Bohnen, I. Yu. Sklyadneva, and E. V. Chulkov, *Phys. Rev. B* **81**, 174527 (2010).
- [49] L. Hedin and B. I. Lundqvist, *J. Phys. C* **4**, 2064 (1971).
- [50] S. Baroni, S. de Gironcoli, A. D. Corso, and P. Giannozzi, *Rev. Mod. Phys.* **73**, 515 (2001).
- [51] N. E. Zein, *Fiz. Tverd. Tela (Leningrad)* **26**, 3028 (1984); *Sov. Phys. Solid State* **26**, 1825 (1984).
- [52] R. Heid and K. P. Bohnen, *Phys. Rev. B* **60**, R3709 (1999).
- [53] I. Yu. Sklyadneva, R. Heid, P. M. Echenique, K.-B. Bohnen, and E. V. Chulkov, *Phys. Rev. B* **85**, 155115 (2012).
- [54] D. Pines and P. Nozieres, *The Theory of Quantum Liquids* (Benjamin, New York, 1966).
- [55] T. Brumme, M. Calandra, and F. Mauri, *Phys. Rev. B* **91**, 155436 (2015).

- [56] P. B. Allen and R. C. Dynes, *Phys. Rev. B* **12**, 905 (1975).
- [57] J. E. Hirsch, *Phys. Rev. B* **70**, 226504 (2004).
- [58] S. Basavaiah, J. M. Eldridge, and J. Matisoo, *J. Appl. Phys.* **45**, 457 (1974).
- [59] S. Bose, C. Galande, S. P. Chockalingam, R. Banerjee, P. Raychaudhuri, and P. Ayyub, *J. Phys.: Condens. Matter* **21**, 205702 (2009).
- [60] W. L. McMillan and J. M. Rowell, in *Superconductivity*, edited by R. D. Parks (Marcel Dekker, New York, 1969), Vol. 1.