

Electronic structure, magnetic ordering and X-ray magnetic circular dichroism in $\text{La}_{1-x}\text{Pr}_x\text{Co}_2\text{P}_2$ phosphides

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The electronic structure and magnetic ordering in $\text{La}_{1-x}\text{Pr}_x\text{Co}_2\text{P}_2$ ($x = 0, 0.25, \text{ and } 1$) phosphides have been studied theoretically using the fully relativistic spin-polarized Dirac linear muffin-tin orbital (LMTO) band-structure method. The X-ray absorption and X-ray magnetic circular dichroism spectra at the $\text{Co}L_{2,3}$ and $\text{Pr}M_{4,5}$ edges have been investigated theoretically within the framework of the LSDA+ U method. The core-hole effect in the final state as well as the effects of the electric quadrupole E_2 and magnetic dipole M_1 transitions have been investigated. Good agreement with experimental measurements has been found.

Key words: *strongly correlated systems, band structure, magnetic moments, X-ray magnetic circular dichroism*

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1. Introduction

Ternary intermetallics AT_2X_2 , where A – alkali, alkali-earth, rare-earth, or actinide metal, T – transition metal, and X – nonmetal, often demonstrate intrinsically complex magnetic structures and a wide variety of physical properties. They belong to the ThCr_2Si_2 structure type and can be conveniently represented as materials built by stacking covalently bonded transition metal-metalloid T_2X_2 layers, made of edge-sharing TX_4 tetrahedra, with ionic A atoms. This body-centered tetragonal structure is rather simple with only one crystallographic site for each atomic species and only one variable positional parameter, the z value of the X site. Due to the great variety of possible elements in this structure, over 1000 different compounds of this type are known already [1]. Some of these compounds exhibit fascinating physical properties such as superconductivity, valence fluctuations, local and itinerant magnetism. Interest to these materials has been reinvigorated recently due to the discovery of non-Fermi-liquid behavior in YbRh_2Si_2 [2] and high-temperature superconductivity in K-doped BaFe_2As_2 [3, 4]. In addition, materials with layered arrangements of magnetic moments are also of great interest because they tend to exhibit a highly anisotropic magnetic behavior. The magnetic properties of the silicides and germanides possessing this structure were studied by many groups [5–11]. Besides superconductivity, AT_2X_2 phases exhibit a plethora of other phase transition phenomena. Thus, SmMn_2Ge_2 becomes ferromagnetic below 348 K and then undergoes a transition to an antiferromagnetic state at 196 K, followed by a re-entrant ferromagnetic transition at 64 K [7]. This peculiar magnetic behavior stems from the layered structure and the presence of two magnetic sublattices in these materials and was later observed for a number of rare-earth manganese germanides and silicides [8–11]. Such sequential magnetic transitions have not been observed in isostructural phosphides which might explain why their magnetic properties have received somewhat less attention than the tellurides. The isotypic phosphides were prepared much later [12] and the investigations of their magnetic properties have started only recently.

In the compounds RT_2P_2 (R = lanthanides, T = Fe, Co, Ni) with the trivalent rare earth elements, magnetic moments at the transition metal sites were observed only for the cobalt containing compounds

[13–15]. Reehuis et al. carried out a comprehensive study of the structural and magnetic properties of ternary RCo_2P_2 materials [15–17]. Recently, Kovnir et al. [18–20] and other groups [21, 22] have demonstrated that the magnetic properties of the ThCr_2Si_2 type phosphides can be as rich and diverse as those of the aforementioned silicides and germanides, provided that proper iso- and aliovalent substitutions are made to modify the crystal and electronic structures of the materials. Ternary cobalt phosphides RCo_2P_2 appear to be on the verge of magnetic instability [20]. Indeed, LaCo_2P_2 is characterized by ferromagnetic ordering of Co magnetic moments at 132 K [14, 18], while other representatives of this family ($\text{R} = \text{Ce}, \text{Pr}, \text{Nd}, \text{Sm}$) order antiferromagnetically at around room temperature [15] (with the exception of the Ce-containing compound, which shows an antiferromagnetic transition at 440 K purportedly due to the mixed valence of Ce). A substantial difference is also observed in the crystal structures of these compounds [23]. In LaCo_2P_2 , the $[\text{Co}_2\text{P}_2]$ layers are far from each other, with the interlayer P–P distance of 3.16 Å indicating essentially no bonding between the phosphorus atoms. By contrast, other RCo_2P_2 structures show a weakly covalent P–P interaction at relatively small interplanar P–P distances (2.57 Å in PrCo_2P_2 [18]). Thus, a drastic change in the magnetic behavior is accompanied by the formation of the P–P bonding pathway between the $[\text{Co}_2\text{P}_2]$ layers.

Kovnir et al. [18] embarked on the study of structural, magnetic, and electronic properties of quaternary compositions $\text{La}_{1-x}\text{Pr}_x\text{Co}_2\text{P}_2$ ($0 \leq x \leq 1$). The choice of Pr was based on the fact that it does not exhibit a mixed valence like Ce, and, therefore, is the closest rare-earth neighbor of La in both the size and the formal ionic charge. They focused on the drastically different magnetic properties of LaCo_2P_2 and PrCo_2P_2 , which exhibit ferromagnetic ($T_C = 132$ K) and antiferromagnetic ($T_N = 305$ K) ordering in the Co sublattice, respectively [18]. In both cases, a ferromagnetic arrangement of magnetic moments within the square plane of Co atoms is observed. In LaCo_2P_2 , the Co moments are aligned in-plane and parallel to the moments in the other layers, leading to the ferromagnetic ground state [17]. In PrCo_2P_2 , they are aligned perpendicular to the plane (along the c axis), but the antiparallel magnetic coupling between the neighboring planes results in antiferromagnetism [16]. In [24], magnetic behavior of $\text{La}_{0.75}\text{Pr}_{0.25}\text{Co}_2\text{P}_2$ was investigated by a combination of magnetic measurements, magneto-optical imaging, neutron diffraction, and X-ray absorption spectroscopy, including X-ray magnetic circular dichroism.

The aim of this paper is a theoretical study, from the first principles, of the electronic structure, magnetic ordering and X-ray magnetic circular dichroism in $\text{La}_{1-x}\text{Pr}_x\text{Co}_2\text{P}_2$ ($x = 0, 0.25, \text{ and } 1$) compounds. The energy band structure of $\text{La}_{1-x}\text{Pr}_x\text{Co}_2\text{P}_2$ ($x = 0, 0.25, \text{ and } 1$) compounds is calculated within the *ab initio* approach taking into account strong electron correlations by applying the local spin-density approximation (LSDA) to the density functional theory supplemented by a Hubbard U term (LSDA+ U) [25]. The paper is organized as follows. The computational details are presented in section 2. Section 3 presents the electronic structure, XAS and XMCD spectra of $\text{La}_{1-x}\text{Pr}_x\text{Co}_2\text{P}_2$ ($x = 0, 0.25, \text{ and } 1$) compounds calculated in the LSDA+ U approximation. The theoretical results are compared to experimental measurements. Finally, the results are summarized in section 4.

2. Computational details

X-ray magnetic circular dichroism. The absorption coefficient $\mu_j^\lambda(\omega)$ for incident X-ray of polarization λ and photon energy $\hbar\omega$ can be determined as the probability of electronic transitions from initial core states with the total angular momentum j to final unoccupied Bloch states:

$$\mu_\lambda^j(\omega) = \sum_{m_j} \sum_{n\mathbf{k}} \left| \langle \Psi_{n\mathbf{k}} | \Pi_\lambda | \Psi_{jm_j} \rangle \right|^2 \delta(E_{n\mathbf{k}} - E_{jm_j} - \hbar\omega) \theta(E_{n\mathbf{k}} - E_F), \quad (2.1)$$

where Ψ_{jm_j} and E_{jm_j} are the wave function and the energy of a core state with the projection of the total angular momentum m_j ; $\Psi_{n\mathbf{k}}$ and $E_{n\mathbf{k}}$ are the wave function and the energy of a valence state in the n -th band with the wave vector \mathbf{k} ; E_F is the Fermi energy. Π_λ is the electron-photon interaction operator in the dipole approximation

$$\Pi_\lambda = -e\boldsymbol{\alpha}\mathbf{a}_\lambda, \quad (2.2)$$

where $\boldsymbol{\alpha}$ are the Dirac matrices, \mathbf{a}_λ is the λ polarization unit vector of the photon vector potential, with $a_\pm = 1/\sqrt{2}(1, \pm i, 0)$, $a_\parallel = (0, 0, 1)$. Here, + and – denotes, respectively, left and right circular photon polarizations with respect to the magnetization direction in the solid. The X-ray magnetic circular and linear

dichroism are given by $\mu_+ - \mu_-$ and $\mu_{\parallel} - (\mu_+ + \mu_-)/2$, respectively. Detailed expressions of the matrix elements in the electric dipole approximation may be found in [26–29]. The matrix elements due to the magnetic dipole and electric quadrupole corrections are presented in [29].

Magnetocrystalline anisotropy energy. The internal energy of a ferromagnetic material depends on the direction of spontaneous magnetization. The magnetocrystalline anisotropy energy (MAE), which possesses the crystal symmetry of a material, is a part of this energy. The MAE is an important property that describes the tendency of the magnetization to align along specific spatial directions rather than to randomly fluctuate over time. The MAE determines the stability of magnetization in bulk as well as nanoparticle systems. Extensive studies on ferromagnetic bulk materials and thin films have highlighted the MAE dependence on crystal symmetry and atomic composition. While the exchange interaction among electron spins is purely isotropic, the orbital magnetization connects the spin magnetization to the atomic structure of a magnetic material via the spin-orbit interaction, giving rise to the magnetic anisotropy [30]. The calculation of the magnetocrystalline anisotropy energy has been a long-standing problem. A first theory of the MAE in Fe and Ni was formulated by Brooks [31] and Fletcher [32], who emphasized that the energy band picture in which the effect of spin-orbit (SO) coupling is taken into account in a perturbative way could provide a coupling of the magnetization orientation to the crystallographic axes of approximately the right order of magnitude. In this pioneering work, the band structure was oversimplified to three empirical bands [31, 32]. Recent investigations [33–37] elaborated the MAE problem using *ab initio* calculated energy bands obtained within the local-spin density approximation to the density functional theory. Although it is beyond doubt that LSDA energy bands are superior to empirical bands, it turned out that the calculation of the MAE from first principles poses a great computational challenge. The prime obstacle is the smallness of the MAE, which is of only a few meV/atom, a value that ought to result from the difference of two total energies for different magnetization directions, which are both of the order of 10^4 eV/atom. Due to this numerical problem, it remained at first unclear if the LSDA could at all describe the MAE correctly, since the wrong easy axis was obtained for hcp Co and fcc Ni [33]. Recent contributions aimed consequently at improving the numerical techniques [35, 38], with the result that the correct easy axis was obtained for hcp Co, but not for fcc Ni [35]. Halilov et al. [36] reported an *ab initio* investigation of the magnetocrystalline anisotropy energy in bcc Fe and fcc Co and Ni. They introduced a spin-orbit scaling technique that yielded the correct easy axis for Fe and Co, but a vanishing MAE for Ni.

For the material exhibiting uniaxial anisotropy, such as a hexagonal crystal, the MAE can be expressed as [39]

$$E(\theta) = K_1 \sin^2 \theta + K_2 \sin^4 \theta + K_3' \sin^6 \theta + K_3 \sin^2 \theta \cos [6(\phi + \psi)] + \dots, \quad (2.3)$$

where K_i is the anisotropy constant of the i th order, θ and ϕ are the polar angles of the Cartesian coordinate system where the c axis coincides with the z axis (the Cartesian coordinate system is chosen so that the x axis is rotated through 90° from the a hexagonal axis) and ψ is a phase angle.

Both the magnetic dipole interaction and the SO coupling give rise to the MAE, the former contributing only to the first-order constant K_1 . Here, we deal with the MAE caused only by the SO interaction. Both the magneto-optical effects and the MAE have a common origin in the SO coupling and exchange splitting. Thus, a close connection between the two phenomena seems plausible.

In this paper, the MAE is defined as the difference between two self-consistently calculated fully relativistic total energies for two different magnetic field directions, $E(\theta) - E_{(001)}$.

Crystal structure. LaCo₂P₂ and PrCo₂P₂ belong to the ThCr₂Si₂ body-centered tetragonal structure type (figure 1). The space group is I4/mmm (No. 139) with Pr (La) at the $2a$ positions (0, 0, 0), Co at the $4a$ positions (0, $\frac{1}{2}$, $\frac{1}{4}$) and P at the $4e$ positions (0, 0, z); $z = 0.3697$ and 0.3568 in PrCo₂P₂ and LaCo₂P₂, respectively [16, 17]. In this structure, the transition metal atoms (Co) form planar square nets, and the nonmetal atoms (P) cap the centers of the squares above and below the planes in a chessboard-like fashion. The resulting [Co₂P₂] layers are separated by layers of rare-earth metal cations (Pr or La). The Pr atom in PrCo₂P₂ has 8 P nearest neighbors at the distance of 3.0329 Å and 8 Co atoms at the 3.1094 Å distance, the shortest P–P distance is equal to 2.5247 Å. In LaCo₂P₂, the La atom has 8 P nearest neighbors

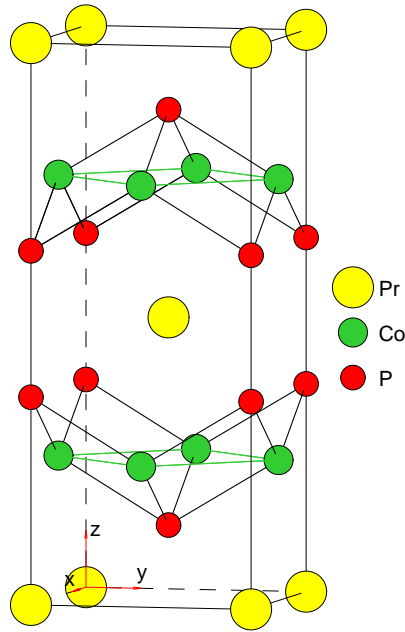


Figure 1. (Color online) Schematic representation of the PrCo_2P_2 structure.

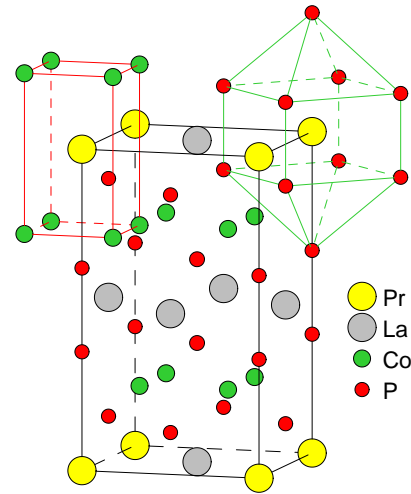


Figure 2. (Color online) Schematic representation of the $\text{La}_{1-x}\text{Pr}_x\text{Co}_2\text{P}_2$ ($x = 0.25$) structure.

at the distance of 3.1265 Å and 8 Co atoms at the 3.3551 Å distance, the P–P distance in LaCo_2P_2 is much larger and is equal to 3.1621 Å.

The calculations of the energy band structure of $\text{La}_{1-x}\text{Pr}_x\text{Co}_2\text{P}_2$ ($x = 0, 0.25$, and 1) compounds were performed for $a \times a \times 2c$ supercells of the tetragonal structure with space group of $P4/mmm$ (No. 123). The structure refinement parameters of $\text{La}_{1-x}\text{Pr}_x\text{Co}_2\text{P}_2$ compounds are presented in [18]. The crystal structure of $\text{La}_{0.75}\text{Pr}_{0.25}\text{Co}_2\text{P}_2$ is presented in figure 2. The Pr atom in this compound has 8 P nearest neighbors at the distance of 3.0329 Å and 8 Co atoms at the 3.1094 Å distance. The Co atoms are surrounded by 4 P atoms at the distance of 2.2688 Å and 4 Co atoms at the distance of 2.7577 Å, 4 La atoms are at the 3.1094 Å distance from Co.

Calculation details. The calculations presented in this work were performed using the spin-polarized fully relativistic linear muffin-tin orbital (SPR LMTO) method [40–42] for the experimentally observed lattice constants: $a = 3.6$ Å, $c = 9.688$ Å for PrCo_2P_2 [16], $a = 3.8145$ Å, $c = 11.041$ Å for LaCo_2P_2 [17], and $a = 3.8260$ Å, $c = 10.9031$ Å for $\text{La}_{1-x}\text{Pr}_x\text{Co}_2\text{P}_2$ ($x = 0.25$) [24]. The basis consisted of the Pr (La) s , p , d , and f ; Co s , p , and d ; P s , p , and d LMTO's. The \mathbf{k} -space integrations were performed with the improved tetrahedron method [43], and the self-consistent charge density was obtained with 1063 irreducible \mathbf{k} -points. To attain good convergence in total energy, a large number of \mathbf{k} points should be used in calculations. To resolve the difference in total energies and to investigate the magnetocrystalline anisotropy, we used 13824 \mathbf{k} points in the irreducible part of the Brillouin zone, which corresponds to 82944 tetrahedra in the full zone.

The X-ray absorption and dichroism spectra were calculated taking into account the exchange splitting of core levels. We also take into account the core-hole effect in the final state using the supercell approximation. The similar approximation has been used by several authors [44, 45]. The finite lifetime of a core hole was accounted for by folding the spectra with a Lorentzian. The widths of core level spectra $\Gamma_{L_2} = 1.13$ eV and $\Gamma_{L_3} = 0.47$ eV for Co, and $\Gamma_{M_{4,5}} = 0.75$ eV for Pr were taken from [46]. The finite apparatus resolution of a spectrometer was accounted for by a Gaussian with the width of 0.6 eV.

It is well known that the LSDA fails to describe the electronic structure and properties of the systems in which the interaction among the electrons is strong. In recent years, more advanced methods of electronic structure determination such as the LSDA plus self-interaction corrections (SIC–LSDA) [29, 47], the

LSDA+ U [25] method, the GW approximation [48] and the dynamical mean-field theory (DMFT) [49–51] have sought to remedy this problem and have met considerable success. The LSDA+ U method is the simplest among them and most frequently used in the literature. We used the “relativistic” generalization of the LSDA+ U method which takes into account the spin-orbit coupling so that the occupation matrix of localized electrons becomes non-diagonal in spin indexes. This method is described in detail in our previous paper [52] including the procedure to calculate the screened Coulomb U and exchange J integrals, as well as the Slater integrals F^2 , F^4 , and F^6 .

The screened Coulomb U and exchange J integrals enter the LSDA+ U energy functional as external parameters and should be determined independently. The value of U can be estimated from the photoemission spectroscopy (PES) and X-ray Bremsstrahlung Isochromat Spectroscopy (BIS) experiments. Because of the difficulties with unambiguous determination of U , it can be considered as a parameter of the model. Then, its value can be adjusted to achieve the best agreement of the results of LSDA+ U calculations with PES or optical spectra [53]. While the use of an adjustable parameter is generally considered an anathema among first principles practitioners, the LSDA+ U approach does offer a plausible and practical method to treat approximately strongly correlated orbitals in solids. The Hubbard U and exchange parameter J can be determined from supercell LSDA calculations using the Slater’s transition state technique [54, 55] or from constrained LSDA calculations [55–57]. Cococcioni and Gironcoli [58] have also provided an internally consistent, basis-set independent method based on the linear response approach for the calculation of the effective interaction parameters in the LSDA+ U method. The constrained LSDA calculations produce $U = 6.17$ eV, $J = 0.87$ eV for Pr and $U = 4.1$ eV, $J = 0.81$ eV for Co in PrCo₂P₂, as well as $U = 6.08$ eV, $J = 0.83$ eV for La and $U = 4.0$ eV, $J = 0.8$ eV for Co in LaCo₂P₂. These values of U and J were used in our calculations presented below.

3. Results

3.1. Electronic structures of LaCo₂P₂ and PrCo₂P₂

Figure 3 presents the partial density of states (PDOS) for LaCo₂P₂ and PrCo₂P₂. The P 3s states are located mostly between -13.6 eV and -10.0 eV below the Fermi level (not shown), while the 3p states of P are found between -6.8 eV and 6.1 eV in LaCo₂P₂ and between -7.5 eV and 5.8 eV in PrCo₂P₂. The spin splitting of the P 3p states is quite small. The Co 3d states occupy the energy interval between -6.9 eV and 5.5 eV and hybridize strongly with the P 3p states. The La 4f empty states occupy the $3.2 - 4.5$ eV energy interval in LaCo₂P₂. The Pr 4f empty states are from 1.5 eV to 4.8 eV above the Fermi level. Two Pr 4f spin-up electrons occupy a small energy interval around -3 eV.

The crystal field at the Co site (D_{2d} point symmetry) in both compounds causes the splitting of Co 3d orbitals into three singlets a_1 (d_{3z^2-1}), b_1 (d_{xy}), b_2 ($d_{x^2-y^2}$) and a doublet e (d_{yz} and d_{xz}). A two peak structure of the minority- and majority-spin Co d states is found in the close vicinity of the Fermi energy. We found that the two peaks (occupied in the majority-spin channel at -0.5 eV and

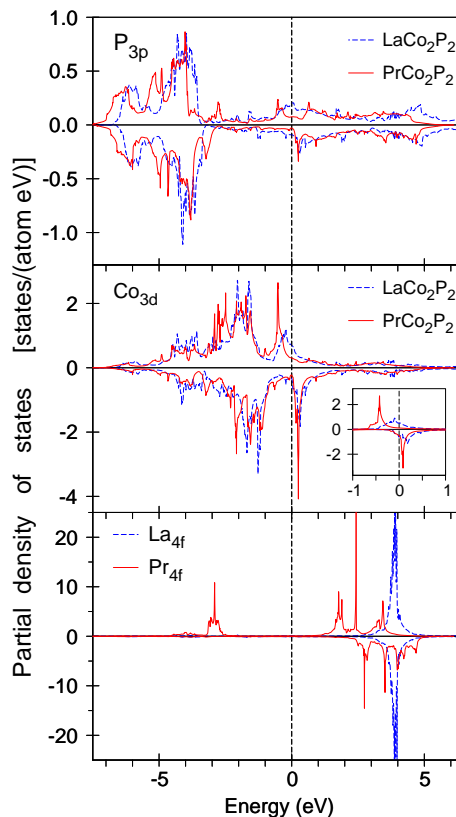


Figure 3. (Color online) Partial density of states of LaCo₂P₂ and PrCo₂P₂. The insert in the middle panel shows the energy distribution of the xy orbitals.

empty in the minority-spin channel at 0.3 eV) are of the xy character. These peaks are rather broad in LaCo_2P_2 and very narrow and intensive in PrCo_2P_2 . Such a difference can be explained by different nearest-neighbors interatomic Co–Co and Co–P distances. The Co–Co distance is equal to 2.697 Å and 2.757 Å in LaCo_2P_2 and PrCo_2P_2 , respectively. Also, the Co–P distance is larger in LaCo_2P_2 than in PrCo_2P_2 by 0.027 Å.

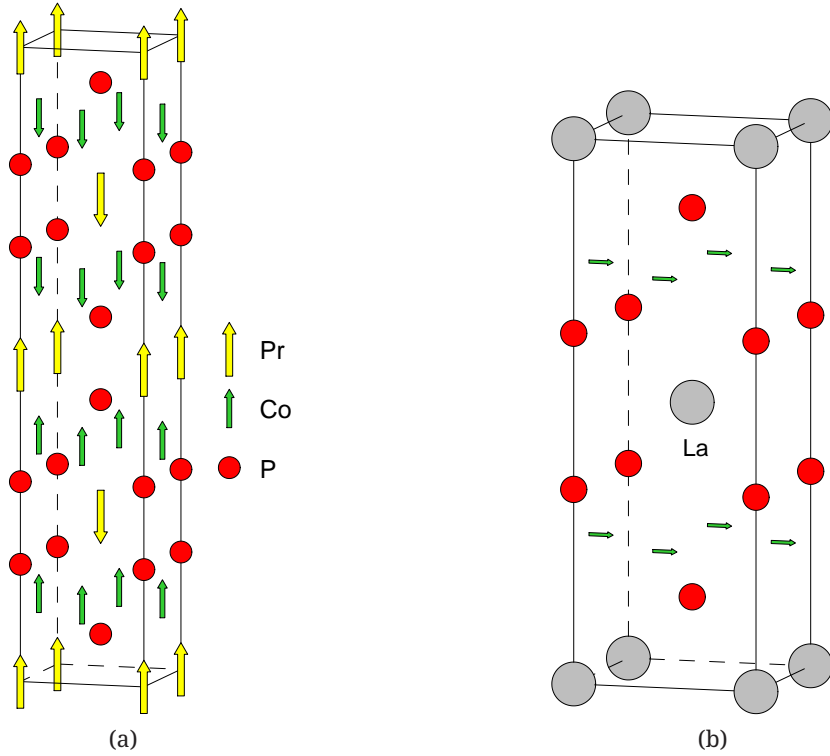


Figure 4. (Color online) Schematic representation of the magnetic ordering in PrCo_2P_2 (left-hand column) and LaCo_2P_2 (right-hand column).

Our self-consistent calculations reveal a ferromagnetic arrangement of magnetic moments in PrCo_2P_2 . The spin magnetic moments at the Pr and Co sites are aligned along the c axis with the antiparallel magnetic coupling between the neighboring planes resulting in antiferromagnetism (left-hand column of figure 4). The similar magnetic arrangement was observed experimentally by Reehuis et al. [16]. The theoretical calculations give a ferromagnetic arrangement of magnetic moments in LaCo_2P_2 , the Co moments are aligned in-plane and parallel to the moments in the other layers (right-hand column of figure 4). Figure 5 (upper panel) shows the MAE as a function of the polar angle θ . The minimum of the total energy corresponds to the magnetic configuration with the Co moments aligned in-plane in agreement with the experimental observation [17]. The theory produces quite a small value of MAE of around 0.28 meV per formula unit in LaCo_2P_2 . The lower panel of figure 5 shows the anisotropy of spin magnetic moments at the Co site in LaCo_2P_2 as a function of the polar angle θ .

The spin m_s and orbital m_l magnetic moments at the Co site in PrCo_2P_2 are larger than in LaCo_2P_2 . Our band structure calculations yield the magnetic moments for the Co atoms $m_s = 0.825 \mu_B$, $m_l = 0.120 \mu_B$ in PrCo_2P_2 and $m_s = 0.665 \mu_B$, $m_l = 0.066 \mu_B$ in LaCo_2P_2 . The induced spin magnetic moments at the P site are of $0.011 \mu_B$ and $0.016 \mu_B$ for the PrCo_2P_2 and LaCo_2P_2 , respectively. The orbital moments at the P sites are small in both compounds ($m_l^P = 0.005 \mu_B$). The orbital magnetic moment at the La site is extremely small $m_l^{\text{La}} = -0.002 \mu_B$, however, we found quite a large orbital moment at the Pr site in PrCo_2P_2 ($m_l^{\text{Pr}} = 0.509 \mu_B$).

3.2. Electronic structure, X-ray absorption and XMCD spectra in La_{1-x}Pr_xCo₂P₂ compounds

Figure 6 presents the partial density of states in La_{1-x}Pr_xCo₂P₂ for $x = 0.25$. The energy position and shape of PDOS slightly differ from the corresponding PDOS of LaCo₂P₂ and PrCo₂P₂. The P 3*p* PDOS has more pronounced peaks near the Fermi level due to the larger P 3*p* – Co 3*d* hybridization. The positions of the empty La and Pr 4*f* states are shifted closer to the Fermi level in comparison with the reference compounds. Also, the shape of the energy distribution of Pr 4*f* states slightly differ from the corresponding 4*f* PDOS in PrCo₂P₂ and LaCo₂P₂. Such differences might be explained by different lattice constants and different numbers and distances of the nearest neighbors.

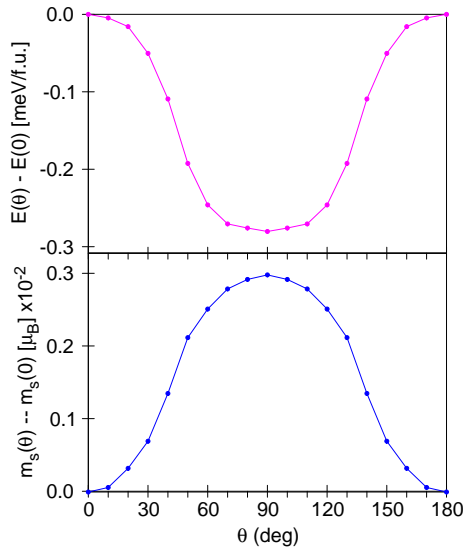


Figure 5. (Color online) MAE (upper panel) and anisotropy of spin magnetic moments at the Co site (lower panel) in LaCo₂P₂ as functions of the polar angle θ .

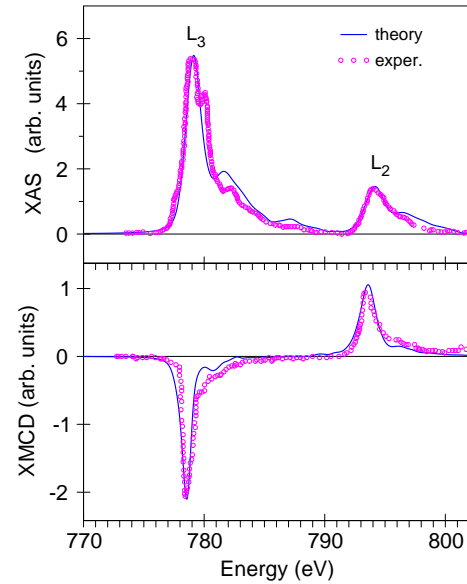


Figure 6. (Color online) Partial density of states in La_{1-x}Pr_xCo₂P₂ for $x = 0.25$.

The spin magnetic moments at the Pr and Co sites ($1.955 \mu_B$ and $0.815 \mu_B$, respectively) are aligned along the c axis in La_{0.75}Pr_{0.25}Co₂P₂ with antiparallel magnetic moments at the La sites ($m_s = -0.002 \mu_B$ and $-0.007 \mu_B$ for two nonequivalent La sites). The orbital magnetic moments are equal to $0.054 \mu_B$, $0.109 \mu_B$, and $0.020 \mu_B$ for Pr, Co, and La sites, respectively. The orbital moments at the P sites are small ($m_l^P = 0.001 \mu_B$).

Figure 7 (upper panel) shows the X-ray absorption spectra at the Co $L_{2,3}$ edges in La_{0.75}Pr_{0.25}Co₂P₂ measured at 5 K [24] with a 1 T magnetic field applied along the c axis compared with the theoretically calculated ones in the LSDA+ U approximation. The Co L_3 X-ray absorption spectrum possesses four major fine structures: a major peak at 779 eV and three high energy shoulders at 780.5 eV, 782.5 eV and 787.5 eV. The theory describes reasonably well the energy position and relative intensity of all the fine structures except the shoulder at 780.5 eV. Due to the electric dipole selection rules ($\Delta l = \pm 1$; $\Delta j = 0, \pm 1$), the major contribution to the absorption at the L_3 edge stems from the transitions $2p_{3/2} \rightarrow 5d_{5/2}$, with a weaker contribution from the $2p_{3/2} \rightarrow 5d_{3/2}$ transitions. For the latter case, the corresponding $2p_{3/2} \rightarrow 5d_{3/2}$ radial matrix elements are only slightly smaller than for the $2p_{3/2} \rightarrow 5d_{5/2}$ transitions. The angular matrix elements, however, strongly suppress the $2p_{3/2} \rightarrow 5d_{3/2}$ contribution. Therefore, the contribution to the XAS spectrum at the L_3 edge from the transitions with $\Delta j = 0$ is one order of magnitude smaller than the transitions with $\Delta j = 1$ [28].

The lower panel of figure 7 presents the XMCD experimental spectra of $\text{La}_{0.75}\text{Pr}_{0.25}\text{Co}_2\text{P}_2$ at the Co $L_{2,3}$ edges and the theoretically calculated ones. The LSDA+ U calculations describe reasonably well all the features of the experimental XMCD spectra.

A study of the $4f$ electron shell in rare earth compounds is usually performed by tuning the energy of X-ray close to the $M_{4,5}$ edges of rare-earth where the electronic transitions between $3d_{3/2,5/2}$ and $4f_{5/2,7/2}$ states are involved. Figure 8 shows the XAS and XMCD spectra at the Pr $M_{4,5}$ edges in $\text{La}_{0.75}\text{Pr}_{0.25}\text{Co}_2\text{P}_2$ measured at 5 K [24] with a 1 T magnetic field applied along the c axis compared with the theoretically calculated ones in the LSDA+ U approximation. The Dirac-Hartree-Fock-Slater single-particle approximation used in this work to calculate the core states is not capable of producing a correct energy position of the spectra (because the self-interaction correction, different kinds of relaxation processes and other many-particle effects were not taken into account). Therefore, we used the experimentally measured positions of the spectra. The theoretically calculated XAS spectra have a rather simple line shape composed of two white line peaks at the M_5 and M_4 edges. The theory reproduces the shape of the M_5 XAS spectrum very well. However, the experimentally measured M_4 spectrum has a well pronounced fine structure at the low energy part that is not reproduced by the theory. We should mention here that a major shortcoming in the band structure approximation is that the multiplet structure is not included. For the Co $L_{2,3}$ edges, this is not the major problem. However, for the Pr $M_{4,5}$ edges, the core-valence electrostatic interactions can significantly effect the line shape of the XAS and XMCD spectra. The fine structure on the low energy side of Pr M_4 XASs is believed to be due to the multiplet structure, which is not included in our calculations. A theoretical method that consistently includes both the band structure and the atomic-like multiplet structure of rare earth metals and compounds is highly desired. However, we should mention that it is still not clear why the multiplet structure is pronounced only at the Pr M_4 edge and is not seen at the M_5 edge.

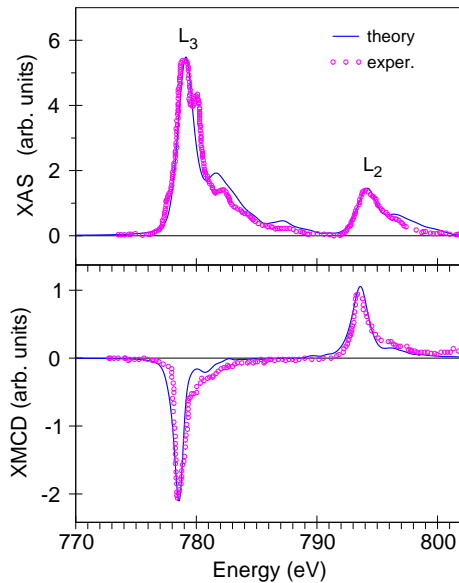


Figure 7. (Color online) Experimental and theoretically calculated X-ray absorption (upper panel) and XMCD (lower panel) spectra of $\text{La}_{0.75}\text{Pr}_{0.25}\text{Co}_2\text{P}_2$ at the Co $L_{2,3}$ edges.

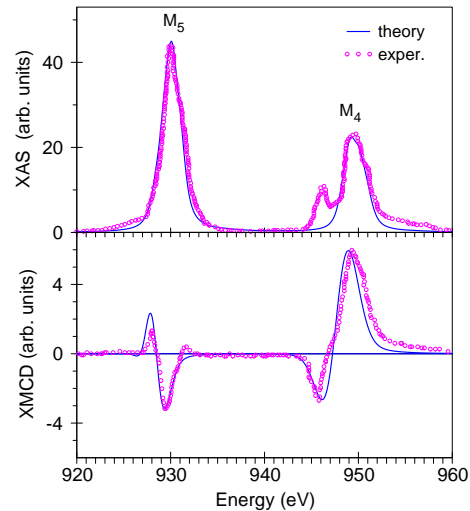


Figure 8. (Color online) Experimental and theoretically calculated X-ray absorption (upper panel) and XMCD (lower panel) spectra of $\text{La}_{0.75}\text{Pr}_{0.25}\text{Co}_2\text{P}_2$ at the Pr $M_{4,5}$ edges.

Figure 8 (lower panel) shows the calculated Pr $M_{4,5}$ XMCD spectra in the LSDA+ U approximation for $\text{La}_{0.75}\text{Pr}_{0.25}\text{Co}_2\text{P}_2$ together with the corresponding experimental data [24]. The experimentally measured dichroism is rather large. The XMCD spectra at the M_5 and M_4 edges have a two-peak structure. The dichroism is positive at a lower energy and negative at a higher energy at the Pr M_5 edge, while the Pr M_4 XMCD spectrum has a negative minimum at a lower energy and a positive maximum at a higher

energy. The LSDA+*U* calculations quite well describe all the features of the experimental XMCD spectra at the $M_{4,5}$ edges. The theory also correctly reproduces the relative intensities of the XAS and XMCD spectra at the M_5 and M_4 edges, namely, the XAS is larger at the M_5 edge than at the M_4 edge, while the XMCD spectra show the opposite behavior with larger dichroism at the M_4 edge in comparison with the M_5 edge.

When a $3d$ core-electron is photo-excited to an unoccupied $4f$ state, the distribution of the charge changes to account for the created hole. We investigated this core-hole effect in the final state using the supercell approximation. We found that the final-state interaction has little effect on the shape of the XAS and XMCD spectra at the Pr $M_{4,5}$ edges. We also investigated the effect of the electric quadrupole E_2 and magnetic dipole M_1 transitions. We found that the M_1 transitions are extremely small in comparison with the E_2 transitions and can be neglected. The E_2 transitions are much weaker than the electric dipole transitions E_1 . They are almost invisible in the XAS and have a very small effect on the XMCD spectra at the Pr $M_{4,5}$ edges.

4. Summary

The electronic structure and magnetic ordering in La_{1-x}Pr_xCo₂P₂ ($x = 0, 0.25, \text{ and } 1$) phosphides have been studied theoretically using the fully relativistic spin-polarized Dirac LMTO band-structure method. The self-consistent calculations reveal a ferromagnetic arrangement of magnetic moments in PrCo₂P₂. The spin magnetic moments at the Pr and Co sites are aligned along the c axis with the antiparallel magnetic coupling between the neighboring planes resulting in antiferromagnetism. LaCo₂P₂ possesses a ferromagnetic arrangement of magnetic moments where the Co moments are aligned in-plane and parallel to the moments in the other layers. The theory gives quite a small value of MAE in LaCo₂P₂ (around 0.28 meV per formula unit).

We have studied the X-ray magnetic circular dichroism at the Co $L_{2,3}$ and Pr $M_{4,5}$ edges in La_{0.75}Pr_{0.25}Co₂P₂. The calculations show good agreement with the experimental measurements. The core-hole effect was found to be very small on the shape of the XAS and XMCD spectra at the Pr $M_{4,5}$ edges. We found that the magnetic dipole M_1 transitions are extremely small in comparison with the electric quadrupole E_2 transitions and can be neglected. The E_2 transitions were found to be weaker than the electric dipole transitions E_1 . They are almost invisible in the XAS and have a very small effect on the XMCD spectra at the Pr $M_{4,5}$ edges.

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Електронна структура, магнітне впорядкування та рентгенівський магнітний циркулярний дихроїзм у фосфідах La_{1-x}Pr_xCo₂P₂

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На основі зонних розрахунків повністю релятивістським спіно-поляризованим лінійним методом МТ-орбіталей (ЛМТО) теоретично вивчені електронна структура і магнітне впорядкування у фосфідах La_{1-x}Pr_xCo₂P₂ ($x = 0, 0.25, \text{ та } 1$). В рамках методу LSDA+ U теоретично досліджені рентгенівські спектри поглинання та спектри рентгенівського циркулярного дихроїзму на краях поглинання Co $L_{2,3}$ і Pr $M_{4,5}$. Вивчено вплив остовної дірки в кінцевому стані, а також електричних квадрупольних E_2 і магнітних дипольних M_1 переходів. Отримано добре узгодження з експериментальними результатами.

Ключові слова: *сильно скорельовані системи, зонна структура, магнітні моменти, рентгенівський магнітний циркулярний дихроїзм*
