## Multipole character of the proposed 220 eV transition in <sup>229</sup>Pa

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Internal conversion coefficients (ICC's) have been calculated for protactinium and transition energies between 170 eV and 10 keV. The ICC's for E1 multipolarity show an unusual behavior, which cannot be approximated by an exponential dependence on the transition energy, whereas the ICC's for M1 and E2multipolarities closely follow such a dependence. Using the newly calculated ICC's the unusually strong "enhancement" of a possible 220 eV E1 transition in <sup>229</sup>Pa proposed earlier is reduced by a factor of  $\sim 5$ , yielding an induced electric dipole moment similar to that observed in the neighboring octupoledeformed isotopes.

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An excited level was proposed by Ahmad et al. [1] at 220 eV in <sup>229</sup>Pa, with spin parities of  $5/2^+$  and  $5/2^-$  assigned to the ground state and the 220-eV state, respectively. These levels would thus form an almost degenerate parity doublet, as is characteristic for nuclei with octupole deformation, a nuclear shape that is expected theoretically in the mass region around A = 225. Ahmad et al. also measured a half-life of  $420\pm30$  ns, assigned to the 220 eV level, from which they derive an E1 transition rate of  $2.5 \times 10^{-2}$  W.u. for the proposed  $5/2^- \rightarrow 5/2^+$ transition. This transition rate is very fast compared to the E1 rates usually observed between one-quasiparticle states. This led Ahmad et al. in a later publication [2] to propose a collective enhancement of the E1 transitions in octupole-deformed nuclei. This idea has turned out to be very fruitful (see, e.g., Ref. [3]), and in fact such "enhanced" E1 transitions are now considered as fingerprints for strong octupole correlations.

Recently Grafen et al. [4] have shown that part of the experimental evidence that led to the spin-parity assignment of  $5/2^{-1}$  to the proposed 220 eV level was wrong. It was also argued in this paper that the proposed enhancement of the  $5/2^- \rightarrow 5/2^+$  transition is much larger than expected from octupole correlations. The collective E1transitions in octupole-deformed nuclei are usually interpreted as resulting from an intrinsic electric dipole moment  $\mathbf{e} \cdot \mathbf{D}_0$  induced by the polarizing electric field of the nuclear octupole deformation. For the proposed 220-eV E1 transition in <sup>229</sup>Pa one has  $B(E1, 5/2^{-} \rightarrow 5/2^{+}) = (15/28\pi)e^2D_0^2$ , and with the transition rate given by Ahmad et al. a value of  $D_0 \approx 0.6$  fm is obtained. This would be the largest intrinsic electric dipole moment observed in the region of octupole deformation around A = 225, and it would be ~5 times larger than the  $D_0$  for the even-core nucleus <sup>228</sup>Th [3]. Such a large polarization of the nucleus <sup>229</sup>Pa, as compared to its even core, seems unusual, although it might possibly be expected on theoretical grounds [5].

Since the identification of the parity doublet in <sup>229</sup>Pa is

not established experimentally, it is important to consider the experimental evidence in favor of such a doublet. In the present work we address the extraction of the E1transition rate from the measured half-life. The problem here is the internal conversion rate, which dominates the transition. From the E1 rate quoted by Ahmad *et al.* we deduce that these authors used an internal conversion coefficient (ICC) of  $\alpha_{tot}(E1) \approx 1600$ , although it is not clear from their paper how this value was obtained.

The ICC's for very low transition energies are not available in current tabulations. In the most recent tables [6] the conversion coefficients are listed for energies above 2.2 keV. Since for E1 multipolarity  $\alpha_{tot}$  varies roughly as  $E_{\gamma}^{-3}$ , an extrapolation over three orders of magnitude is needed in the present case. Moreover, it is known that the ICC's often exhibit "strange" behavior when the transition energy approaches threshold. It is thus clear that a direct calculation of the conversion coefficients in the low-energy region is necessary to obtain reliable values for the total ICC of the 220-eV transition in <sup>229</sup>Pa. To our knowledge the internal conversion coefficients were never calculated for such low energies, except for the 77-eV E3 transition in <sup>235</sup>U [7,8].

The following physical assumptions were used in our calculations. Both the bound-electron and free-electron wave functions are solutions of the Dirac equation with the relativistic Hartree-Fock-Slater potential [9]. The nucleus is described by a Fermi distribution of the charge with R = 6.7 fm. The kinetic energies of the conversion electrons were determined from the experimental binding energies [10]. Special attention was paid to correct normalizations of the wave functions for the free electrons with very low kinetic energies. The numerical accuracy of the calculated ICC's is estimated to be about 6%. In accordance with Rösel *et al.* [6], our calculation corresponds to the lowest-nonvanishing order of the perturbation theory of quantum electrodynamics.

The results for the  $O_4$  subshell and the total ICC's for E1 multipolarity are shown in Fig. 1. The fine energy

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FIG. 1. Internal conversion coefficients for low-energy transitions of E1 multipolarity in protactinium. The points indicate the energies for which the coefficients were calculated in the present work, the horizontal line denotes the energy region covered by the tables of Rösel *et al.* [6]. Displayed are (a) the total ICC's, (b) the sum over the subshells  $O_2$  through  $Q_1$  playing the role of the total ICC's for  $\gamma$ -ray energies between 245 and 305 eV, and (c) the ICC's for the  $Q_4$  subshell. The bump on the curve for the total ICC's at ~0.5 keV is due to internal conversion in the  $N_6$  and  $N_7$  subshells (4*f* electrons) which is important only for low-energy E1 transitions [11].

mesh of our calculations enabled us to follow a complex behavior of the conversion coefficients to energies far below the 2.2 keV threshold of existing tabulations. We extended the calculations for E1 multipolarity, and for the  $M_1$  through  $Q_1$  atomic subshells, up to 10 keV to compare our values to those of Rösel et al. [6]. In all cases the conversion coefficients agreed within a few percent except for the  $N_5$  subshell, where the deviation was 11%. As demonstrated in Fig. 1 for the  $O_4$  subshell, the ICC's exhibit minima around  $E_{\gamma} \sim 1$  keV which are most shallow for the  $s_{1/2}$  subshells and sharpen with increasing angular momentum of the subshell. We verified that these resonancelike structures are due to cancellations in the leading matrix elements demonstrated earlier for E2multipolarity at higher energies [12]. It is also obvious from our calculation that an extrapolation of ICC's from existing tabulations to energies below  $\sim 1 \text{ keV}$  does not lead to reliable values for E1 multipolarity (see Fig. 1).

In the region of interest,  $170 \le E_{\gamma} \le 270$  eV, the calculated total ICC's for *E*1 multipolarity vary approximately as  $E_{\gamma}^{-5}$  (to  $\le 2\%$ ) with proportionality constants as given in Table I. With the 420-ns half-life of the proposed 220-eV level we derive an induced electric dipole moment of  $D_0 = 0.27 \pm 0.04$  fm, where the error includes uncer-

TABLE I. Calculated ICC's for the 220 $\pm$ 50 eV transition in  $^{229}\text{Pa}.$ 

$E_{\gamma}$ (keV)	$E_{\gamma}^5 \alpha_{\rm tot}(E1)$	$E_{\gamma}^5 \alpha_{\rm tot}(E2)$	$E_{\gamma}^{3}\alpha_{\rm tot}(M1)$
0.170-0.188	2.61	$2.22 \times 10^{8}$	$1.18 \times 10^{4}$
0.188-0.245	3.87	$8.89 \times 10^{8}$	$1.22 \times 10^{4}$
0.245-0.270	5.55	$14.2 \times 10^{8}$	$1.74 \times 10^{4}$

tainties in both the half-life and the transition energy as well as numerical uncertainty of the ICC's. This value is much closer to the experimental result of  $0.11\pm0.02$  fm of the <sup>228</sup>Th core nucleus than that given by Ahmad *et al.* [1].

As pointed out by Ahmad *et al.* [1] the observed halflife is consistent with M1 multipolarity for the 220-eV transition. We have therefore also calculated the conversion coefficients for M1 and E2 multipolarity. In these cases the calculated ICC's closely follow a straight line in the log-log plot, even in the very-low-energy region. The total conversion coefficients are proportional to  $E_{\gamma}^{-3}$  and  $E_{\gamma}^{-5}$  for M1 and E2 multipolarity, respectively. The results of the calculation in the energy region of interest in the present work are included in Table I. For the reduced transition probability of a  $220\pm50$ -eV transition one would obtain  $B(M1)=(6.6\pm1.7)\times10^{-3}$   $(eh/2Mc)^2$ and  $B(E2)=(6.1\pm0.8)\times10^3 e^2$  fm<sup>4</sup> or  $(1.2\pm0.3)\times10^3 e^2$ fm<sup>4</sup> for  $E_{\gamma}$  below or above the  $O_3$  binding energy, respectively.

In summary, our calculations show that for E1 transitions with energies below  $\sim 1$  keV the ICC's cannot be derived by an extrapolation from higher energies, whereas such an extrapolation yields reasonable values for M1 and E2 multipolarities, at least for heavy elements. For a possible 220-eV E1 transition in <sup>229</sup>Pa the half-life of 420 ns gives an enhancement of a factor of  $\sim 2$ for the induced electric dipole moment, as compared to the <sup>228</sup>Th core nucleus, which might be expected on theoretical grounds due to the polarization of the core by the extra proton [5].

Using modern electrostatic spectrometers one should be able to examine the conversion-electron spectrum of the <sup>229</sup>U $\rightarrow$ <sup>229</sup>Pa decay at very low energies. An observation of the conversion electrons of the proposed 220-eV transition would not only provide uncontested evidence for its existence, but would also allow one to determine its multipolarity, even from a rough measurement of conversion-electron intensity ratios. For example, for  $E_{\gamma} \sim 220$  eV, the  $(P+Q)/(O_4+O_5+O_6)$  ICC ratio takes on values of ~0.3, 11.3, and 114 for E1, E2, and M1 multipolarity, respectively. Moreover, a precise measurement of the conversion-line intensities could serve as a test for some of the higher-order effects in the internal conversion process [8,13].

After completion of the present work we learned that Band *et al.* [14] calculated ICC's for 220-eV E1, M1, and E3 transitions in protactinium using several atomic models. Their total ICC's corresponding to our approach agree with our values to within 9%.

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