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Photoemission spectra of CeAl$_2$, CeBe$_{13}$, CeSi$_2$, and CeCu$_2$Si$_2$: Weights and widths of the 4f emission features

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We present valence-band photoemission spectra for CeAl$_2$, CeBe$_{13}$, CeSi$_2$, CeCu$_2$Si$_2$, and related conventional rare-earth counterpart compounds, taken at photon energies corresponding to the giant 4d resonance with resolution $\sim 150$ meV. We take into account the 5d emission, which comprises 30% of the valence-band emission at the 4d resonance. We compare the resulting 4f emission to the predictions of the Anderson impurity model calculated in a low-order 1/N expansion, and including spin-orbit, crystal-field, and finite Coulomb correlation effects. The calculation gives order-of-magnitude fits to the data, but underestimates the spectral weight near the Fermi level by a factor of 2 and overestimates the width of the main f emission at 2 eV by a factor of 4. We argue that Ce photoemission remains an open problem and discuss several experimental and theoretical issues which need to be resolved to make further progress.

I. INTRODUCTION

In this paper we address the question$^{1-6}$ of whether the Anderson impurity model adequately describes valence-band 4f photoemission in cerium compounds. The comparison is made over the scale of the bandwidth at moderate resolution (150 meV). There are several reasons for undertaking this: our resolution is significantly better than that used in similar earlier studies;$^4$ in some cases single crystals are now available where older studies were performed on polycrystals; we have made a special effort to subtract the 5d emission, which we will argue was done incorrectly in the past; and we have developed a calculation which simultaneously includes spin-orbit and crystal-field splitting, and finite-Coulomb correlation energy.

We present ultraviolet photoemission spectra for four compounds (CeAl$_2$, CeCu$_2$Si$_2$, CeSi$_2$, and CeBe$_{13}$) chosen so that comparison to theory can be made over two orders of magnitude of Kondo temperature ($T_K = 4$, 8, 40, and 400 K, respectively). The experiments utilized the giant 4d resonance at the photon energy $h\nu = 120$ eV to enhance 4f emission;$^7$ the resolution was 150 meV. We estimate the background emission (primarily 5d) by comparing to spectra of La and Pr compounds taken at their respective 4d resonance energies, showing that the 5d emission forms an appreciable fraction ($\approx 30\%$) of the 4d resonance spectra in these Ce compounds. We then compare the spectra to the predictions of the Anderson impurity model, calculated via the large $N$ expansion method.$^{1,8}$ As mentioned, the calculation includes finite Coulomb correlation $U$, with spin-orbit and crystal-field splittings constrained to their (experimentally) known values. For a fixed background bandwidth $W$ and Coulomb correlation $U$ we then constrain the hybridization $\Delta = V^2\rho$ and the bare $f$ level energy $E_f$ to give the experimentally known Kondo temperature and measured ("renormalized") $f$ level energy $\tilde{E}_f$. The spectra were finally subjected to a degree of broadening equivalent to our experimental instrumental resolution. Our basic findings are that the weight near the Fermi energy ($\epsilon_F$) observed experimentally is at least two times greater than predicted for the Kondo resonance and its crystal-field and spin-orbit sidebands and that the width of the $f^1 \rightarrow f^0$ emission peak near 2 eV is three to four times smaller in the experimental spectra than predicted.

In the discussion we propose several possibilities to explain these discrepancies between experiment and theory. First, we consider experimental difficulties in determining the bulk 4f spectra. These include uncertainties due to our method of subtracting the bulk (mainly 5d) background; the lack of knowledge of the photon energy dependence of the branching ratio of the $f^{n-1} \rightarrow f^0$ peak to the near-$\epsilon_F$ peak; and the problem of separating out the surface emission. Although the latter is clearly important for 120-eV photoemission spectroscopy (PES), we have not attempted it; but we argue that were we to do so, then in most instances the disagreement between experiment and theory would be worse. We then consider the theoretical difficulties of determining the energy-dependent hybridization $\Delta(\epsilon) = V^2(\epsilon)\rho(\epsilon)$. While use of smaller values of bandwidth $W$ than are expected for these compounds can alleviate the above-mentioned discrepancies somewhat, we argue that the resulting spec-
SPECTRA still do not resemble the experimental data in significant respects. The large N approximation scheme may also affect the predictions of the Anderson model. Finally, we discuss the possibility that the discrepancies may arise from a more fundamental source, and that resolution of the difficulties may require (i) inclusion in the model of Coulomb screening of the 4f hole, and (ii) recognition that these cerium compounds may be closer to the band limit than implied by the model.9,10 All of this suggests that photoemission in cerium compounds is an open problem.

II. EXPERIMENTAL TECHNIQUE

The samples consisted of flux-grown single crystals of CeCu2Si2, CeBe13, and LaBe13 and arc-melted polycrystals of PrBe13, RA13 (R = La, Ce, and Pr) and RSi2 (R = La and Ce). The experiments were performed primarily at the U3C beamline at NSLS; additional experiments were performed at the University of Wisconsin SRC. Both beamlines utilize extended range grasshopper (ERG) monochromators; at lower photon energies at the SRC a Seya monochromator was used. For most of the spectra reported here the photon energy was 116, 120, and 126 eV for the La, Ce, and Pr compounds, respectively, corresponding to the energies of the giant 4d resonance for these three elements. An HA-50 hemispherical analyzer coupled to a multichannel detector provided the electron counting. For the 4d resonance spectra the total resolution, as determined by numerous measurements of the Fermi edge of Au or Pt, was in the range of 120–180 meV. The samples were cleaved or fractured in an ultrahigh vacuum; the base pressure was in the range 5–10×10−11 torr for all experiments. The samples were in direct contact with a Cu cold finger which could be cooled by liquid N2 (T ≈ 80 K) or liquid He (T ≈ 20 K); a thermocouple attached to the cold finger in close proximity to the sample provided the temperature measurement. Changes in the spectra with temperature were minor. For all the spectra reported here we have subtracted the inelastic background under the conventional assumption that the background at energy E is proportional to the area bounded by the background and the spectrum between E and $\varepsilon_F$.

A problem which plagues the whole area of heavy fermion photoemission is that, with very few exceptions,11 the samples utilized are polycrystals, and rather poorly characterized by the standards of modern surface science. The spectra can vary somewhat from cleave to cleave and are extremely sensitive to oxidation, which occurs rapidly. Hence, cleanliness and reproducibility are key issues. To study these issues, we cleaved each sample three to six times, under a variety of conditions, and studied the evolution of the spectra with time, as the degree of oxidation increased. As examples of the latter effect we show in Fig. 1 room-temperature spectra of CeAl3 and CeBe13, where following the initial spectra the samples were exposed to the base pressure for periods of 6 and 2 h, respectively. It is clear that both the line widths and the fractional weight in the feature near the Fermi level increase markedly with continued exposure. At the same time, a maximum develops near 6-eV binding energy, which is characteristic of O 2p emission. For our cleanest spectra, which showed no sign of oxygen emission, we found a smaller degree of cleave dependence, which gives an uncertainty in the linewidths and fractional spectral weights reported below of order 10%. It is worth pointing out that the effects of oxidation and cleave dependence (as well as the uncertainties in determining the background non-4f emission, discussed below) are less of a problem in this study, where our effort is to examine overall features (linewidths and relative spectral weights) at moderate resolution, than for detailed high-resolution studies of the line shape and temperature dependence near $\varepsilon_F$.

III. EXPERIMENTAL RESULTS AND ANALYSIS

The spectrum of the valence-band region of CeAl3, taken at the giant 4d resonance at $h\nu = 120$ eV, is shown in Fig. 2(a). Both the 4f and 5d electrons are known7 to contribute to this resonance spectrum. To determine the 5d contribution, we measured the spectra of the conventional rare-earth counterparts LaAl3 and PrAl3 at their 4d resonances at 116 and 126 eV, respectively [Figs. 3(b) and 3(c)]. Simultaneously, we measured the Al 2p spectra of all three compounds at the same photon energies [Fig. 3(a)]. All spectra presented are normalized so as to have equal area in the Al 2p peak; this assumes that the reso-
nance does not affect the emission from the Al core level, and ignores the photon energy dependence of the Al 2p emission in the region 116–126 eV. We then fit the valence band of LaAl₃ to a Gaussian [Fig. 3(b)]; we assume this represents the 5d emission. The spectrum of PrAl₃ can be fit to the sum of a Lorentzian representing the Pr 4f emission [and whose full width at half maximum (FWHM) is 1.5 eV] and a Gaussian with exactly the same binding energy, width, and height as for LaAl₃; this supports the assumption that the 5d emission will be essentially identical in all the light rare-earth compounds RAl₃. We thus estimate the 5d emission in CeAl₃ with precisely the same Gaussian [Fig. 2(a)] noting that the 5d emission forms 28% of the total valence-band emission in CeAl₃ at 120 eV. After subtracting off the 5d emission we obtain our estimate of the 4f emission [Fig. 2(b)]. We note that the ratio of the (normalized) 4f emission in PrAl₃ to that of CeAl₃ is 1.8, or approximately 2, as expected since there are twice as many 4f electrons in Pr as in Ce; this lends added confidence to our procedure. Finally we fit the 4f emission in CeAl₃ to the sum of a Lorentzian, representing emission from a local 4f state, which we call the “$4f^1 \rightarrow 4f^{0*}$” emission, and a remainder, which we label as the “near-$\varepsilon_F$” emission. The $4f^1 \rightarrow 4f^{0*}$ peak has a width (FWHM) of 0.9 eV; the fractional weight in the near-$\varepsilon_F$ emission is 37%.

The 4d resonance spectra of LaSi₂ and CeSi₂, normalized to equal area in the Si 2p peaks [Fig. 4(a)] are shown in Fig. 4(b). The LaSi₂ spectrum has an area equal to 30% of the total CeSi₂ spectrum; this is very close to the value 28% for the 5d–valence-band fraction given above for CeAl₃ and increases our confidence that the LaSi₂ spectrum can be taken as the 5d emission in CeSi₂, and is appropriately normalized. After subtracting the 5d emission we obtain the 4f emission in CeSi₂ [Fig. 4(c)]; this is again fit to the sum of a Lorentzian, representing the $4f^1 \rightarrow 4f^{0*}$ peak, and a remainder. This latter near-$\varepsilon_F$ peak has a fractional area of 33%; the $4f^1 \rightarrow 4f^{0*}$ peak has a width (FWHM) of 1.5 eV.

Additional information concerning the background emission can be obtained by studies at photon energies below the threshold for the onset of f emission; such spectra are often taken as representing the background emission from 5d and other (e.g., Si 3p) states. In Fig. 5 we compare such spectra for LaSi₂ and CeSi₂, taken at 26 eV, to the spectra of LaSi₂ taken at the 4d resonance (116 eV) and of CeSi₂ taken at the Fano minimum of the 4d resonance (112 eV). Both pairs of spectra have three peaks (at −4, −2, and 0 eV) but the branching ratios of
the three peaks are clearly different at the two photon energies; in particular the peak near the Fermi edge is enhanced and the peak at 4 eV is suppressed in the 26-eV spectra. These differences arise from relative changes in the optical-absorption matrix elements for the three peaks as the photon energy varies. Since our CeSi₂ valence-band spectrum was obtained at \( h\nu = 120 \text{ eV} \), it is clearly more appropriate to determine the strength of the 5d emission at an approximately equal photon energy (as we have done by use of the LaSi₂ 4d resonance spectrum) than at the lower energies (\( h\nu = 20–30 \text{ eV} \)). We did not measure a core level for normalization in the \( \text{RBe}_{13} \) series. Instead, we proceeded as follows: We measured the LaBe₁₃ spectrum at the 4d resonance, and the spectra of CeBe₁₃ and PrBe₁₃ at a low photon energy (27 eV) chosen to lie below the threshold for the onset of \( f \) emission [Fig. 6(a)]. All three spectra have the same three-peaked structure and furthermore the branching ratios are the same at both photon energies; hence, we take the 5d emission as proportional to the LaBe₁₃ spectrum, but with as yet undetermined overall scale factor. Two ways to choose the scale factor are shown in Fig. 7 for CeBe₁₃. In the top panel we have chosen the scale factor to be sufficiently large that when 5d is subtracted there is no trace of the middle peak near 1.5-eV binding energy, which is the peak that corresponds to the largest peak in the 5d emission. In doing this, however, we find that the resulting 5d emission is nearly 50% of the total valence-band emission in CeBe₁₃ and the resulting fractional weight in the near-\( \varepsilon_F \) emission is only 11% of the 4f emission, as opposed to the values of 30% and 35% observed in CeAl₁₃ and CeSi₂. In the lower panel we have chosen the scale factor so that the 5d emission is 30% of the valence band. For this choice, after subtraction of the \( 4f^1 \rightarrow 4f^0 \) Lorentzian (FWHM = 1.2 eV), the remaining near-\( \varepsilon_F \) emission has a fractional weight of 28%, now comparable to that for CeAl₁₃ and CeSi₂. We take the latter choice of the 5d emission as the more probable. We have also attempted to determine the correct choice of scale factor for PrBe₁₃. However, for this compound the valence-band spectra cannot be neatly decomposed into the sum of a Lorentzian plus the 5d emission, as for PrAl₁₃. The choice of 5d emission shown in Fig. 6(b) has the virtue that it fits the Fermi edge very well; a greater choice of scale factor would overestimate the emission at \( \varepsilon_F \). This leads to the result, however, that there is a large remaining peak [the long-dashed line in Fig. 6(b)] in the valence band of PrBe₁₃.

The spectra of CeCu₂Si₂ at the 4d resonance (\( h\nu = 120 \text{ eV} \)) as well as off-resonance (\( h\nu = 112 \text{ eV} \)) are shown in

![FIG. 4](image)

**FIG. 4.** (a) Si sp core levels, normalized to equal area, for LaSi₂ (\( h\nu = 116 \text{ eV}, T = 80 \text{ K} \)) and CeSi₂ (\( h\nu = 120 \text{ eV}, T = 20 \text{ K} \)). (b) Valence-band spectra, normalized to the Si 2p core levels, for CeSi₂ (\( h\nu = 120 \text{ eV}, T = 20 \text{ K} \)) and LaSi₂ (\( h\nu = 116 \text{ eV}, T = 80 \text{ K} \)). (c) 4f emission for CeSi₂ (solid circles), obtained by subtraction of the LaSi₂ emission. The local \( f^1 \) peak is represented as a Lorentzian with binding energy \( E_B = 2.35 \text{ eV} \) and FWHM 1.46 eV. The open triangles are the near-\( \varepsilon_F \) emission, obtained by subtracting the Lorentzian.

![FIG. 5](image)

**FIG. 5.** (a) Off-resonance spectrum for CeSi₂ (\( T = 20 \text{ K} \)) and on-resonance spectrum for LaSi₂ (\( T = 80 \text{ K} \)). (b) Spectra for CeSi₂ (20 K) and LaSi₂ (80 K) taken below the threshold (\( h\nu = 30 \text{ eV} \)) for the onset of \( f \) emission. This figure shows two methods of estimating the 5d emission, but demonstrates that the branching ratios between the three peaks in the 5d spectra vary with photon energy. (All spectra are normalized to unit height near 2 eV.)
Fig. 8(a); the spectra are normalized to unity at the copper 3d peak at 4 eV. For this case we do not at present have data for the La counterpart, so we proceed as follows. An examination of Fig. 8 shows a clear peak near 1.67 eV in both the on- and off-resonance spectra. In Fig. 8(b) we fit the off-resonance data in the interval 3.5 eV below the Fermi level to the sum of a Lorentzian representing the tail of the Cu peak at 4 eV, a second Lorentzian for the peak at 1.67 eV, and a reversed arc tangent for the Fermi step. We then assume that the emission in the Fermi step and the peak at 1.67 eV represent the 5d emission, and hence at resonance will be enhanced to a magnitude equal to 30% of the total emission in the valence band, as for the other compounds. To determine the 4f emission, we first subtract the peak representing the copper emission from the on-resonance data, assuming that it should not resonate. We then scale the sum of the peak at 1.67 eV and the Fermi step so that its weight is 30% of the resulting valence-band emission. The 4f emission obtained after subtraction [Fig. 9(a)] shows no hint of the peak at 1.67 eV. The resulting 4f$^1$→4f$^0$ peak has a width 1.12 eV (FWHM) and the near-$\epsilon_F$ peak comprises 49% of the total 4f emission.

![Graph](image_url)

**FIG. 6.** (a) Un-normalized spectra for LaBe$_{13}$ taken on resonance (300 K) and for CeBe$_{13}$ (20 K) and PrBe$_{13}$ (80 K) taken below the threshold for 4f emission. For this material the branching ratio for the non-4f emission appears to be constant with photon energy. (b) Spectrum for PrBe$_{13}$ taken on resonance. The dotted curve, proportional to the curves in (a), and normalized as discussed in the text, represents the non-4f emission. The Lorentzian (E$_g$ = 3.9 eV, FWHM = 1.20 eV) represents the local f$^3$→f$^1$ peak. The long-dashed curve is the extra 4f emission.

![Graph](image_url)

**FIG. 7.** (a) The on-resonance spectrum of CeBe$_{13}$ (hv=120 eV, $T=20$ K) and LaBe$_{13}$ (hv=117 eV, $T=300$ K). The latter is normalized so that when subtracted, no vestige of the peak near 1 eV remains in the 4f spectra (solid triangles). (b) 4f emission for CeBe$_{13}$ (solid circles) obtained by subtracting the LaBe$_{13}$ spectrum, normalized so as to give the same fraction of 5d emission on resonance as in CeAl$_3$ and CeSi$_3$. For reasons stated in the text, we believe this is a better method of normalizing than in (a). The parameters of the Lorentzian (solid line) are E$_g$=2.6 eV, FWHM=1.2 eV. The remainder after subtracting the Lorentzian is the near-$\epsilon_F$ emission (open triangles).

(Variants on our procedure, i.e., other choices of the ratio of the weight in the peak at 1.67 eV to the weight in the Fermi step, give similar final results for the near-$\epsilon_F$ weight and f$^1$→f$^0$ width.)

**IV. THEORETICAL METHODS**

The photoemission spectrum was fit using the results of a variational 1/N expansion for the single-impurity Anderson model. The calculational method used was that of Refs. 12 and 13. We will give full details in a future review; here we focus on similarities and differences between our calculation and those of Refs. 12 and 13. The calculation included the effects of splitting the full fourteenfold degeneracy of the 4f$^1$ configuration by spin-orbit and crystal-field interactions. That is, we consider f$^1$ multiplets $\{\alpha\}$ with energies $E_{f|\alpha\rangle}$ and degeneracies $N_{\alpha|\alpha\rangle}$. The restriction to the $U\rightarrow\infty$ limit of the Anderson model was partially lifted by including states incorporating the f$^2$ configuration into the set of variational basis states. (To the best of our knowledge, this is the first calculation of photoemission spectra using a 1/N treatment of the Anderson model to simultaneously include crystal-field, spin-orbit, and finite-$U$ effects.) The f$^2$ configuration was assumed not to be split by spin-orbit and crystal-field interactions. The hybridization matrix elements were assumed to be independent of the particular f$^1$ multiplet $\{\alpha\}$ involved, and also independent of energy. The variational basis states used for the ground-
state wave function consisted of those specified in Eqs. (3.1), (3.2), and (3.4) of Ref. 12, which incorporate terms of leading order in \( 1/N \).

A set of coupled equations for the variational coefficients for the ground-state wave function and the ground-state energy were found. These were reduced to an integral equation for an amplitude \( d_{k,\alpha} \) which, when solved, yields a transcendental equation for the ground-state energy. This procedure is analogous to that used in Refs. 12 and 13. The integral equation [see Eq. (4.3)] is not separable, except in the \( U \to \infty \) limit. Following the procedure outlined in Ref. 13 [see Eq. (4.6)], the integral equation can be decoupled by neglecting the kinetic-energy dependence of the denominator. This approximation is expected to be reasonable in the large \( U \) limit, i.e., \( U \gg W \). (For the purposes of fitting spectra in real materials, other authors\(^{1,8} \) have solved the integral equation numerically, without resort to this approximation. We argue, however, that use of this decoupling procedure should lead to no greater inaccuracy than the neglect of configurations \( f^n \) with \( n > 2 \).) Unlike the \( N=14 \) case, the resulting transcendental equation for the bound-state energy [a generalization of Eq. (4.7) of Ref. 13] may possess more than one solution. The lowest energy solution corresponds to the ground-state energy.

In the variational ground state, the Kondo singlet is formed by binding a conduction electron to an electron predominantly in the lowest \( f^1 \) multiplet. The resulting complex, consisting of a local \( f \) moment and the screening electron, produces a nonmagnetic singlet. The higher energy solutions of the integral equation, when they exist, also correspond to a Kondo singlet in which the \( f^1 \) configuration is a superposition consisting mostly of the higher multiplets. These excited spin-orbit and crystal-field-split Kondo states have characteristic binding energies or Kondo temperatures \( T_K \) which are different\(^2 \) from the ground-state Kondo temperature \( T_K \). These higher energy Kondo singlets and the resulting features in the photoemission spectra are well separated when the splittings between \( f^1 \) multiplets are greater than the relevant Kondo temperatures.

In the sudden approximation, the zero-temperature limit of the photoemission spectrum is approximated by the density of states for removal of one \( f \) electron. The matrix element of the resolvent, taken between states where one \( f \) electron has been removed, is calculated by a method analogous to that of Ref. 13. The appropriate

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**FIG. 8.** (a) The on-resonance (\( h\nu=120 \) eV) and off-resonance (\( h\nu=112 \) eV) spectra of CeCuSi\(_2\) at \( T=80 \) K, normalized to unity at the Cu emission peak at 4 eV. (b) The off-resonance spectrum, fit to the sum of (i) a Lorentzian (dotted line) with \( E_g=3.7 \) eV and FWHM=0.58 eV, representing the tail of the Cu emission peak; (ii) a Lorentzian (medium-dashed line) with \( E_g=1.66 \) eV and FWHM=0.86 eV, representing the 5d emission; and (iii) a reversed arc tangent with width 0.5 eV (long-dashed line) representing the Fermi step. The solid line is the sum of the three contributions. A small amount of \( 4f \) emission is apparent near 2.5 eV. (c) The on-resonance spectrum. The three fit curves are the same as in (b), only enhanced as explained in the text to give a non-\( 4f \) contribution to the valence band of 30%, as for the other compounds.

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**FIG. 9.** (a) The \( 4f \) emission for CeCuSi\(_2\) (solid circles) obtained after subtracting the fit curves of Fig. 8(c) from the total spectrum. The parameters of the \( 4f \) Lorentzian are \( E_g=2.44 \) eV and FWHM=1.12 eV. The open circles are the remaining near-\( 5d \) emission, after subtraction of the Lorentzian. (b) Calculation of the \( 4f \) emission for CeCuSi\(_2\) for \( W=5 \) eV and \( U=8 \) eV. (c) Calculation of the spectrum assuming \( W=10 \) eV and \( U=14 \) eV. (For further explanation of the theory curves, see the caption for Fig. 10 and the text of Sec. V.)
basis states [(5.17), (5.18), and (5.20)] have been modified to include the splitting of the $f^1$ configuration into multiplets. The set of coupled equations for the resolvent can be separated by using a decoupling approximation similar to that used in Ref. 13. The decoupling approximation is essentially the same as the one that occurs in the calculation of the ground state, and is also expected to produce reasonable results in the limit $U \gg W$.

The photoemission spectrum is then obtained by taking matrix elements of the resolvent and the state in which an $f$ electron has been removed from the ground state. The resulting expression for the one-electron removal spectrum contains

$$
|\langle \omega \rangle | = \Re^{-4} \sum_{[a]} \left( \frac{U}{U+E_f-\varepsilon_F} \right)^2 \left( N_{[\alpha]} |\Delta| \pi \right)^{1/2} \left[ \frac{U}{U+E_f-\varepsilon_F} \right]^2 \left( N_{[\beta]} |\Delta| \pi \right)^{1/2} \left( \frac{1}{U+E_f-k_B T_K} \right) - \left( \frac{1}{U+E_f} \right) \right|^2 ,
$$

where $\Re$ is the renormalization of the ground state, $E_F < 0$ for cerium, $\Delta = V^2 r$, and $a_{k,[\alpha]}$ depends on $\varepsilon_k = h\omega + \mu$ through the relation

$$
a_{k,[\alpha]} = -\left( N_{[\alpha]} |\Delta| \pi \right)^{1/2} \left( \frac{U+2U_2}{U+2U_2+E_f-\varepsilon_F} \right) \left( E_{f,[\alpha]} - E_f + k_T + T_K \right)^{-1} .
$$

Here, $\Gamma \approx (14|\Delta|/\pi)(W/10E_f-\varepsilon_F)$ is small (for $\Delta = 0.1$ eV it is of order 0.4 eV) relative $U$. Each multiplet contributes three terms to the photocurrent. The first, proportional to $a_{k,[\alpha]}$, is already present for $U = \infty$; it represents the rising tail of the ground multiplet, crystal-field, or spin-orbit Kondo resonance, with apparent divergence above $\varepsilon_F$ at $k_T$, $\Delta_{cr}+k_T$, or $\Delta_{SO}+k_T$, respectively. Physically, these tails represent the energy distribution of the conduction electrons involved in the screening of the $f$ moment. It is seen that the inclusion of the $f^2$ component leads to the addition of an energy-independent constant to the photoemission amplitude near the Fermi level, as evidenced by the last term in the bold parentheses of Eq. (1). Finally, cross terms give rise to a weakly energy-dependent constructive interference, which is smaller by a factor $|\Delta|/(U+E_f)$ than the leading term for the tail of the resonance. (The situation is very similar to Fig. 3 of Ref. 13, except that there are separate contributions for each multiplet.)

For energies further away from the Fermi energy, both the Dirac $\delta$ function and the continuum part of the resolvent contribute terms to the spectrum. The resulting convolution is evaluated numerically. The calculated spectrum has features similar to those found by Bickers, Cox, and Wilkins in their evaluation of the $f$ electron density of states for the $U \rightarrow \infty$ limit of the single-impurity Anderson model in the noncrossing approximation (NCA). That is, the spectrum has a broad continuum of $f$ states stretching down at least to the lower edge of the conduction band. The precise shape, including the width and peak energy position, is strongly influenced by the choice of the conduction-band density of states, and the magnitude of the width $W + \varepsilon_F$ of the occupied portion of the conduction band. Closer to the Fermi energy, the spectrum shows Kondo sidebands, due to the existence of the Kondo spin singlets involving the higher energy $f^1$ multiplets. In the theory, these split Kondo singlet states may occur as final states.

In addition to the features that are also found in the factors which depend on both the proportions of the $f^1$ and the $f^2$ configuration in the ground state, as well as cross terms.

For energies within $T_K$ of the Fermi level $\varepsilon_F$, the only contribution to the spectrum comes from a Dirac $\delta$-function component of the resolvent. In this energy regime, the integral over the Dirac $\delta$ function can be performed, and the photoemission spectrum is of the form of Eq. (5.30) of Ref. 13. However, use of the same large $U/W$ approximation as made in the decoupling procedures described above, removes the integrals to yield the analytic expression

$$
|\langle \omega \rangle | = \Re^{-4} \sum_{[a]} \left( \frac{U}{U+E_f-\varepsilon_F} \right)^2 |N_{[\alpha]} |\Delta| \pi \right)^{1/2} \left[ \frac{U}{U+E_f-\varepsilon_F} \right]^2
$$

where $\Re$ is the renormalization of the ground state, $E_F < 0$ for cerium, $\Delta = V^2 r$, and $a_{k,[\alpha]}$ depends on $\varepsilon_k = h\omega + \mu$ through the relation

$$
a_{k,[\alpha]} = -\left( N_{[\alpha]} |\Delta| \pi \right)^{1/2} \left( \frac{U+2U_2}{U+2U_2+E_f-\varepsilon_F} \right) \left( E_{f,[\alpha]} - E_f + k_T + T_K \right)^{-1} .
$$

for $\Gamma = 42$ K for CeAl$_3$, CeCu$_2$Si$_2$, and CeSi$_3$, respectively. Use of the rule$^{18}$ that $T_K$ equals the neutron-scattering quasieleastic linewidth $\omega_0$ gives $T_K = 6$, 6–10, and 290 K for CeAl$_3$, CeCu$_2$Si$_2$, CeSi$_3$, and CeBe$_{13}$, respectively. Use of the rule$^{18}$ that $T_K = 3.5 T_{\text{max}}$, where $T_{\text{max}}$ is the temperature of the broad maximum in susceptibility in mixed valent compounds gives $T_K = 490$ K for CeBe$_{13}$. We therefore set $T_K = 4, 8, 40$, and 400 K for the four compounds. (In Ref. 15, inclusion of an extra factor of $\pi$ in the formula relating $T_K$ and $\gamma$ leads to a correspondingly larger value of $T_K$ for CeSi$_3$. Our choice gives consistency between the value deduced from $\gamma$ and that deduced from $\omega_{\text{opt}}$. Next, the crystal-field splittings are set at 5 and 8 meV for CeAl$_3$, at 20 and 32 meV for CeCu$_2$Si$_2$, and at 25 and 48 meV for CeSi$_3$, using neutron-scattering results$^{21,22}$ For CeBe$_{13}$ only quasielastic scattering (i.e., no crystal-field excitations) was observed in neutron scattering$^{20}$ and the susceptibility$^{23}$ has a broad maximum characteristic$^2$ of mixed valent com-
Photoemission with the spectra. We have observed a main line for CeCu$_2$Si$_2$ Ef = -8 eV, and a value of 2.8 eV in compounds, mainly observed in conduction bands. We have used the value 0.28 eV for all four cases, as this value is consistent with the splitting observed in higher resolution photoemission spectra and is typical for cerium compounds.

As we will see, a key parameter for determining the spectral weights and $f^1$ linewidth is the bandwidth of the backconduction band. For all four compounds, the cerium hybridizes with the $s, p$ electrons on the nearest neighbor (Al, Be, or Si). Band calculations for CeAl$_3$ and CeCu$_2$Si$_2$ show that the bottom of the $s, p$ band lies at a distance $W = 9-10$ eV below the Fermi level $\epsilon_F$. We do not know of calculations for the other two compounds, but calculations for UBe$_{13}$, which should have a similar $s, p$ band to CeBe$_{13}$, give $W = 10$ eV; and we assume that the Si $s, p$ band in CeSi$_2$ should have a similar width as in CeCu$_2$Si$_2$. Therefore we have set $W = 10$ eV for all four cases, as this appears to be a characteristic value for compounds of rare earths with these $s, p$ elements. The conduction band is taken as rectangular (constant $\rho = 1/2W$) with $\epsilon_F = 0$. We fix the Coulomb correlation energy as $U = 14$ eV; this value which is somewhat larger than the accepted value $U = 6$ eV, is taken as small as possible, but larger than $W$ since the calculation requires $U > W$, as discussed above.

Finally we adjust both the bare value of the energy $E_f$ of the $f^1$ level and the average hybridization width $\Delta = Vp$ to simultaneously yield the observed (“renormalized”) $f^1$ level energy $E_f$ and the Kondo temperature. Following this we convolute the spectra with the known resolution (150 meV) using a three-point smoothing method (which correctly convolutes 6 functions and step functions). The resulting spectra are shown in Figs. 9(c) and 10. To determine the $f^1 \rightarrow f^0$ linewidth and the relative spectral weights in the two peaks, we fit the high binding energy part of the spectra to Lorentzians [dashed lines in Figs. 9(c) and 10] in precisely the same manner as for the experimental spectra. The resulting linewidths and fractional weights in the near-$\epsilon_F$ peaks are given in Table I where it can be seen that the experimental widths of the $f^1 \rightarrow f^0$ peaks are 1.5 to 2.0 times narrower than the theoretical widths, and the fractional weight in the near-$\epsilon_F$ peak is 1.65, 1.85, and 2.33 times greater in the experimental spectra for CeSi$_2$, CeAl$_3$, and CeCu$_2$Si$_2$ than in the theoretical spectra; for CeBe$_{13}$, the agreement is much better.

Somewhat better agreement with experiment can be obtained by choosing a more realistic value of $U$ (8 eV) but a less realistic value of $W$ (5 eV). The results are shown in Figs. 9(b) and 11 and Table I. For CeSi$_2$ and CeBe$_{13}$ the fractional weights in the near-$\epsilon_F$ peaks are closer to the experimental values; however, the theory continues to give too little weight for CeAl$_3$ and CeCu$_2$Si$_2$ and too large a value for the width of the $f^1 \rightarrow f^0$ excitation for all four compounds.

VI. DISCUSSION

Comparison of the experimental $4f$ spectra [Figs. 2(b), 4(c), 7(b), and 9(a)] to the theoretical spectra (Figs. 9–11) show that on the scale of the bandwidth and for the resolution utilized here, the theory underestimates the spectral weight in the near-$\epsilon_F$ peak, and it overestimates the width of the main $f^1 \rightarrow f^0$ peak (see also Table I). Visual comparison of the spectra for CeCu$_2$Si$_2$ in Fig. 9 makes this particularly clear. Reduction of $U$, with a concomitant reduction of $W$, can improve the fractional weights somewhat, but overall agreement remains visually poor.

We note in passing that our spectrum, taken for a cleaved single crystal of CeCu$_2$Si$_2$ at resolution $\sim 150$ meV, is significantly different from the low resolution ($\sim 600$ meV) spectrum$^{27}$ of polycrystalline CeCu$_2$Si$_2$ used recently by Kang et al.$^{28}$ Adding additional rounding to our spectra to simulate 600 meV resolution does not remove the discrepancy, which is mainly that our data show more weight in the near-$\epsilon_F$ peak than do the older data. It is our experience that single-crystal data are more reliable. Thus, the analysis of Ref. 28 needs to be reconsidered.

The experimental $f^1 \rightarrow f^0$ line widths all have comparable magnitude and also are comparable to those of normal rare-earth core levels (for example, the $4f$ level of

![Figure 10](image_url)

FIG. 10. Calculations of the $4f$ emission for CeAl$_3$, CeSi$_2$, and CeBe$_{13}$ using the low-order $1/N$ expansion of the Anderson impurity model. The bottom of the conduction band is set at $W = 10$ eV in accordance with known $s, p$ bandwidths of conventional rare-earth counterparts, and the Coulomb correlation is set at $U = 14$ eV. Other parameters are chosen, as described in the text, in accord with experimentally known Kondo temperatures, crystal-field and spin-orbit splittings, and final position of the $f^1 \rightarrow f^0$ peak. The dashed lines are Lorentzians, representing the $f^1 \rightarrow f^0$ emission; the binding energies are the same as for the experimental case and the widths and spectral weights are given in Table I.
TABLE I. Widths of the $f^1$ Lorentzian and ratio of the spectral weight in the near-$e_F$ emission to the total $4f$ emission for the experimental spectra and for the theoretical spectra of Figs. 9–11. The values given for the experimental spectral weight are lower limits; subtraction of surface emission would decrease the relative $f^1 \to f^0$ weight. The parameters $W$ and $U$ used in the calculation are given for each case, as are $E_f$ (bare $f$-level energy) and $\Delta$, which are adjusted to yield the experimental ("renormalized") $f$-level energy $E_f$ and Kondo temperature $T_K$. All energies are given in eV.

<table>
<thead>
<tr>
<th>Experiment</th>
<th>Width FWHM $f^1 \to f^0$</th>
<th>Fractional weight near $e_F$</th>
<th>Width FWHM $f^1 \to f^0$</th>
<th>Theory</th>
<th>Fractional weight near $e_F$</th>
</tr>
</thead>
<tbody>
<tr>
<td>CeAl$_3$</td>
<td>0.9</td>
<td>&gt; 37%</td>
<td>$W = 10$</td>
<td>$U = 14$</td>
<td>$E_f = -2.65$</td>
</tr>
<tr>
<td>$T_K = 4$ K</td>
<td></td>
<td></td>
<td>$W = 4.7$</td>
<td>$U = 8$</td>
<td>$E_f = -2.00$</td>
</tr>
<tr>
<td>CeCu$_2$Si$_2$</td>
<td>1.1</td>
<td>&gt; 49%</td>
<td>$W = 5$</td>
<td>$U = 14$</td>
<td>$E_f = -2.70$</td>
</tr>
<tr>
<td>$T_K = 8$ K</td>
<td>$E_f = -2.45$</td>
<td></td>
<td>$W = 4.0$</td>
<td>$U = 8$</td>
<td>$E_f = -2.20$</td>
</tr>
<tr>
<td>CeSi$_2$</td>
<td>1.5</td>
<td>&gt; 33%</td>
<td>$W = 10$</td>
<td>$U = 14$</td>
<td>$E_f = -2.70$</td>
</tr>
<tr>
<td>$T_K = 40$</td>
<td>$E_f = -2.35$</td>
<td></td>
<td>$W = 4.7$</td>
<td>$U = 8$</td>
<td>$E_f = -2.20$</td>
</tr>
<tr>
<td>CeBe$_{13}$</td>
<td>1.2</td>
<td>&gt; 28%</td>
<td>$W = 10$</td>
<td>$U = 14$</td>
<td>$E_f = -2.75$</td>
</tr>
<tr>
<td>$T_K = 400$ K</td>
<td></td>
<td></td>
<td>$W = 4.7$</td>
<td>$U = 8$</td>
<td>$E_f = -1.95$</td>
</tr>
</tbody>
</table>

PrAl$_3$ in Fig. 3 has a FWHM of 1.5 eV; the Lorentzian component of the $4f$ spectra for PrBe$_{13}$ has a FWHM of 1.2 eV. Such small observed linewidths are disturbing in that the prediction of a large hybridization width (essentially $28\Delta$ FWHM) for the $f^1 \to f^0$ transition is one of the most basic features of the model. [Kang et al. 38] calculated a narrow width for CeCu$_2$Si$_2$ by ignoring crystal-field splitting; in the calculation they use $\Delta \approx 0.06$ eV. For our calculation, where crystal-field splitting is not ignored, a larger value of $\Delta$ (0.09–0.10 eV) is required to reproduce $T_K$; hence, our value of the $f^1 \to f^0$ width is also larger.] It is as though the decay of the $4f$ core hole (which, given the narrow experimental width, is slower than predicted) has nothing to do with hybridization. Concerning the spectral weights, not only are the theoretical predictions too small, but the systematics are wrong. The theory predicts (Table I) a nearly linear increase in the near-$e_F$ spectral weight as the hybridization $\Delta$ increases at fixed $W$ and $U$; but the data show an inverse correlation between $\Delta$ and spectral weight near the Fermi level.

The question is, how significant are these disagreements? On the one hand, the calculation does fit the data to within an order of magnitude. Furthermore, the Anderson model is simply that—a model which is designed to give the correct low-energy behavior (which behavior exhibits universality, and hence has a certain degree of independence of the high-energy details), but which perhaps should not be expected to give good agreement on large energy scales. From this point of view, a prediction of two peaks with the correct order of magnitude for the widths and weights is sufficient. Of course, such a relaxed requirement would apply equally to any competing model of cerium photoemission. On the other hand, a good deal of emphasis$^2$–$^4$ has been given as to the degree to which the predictions of the model fit the experimental photoemission spectra. In addition, we wish to know whether the model ignores any fundamental physics; in particular, whether final-state Coulomb screening and banding effects play a more significant role in the spectra than implied by use of the impurity model. For
these reasons, we take the disagreements seriously; and in what follows, we examine several experimental and theoretical problems which will need to be resolved in order to make progress on these issues.

On the experimental side we first consider the method of determining the non-4f background. There are at least three methods that are widely used. The first is to compare spectra taken on (120 eV) and off (112 eV) the giant 4f resonance; the second is to compare spectra taken above (e.g., at $h\nu=40$ eV) and below (e.g., at $h\nu=20$ eV) the presumed threshold (near $h\nu=30$ eV) for the onset of Ce 4f emission; the third is to compare the emission for the Ce compound to that of a La compound. As we have shown, the first method ignores the fact that the 5d electrons contribute at least 30% of the spectrum at 120 eV. Applying this approach to CeCu$_2$Si$_2$, by simply subtracting the two spectra in Fig. 8(a), gives a non-4f fraction of 17%; it also leaves a significant peak at 1.67 eV in the resulting spectrum, which we believe is due to non-4f emission, rather than 4f emission as implied by this method. The second method can underestimate the amount of 4f emission at 20 eV (it does not actually vanish) and 5d emission at 40 eV. The third (which is the method we use) overlooks possible differences between the Ce and La 5d spectra.

All three methods require additional input as to how to correctly normalize between comparison spectra. We have chosen to normalize our spectra to core levels whenever possible; in doing so we have ignored possible photon energy dependence of the core-level emission over the limited range 116–126 eV. An alternative method is to normalize to incident photon flux (by measuring mesh current); however, in polycrystals this can give very bad results, due to unpredictable cleave dependencies.

A second source of uncertainty is that the photon energy dependence of the matrix elements for optical absorption may differ for different energy regions of the 4f spectrum. In particular, the branching ratio of the near-$e_F$ peaks to the $f^1\rightarrow f^0$ peak may vary with photon energy; indeed, this is known to occur in the vicinity of the 4d resonance. Examination of Fig. 5 demonstrates clearly that the branching ratio for the 5d states varies considerably between 26 and 120 eV. Hence, there is no a priori reason for assuming that the 4f branching ratio is constant over any energy range. This means that the spectra taken at a particular photon energy may not be simply proportional to the spectral density.

At the photon energy that we use (120 eV) the spectra are highly surface sensitive. In our analysis, however, we have ignored the distinction between surface and bulk emission. Recently it has been shown that the surface emission occurs primarily in the $f^1\rightarrow f^0$ peak near 2 eV. For the compound CeRh$_3$ (which has a very large $T_K>1000$ K) there is substantial weight in the 2-eV feature at 120 eV, but spectra taken at $h\nu=848$ eV show very little weight in the $f^1\rightarrow f^0$ peak. The argument is that for the larger photon energy the spectra should be dominated by bulk emission. If all this is true, then were we to account for surface emission in our spectra, there would be less weight in the bulk 2-eV feature, and hence the fractional weight in the near-$e_F$ peak would be even larger than our results suggest. In that sense, our results are lower limits for the fractional spectral weight. Hence, the disagreement with theory is even greater than we have stated above. It should be pointed out that, given the lack of knowledge of the 4f branching ratio discussed above, it is possible that some of the differences between the 120- and 844-eV spectra may not be due to such a surface effect.

The greatest uncertainty in theoretical spectra arises from the lack of knowledge of the energy dependence of the combined hybridization matrix element and conduction-band density of states $\Delta(E)=V^2(E)p(E)$. The line shape of the 4f spectrum over the full energy range is extremely sensitive to the choice of the energy dependence of this parameter. (For this reason, it is all the more surprising that the $f^1\rightarrow f^0$ peak observed experimentally is for all compounds a Lorentzian with FWHM $\approx 1$ eV.) We have chosen to ignore the energy dependence $\Delta(E)=\Delta$ independent of energy and have chosen to cut the conduction band off 10 eV below the Fermi level, since this is to be the appropriate bandwidth for compounds of rare earths with the $s,p$ elements Al, Be, and Si. A smaller choice of bandwidth causes more $f^2$ weight to pile up near the Fermi energy and changes the ratio of near-$e_F$ emission to 2-eV emission. In addition, if the bandwidth is made so small that the bare 4f level resides close to the bottom of the band, then the $f^1\rightarrow f^0$ line shape alters dramatically. In this model $U=\infty$ NCA calculation? for CeSi$_2$, the background band was assumed to be Gaussian, with a small bandwidth (3 eV FWHM); the final $f$ level is pushed well out in the tail where the density of states is low and hence the hybridization is weak. For this choice of band shape, the resulting $f^1\rightarrow f^0$ width is small and comparable to experiment. However, in our opinion, this is a very unrealistic choice of band shape for CeSi$_2$, and it strengthens our point that fits in the literature are strongly dependent on the choice of $\Delta(E)$, which is essentially unknown.) A variety of recent studies$^{28,32-35}$ attempt to derive $\Delta(E)$ and other parameters of the model from first-principles local-density approximation (LDA) electronic structure calculations. However, there is no general consensus on how to achieve this, since there is uncertainty about the extent to which the LDA already includes the many-body effects which are to be calculated from the resulting model. The issues at hand (concerning the line shape on the scale of the bandwidth) cannot be resolved without solution of this problem.

A second source of uncertainty is the convergence of the large $N$ expansion and the degree to which the truncated expansion adequately represents the spectrum of the Anderson impurity model. Furthermore, the generalization from the infinite $U$ limit to the finite values of $U$ also raises questions; this generalization does not include the effects of configurations $f^n$ with $n>2$, and the nature of the errors introduced by the decoupling approximation remains unknown.

Given all this, the possibility remains that adequate treatment of the photoemission spectra may require inclusion of other physics. Coulomb screening of the final-state hole has been utilized in an attempt to explain the
large differences between Kondo temperatures determined from thermodynamic measurements and alternatively from the photoemission spectra. In addition, it is known that in certain limits a two-peaked spectrum ("well-screened" and "poorly screened" final states) is expected when such screening is present. When such screening is added to the Anderson model, the relation between high-energy scales and low-energy scales is known to be affected. A possibility is that some of the extra weight that we observe in the near-$e_F$ peak arises from this source. It is very probable that the extra width in the near-$e_F$ peak, which is apparent in higher resolution spectra, arises from this source. In this regard the extra $f$ emission seen in PrBe$_3$ [Fig. 6(b), long-dashed line] is suggestive. Similar peaks in several other Pr compounds have been interpreted as evidence for the existence of two final-state screening channels. Indeed, the existence of two $4f$ peaks in Pr compounds with large hybridization (as estimated by the large Kondo temperatures $T_K > 100$ K of the corresponding Ce compounds) now appears to be more the rule than the exception; and this supports the view that the second peak cannot be universally obtained from the Anderson model as suggested in Refs. 1 and 8.

Finally, we raise the possibility that the enhanced spectral weight at the Fermi level, which approaches 50% in these compounds, may mean that the Ce compounds reside closer to the band limit than is implied by use of an essentially local impurity model. In this approach, the 2-eV peak would arise in the same way as the 6-eV satellite in nickel. While the physics of this satellite still involves the Coulomb correlation, there are significant differences in the way this approach (as opposed to the Anderson model approach) generates the satellite. The possibility that the band limit is more appropriate is supported by the recent work mentioned above for CeRh$_3$ where subtraction of surface emission so drastically reduces the weight in the 2-eV peak that the resulting spectrum is simply that expected for an $f$ band. While the compounds studied here have smaller Kondo temperatures than that of CeRh$_3$, it is quite plausible that correct subtraction of the surface peak weight would lead to spectra with a much larger peak at $e_F$ and a much smaller satellite at 2 eV.

All of this suggests that photoemission in cerium compounds is very much an open problem, and that future work, both experimental and theoretical, is in order. On the experimental side, the greatest need is for use of high-quality, well characterized single-crystal surfaces. Studies over a broad range of photon energies, coupled with studies of the take-off angle dependence of the emission, are needed to help determine the degree of surface and bulk contribution to the spectra. For similar reasons studies of the photon energy dependence of both the $4f$ and non-$f$ emission over a broad (20–1000 eV) photon energy range in both Ce and related La and Pr compounds are needed to determine the non-$f$ emission and the $4f$ branching ratios. Further thought should be given to normalization. Ongoing improvements in experimental resolution can also be expected to be important.

On the theoretical side, the problem of determining the parameters of the model [such as $\Delta(\epsilon)$] from first principles presents an important and difficult challenge. Improvements in our understanding of the effect of approximations on the predictions of the Anderson model would also be helpful; in this regard, the recent Monte Carlo work is very promising. Further work to assess the importance of Coulomb screening and to determine its effect on the predictions of the Anderson model is necessary.

Finally, use of many-body theories which use band theory as a starting point may well be necessary to understand valence-band photoemission in cerium compounds.

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PHOTOEMISSION SPECTRA OF CeAl$_3$, CeBe$_{13}$, CeSi$_2$ . . .