



Hidden magnetic order in plutonium dioxide nuclear fuel†

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A thorough understanding of the chemistry of PuO₂ is critical to the design of next-generation nuclear fuels and the long-term storage of nuclear materials. Despite over 75 years of study, the ground-state magnetic structure of PuO₂ remains a matter of much debate. Experimental studies loosely indicate a diamagnetic (DM) ground-state, whereas theoretical methods have proposed either a collinear ferromagnetic (FM) or anti-ferromagnetic (AFM) ground-state, both of which would be expected to cause a distortion from the reported *Fm* $\bar{3}m$ symmetry. In this work, we have used accurate calculations based on the density functional theory (DFT) to systematically investigate the magnetic structure of PuO₂ to resolve this controversy. We have explicitly considered electron-correlation, spin-orbit interaction and noncollinear magnetic contributions to identify a hereto unknown longitudinal 3k AFM ground-state that retains *Fm* $\bar{3}m$ crystal symmetry. Given the broad interest in plutonium materials and the inherent experimental difficulties of handling this compound, the results presented in this paper have considerable implications for future computational studies relating to PuO₂ and related actinide structures. As the crystal structure is coupled by spin-orbit interactions to the magnetic state, it is imperative to consider relativity when creating computational models.

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1 Introduction

In the nuclear fuel cycle, PuO₂ plays a significant role as a fissile material and a thorough understanding of its fundamental properties is therefore important. To characterise the nature of condensed matter systems, it is imperative to identify coupled and competing magnetic interactions. An accurate description of the underlying physics, including the magnetic ground-state, is key to the design of reliable computational models. A controversy exists between experimental and theoretical methods,¹ which affects the calculation of phonon frequencies, thermodynamic properties and excited electronic states,^{1,2} and has implications for the validation of basic models of reactivity, the design of surface models and the interaction of adsorbents.³ A singlet diamagnetic (DM) ground-state is inferred

by experimental measurements and the one-electron interpretation of the crystal electric field,^{4–8} whereas a ferromagnetic (FM) or an anti-ferromagnetic (AFM) ground-state is calculated by first-principle methods.^{2,3,9–21} In this paper, the controversy detailed in the literature is addressed by high-end first-principles methods and, for the first time, a fully relativistic noncollinear investigation of magnetic order in PuO₂ has been conducted. As a key computational consideration, the spin-orbit interaction (SOI, which couples the lattice structure to the magnetic state) is fully considered in this study.²² A number of experimental measurements on PuO₂ report *Fm* $\bar{3}m$ (no. 225) crystal symmetry (Fig. 1),²³ but suggestions of low-temperature ($T < 30.8$ K) lattice anomalies (reminiscent of transverse 3k AFM behaviour in UO₂ with *Pa* $\bar{3}$ (no. 205) crystal symmetry) remain uninvestigated.²⁴

The low-temperature characterisation of the PuO₂ magnetic ground-state is challenging, due to: nucleonic radioactive decay,²⁵ the toxicity of the metals,²⁶ and the inhomogeneity of samples.⁷ A number of inconsistencies in the literature have been recorded.²⁷ For example, a singlet Γ_1 DM ground-state (in accordance with the one-electron crystal-field model) has been inferred by: magnetic susceptibility ($T = 4$ K),⁴ inelastic neutron scattering (INS) ($T > 30$ K),^{5,6} and nuclear magnetic resonance (NMR) ($T > 4$ K)^{7,8} measurements. Although a number of inconsistencies in the literature have been recorded,²⁷ a singlet Γ_1 ground-state has been inferred from the temperature-independent magnetic susceptibility of PuO₂ (5.4×10^{-4} cm³ mol⁻¹).¹

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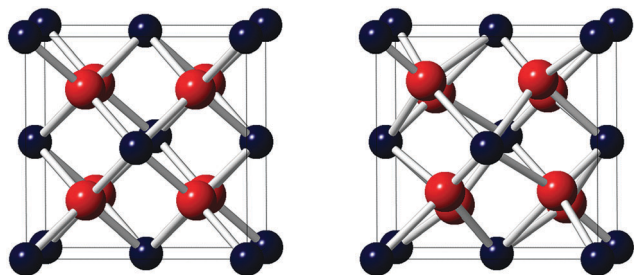


Fig. 1 The crystal structure of PuO_2 with; (left) $Fm\bar{3}m$ (no. 225) (right) $Pa\bar{3}$ (no. 205) crystal symmetry. The colours in the parentheses indicate the Pu^{4+} (blue) and O^{2-} (red) ions. Note, in $Pa\bar{3}$ symmetry, the displacement of the O^{2-} ions is exaggerated.

In contrast, the temperature-dependent magnetic susceptibility of $\text{Th}_{1-x}\text{Pu}_x\text{O}_2$ ($\sim 1.1 \times 10^{-3} \text{ cm}^3 \text{ mol}^{-1}$) indicates a magnetic response that is inconsistent with earlier results.²⁸ INS measurements of PuO_2 indicate a singlet Γ_1 to triplet Γ_4 transition of 123 meV, which corresponds to a temperature-dependent magnetic susceptibility of $\sim 1 \times 10^{-3} \text{ cm}^3 \text{ mol}^{-1}$ when $T > 400$ K. The magnetic coupling of proximal cations by intermediary DM anions is described by magnetic superexchange.²⁹ The full width at half maximum (FWHM) INS peak results of 25 meV is indicative of AFM superexchange.^{1,30} In addition, magnetic susceptibility measurements of $\text{U}_{1-x}\text{Pu}_x\text{O}_2$ indicate AFM superexchange with high concentrations of plutonium.³¹ Finally, the low-temperature ^{17}O NMR measurements of PuO_2 result in an apparent DM state at 6 K and it is conceivable that even lower temperatures may result in a different magnetic ground-state. For instance, low temperature measurements of $\text{U}_{1-x}\text{Pu}_x\text{O}_2$ indicate a Néel temperature under 5 K when $x = 0.5$.³¹ To compensate for experimental issues (thermal energy released by the decay of actinide nuclei), theoretical methods offer an alternative and complementary means of investigation.

A computational investigation of the PuO_2 systems is highly complicated. This is due to exchange–correlation effects, relativistic influences and magnetic behaviour. It is now well recognised that in highly correlated f-electron systems, conventional DFT^{32,33} methods fail to reproduce the correct electronic structure. For instance, identifying finding well-known AnO_2 insulators as metallic by underestimating the band-gap.^{17,21} To calculate highly-correlated materials, a number of methods offer a correction to the self-interaction error. These include the self-interaction correction (SIC) method,³⁴ modified density functional theory (DFT+ U),^{32,33,35–37} dynamic mean field theory (DMFT),^{38,39} and hybrid density functionals.^{12,40,41} In condensed matter physics, the DFT+ U method offers a computationally tractable means of study, where the on-site Coulomb repulsion of the f-electrons is treated by tuneable U and J modifiers. In contrast, although the use of hybrid functionals are computationally expensive, they represent one of the more accurate methods available.

DFT studies have consistently reported either a collinear ferromagnetic (FM) or a collinear antiferromagnetic (AFM) ground-state.³ The ordered magnetic states have been attributed

to approximations inherent to DFT methods.⁴² In response to the highly-controversial ordered magnetic structures, a number of investigations have focused on the DM solution.^{42,43} As a combination of the exact diagonalization of the Anderson impurity model and LDA+ U ($U = 6.5$ eV), correlated band theory yielded a DM solution with an f-shell occupation of ~ 4.5 at the Pu atoms. The fully spherical symmetric Coulomb interaction, crystal field, and SOI were included in the multiorbital impurity model.⁴³ The authors found an excited triplet state at 126 meV; however, a complete investigation of ordered magnetic states is needed.

In cases where an ordered magnetic state has been incorporated within the DFT methodology, the investigators have limited the discussion to FM(001) and collinear 1k AFM states.^{2,3,9–21} As the actinide elements are heavy-fermion systems, SOI is imperative for the computational treatment of nuclear materials.⁴⁴ A limited number of studies have included SOI (due to the high computational cost) in their calculations.^{18–21,42,43} The inclusion of noncollinear magnetic behaviour PuO_2 has been ignored.

In addition to general complications associated with highly-correlated systems, the magnetic ground-state of the AnO_2 is influenced by superexchange, multipolar and noncollinear interactions.^{1,27} Other methods have been used to model PuO_2 systems. The complete active space (CAS) wavefunctions to correct the exchange–correlation errors common to DFT can be used to achieve a DM ground-state.⁴⁵ A limitation of the embedded cluster method, however, is that the quantum cluster may not be sufficiently large enough to capture the superexchange interactions in PuO_2 , and therefore the findings are incomplete. In addition, the ground-state electronic structure and crystal-field levels of PuO_2 have been calculated by DFT in terms of the total energy differences.³⁰ The total energy of the crystal-field states are calculated by the Clebsch–Gordon method from a standard model of rare-earth elements and as a function of their single-electron components. The calculated ground Γ_1 state-excited Γ_4 state transition of 99 meV compares with the measured INS result of 123 meV.

In the current study, the magnetic structure of PuO_2 has been investigated by noncollinear relativistic calculations. We have implemented three levels of theory. These include density functional theory (PBEsol),^{32,33} DFT with the Hubbard correction (PBEsol+ U),^{32,33,35–37} and hybrid (HSE06) functionals.^{12,40,41,46} To date, noncollinear contributions to the magnetic ground-state have remained unexplored, and we have therefore investigated the noncollinear interactions illustrated in Fig. 2. To highlight the importance of SOI, we have compared with non-SOI HSE06 calculations (see additional information, ESI†).

2 Computational methodology

A noncollinear relativistic computational study of the PuO_2 magnetic structure by the Vienna Ab initio Simulation Package (VASP) code has been conducted.^{34,38,47} The code employs relativistic effective core potentials (ECPs), the frozen-core projector-augmented wave (PAW) method and a planewave



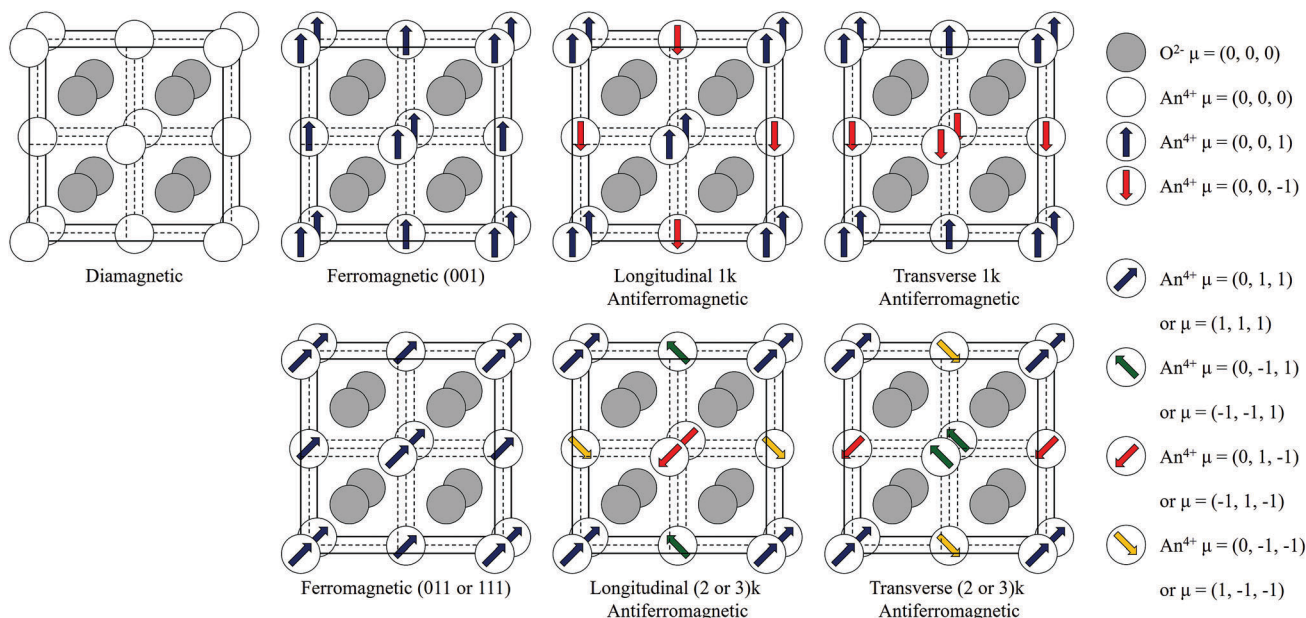


Fig. 2 The magnetic structure of PuO_2 for the diamagnetic (DM), ferromagnetic (FM) and antiferromagnetic (AFM) states. The direction of the effective magnetic moment (μ) of each ion is described by the $\mu = (a, b, c)$ wave-vector matrix. Here, transverse (T) and longitudinal (L) domains are considered for the AFM state. The number of independent wave vectors is denoted by the k prefix.

basis set. The cut-off energy of the planewave basis set is 500 eV. The cut-off energy and k -point mesh have been checked for convergence. The following explicit electrons for each atomic species are considered: O ($1s^2 2s^2 2p^2$) and Pu ($5f^6 6s^2 6p^6 6d^1 7s^2$). The ion–electron interaction was described using the projector-augmented wave (PAW)^{41,48} approximation, considering spin–orbit interactions (SOI)⁴⁹ and noncollinear magnetic wave vectors. The spin quantisation axis is defined by $(0, 0, 1)$ plane, from which magnetic and spinor-like values are calculated. Ionic forces have been evaluated using the Hellmann–Feynman theorem⁵⁰ and conjugate gradient algorithm.⁵¹ The space group has been evaluated to 10^{-5} Å based on a symmetry analysis of the unit cell.

As a computationally intensive method, hybrid functionals incorporate Hartree–Fock (HF) exchange energy into the DFT formalism. In this study, the hybrid Heyd–Scuseria–Ernzerhof (HSE06) functional is employed.^{12,40,41,46} The functional offers improved: thermochemical data, band-gaps (for low-medium semi-conductors) and lattice constants.

$$E_{\text{XC}}^{\text{HSE}} = (a)E_{\text{X}}^{\text{HF,SR}}(\mu) + (1 - a)E_{\text{X}}^{\text{PBE,SR}}(\mu) + E_{\text{X}}^{\text{PBE,LR}}(\mu) + E_{\text{C}}^{\text{PBE}}. \quad (1)$$

The terms in the parentheses define; the exchange–correlation HSE06 energy ($E_{\text{XC}}^{\text{HSE}}$), an adjustable constant (a), the short-range (SR) interaction energy and the long-range (LR) interaction energy. The adjustable screening (μ) modifier, in the HSE06 functional, is 0.207 \AA^{-1} . The integration of reciprocal space was completed with a Γ -centred $(4 \cdot 4 \cdot 4)$ k -point mesh⁵² by the conventional Gaussian or Blöchl tetrahedron method.⁵³ The iteration threshold for electronic and ionic convergence is 1×10^{-6} eV and 1×10^{-2} eV \AA^{-1} , respectively.

As a computationally more tractable means of investigation, the DFT+ U ^{32,33,35–37} method addresses the on-site Coulombic repulsion of the Pu f -electrons. The correction of the electron self-interaction error (SIE) implicit in DFT calculations, is treated by the rotationally invariant Liechtenstein *et al.*³⁶ formalism. In the rotationally invariant Liechtenstein *et al.*³⁶ formalism, the on-site Coulombic (U) and exchange (J) modifiers are treated as independent variables.

$$E_{\text{dc}}(\hat{n}) = \frac{U}{2} \hat{n}_{\text{tot}} (\hat{n}_{\text{tot}} - 1) \frac{J}{2} \sum_{\sigma} \hat{n}_{\text{tot}}^{\sigma} (\hat{n}_{\text{tot}}^{\sigma} - 1). \quad (2)$$

The terms in the parentheses indicate the; double counting energy term (E_{dc}), the main quantum number (n) and the spin index (σ). These parameters are applied to the f -orbitals of plutonium. Thus, in the DFT+ U modification, the electrons are correctly localised to their specific lattice sites.

In the past, the influence of J on noncollinear magnetic materials has been investigated.^{44,54} The introduction of J increased the anisotropic nature of the f -states.⁴² As such, in the current study, the J modifier is kept at a constant value of 0.00 eV.⁴⁴ By comparison, when $J = 0.00$ eV, the Dudarev *et al.*³⁵ formalism and Liechtenstein *et al.*³⁶ formalism are equivalent.⁵⁴ The integration of reciprocal space was completed with a Γ -centred k -point $(5 \cdot 5 \cdot 5)$ mesh⁵² by Blöchl tetrahedron method.⁵³ The iteration threshold for electronic and ionic convergence is 1×10^{-8} eV and 1×10^{-3} eV \AA^{-1} , respectively. The DFT^{32,33} and DFT+ U ^{35–37} calculations are performed with PBE revised for solids (PBEsol)⁵⁵ exchange–correlation functional. In the calculation of actinide materials, the functional has shown improved results.⁴⁴ For the optical absorbance calculation, the k -point mesh is $15 \times 15 \times 15$. For the band structure calculations, the $Fm\bar{3}m$ k -point pathway of high



symmetry points in the Brillion zone is defined as $\Gamma \rightarrow L \rightarrow W \rightarrow X \rightarrow \Gamma$. The density of states and band structure have been illustrated using the SUMO code, a command-line plotting tool for *ab initio* calculations.⁵⁶

2.1 Noncollinear magnetic behaviour

In the high-temperature PM state, the AnO₂ crystallises in the calcium fluorite (CaF₂) motif with $Fm\bar{3}m$ (no. 225) cubic symmetry. In this structure, the An⁴⁺ cations occupy octahedral (4a) sites; whereas, the O²⁻ anions occupy tetrahedral (8c) sites. As the magnetic planes are indistinct, the implicit PM disorder stabilizes the high symmetry $Fm\bar{3}m$ crystal structure. At low-temperature, the AnO₂ exhibit either DM or AFM behaviour. In addition, multiple first-principle methods calculate a FM ground-state under certain constraints. In the DM and FM states, the implications of noncollinearity are irrelevant. In the AFM state, noncollinear contributions result in 1k (one wave-vector), 2k (two independent wave-vectors) or 3k (three independent wave-vectors) configurations. These configurations are further differentiated when considering longitudinal and transverse domains (Table 1).

In the Russell–Saunders coupling scheme, under the cubic crystal-field, the degeneracy of the ⁵I₄ multiplet is split into ascending Γ_1 (1), Γ_4 (3), Γ_3 (2), and Γ_5 (3) crystal-field levels (the number in the parentheses denotes the degeneracy).^{30,42,43} The calculated singlet Γ_1 ground-state to triplet Γ_4 excited state transition is 123 meV as measured by neutron inelastic scattering.⁶ The crystal-field model is technically only valid for the one-electron case; nevertheless, the DM state inferred by experiment is often justified by crystal-field theory.

The majority of studies on the AnO₂ are often limited by the computational expense of the systems and restricted to discussions of collinear 1k magnetism.^{2,3,9–21,57} In general, the actinide dioxides (AnO₂) exhibit low-temperature noncollinear 3k AFM or PM behaviour (Table S1, ESI†).^{7,24,58–64} The low-temperature magnetic state can result in a distortion of the crystal structure.^{1,27,44,65} For instance, in uranium dioxide (UO₂) transverse 3k AFM order results in $Pa\bar{3}$ symmetry; whereas, in neptunium dioxide (NpO₂) longitudinal 3k AFM order preserves $Fm\bar{3}m$ symmetry. As the crystal structure is coupled by spin–orbit interactions (SOI) to the magnetic state, it is imperative to incorporate relativistic influences when creating computational models.

3 Results & discussions

3.1 Magnetic structure

In the current study, the magnetic structure of PuO₂ has been investigated by first-principle methods. The magnetic state clearly influences: relative energetics, band structure, magnetic moment, and crystal symmetry. In our HSE06 and PBEsol+*U* calculations, the DM state of PuO₂ is the highest energy magnetic configuration. In each instance, a longitudinal 3k AFM ground-state with $Fm\bar{3}m$ cubic symmetry has been calculated. In the absence of SOI, a degenerate FM(011)–FM(111) ground-state has been calculated (see additional information, Table A1, ESI†). The result highlights the need (when modelling PuO₂) to consider: relativistic effects, exchange–correlation, and noncollinear magnetism. In our hybrid (HSE06) calculations (compared to the DM configuration) the ordered FM and AFM configurations are ~ 1.01 eV per F.U. (formula unit) lower in energy (Table 2). The finding is inconsistent with the DM ground-state indicated by current experimental data.^{4–8}

It is noted that the experimental optical band-gap (2.80 eV) has been measured by epitaxial thin film absorbance.⁶⁶ As the band-gap is influenced by the magnetic state, the experimental PM (high-temperature, $T \sim 298$ K) result and computational longitudinal 3k AFM result (low-temperature, $T \sim 0$ K) cannot be directly compared. In addition, the magnetic structure is coupled by SOI to the crystal symmetry. In our calculations, the DM and longitudinal 3k AFM states retain $Fm\bar{3}m$ crystal symmetry, only. The collinear 1k AFM states (used in past studies) calculate an incorrect crystal structure. A tetragonal $I4/mmm$ (no. 139) (longitudinal 1k AFM) or orthorhombic $Fmmm$ (no. 69) (transverse 1k AFM) crystallographic distortion results.^{2,3,9–21} This inconsistent with experimental results. In addition, transverse 3k AFM order results in $Pa\bar{3}$ crystal symmetry (as observed in UO₂ and AmO₂).⁴⁴

As a more computationally tractable method of theory (compared with the hybrid functionals), the results obtained from PBEsol+*U* are reported for different values of *U* (Fig. 3). In an identical manner to hybrid (HSE06) calculations, the magnetic state influences the electronic and crystallographic structure. PBEsol ($U = 0$ eV) calculates a degenerate metallic FM (111), (011), (001) ground-state; however, the metallic ground-state is inconsistent with the known insulating nature. The result exemplifies the failure of conventional DFT methods

Table 1 A list of magnetic wavevectors in PuO₂ for the ordered states

Ion	Ferromagnetic			Antiferromagnetic					
	(001)	(011)	(111)	Longitudinal domain			Transverse domain		
				1k	2k	3k	1k	2k	3k
(0, 0, 0)	(0, 0, 1)	(0, 1, 1)	(1, 1, 1)	(0, 0, 1)	(0, 1, 1)	(1, 1, 1)	(0, 0, 1)	(0, 1, 1)	(1, 1, 1)
$\left(\frac{1}{2}, \frac{1}{2}, 0\right)$	(0, 0, 1)	(0, 1, 1)	(1, 1, 1)	(0, 0, 1)	(0, $\bar{1}$, 1)	($\bar{1}$, $\bar{1}$, 1)	(0, 0, $\bar{1}$)	(0, $\bar{1}$, $\bar{1}$)	(1, $\bar{1}$, $\bar{1}$)
$\left(\frac{1}{2}, 0, \frac{1}{2}\right)$	(0, 0, 1)	(0, 1, 1)	(1, 1, 1)	(0, 0, $\bar{1}$)	(0, 1, $\bar{1}$)	($\bar{1}$, 1, $\bar{1}$)	(0, 0, 1)	(0, $\bar{1}$, 1)	($\bar{1}$, $\bar{1}$, 1)
$\left(0, \frac{1}{2}, \frac{1}{2}\right)$	(0, 0, 1)	(0, 1, 1)	(1, 1, 1)	(0, 0, $\bar{1}$)	(0, $\bar{1}$, $\bar{1}$)	(1, $\bar{1}$, $\bar{1}$)	(0, 0, $\bar{1}$)	(0, 1, $\bar{1}$)	($\bar{1}$, 1, $\bar{1}$)



Table 2 The relative energy (eV per F.U.), band-gap (eV), magnetic moment (μ_B /Pu ion), lattice volume (\AA^3) and space group (number) for each PuO₂ magnetic configuration (calculated by the HSE06 functional). The energetics of the magnetic configurations are calculated relative to the longitudinal 3k antiferromagnetic ground-state

Magnetic configuration	Relative energy (eV per F.U.)	Band-gap (eV)	Magnetic moment (μ_B /Pu ion)	Lattice volume (\AA^3)	Space group (number)
Diamagnetic	1.069	2.92	0.00	155.37	$Fm\bar{3}m$ (no. 225)
Ferromagnetic	(001)	0.074	1.91	155.88	$I4/mmm$ (no. 139)
	(011)	0.037	1.79	156.03	$Immm$ (no. 71)
	(111)	0.027	1.68	155.99	$R\bar{3}m$ (no. 166)
Antiferromagnetic Longitudinal	1k	0.058	2.83	155.69	$I4/mmm$ (no. 139)
	2k	0.011	3.05	155.65	$I4/mmm$ (no. 139)
	3k	0.000	3.04	155.65	$Fm\bar{3}m$ (no. 225)
	Transverse	1k	0.062	2.50	155.64
	2k	0.020	2.93	155.73	$Pbca$ (no. 61)
	3k	0.009	2.68	155.72	$Pa\bar{3}$ (no. 205)
Experimental	—	2.80 ⁶⁶	0.00 ^{4–8}	157.25 ²³	$Fm\bar{3}m$ (225) ²³

with highly-correlated systems. The band-gap is only replicated by the AFM and DM states when $U = 5–7$ eV.⁶⁶ In particular, the band-gap of the experimental DM state and theoretical longitudinal 3k AFM ground-state is reproduced when $U = 6.35$ eV and $U = 6.00$ eV, respectively. In contrast, when calculating the band-gap for the FM states, at every value of U , the band-gap is underestimated relative to experimental values.

In experimental studies, DM order has been inferred from the absence of an effective Pu⁴⁺ magnetic moment.^{4–8} In this study, the effective magnetic Pu⁴⁺ moment of the ordered FM and AFM states is reported. In the FM states, the effective magnetic moment decreases from 3.85 to 3.75 μ_B /Pu ion in the $U = 0–2$ eV range; however, increases from 3.75 μ_B /Pu ion to 3.83 μ_B /Pu ion when $U = 3–7$ eV. In contrast, the effective magnetic moment of the AFM states continually increases from 3.55 μ_B /Pu ion to 3.80 μ_B /Pu ion over the range of $U = 0–7$ eV. However, when $U = 6$ eV, the effective magnetic moment converges at 3.80 μ_B /Pu ion for all magnetic states.

3.2 Longitudinal 3k antiferromagnetic ground-state

In this paper, a theoretical longitudinal 3k AFM ground-state for PuO₂ has been calculated by PBESol+ U ($U = 6.00$ eV) and hybrid (HSE06) functionals. The noncollinear magnetic ground-state of UO₂, NpO₂ (established by experimental means) and PuO₂ are highly comparable. The optical absorbance, band structure and density of states for the longitudinal 3k AFM ground-state have been calculated by PBESol+ U ($U = 6$ eV) (Fig. 4 and Table 3).

A charge-transfer insulator with an indirect bandgap of 2.80 eV has been calculated; this contrasts against the Mott-insulator calculated by the fully relativistic linear combinations of Gaussian-type orbitals-fitting function (LCGTO-FF) method.¹⁹ The respective X- Γ valence band maximum (VBM) to L conduction band minimum (CBM) is comprised predominately of O p- and Pu f-states. The degeneracy of the bands in the longitudinal 3k AFM ground-state is perturbed by the presence of magnetic order. This key finding was made possible only through the inclusion of noncollinear and spin-orbit interactions.

The fundamental band-gap (the difference between the ionisation potential and the electron affinity, the valence band maximum (VBM) to conduction band minimum (CBM) transition) and optical band-gap (transitions are restricted by orbital symmetry) are influenced by the magnetic state. In our calculations, the longitudinal 3k AFM electronic structure is in excellent agreement with optical absorbance measurements.⁶⁶ The direct and optical band-gaps differ by 0.17 eV, although both are in excellent agreement with the experimental band-gap of 2.80 eV (Table 3).⁶⁶

3.3 Antiferromagnetic superexchange & small-moment systems

The experimental low-temperature structure of PuO₂ (not necessarily indicative of DM order) is characterised by the absence of a lattice distortion and of a magnetic moment. The ambient PM, experimental DM ground-state, and theoretical longitudinal 3k AFM ground-state retain $Fm\bar{3}m$ crystal symmetry. The absence of any change in crystal structure between the PM and the longitudinal 3k AFM phase is presumed to have helped the ground-state elude detection. A complete analysis can be conducted by comparing other f-electron materials. The results are rationalised by considering magnetic anomalies in NpO₂, URu₂Si₂, UPt₃, and UPd₃, which are small-moment materials.¹

In the absence of an ordered moment and of noticeable lattice anomalies, a phase-transition could be driven by an ordered parameter no longer invariant under time-reversal symmetry.¹ The interaction leading to a phase-transition would have to arise from a purely electronic mechanism, as phonons may only carry interactions between time-reversal-invariant multipoles. A magnetic multipole (for which Kramers' theorem would no longer apply) by AFM superexchange is a candidate for an ordered mechanism.^{1,27,67} As an indication of superexchange, the magnetic susceptibility measurements for PuO₂ and Th_{1-x}Pu_xO₂ are inconsistent.^{4,28,31} The interaction of the Pu⁴⁺ ions in Th_{1-x}Pu_xO₂ is reduced by dilution; this suppresses the superexchange mechanism and increases the magnetic susceptibility.¹



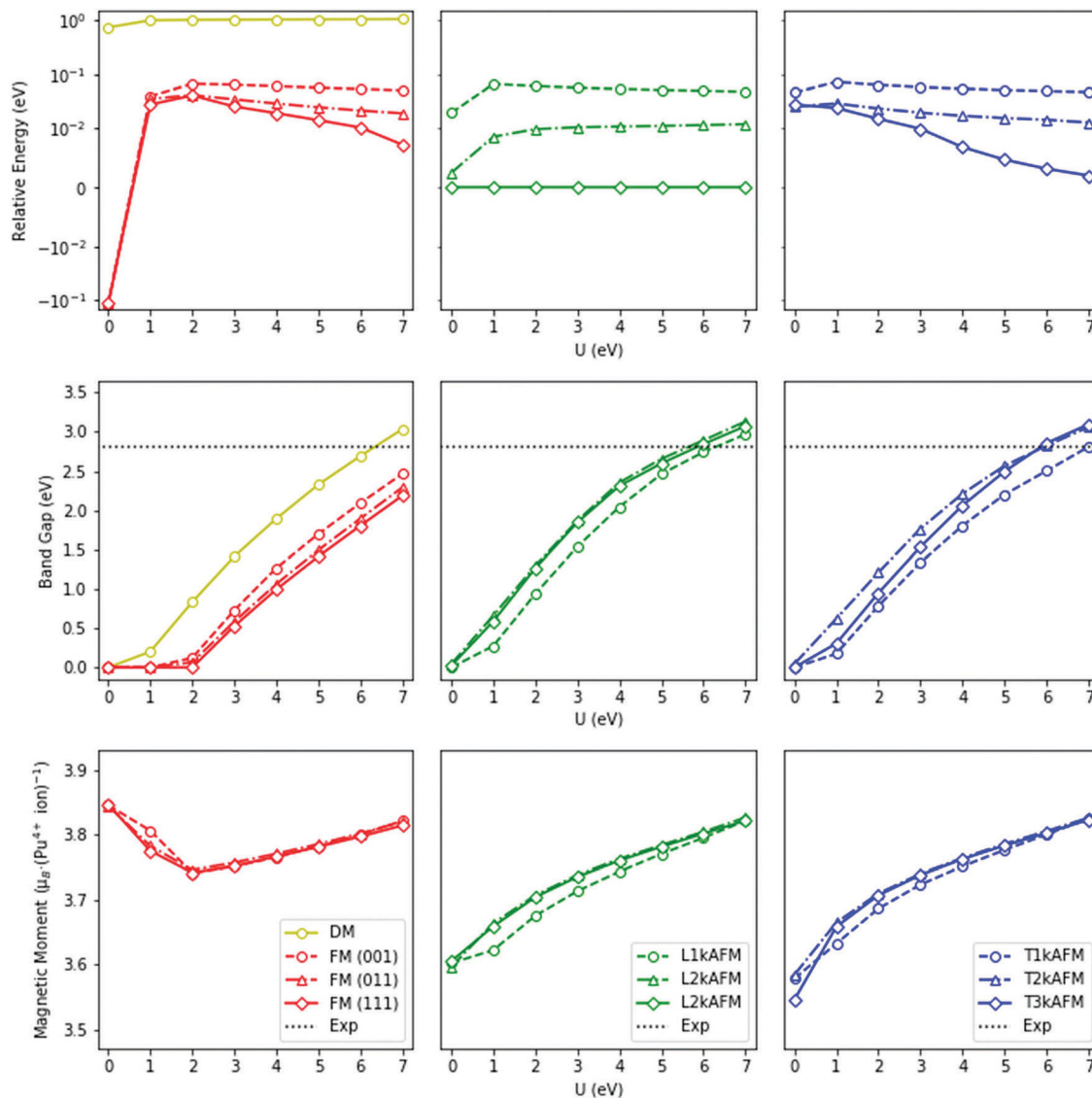


Fig. 3 The relative ground-state energies, band-gaps, and effective magnetic moments against the Coulomb modifier (U) for diamagnetic (DM), ferromagnetic (FM), and antiferromagnetic (AFM) states of PuO_2 , calculated with PBEsol ($U = 0$ eV) and PBEsol+ U . The antiferromagnetic transverse (T) and longitudinal (L) domains are additionally represented. The k -prefix denotes the number of independent wave vectors: (upper row) the calculated energy of magnetic states relative to the longitudinal $3k$ antiferromagnetic ground-state; (middle row) the direct band-gap; (lower row) the effective Pu magnetic moment. The DM (yellow), FM (red), longitudinal AFM (green) and transverse AFM (blue) states are denoted by the colours in the parentheses.

In addition, anomalies in the INS measurements of PuO_2 have been attributed to Jahn–Teller distortions and superexchange interactions.^{5,6} As a comparison, a superexchange interaction is theorised to occur in NpO_2 where a magnetic octupole has been established.⁶⁸

In another interpretation, PuO_2 may be a small-moment system. This is observed in other actinide materials where the AFM response of URu_2Si_2 ($T_N = 17.5$ K), UPt_3 ($T_N = 5$ K) and UPd_3 ($T_N = 4.5$ K) is 0.02 – $0.03 \mu_B/\text{U}$ ion.¹ Indeed, a small-moment magnetic state has been theorised for NpO_2 where muon spin rotation measurements of the magnetic moment is 0.06 – $0.15 \mu_B/\text{Np}$ ion.^{1,69} The nature of small-moment systems is unclear.¹ The moment of a system may be reduced by competing magnetic interactions. In the localised PuO_2

f -electron system, the interaction may be Rudermann, Kittel, Kasuya, Yoshida (RKKY) or Kondo in nature. If a small-moment in the PuO_2 system exists, establishing the magnetic ground-state is a considerable challenge. In our results, the magnetic moment of the longitudinal $3k$ AFM ground-state is $\sim 3.65 \mu_B/\text{Pu}$ ion. In this regard, first-principle DFT methods are known to overestimate small-moment actinide materials.¹ It is also conceivable that PuO_2 undergoes a second low-temperature phase transition, which is known to occur in other actinide compounds. For instance, the specific heat capacity of UPt_3 has two distinct anomalies at $T_{C1} = 0.49$ K and $T_{C2} = 0.44$ K indicative of changing magnetic order. A detailed low-temperature experimental study of magnetic order in PuO_2 is needed.



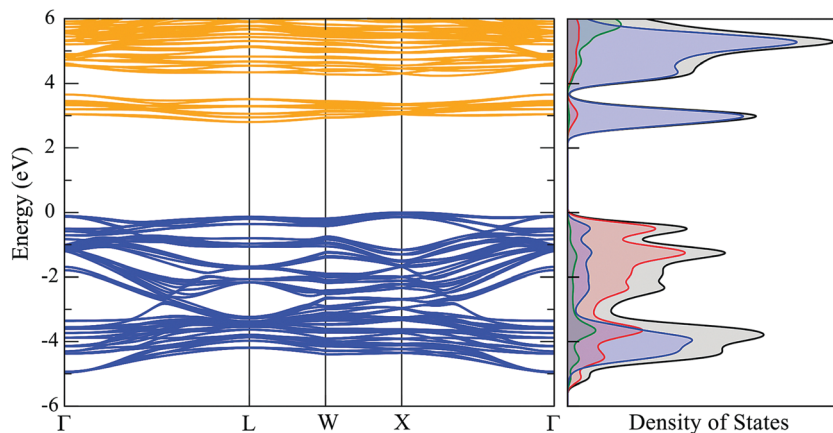


Fig. 4 The electronic structure of the longitudinal 3k antiferromagnetic ground-state of PuO₂ calculated by PBEsol+*U* where *U* = 6 eV; (left) band structure, (right) density of states. The colours in the parentheses indicate the Pu f- (blue), Pu d- (green) and O p- (red) states.

Table 3 The fundamental band-gap (eV), optical band-gap (eV), lattice constant (Å), bulk modulus (GPa) and magnetic moment (μ_B /Pu ion) for the diamagnetic (DM, *U* = 6.35 eV) and longitudinal 3k antiferromagnetic (AFM, *U* = 6.00 eV) state of PuO₂ (calculated by PBEsol+*U*)

DFT+ <i>U</i>	Band-gap (eV)		Lattice constant (Å)	Bulk modulus (GPa)	Magnetic moment (μ_B /Pu ion)	Crystal symmetry	Magnetic configuration
	Fundamental	Optical					
6.35	2.81	2.82	5.411	217	0.00	<i>Fm</i> $\bar{3}$ <i>m</i> (225)	DM ⁴⁴
6.00	2.80	2.97	5.415	215	3.80	<i>Fm</i> $\bar{3}$ <i>m</i> (225)	Longitudinal 3k AFM
—	—	2.80 ⁶⁶	5.395–5.398 ²³	178–379	0.00	<i>Fm</i> $\bar{3}$ <i>m</i> (225)	Exp.

4 Conclusions

In conclusion, the magnetic structure of PuO₂ has been investigated by computational means. As a highly-correlated and heavy-fermion system, the electronic structure of PuO₂ is notoriously difficult to calculate. Investigations of surface reactivity and bonding often rely on exact calculations of the electronic structure. A key finding is the impact of transverse and longitudinal domains on the band structure. In the case of collinear 1k AFM order, the electronic structure is significantly different depending on the domain employed, which is important as most investigations on AnO₂ do not differentiate between collinear magnetic structures.

In our calculations, the metastable DM state (in contrast with the longitudinal 3k AFM ground-state) is inconsistent with current experimental information. In this paper, we have established a longitudinal 3k AFM ground-state with *Fm* $\bar{3}$ *m* crystal symmetry. The fact that this magnetic ground-state has not been experimentally observed may be explained in two possible ways. First, PuO₂ may be a small-moment material for which current experimental resolution is insufficient to identify an ordered magnetic state. If this is the case, dynamic mean field-theory (DMFT) may be needed to resolve the magnetic moment. Second, an AFM-DM phase transition may occur outside of the temperature range typically studied in the available literature. In theory a superexchange mechanism might result in a hidden magnetic phase transition. This paper calls for further experimental studies to resolve outstanding issues, in particular low-temperature specific heat capacity and muon spin relaxation measurements are needed.

Furthermore, we have shown how the crystal structure is influenced by the magnetic state. For instance, transverse 3k AFM behaviour results in *Pa* $\bar{3}$ crystal symmetry,^{22,44} synonymous with calculations of UO₂ and AmO₂ when SOI are considered. Finally, the electronic structure of PuO₂ is highly influenced by the magnetic state employed. These results have profound implications for investigations of bonding, reactivity, and surface structure. The re-orientation of magnetic vectors in surface calculations is often neglected (the direction of magnetism is usually defined by the z-axis), resulting in incomparable magnetic configurations between low-index surfaces. As the magnetic structure has a clear impact of the energetics, it is imperative that the re-orientation of magnetic vector is accounted for in computational surface investigations.

Conflicts of interest

The authors declare no competing financial interests.

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