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High-energy x-ray core-level spectroscopy for $PrFe_4P_{12}$

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In our previous study, we observed a very strong Pr 4f spectral intensity just below the Fermi level in PrFe₄P₁₂ by means of the bulk-sensitive high-resolution photoemission spectroscopy [1]. The strong spectral intensity is interpreted as a Kondo resonance arising from the hybridization between the Pr 4f and P 3p states. The rare-earth 3d core-level spectroscopy by means of the hard x-ray light source enables us to investigate not surface-sensitive but bulk-sensitive Pr 4f electronic structures because of the large escape depth of photoelectrons in solid as well as the valence band photoemission spectroscopy by means of the soft x-ray light source. In order to obtain a deeper insight into the bulk electronic structures of PrFe₄P₁₂, we have carried out the x-ray core-level spectroscopy using the high-energy synchrotron radiation light source up to $h\nu = 5450$ eV at European Synchrotron Radiation Facility (ESRF).

The contribution of $|f^3\rangle$ configuration is obviously seen in Pr 3d core-level spectra of PrFe₄P₁₂, PrSn₃, and Pr metal. Among them, PrFe₄P₁₂ has much larger $|f^3\rangle$ intensity than others due to the stronger *c*-*f* hybridization. Configuration interaction cluster model calculations reproduce the Pr 3d core-level spectra of both well hybridized and localized Pr compounds. Difference of the estimated 4*f* electron numbers between high and low h ν s indicates the larger *c*-*f* hybridization strength in the bulk than near the surface. Furthermore, the results of the calculations predict that the Kondo effect observed in PrFe₄P₁₂ and PrSn₃ originates from the mixing of $|4f^n\rangle$ and $|4f^{n+1}\rangle$ configurations in contrast with the mixing of $|4f^n\rangle$ and $|4f^{n-1}\rangle$ configurations in Ce (*n*=1) compounds.

 A. Yamasaki, S. Imada, T. Masuda, T. Nanba, A. Sekiyama, H. Sugawara, T. D. Matsuda, H. Sato, C. Sekine, I. Shirotani, H. Harima, and S. Suga, Acta phys. Polonica B 34, 1035 (2003).