



Cite this: *J. Mater. Chem. A*, 2024, 12, 29645

Spinel ferrites MFe_2O_4 ($M = Co, Cu, Zn$) for photocatalysis: theoretical and experimental insights†

Charlotte A. Hall,^{ab} Pilar Ferrer,^{id}*^b David C. Grinter,^{id}^b Santosh Kumar,^b Ivan da Silva,^{id}^c Juan Rubio-Zuazo,^{de} Peter Bencok,^b Frank de Groot,^f Georg Held,^{id}^b and Ricardo Grau-Crespo,^{id}*^a

Spinel ferrites exhibit significant promise in photocatalysis and other applications due to their compositional diversity and favourable electronic structure, magnetism, and partially tuneable cation distribution. However, their complex properties, for example, the different behaviour of bulk and nanostructured materials, are not well understood. Here, we combine advanced computational and experimental methods with reactivity measurements to explore the inversion degrees, electronic structures, and photocatalytic activities of MFe_2O_4 spinels ($M = Co, Cu, Zn$). X-ray diffraction and anomalous X-ray scattering measurements determined bulk inversion degrees of 0.81, 0.91, and 0.26 for $CoFe_2O_4$, $CuFe_2O_4$, and $ZnFe_2O_4$, respectively. Photocatalytic tests showed that only $ZnFe_2O_4$ is active in the oxygen evolution reaction (OER), which correlates with its favourable band alignment, as determined through electronic structure simulations. Surface-sensitive X-ray Absorption Spectroscopy (XAS) measurements provided insights into the cation distributions at the surfaces, showing significant deviations from bulk properties, particularly in $ZnFe_2O_4$ in which 52% of the near-surface tetrahedral sites are occupied by Fe cations, compared to 26% in the bulk. DFT simulations of $ZnFe_2O_4$ illustrated how the surface terminations can alter the thermodynamic preference for cation distribution in comparison with the bulk. Our findings illustrate the complex interplay between surface and bulk properties in spinel ferrites.

Received 16th July 2024
Accepted 30th September 2024

DOI: 10.1039/d4ta04941a

rsc.li/materials-a

1. Introduction

Spinel ferrites constitute a versatile family of materials with important applications in photocatalysis,^{1,2} water purification,³ biomedicine,⁴ and other fields.⁵ They are metal oxides with composition MFe_2O_4 , where M represents a divalent metal cation (e.g. Co^{2+} , Cu^{2+} , Zn^{2+}) and iron is present in trivalent form, Fe^{3+} . In a “normal” spinel the M^{2+} and Fe^{3+} cations occupy the tetrahedral (Td) and octahedral (Oh) positions, respectively, of the spinel structure (Fig. 1). But for some compositions, the

cations are redistributed across the Td and Oh sites, therefore the formula can be written as $(M_{1-x}Fe_x)[M_xFe_{2-x}]O_4$ where $()$ represents the Td sites and $[]$ represents the Oh sites. The degree of inversion ($0 \leq x \leq 1$) is defined here as the fraction of Fe^{3+} cations occupying the Td site.

Spinel ferrites are attractive materials for photocatalytic applications for several reasons. First, their electronic structure and optical properties can be tuned *via* their composition and/or cation distribution, which allows optimising light absorption or targeting specific band alignments.^{6–8} Second, they tend to be chemically stable under a wide range of temperatures and pH levels.⁹ Third, they are magnetic, which allows for easy recovery and reuse of the photocatalyst from the reaction mixture, reducing waste and improving process efficiency.⁵ Spinel ferrites are also relatively cheap catalysts and can be made up of naturally abundant metals.^{10,11} One potential application of spinel ferrite photocatalysts is water splitting to produce renewable hydrogen gas, H_2 .¹² Spinel ferrites such as $CoFe_2O_4$ and $CuFe_2O_4$ have been reported as potential photocatalysts for water splitting; however, they are typically used as part of composite photocatalysts.^{10,13} Drawbacks of cobalt and copper ferrites include poor conductivity, insufficient active sites, rapid charge carrier recombination, and irregular morphology.^{14,15} In

^aDepartment of Chemistry, University of Reading, Whiteknights, Reading RG6 6DX, UK. E-mail: r.grau-crespo@reading.ac.uk

^bDiamond Light Source, Harwell Science and Innovation Campus, Didcot OX11 0DE, UK. E-mail: pilar.ferrer-escorihuela@diamond.ac.uk

^cISIS Neutron and Muon Source, STFC, Rutherford Appleton Laboratory, Didcot, OX11 0QX, UK

^dBM25-SpLine, European Synchrotron Radiation Facility, CS40220, F-38043 Grenoble Cedex 9, France

^eICMM-CSIC, Sor Juana Inés de la Cruz 3, Cantoblanco, Madrid, Spain

^fDebye Institute for Nanomaterials Science, Utrecht University, 3584 CA Utrecht, The Netherlands

† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d4ta04941a>

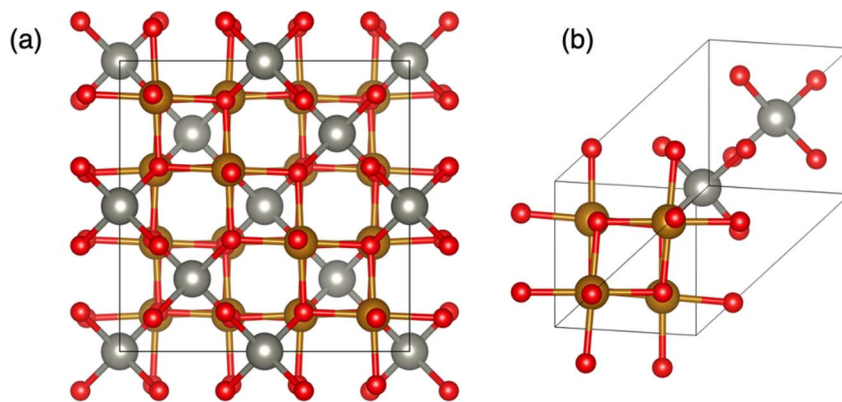


Fig. 1 Crystal structure of MFe_2O_4 spinels represented by (a) the conventional cubic cell and (b) the primitive unit cell. Colour scheme: tetrahedral (Td) sites = silver; octahedral (Oh) sites = gold; oxygen = red.

many studies where activity has been observed, a co-catalyst or sacrificial agent has been used alongside the spinel-based catalyst; the use of different sacrificial agents, such as oxalates^{16,17} or methanol,^{14,18} can have a significant effect on the efficiency of the catalyst. Another common strategy for improving the activity of $CoFe_2O_4$ or $CuFe_2O_4$ is constructing heterojunctions with other wide band gap materials, such as Fe_2O_3 (ref. 19 and 20) or C_3N_4 .^{21,22} Compared to cobalt and copper ferrites, zinc ferrite, $ZnFe_2O_4$, is a more efficient photocatalyst.²³ In a study by Rodríguez *et al.*,²⁴ more than twice the amount of H_2 was produced by $ZnFe_2O_4$ compared to $CoFe_2O_4$ over 8 hours with a methanol sacrificial agent. However, $ZnFe_2O_4$ is also still often used in conjunction with a co-catalyst or as a hetero-junction.^{23,25,26} Intrinsically altering the spinel by cation substitution, such as Ga in place of Fe, has proved effective in improving zinc ferrites performance as a photocatalyst.²⁷ To overcome the limitations of spinel ferrites for water splitting, a fundamental understanding of their electronic and photocatalytic properties is required.

Previous work has suggested substantial differences in the properties of spinel ferrites between bulk crystals and nanoparticles.^{28,29} For example, the cation distribution in nanoparticles can be significantly different from that in the bulk, and is heavily influenced by factors such as preparation method³⁰ and/or thermal treatment.³¹ $CoFe_2O_4$ is known to have fully inverse cation distribution ($x \approx 1$) in the bulk,^{32,33} whereas in nanoparticles lower degrees of inversion in the range of $x = 0.66$ – 0.68 are observed.^{28,34} $CuFe_2O_4$ also displays a high bulk inversion degree; Siddique *et al.*²⁹ report $x = 0.88$ in the bulk compared to $x = 0.80$ in nanoparticle form. However, inversion degrees as low as $x = 0.57$ have been observed in copper ferrite samples with a particle size of less than 10 nm.³⁵ In contrast, bulk $ZnFe_2O_4$ has a very low inversion degree of $x \approx 0$,³⁶ whereas $ZnFe_2O_4$ nanoparticles display a higher degree of inversion of up to $x = 0.4$, depending on particle size and thermal history.^{30,37} Understanding the properties that are inherent to the bulk materials and differentiating them from the effects of the surface is important in applications including photocatalysis, because both the bulk and the surface participate with different roles in the photocatalytic process.

In this study we have used a combination of computational simulations, X-ray absorption spectroscopy, X-ray diffraction, and photocatalytic activity measurements to investigate bulk and surfaces properties of MFe_2O_4 ($M = Co, Cu$ or Zn) in an effort to rationalise the bulk/surface behaviour of these metal ferrites nanoparticles (particle sizes < 35 nm). In addition to the characterisation of structural, photocatalytic and electronic properties, our modelling offers insights on the departures from bulk behaviour seen in small nanoparticles due to different behaviour of bulk and surfaces in terms of degree of inversion.

2. Methods

2.1 *Ab initio* simulations of bulk and surface models

The calculations were performed using density functional theory (DFT) as implemented in the VASP code.^{38,39} Geometry optimisations were performed using the generalised gradient approximation (GGA) with the Perdew–Burke–Ernzerhof (PBE) exchange–correlation functional.⁴⁰ Hubbard (GGA + U) corrections with U_{eff} values of 3.3 eV and 4.0 eV were applied to the Co and Cu/Fe d orbitals, respectively; these values were obtained by Wang *et al.*⁴¹ via fitting to the experimental oxidation enthalpies of the corresponding binary metal oxides, and have been found to transfer well to the study of more complex oxides (*e.g.* $FeSbO_4$,⁴² $LaCoO_3$ and $LaFeO_3$,⁴³ $YBa_2Fe_{3-x}Co_xO_8$,⁴⁴ and $BiFe_{1-x}Co_xO_3$ (ref. 45)). The interaction between the valence and core electrons was described with the projector augmented wave (PAW) method.⁴⁶ An energy cut-off of 520 eV, 30% above the recommended value for the PAW potentials, was used for all geometry optimisations involving cell volume charges, to decrease the Pulay errors.

For the bulk calculations, primitive unit cells of MFe_2O_4 ($M = Co, Cu, Zn$) containing two formula units were modelled with inversion degrees of $x = 0, 0.5$ and 1. For each inversion degree there is only one symmetrically different configuration in the primitive cell, therefore allowing us to calculate the inversion energy (configurational contribution only) as $\Delta E_{\text{conf}}(x) = E(x) - E(0)$. The electronic structure calculations to determine the band gap and alignment of the most stable configurations were



completed using the hybrid functional by Heyd, Scuseria and Ernzerhof (HSE06),⁴⁷ which includes 25% of Hartree–Fock exchange as well as range-separated screening with an attenuation parameter of $0.2 \text{ \AA}^{-1}$. The HSE06 calculations used the geometries optimised at GGA + U level, *i.e.* we did not re-optimize the structures at HSE06 level (but we checked, using ZnFe_2O_4 as a test, that this is an acceptable approximation, leading to both cell parameters and band gaps very close, within 0.5%, to those obtained with the full HSE06 approach). The band alignment was calculated using the top of the valence band and the bottom of the conduction band in the HSE06 band structure.

In all calculations, the magnetic moment for Fe^{3+} (and for Co^{2+} in the case of CoFe_2O_4) was initialised in high-spin state.^{48,49} For CoFe_2O_4 there is an additional degree of freedom: both cobalt and iron cations can exist in +2 or +3 oxidation states; therefore, charge transfer could occur resulting in Co^{3+} and Fe^{2+} being present. Test calculations were completed in which charge transfer and different spin states of the cations were considered. In all cases, the structures were either higher in energy or converged back to the more stable $\text{Co}^{2+}/\text{Fe}^{3+}$ high-spin configuration. Magnetic moment orientations were initialised at the magnetic ground state which was found by considering all the possible orientations of the moments. For ZnFe_2O_4 with normal distribution, an antiferromagnetic configuration of the Fe^{3+} in Oh sites had the lowest energy. However, a ferromagnetic orientation of Fe^{3+} in Oh sites was most stable for normally distributed CoFe_2O_4 and CuFe_2O_4 . For the ground state of all three spinels structures, when partially or fully inverse, the Fe^{3+} in Td sites had magnetic moments opposite to the moments of the Fe^{3+} in the Oh sites.

To simulate the (100) and (111) ZnFe_2O_4 terminated surfaces, four periodic slab models of ZnFe_2O_4 with different terminations separated by a vacuum gap of 10 Å were optimised using the same parameters outlined for the bulk GGA + U calculations. These surfaces of the spinel are Tasker type-III surfaces, in which there is a dipole moment perpendicular to the surface which can only be eliminated by surface reconstruction.⁵⁰ Therefore, we need to modify the slabs to build stoichiometric, non-polar surface models, which are generally expected to be stable under neutral (not very reducing or very oxidising) conditions. Having stoichiometric and non-polar slabs is also important in our work because they are used to estimate the offset of the macroscopic electrostatic potential between the bulk and the vacuum level. However, it is generally possible to study deviations in stoichiometry, as done elsewhere for spinel oxides,^{51–55} to understand the variation in surface stoichiometry with external conditions, such as the oxygen partial pressure; we have not conducted such analysis here. The details of reconstructions of the surfaces for our study are shown in the ESI Fig. 5,[†] and the notation for the stoichiometric non-polar surfaces follows the one used in ref. 52. As seen in Fig. 2, reconstructions A and B of the (100) surface terminate on Zn and Fe/O respectively. Building the (111) presented a greater challenge, as the unit cell needed to be expanded in both lateral directions. The (111) surfaces are Fe-terminated, but the (111)_B reconstruction also has Zn exposed at the surface, from the

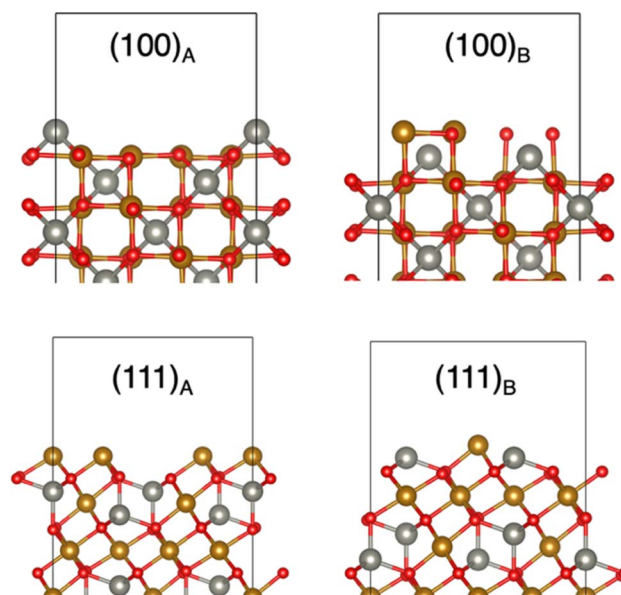


Fig. 2 Surface structures of ZnFe_2O_4 (100) and (111) surface reconstructions leading to stoichiometric and non-polar terminations. Only one side of the slab is shown but the other side of the slab is equivalent by symmetry. Colour scheme: Zn = silver; Fe = gold; oxygen = red.

second atomic layer. An inversion on the surface was modelled by switching one Fe atom in an Oh site with one Zn atom in a Td site on either side of the slabs to maintain the stoichiometry and symmetry. For the most stable surface termination further inversions were created propagating into the surface.

Surface energies (γ) for the different surface terminations were obtained from the equation:

$$\gamma = \frac{E_{\text{slab}} - E_{\text{bulk}}}{2A},$$

where E_{slab} is the total energy of the relaxed slab, E_{bulk} is the energy of the bulk with the same number of formula units as the respective slab and A is the surface area of one side of the slab. In the calculation both sides of the symmetric slab are allowed to relax and therefore both must be considered in the surface energy calculation.

2.2 Semi-empirical simulations of core-level spectra

The semi-empirical quantum many-body program QUANTY,^{56–58} within the graphical user interface CTM4XAS,⁵⁹ was used to simulate the Fe $L_{2,3}$ edges. This semi-empirical approach considers Coulomb interaction, spin–orbit coupling and crystal-field splitting around a given species, without consideration of the crystalline structure. Independent calculations were completed for the three Fe species observed in the spinels: Fe^{3+} in Oh and Td symmetry and Fe^{2+} in Oh symmetry. Based on experience in modelling similar systems, the Coulomb interaction was scaled to 94% and 88% of the Hartree–Fock values of the Slater integrals, whereas the spin–orbit coupling parameter were kept at 1.0 (no screening) for both core and valence levels.⁵⁹ A broadening of 0.1 eV was used for Gaussian functions and broadenings of 0.2–0.4 eV were used for Lorentzian functions in both the Near-Edge X-ray Absorption Fine Structure (NEXAFS)



and X-Ray Magnetic Circular Dichroism (XMCD) simulations. The broadening values for each Fe species have been outlined in Table 1, alongside the crystal field parameters. The integrated Fe^{2+} and Fe^{3+} spectra of the were normalised to the corresponding numbers of d electrons (6 and 5, respectively). The energies of the simulated spectral shapes were aligned by fitting to the experimental XMCD signals.

2.3 X-ray diffraction

Powder X-ray diffraction (PXRD) patterns from CuFe_2O_4 and ZnFe_2O_4 samples were collected on a Rigaku SmartLab diffractometer, using $\text{CuK}\alpha_1$ radiation, in reflection mode and at room temperature, over a 2θ range of 10 – 80° .

In the case of the CoFe_2O_4 spinel, the PXRD technique is limited because Co and Fe have similar atomic numbers (27 and 26, respectively), leading to similar scattering factors, and making it difficult to distinguish between these cations when they share a given spinel site. For this sample, an Anomalous X-Ray Scattering (AXRS) experiment was carried out on the multipurpose six-circle geometry diffractometer of SpLine BM25 Beamline at the European Synchrotron Radiation Facility (Grenoble, France). The CoFe_2O_4 sample was loaded in a 0.5 mm diameter borosilicate capillary, and the recorded diffraction patterns were collected for sample CoFe_2O_4 : one using a beam energy of 20 000 eV (6 – 60° 2θ range), and another one using an energy of 7097 eV (15 – 67° 2θ range), that is 15 eV below the Fe K-edge absorption edge at 7112 eV.

The Rietveld method was used for fitting the powder diffraction patterns and crystal structure refinements, by means of the Topas Academic v6 software. The structural starting model used for the refinements was the normal spinel structure and, for the case of AXRS data, the diffraction patterns at both incident energies were fitted simultaneously using the same structural parameters for CoFe_2O_4 . In all cases, the A and B cation occupancy at both Td and Oh crystallographic positions were refined, applying the constrain that both sites should be fully occupied, and that the final calculated formula should be AB_2O_4 .

Instrument peak profile parameters, which were calculated from a Silicon NIST-640C standard reference sample measured at the same conditions as the three different samples, were used to calculate broadening effects, due to crystalline size, and perform particle size analysis.

2.4 Catalytic testing

The catalytic testing was undertaken at the Catalysis Hub based at the Research Complex at Harwell. The photocatalytic oxygen

evolution was measured at room temperature in a gas-tight 50 mL quartz photoreactor. The light intensity was adjusted to 1 sun (100 mW cm^{-2}) using an AM 1.5G mass filtered 300 W Xe source. 25 mg of sample was used for each spinel (MFe_2O_4 , M = Co, Cu, Zn) measurement in a 0.5 M AgNO_3 medium to act as the hole scavenger. The system was purged for 1 hour with 1.5 bar Ar gas. The gas composition was monitored by gas chromatography with a barrier ionisation discharge (BID) detector (GC, Shimadzu GC-2010 Plus). The oxygen evolution was measured over 5 hours. Reference measurements for each medium were taken to normalise the oxygen evolution data for the spinel samples; further details of the control measurements can be seen in ESI Fig. 3.†

2.5 Near-edge X-ray absorption fine structure

Near-edge X-ray absorption fine structure (NEXAFS) measurements were carried out on branch B of the B07 (VerSoX) beamline at Diamond Light Source (DLS) using the total electron yield (TEY) mode collected under 1 mbar helium and corrected for the beamline transmission.^{60,61}

2.6 X-ray magnetic circular dichroism

X-ray magnetic circular dichroism (XMCD) measurements were carried out on the I10 beamline at DLS on the electromagnet end station in TEY mode. The measurements were done at normal incidence with a positive helicity at room temperature. Varied external magnetic fields of ± 1.5 to ± 1.9 T were applied to the spinels.⁶²

3. Results

3.1 Bulk structure: DFT simulations and X-ray diffraction

We first discuss the thermodynamics of cation distribution in the three ternary oxides CoFe_2O_4 , CuFe_2O_4 , and ZnFe_2O_4 . To approximate the free energy of inversion as a function of inversion degree and temperature, we interpolate the inversion energies obtained from the three DFT calculations at $x = 0$, 0.5 and 1 using a quadratic dependence, which was originally proposed by O'Neill and Navrotsky⁶³ and has subsequently been used in several investigations of the thermodynamics of inversion in spinels.^{64–66} The interpolated inversion energy functions, ΔE_{conf} , are shown in Fig. 3a. They only depend on the configuration and are therefore independent of the temperature.

From the calculated inversion energies, the configurational free energy of inversion, ΔF_{conf} , can be estimated as:

$$\Delta F_{\text{conf}} = \Delta E_{\text{conf}} - T\Delta S_{\text{conf}},$$

where ΔS_{conf} is the ideal configurational entropy of inversion:^{67,68}

$$\Delta S_{\text{conf}} = -R \left[x \ln x + (1-x) \ln(1-x) + x \ln \frac{x}{2} + (2-x) \ln \left(1 - \frac{x}{2} \right) \right].$$

Table 1 Crystal field parameters of independent Fe species for the CTM4XAS input

Fe species	10 Dq/eV	Exchange field/meV	
		XAS	XMCD
Fe^{3+} (Td)	−0.5	0	−90
Fe^{3+} (Oh)	1.6	0	90
Fe^{2+} (Oh)	1.2	0	90



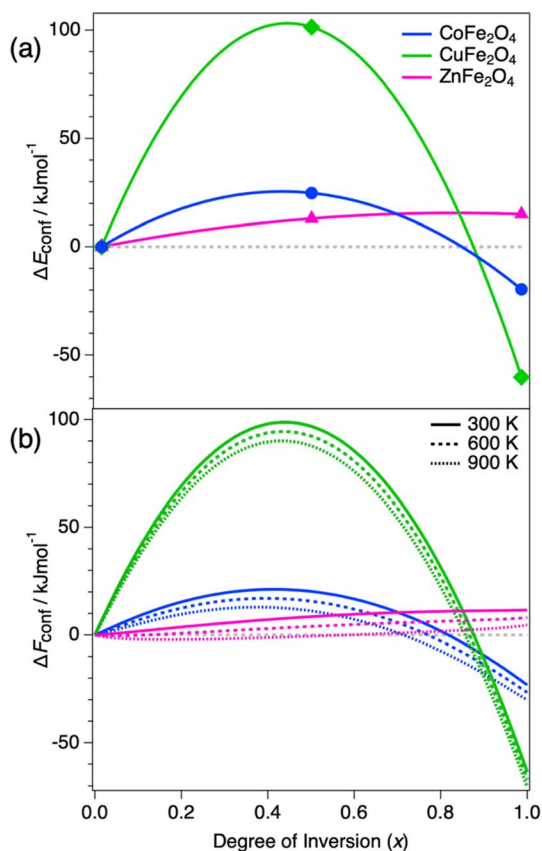


Fig. 3 (a) Inversion energies per formula unit (ΔE_{conf} , configurational contributions only) obtained by DFT, and (b) configurational free energies (ΔF_{conf}) at 300 K (solid line), 600 K (dotted line) and 900 K (dashed line) of CoFe₂O₄ (blue circles), CuFe₂O₄ (green diamonds) and ZnFe₂O₄ (pink triangles).

There might be other (excess) contributions to the inversion entropy, arising from energy differences between configurations at a given inversion degree, or from vibrational contributions. However, previous work⁶⁵ showed that these contributions are relatively small and can be ignored in a first approximation.

The dominant effect is the inversion energy, whereas the entropic term plays a relatively small role. Both CoFe₂O₄ and CuFe₂O₄ reach the minimum inversion energy at $x = 1$, implying a preference to be fully inverse. In contrast, the inversion energy of ZnFe₂O₄ is positive across the full range of x , with the most stable configuration being normal ($x = 0$). These results can be rationalised based on simple physical arguments. In A²⁺B₂³⁺O₄²⁻ spinels, the lattice (Madelung) energy slightly favours the normal cation distribution. Thus, in the absence of crystal field stabilisation energy (CFSE) effects, the normal distribution is preferred, as observed for ZnFe₂O₄ (d¹⁰ cations like Zn²⁺ and d⁵ cations like Fe³⁺ do not have CFSE). In both CoFe₂O₄ and CuFe₂O₄ the divalent cation is a transition metal with higher CFSE in the octahedral than in the tetrahedral site (excess octahedral stabilisation energy is 30.9 kJ mol⁻¹ for Co²⁺ and 63.5 kJ mol⁻¹ for Cu²⁺ ref. 69); therefore, these spinels favour the inverse distribution.

The inclusion of configurational entropy effects permits the consideration of finite temperatures within this simple model, but it does not change the picture considerably. The free energy of inversion variation with x at different temperatures is shown in Fig. 3b. In the cases of CoFe₂O₄ and CuFe₂O₄, the free energy of inversion minima occurs at $x = 1$ even at high temperatures up to 900 K. On the other hand, the inversion free energy minima of ZnFe₂O₄ are close to the normal end, ranging from $x = 0$ at room temperature up to $x = 0.2$ at 900 K.

We now consider how these theoretical bulk values of the inversion degree compare with the observed values for small nanoparticles. The nanoparticle size, inversion degree (x), cell parameter (a) and the oxygen coordinate parameter (u) of the spinel samples, as determined by AXRS or XRD with the Rietveld method, are listed in Table 2. The corresponding XRD and AXRS patterns can be seen in ESI Fig. 1 and 2.† The particle sizes of the cobalt, copper and zinc ferrites are 35, 24, and 22 nm respectively. CoFe₂O₄ and CuFe₂O₄ display high inversion degrees (x) of 0.81 and 0.91 respectively, which are slightly below the DFT-predicted value of 1 (fully inverse) for both systems. In contrast, the degree of inversion of ZnFe₂O₄ (0.26), although much lower compared to the cobalt and copper ferrites, is more inverse than the DFT predicted normal structure. The inversion energy curve, however, shows a very small x -dependence, therefore small additional entropy contribution can alter the position of the minimum significantly. Overall, the general trends of x agree with that of DFT predictions and with previous literature reports.^{29,30,32}

The absolute values of the cell and oxygen parameters of the nanoparticle samples are generally close to those calculated by DFT, with the largest percentage differences between the experimental and calculated a and u (2% and 3% respectively) being observed in the case of CuFe₂O₄. The trend in a values observed in the experimental data (Cu < Co < Zn) is also seen in the simulation results. Factors effecting the accuracy of the simulation prediction include the approximations made in the density functional (the generalised gradient approximation), the absence of nanostructuring effects (calculations are done for the infinite bulk crystal), and the temperature difference (0 K in DFT, room temperature for experiment). Also, a direct comparison between the parameters is difficult as the simulated spinels have extreme inversion degrees (*i.e.* $x = 0$ or 1), from which the nanoparticles deviate.

Table 2 Summary of experimentally determined (AXRS data for CoFe₂O₄ and XRD data for CuFe₂O₄ and ZnFe₂O₄) and DFT-calculated structural parameters x (inversion degree) and a (cell parameter). The u parameter determines the O atoms coordinates (u, u, u) in the standard setting of the $Fd\bar{3}m$ space group

Sample	Size/nm	Experimental (nanoparticles at room temperature)			DFT (bulk at 0 K)		
		x	$a/\text{\AA}$	u	x	$a/\text{\AA}$	u
CoFe ₂ O ₄	35	0.81	8.398	0.246	1	8.423	0.245
CuFe ₂ O ₄	24	0.91	8.371	0.249	1	8.409	0.242
ZnFe ₂ O ₄	22	0.26	8.439	0.243	0	8.481	0.239



3.2 Electronic structure, band alignment, and photocatalysis

The water splitting reaction can be separated into half-reactions, the hydrogen evolution reaction (HER), $2\text{H}^+ + 2\text{e}^- \rightarrow \text{H}_2$, and the oxygen evolution reaction (OER), $2\text{H}_2\text{O} \rightarrow 4\text{H}^+ + 4\text{e}^- + \text{O}_2$. The kinetics of the OER are substantially slower than those of the HER, as evidenced by a large overpotential, making this the rate-limiting step.^{12,70}

The photocatalytic OER using Ag^+ ions as an electron sacrificial agent on all the samples was carried out under simulated solar light (1 sun). As seen in Fig. 4, CoFe_2O_4 and CuFe_2O_4 exhibit very little photocatalytic oxygen evolution across 5 hours of irradiation ($<3 \mu\text{mol g}^{-1}$ of O_2). In contrast, ZnFe_2O_4 produced approximately $65 \mu\text{mol g}^{-1}$ of O_2 after 5 hours.

We now attempt to rationalise the photocatalytic behaviour of the samples in terms of the electronic structures and band alignments. The density of states (DOS), including the partial DOS contributions from the ions, of the spinels can be seen in Fig. 5. In each case, the Fe 3d levels are the main contribution to the conduction band (CB). On the other hand, the character of the valence band (VB) differs among the spinels: the high-lying filled Co 3d levels make the main contribution to the VB of CoFe_2O_4 , leading to the narrowest gap in the series; whereas the absence of d level contributions to the VB of ZnFe_2O_4 leads to the widest gap among the three spinels. The calculated band gaps of CoFe_2O_4 , CuFe_2O_4 and ZnFe_2O_4 are 1.96, 2.17 and 2.84 eV respectively, which are similar to those observed in respective nanoparticle samples in the literature.^{71,72} These band gap values are all sufficient in principle for photocatalysis of the overall water splitting reaction, for which a minimum thermodynamic potential of 1.23 eV is required.^{12,70}

In addition to having a suitable band gap, a semiconductor must also have CB and VB positions straddling the HER and OER levels in order to be a good water splitting photocatalyst (in a single-semiconductor configuration).¹² From the bulk simulations, the CB minimum and VB maximum are calculated with respect to the average electron potential in the solid. To compare these potentials with respect to the HER and OER potentials the electronic structure needs to be aligned relative to the vacuum

level. To do so, a slab calculation can be used to determine the potential difference (ΔV) between the pseudo-bulk average and the vacuum potential. Fig. 6 shows a stoichiometric slab, with a symmetric (100) terminated surface and vacuum level.

The calculated band alignment of the spinels with respect to vacuum scale are shown in Fig. 7, compared with potentials of the water splitting half-reactions. The potentials of the HER and OER in the vacuum scale at pH = 0 are -4.44 and -5.67 eV respectively. These energy levels are shifted up with a pH > 0 at temperature T by $k_{\text{B}}T \times \text{pH} \times \ln 10$.^{45,73} Therefore, at room temperature and pH = 7 the HER and OER potentials are -4.03 and -5.25 eV respectively, corresponding to those seen in Fig. 7. Despite all three spinels having a suitable band gap to catalyse the overall water splitting process, their band alignments do not fit the thermodynamic requirements, due their high-lying CB minima with respect to the HER potential (-4.03 eV). The lack of oxygen evolution displayed by CoFe_2O_4 and CuFe_2O_4 (Fig. 4) could be explained by the VB maxima positions, which lie above the OER potential (-5.25 eV). In contrast, the band alignment of ZnFe_2O_4 meets the thermodynamic requirements for the OER, with a VB potential -5.90 eV.

3.3 Surface effects: NEXAFS and XMCD experiments and slab calculations

While the prior discussion is focused on bulk properties, the behaviour at the oxide surfaces, where the actual catalytic reactions take place, might depart considerably from the bulk behaviour. Understanding the surface properties of these complex oxides is challenging, but some insights can be obtained from using surface-sensitive techniques, such as total electron yield (TEY) NEXAFS and XMCD. These techniques probe approximately 2 nm into the sample surface. The Fe $\text{L}_{2,3}$ edges measured in TEY mode by NEXAFS of CoFe_2O_4 , CuFe_2O_4 , and ZnFe_2O_4 are shown in Fig. 8a. The spectral features are dependent on the relative quantities of each iron species, as calculated in Fig. 8b. The intensity of the feature between the pre- and main edges at 708 eV (indicated by the red arrow in Fig. 8a) can be related to the presence or absence of tetrahedral (Td) Fe^{3+} . The width of the main edge at 709 eV is also affected by the cation distribution. In ZnFe_2O_4 , the intensity of this feature is lower compared to CoFe_2O_4 and CuFe_2O_4 , indicating that there is less Fe^{3+} (Td) (*i.e.* less inversion) in the zinc ferrite surface compared to the surface of the other two spinels. Therefore, the same trend in cation distribution is observed in both the surface and bulk of these materials. However, quantitatively determining the relative amounts of the different Fe species is difficult to do by just fitting NEXAFS spectra; more spectral features or information is required, which can be obtained from XMCD.

The Fe L_3 edges measured by XMCD seen in Fig. 9a–c (also collected in TEY mode, and therefore surface-sensitive) display spectral features that are more distinguishable between the different Fe species, compared to NEXAFS. The features at 707.5 eV, 708.7 eV and 709.4 eV can be attributed mainly to contributions from Fe^{2+} (Oh), Fe^{3+} (Td) and Fe^{3+} (Oh), respectively, with the Oh and Td ions displaying opposite dichroism.

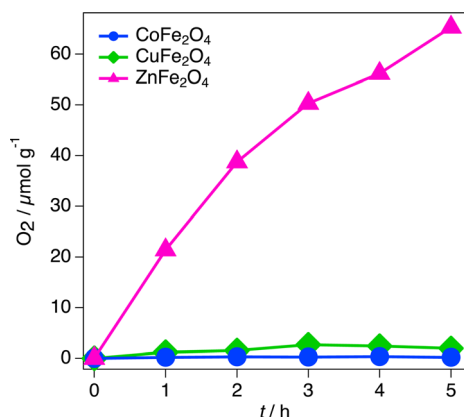


Fig. 4 Photocatalytic oxygen evolution of MFe_2O_4 ($\text{M} = \text{Co}, \text{Cu}$ or Zn) over a time of 5 hours under simulated sun light using AgNO_3 as an electron sacrificial agent.



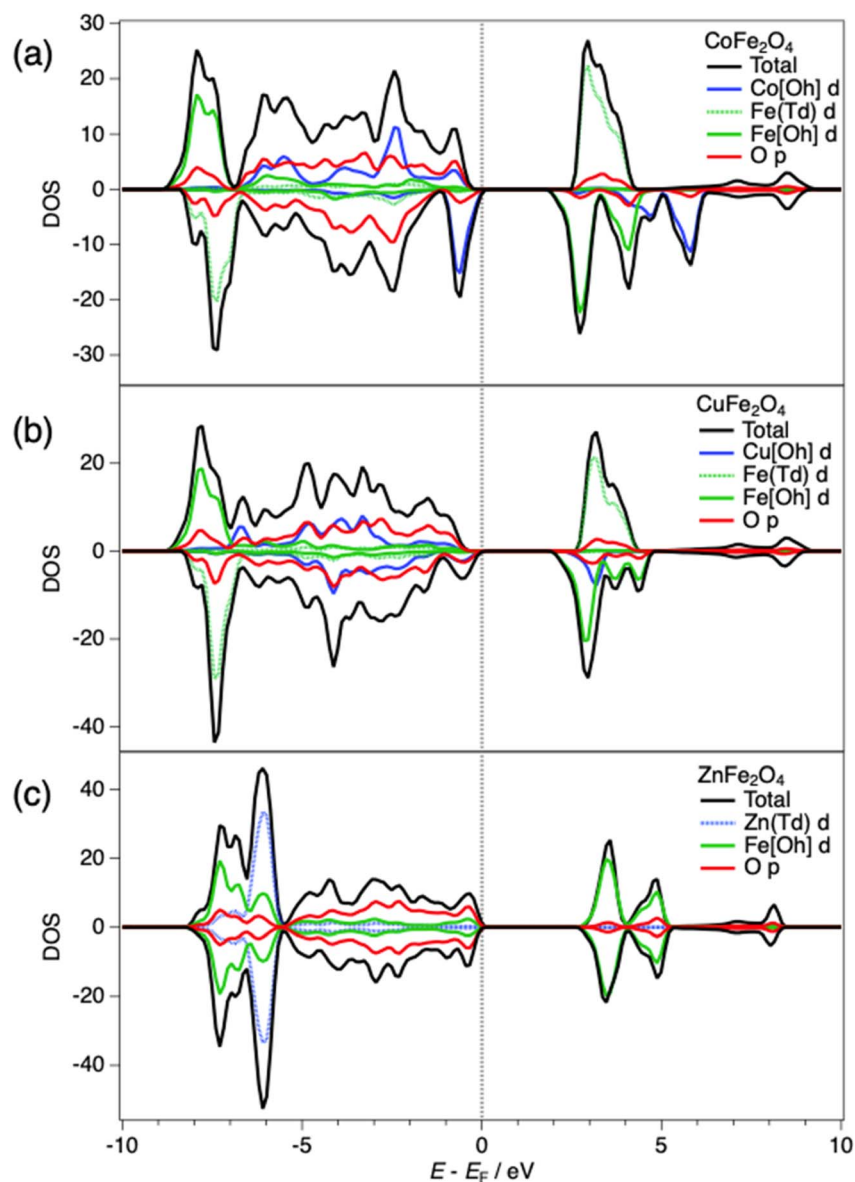


Fig. 5 Density of states (DOS) at HSE06 level for (a) CoFe_2O_4 , (b) CuFe_2O_4 and (c) ZnFe_2O_4 and partial DOS contributions from Co, Cu, Zn and Fe d orbitals and O p orbitals.

The difference of around 2 eV observed in our spinel XMCD spectra between the Fe^{2+} and Fe^{3+} peak maxima in Oh coordination is smaller than that reported for magnetite (Fe_3O_4), a fully inverse spinel (2.5 eV).⁷⁴ However, the relative shifts between the iron species peak maxima observed in Fig. 9a–c are comparable with XMCD shifts reported of spinel ferrite reported in the literature.^{75,76} The relative quantities of the Fe species at the surface can be estimated by fitting a combination of the calculated species-specific spectra (Fig. 9d) to the experimental spectra. Since the XMCD measurements were also collected in TEY mode, the Fe distribution at the near-surface can be determined. The percentages of $\text{Fe}^{2+}/\text{Fe}^{3+}$ in Oh/Td sites, as derived from the XMCD fit, are listed in the ESI Table 1.† The fit of the CoFe_2O_4 signal showed 76% percent of near-surface Td sites are occupied by Fe cations, in contrast to 81%

of Fe occupied Td sites in the bulk. A similar difference was observed in CuFe_2O_4 with 74% of the near-surface Td sites being occupied by Fe cations, compared to 91% in the bulk. ZnFe_2O_4 showed a significant contrast from 26% of Td occupied by Fe in the bulk to 52% at the surface. The XMCD signals also indicate that around 20% of the Fe in Oh was Fe^{2+} in all three of the samples, indicating some level of surface reduction which could be explained by the formation of oxygen vacancies or other surface defects.

To illustrate how the presence of the surface can alter the preferred cation distribution observed in the bulk, we conducted DFT simulations in ZnFe_2O_4 slabs with different surface terminations (Fig. 2) and cation distributions (swapping Zn and Fe cations at the top layer). Only ZnFe_2O_4 was considered for the surface calculations, since for this composition nanoparticles



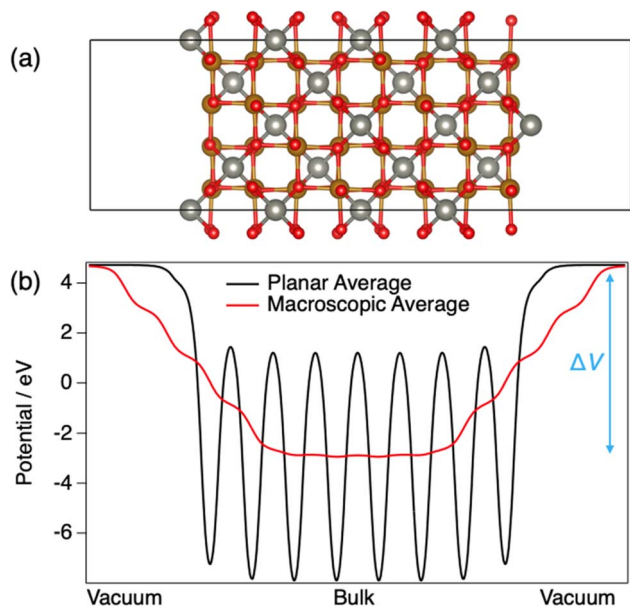


Fig. 6 (a) ZnFe_2O_4 slab with a (100) termination and (b) the planar-averages of the electrostatic potential.

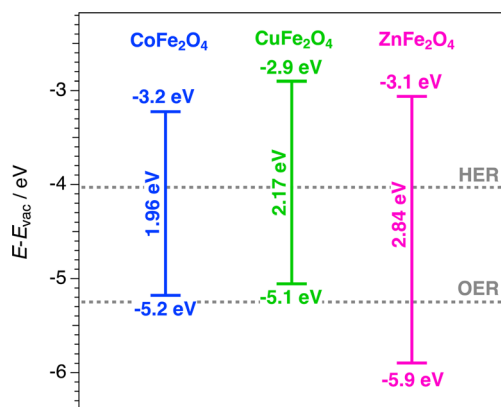


Fig. 7 Calculated CB and VB positions and band gaps of MFe_2O_4 ($\text{M} = \text{Co}, \text{Cu}$ or Zn). Half-reaction potentials for water splitting are represented by dotted lines.

have a significant difference in the inversion degree observed in the surface compared to the bulk. Also, unlike CoFe_2O_4 and CuFe_2O_4 , zinc ferrite was the only sample that demonstrated any catalytic activity for the OER.

The relaxation of the different terminations can be seen in ESI Fig. 6 and 7.† In the $(100)_\text{A}$ and $(111)_\text{A}$ terminated surfaces, the surface cations shift towards the bulk by up to $1.2 \text{ \AA}$, which generates minor distortion or shifting towards surface within their sub-surface layers. The $(100)_\text{B}$ sub-surface layers shift towards bulk, however the top cations remain relatively fixed in the square structure. Minimal cation shifting occurs in the surface and sub-surface layers in $(111)_\text{B}$ structures during relaxation.

The calculated surface energies are summarised in Table 3. The most stable surface is the Zn-terminated $(100)_\text{A}$ surface,

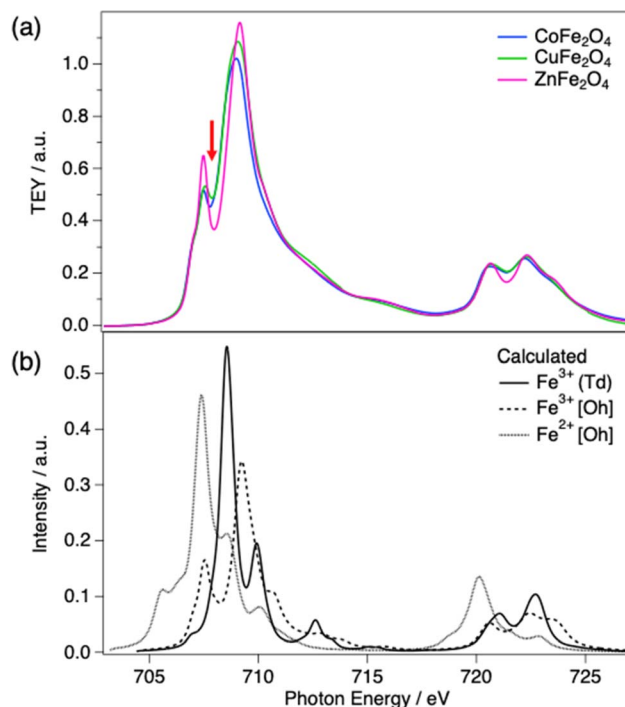


Fig. 8 (a) Fe $\text{L}_{2,3}$ edge NEXAFS spectra of MFe_2O_4 ($\text{M} = \text{Co}, \text{Cu}$ or Zn) and (b) calculated spectra of Fe^{3+} (Td), Fe^{3+} (Oh) and Fe^{2+} (Oh) by CTM4XAS.

whereas the Fe/O-terminated $(100)_\text{B}$ surface is much less stable. This result aligns with a previous *ab initio* study that found that the (100) surface of zinc ferrite is always more stable when Zn-terminated, regardless of the presence of O-rich or O-poor conditions.⁵⁴ We also find that the (111) surface is less stable than the $(100)_\text{A}$ termination. This contrasts with the conclusion from a theoretical study by Guo *et al.*⁵³ that found that the (111) surface is the most stable under the range of chemical potentials at which bulk ZnFe_2O_4 is stable. Since we do not perform an analysis here as a function of chemical potentials, it is difficult to compare with the results of ref. 53. However, for the purpose of this work, we are less interested in the relative stabilities of the surfaces, and more focused on the effect of changes in the cation distribution at the surface with respect to that of the bulk.

The comparison of the normal *vs.* inverted distribution of cations shows that in all cases the surface becomes more stable after the cation inversion at the surface. The $(100)_\text{A}$ surface has the smallest difference in surface energy between the normal and inverted surface (0.02 J m^{-2}). In contrast, the $(100)_\text{B}$ termination was the least stable normal surface, but showed the most stabilisation with the inversion. The $(111)_\text{B}$ termination is more stable than the $(111)_\text{A}$ termination with no inversion; however, when inverted there is only a 0.01 J m^{-2} difference in surface energy between the A and B terminations.

The stabilisation of inversion at the ZnFe_2O_4 surfaces illustrates how surface terminations, which imply a change in cation coordination, can alter the thermodynamic preferences observed in the bulk, which were driven by crystal field effects.

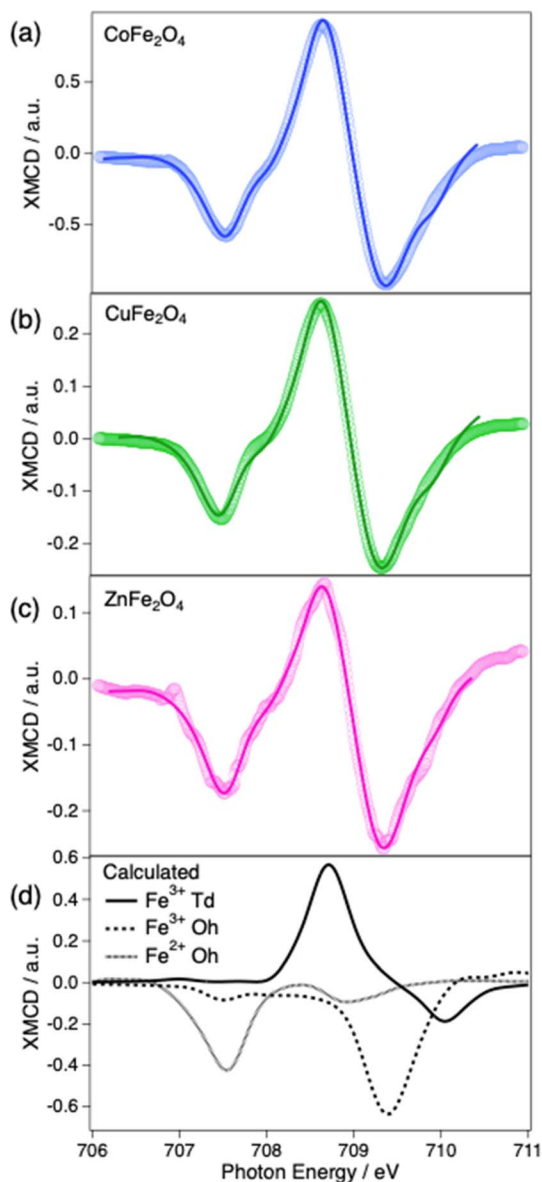


Fig. 9 Fe L_3 edge XMCD spectra of (a) CoFe_2O_4 , (b) CuFe_2O_4 and (c) ZnFe_2O_4 in which the circles are the experimental data and the solid line are the calculated spectra. (d) the calculated Fe L_3 edge of Fe^{3+} (Td), Fe^{3+} (Oh) and Fe^{2+} (Oh) by CTM4XAS.

Table 3 Calculated surface energies (γ) of the relaxed terminations of (100) and (111) surfaces of ZnFe_2O_4

Surface	Termination	Cation distribution at surface	$\gamma/\text{J m}^{-2}$
(100)	A	Normal	1.28
		Inverted	1.26
	B	Normal	2.91
		Inverted	1.91
(111)	A	Normal	2.32
		Inverted	1.62
	B	Normal	1.75
		Inverted	1.61

Table 4 Calculated inversion energies (ΔE) of the relaxed terminations of the (100)_A ZnFe_2O_4 surfaces with 0, 1 and 2 inverted cation pairs

Surface	Number of cation pair inversions on surface	$\Delta E/\text{kJ mol}^{-1}$
(100) _A	0	0.0
	1	−10.6
	2	−16.9

To investigate how deep this effect can propagate from the oxide in the top layer, a second cation pair in the sub surface was inverted in the most stable surface termination, (100)_A. The relaxation of this surface can be seen in ESI Fig. 8.† A similar shift and distortion in the surface and sub-surface layers is observed in both the (100)_A surface with one and two inverted cation pairs. The calculated inversion energies (ΔE) of the normal and inverted (100)_A surfaces are shown in Table 4. A negative inversion energy of $-10.6 \text{ kJ mol}^{-1}$ is observed when one inversion is created on the surface, therefore increasing the stability as seen in the surface energies (Table 3). When creating a second inversion in the sub-surface layer the inversion energy is even lower at $-16.9 \text{ kJ mol}^{-1}$. This demonstrates that it is thermodynamically favourable for inversion to be propagated deeper into top layer of ZnFe_2O_4 . In contrast, creating an inversion in the bulk is an unfavourable process, with a calculated inversion energy of 10.6 kJ mol^{-1} .

Given the small size and high specific surface of the oxide nanoparticles, these surface effects can have a significant impact on the overall cation distributions in the nanoparticles. It is indeed reported that small nanoparticles of ZnFe_2O_4 tend to have higher degree of inversion compared to bulk material.^{30,77,78} Due to the nature of the simulated surfaces, an extensive study of the surface effects, including reduction and oxygen vacancies, have not yet been investigated. However, given the important role of nanostructuring (and surfaces) in photocatalysis, these effects deserve further research attention.

4. Conclusions

Our comprehensive study on MFe_2O_4 ($\text{M} = \text{Co}, \text{Cu}, \text{Zn}$) spinel ferrites, using a combination of theoretical and experimental techniques, provides insights into their structure, electronic properties, and photocatalytic behaviours. The results from DFT simulations align well with experimental findings, revealing distinct inversion degrees and photocatalytic activities across the spinels. The DFT-predicted preference for inverse configurations in CoFe_2O_4 and CuFe_2O_4 and a normal configuration in ZnFe_2O_4 were confirmed by X-ray diffraction and AXRS measurements which showed inversion degrees of 0.81, 0.91, and 0.26, respectively.

Among the three spinel compositions, only ZnFe_2O_4 demonstrated photocatalytic activity for the oxygen evolution reaction (OER), generating $65 \mu\text{mol g}^{-1}$ of oxygen over 5 hours under UV irradiation. This can be attributed to its favourable band alignment, as demonstrated through our electronic structure simulations. CoFe_2O_4 and CuFe_2O_4 do not exhibit



OER activity, which could be explained by their band positions relative to the OER potential.

Surface-sensitive measurements *via* X-ray Absorption Spectroscopy (XAS) reveal significant variations in cation distribution at the surface compared to the bulk, particularly in CuFe_2O_4 (74% of the near-surface Td sites being occupied by Fe cations, compared to 91% in the bulk) and ZnFe_2O_4 (52% of the near-surface Td sites occupied by Fe cations, compared to 26% in bulk). These findings highlight the influence of surface chemistry on the photocatalytic properties of these materials. Further, DFT simulations of surface terminations provided additional understanding of the stability and properties of the surfaces, showing that cation inversion is energetically favourable at the surfaces of ZnFe_2O_4 , even if it is not in the bulk. Our calculations show that the effect of the surface termination in the energetics of cation inversion propagates beyond the top surface layer. The role of surface defects such as oxygen vacancies in the stabilisation of cation inversion at the surface requires investigation in the future.

This study not only deepens the understanding of spinel ferrites but also illustrates the critical role of both bulk and surface properties in determining the photocatalytic performance of these materials. Future work should continue to explore the intricate relationship between structural characteristics and catalytic activities, potentially leading to the development of more effective photocatalytic materials based on spinel ferrites.

Data availability

The experimental and computational data underlying this study are openly available in the University of Reading Research Data Archive at <https://doi.org/10.17864/1947.001342>.

Conflicts of interest

No conflicts of interest.

Acknowledgements

We are grateful to the UK Materials and Molecular Modelling Hub for computational resources, which is partially funded by EPSRC (EP/T022213/1, EP/W032260/1 and EP/P020194/1). C. A. H. acknowledges financial support for her PhD studies by Diamond Light Source (DLS) and the University of Reading. Collaboration with the Utrecht group was funded by COST Action CA18234 (CompNanoEnergy), supported by COST (European Cooperation in Science and Technology) <https://www.cost.eu>. Authors thank D. L. S. for beamtime on B07-B under proposal SI33639 and SI34919 and on I10 under proposal MM36558; ESFR for beamtime on BM25 under proposal A25-2-1023; ISIS Neutron and Muon Source, for access to the facilities at the Materials Characterisation Laboratory. We also thank the Research Complex and the UK Catalysis Hub at Harwell Campus. J. R. Z. acknowledges the Spanish Ministerio de Ciencia, Innovación y Universidades and Consejo Superior de Investigaciones Científicas for financial support and for

provision of synchrotron radiation facilities at BM25-SpLine (PIE 2010 6 0E 013 and PIE 2021 60 E 030).

References

- 1 H. Ma and C. Liu, *Front. Energy*, 2021, **15**, 621–630.
- 2 T. Tatarchuk, B. Al-Najar, M. Bououdina and M. A. A. Ahmed, in *Handbook of Ecomaterials*, 2018, ch. 158-1, pp. 1–50, DOI: [10.1007/978-3-319-48281-1_158-1](https://doi.org/10.1007/978-3-319-48281-1_158-1).
- 3 D. H. K. Reddy and Y.-S. Yun, *Coord. Chem. Rev.*, 2016, **315**, 90–111.
- 4 M. Amiri, M. Salavati-Niasari and A. Akbari, *Adv. Colloid Interface Sci.*, 2019, **265**, 29–44.
- 5 S. J. Salih and W. M. Mahmood, *Heliyon*, 2023, **9**, e16601.
- 6 M. Feng, A. Yang, X. Zuo, C. Vittoria and V. G. Harris, *J. Appl. Phys.*, 2010, **107**, 09A521.
- 7 G. Liu, J. Wang, X. Sheng, X. Xue and Y. Wang, *Ceram. Int.*, 2023, **49**, 29747–29754.
- 8 Z. Szotek, W. M. Temmerman, D. Ködderitzsch, A. Svane, L. Petit and H. Winter, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2006, **74**, 174431.
- 9 V. Tsurkan, H.-A. Krug von Nidda, J. Deisenhofer, P. Lunkenheimer and A. Loidl, *Phys. Rep.*, 2021, **926**, 1–86.
- 10 M. G. Idris, H. Y. Hafeez, J. Mohammed, A. B. Suleiman and C. E. Ndikilar, *Appl. Surf. Sci. Adv.*, 2023, **18**, 100468.
- 11 K. Malaie, Z. Heydari and M. R. Ganjali, *Int. J. Hydrogen Energy*, 2021, **46**, 3510–3529.
- 12 S. A. Razeq, M. R. Popeil, L. Wangoh, J. Rana, N. Suwandaratne, J. L. Andrews, D. F. Watson, S. Banerjee and L. F. J. Piper, *Electron. Struct.*, 2020, **2**, 023001.
- 13 S. Chandrasekaran, C. Bowen, P. Zhang, Z. Li, Q. Yuan, X. Ren and L. Deng, *J. Mater. Chem. A*, 2018, **6**, 11078–11104.
- 14 S. Bera, S. Ghosh, T. Maiyalagan and R. N. Basu, *ACS Appl. Energy Mater.*, 2022, **5**, 3821–3833.
- 15 H. Wang, Q. Hu, J. Qiu, R. Guo and X. Liu, *Catal. Sci. Technol.*, 2023, **13**, 6102–6125.
- 16 M. Benlambarek, N. Salhi, R. Benrabaa, A. M. Djaballah, A. Boulahouache and M. Trari, *Int. J. Hydrogen Energy*, 2022, **47**, 9239–9247.
- 17 H. Yang, J. Yan, Z. Lu, X. Cheng and Y. Tang, *J. Alloys Compd.*, 2009, **476**, 715–719.
- 18 G. He, Y. Wen, C. Ma, X. Li, L. Gao and Z. Sun, *Int. J. Hydrogen Energy*, 2021, **46**, 5369–5377.
- 19 A. Farooq, S. Khalil, B. Basha, A. Habib, M. S. Al-Buriah, M. F. Warsi, S. Yousaf and M. Shahid, *Int. J. Hydrogen Energy*, 2024, **51**, 1318–1332.
- 20 S. Hussain, S. Hussain, A. Waleed, M. M. Tavakoli, Z. Wang, S. Yang, Z. Fan and M. A. Nadeem, *ACS Appl. Mater. Interfaces*, 2016, **8**, 35315–35322.
- 21 S. Bellamkonda, C. Chakma, S. Guru, B. Neppolian and G. R. Rao, *Int. J. Hydrogen Energy*, 2022, **47**, 18708–18724.
- 22 R. Cheng, X. Fan, M. Wang, M. Li, J. Tian and L. Zhang, *RSC Adv.*, 2016, **6**, 18990–18995.
- 23 R. Dillert, D. H. Taffa, M. Wark, T. Bredow and D. W. Bahnemann, *APL Mater.*, 2015, **3**, 104001.
- 24 A. A. Rodríguez-Rodríguez, M. B. Moreno-Trejo, M. J. Meléndez-Zaragoza, V. Collins-Martínez, A. López-



- Ortís, E. Martínez-Guerra and M. Sánchez-Domínguez, *Int. J. Hydrogen Energy*, 2019, **44**, 12421–12429.
- 25 L. Kong, Z. Jiang, T. Xiao, L. Lu, M. O. Jones and P. P. Edwards, *Chem. Commun.*, 2011, **47**, 5512–5514.
- 26 S. Xu, D. Feng and W. Shangguan, *J. Phys. Chem. C*, 2009, **113**, 2463–2467.
- 27 X. Xu, A. K. Azad and J. T. S. Irvine, *Catal. Today*, 2013, **199**, 22–26.
- 28 D. Carta, M. F. Casula, A. Falqui, D. Loche, G. Mountjoy, C. Sangregorio and A. Corrias, *J. Phys. Chem. C*, 2009, **113**, 8606–8615.
- 29 M. Siddique and N. M. Butt, *Phys. B Condens. Matter*, 2010, **405**, 4211–4215.
- 30 F. S. Li, L. Wang, J. B. Wang, Q. G. Zhou, X. Z. Zhou, H. P. Kunkel and G. Williams, *J. Magn. Magn. Mater.*, 2004, **268**, 332–339.
- 31 L. I. Granone, R. Dillert, P. Heitjans and D. W. Bahnemann, *ChemistrySelect*, 2019, **4**, 1232–1239.
- 32 J. Venturini, A. M. Tonelli, T. B. Wermuth, R. Y. S. Zampiva, S. Arcaro, A. Da Cas Viegas and C. P. Bergmann, *J. Magn. Magn. Mater.*, 2019, **482**, 1–8.
- 33 W. H. Wang and X. Ren, *J. Cryst. Growth*, 2006, **289**, 605–608.
- 34 H. L. Andersen, M. Saura-Muzquiz, C. Granados-Miralles, E. Canevet, N. Lock and M. Christensen, *Nanoscale*, 2018, **10**, 14902–14914.
- 35 D. Pajić, K. Zadro, R. E. Vanderberghe and I. Nedkov, *J. Magn. Magn. Mater.*, 2004, **281**, 353–363.
- 36 J. R. Sandemann, K. A. H. Stockler, X. Wang, B. C. Chakoumakos and B. B. Iversen, *J. Am. Chem. Soc.*, 2023, **145**, 21053–21065.
- 37 J. Philip, G. Gnanaprakash, G. Panneerselvam, M. Antony, T. Jayakumar and B. Raj, *J. Appl. Phys.*, 2007, **102**, 054305.
- 38 G. Kresse and J. Furthmüller, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1996, **54**, 11169.
- 39 G. Kresse and J. Furthmüller, *Comput. Mater. Sci.*, 1996, **6**, 15–50.
- 40 J. P. Perdew, K. Burke and M. Ernzerhof, *Phys. Rev. Lett.*, 1996, **77**, 3865.
- 41 L. Wang, T. Maxisch and G. Ceder, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2006, **73**, 195107.
- 42 R. Grau-Crespo, F. Corà, A. A. Sokol, N. H. de Leeuw and C. R. A. Catlow, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2006, **73**, 035116.
- 43 Y.-L. Lee, M. J. Gadre, Y. Shao-Horn and D. Morgan, *Phys. Chem. Chem. Phys.*, 2015, **17**, 21643–21663.
- 44 C. Collins, M. S. Dyer, A. Demont, P. A. Chater, M. F. Thomas, G. R. Darling, J. B. Claridge and M. J. Rosseinsky, *Chem. Sci.*, 2014, **5**, 1493–1505.
- 45 S. Grover, K. T. Butler, U. V. Waghmare and R. Grau-Crespo, *Adv. Theory Simul.*, 2023, **6**, 2200673.
- 46 P. E. Blöchl, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1994, **50**, 17953–17979.
- 47 J. Heyd, G. E. Scuseria and M. Ernzerhof, *J. Chem. Phys.*, 2003, **118**, 8207–8215.
- 48 A. Hossain, M. S. I. Sarker, M. K. R. Khan and M. M. Rahman, *Mater. Sci. Eng. B*, 2020, **253**, 114496.
- 49 Y. H. Hou, Y. J. Zhao, Z. W. Liu, H. Y. Yu, X. C. Zhong, W. Q. Qiu, D. C. Zeng and L. S. Wen, *J. Phys. D: Appl. Phys.*, 2010, **43**, 445003.
- 50 P. W. Tasker, *J. Phys. C: Solid State Phys.*, 1979, **12**, 4977.
- 51 Q. Cai, J.-g. Wang, Y. Wang and D. Mei, *J. Phys. Chem. C*, 2016, **120**, 19087–19096.
- 52 D. Santos-Carballal, A. Roldan, R. Grau-Crespo and N. H. de Leeuw, *Phys. Chem. Chem. Phys.*, 2014, **16**, 21082–21097.
- 53 H. Guo, A. C. Marschilok, K. J. Takeuchi, E. S. Takeuchi and P. Liu, *ACS Appl. Mater. Interfaces*, 2018, **10**, 35623–35630.
- 54 K. L. S. Rodríguez, J. J. M. Quintero, H. H. Medina Chanduvi, A. V. G. Rebaza, R. Faccio, W. A. Adeagbo, W. Hergert, C. E. R. Torres and L. A. Errico, *Appl. Surf. Sci.*, 2020, **499**, 143859.
- 55 R. E. Warburton, H. Iddir, L. A. Curtiss and J. Greeley, *ACS Appl. Mater. Interfaces*, 2016, **8**, 11108–11121.
- 56 M. W. Haverkort, G. Sangiovanni, P. Hansmann, A. Toschi, Y. Lu and S. Macke, *Europhys. Lett.*, 2014, **108**, 57004.
- 57 M. W. Haverkort, M. Zwierzycki and O. K. Andersen, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2012, **85**, 165113.
- 58 Y. Lu, M. Höppner, O. Gunnarsson and M. W. Haverkort, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2014, **90**, 085102.
- 59 E. Stavitski and F. M. F. de Groot, *Micron*, 2010, **41**, 687–694.
- 60 G. Held, F. Venturini, D. C. Grinter, P. Ferrer, R. Arrigo, L. Deacon, W. Quevedo Garzon, K. Roy, A. Large, C. Stephens, A. Watts, P. Larkin, M. Hand, H. Wang, L. Pratt, J. J. Mudd, T. Richardson, S. Patel, M. Hillman and S. Scott, *J. Synchrotron Radiat.*, 2020, **27**, 1153–1166.
- 61 D. C. Grinter, P. Ferrer, F. Venturini, M. A. van Spronsen, A. I. Large, S. Kumar, M. Jaugstetter, A. Iordachescu, A. Watts, S. L. M. Schroeder, A. Kroner, F. Grillo, S. M. Francis, P. B. Webb, M. Hand, A. Walters, M. Hillman and G. Held, *J. Synchrotron Radiat.*, 2024, 578–589, DOI: [10.1107/S1600577524001346](https://doi.org/10.1107/S1600577524001346).
- 62 H. Wang, P. Bencok, P. Steadman, E. Longhi, J. Zhu and Z. Wang, *J. Synchrotron Radiat.*, 2012, **19**, 944–948.
- 63 H. S. C. O'Neill and A. Navrotsky, *Am. Mineral.*, 1983, **68**, 181–194.
- 64 D. Santos-Carballal, A. Roldan, R. Grau-Crespo and N. H. de Leeuw, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2015, **91**, 195106.
- 65 Y. Seminovski, P. Palacios, P. Wahnón and R. Grau-Crespo, *Appl. Phys. Lett.*, 2012, **100**, 102112.
- 66 S.-H. Wei and S. Zhang, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2001, **63**, 075205.
- 67 A. Navrotsky and O. J. Kleppa, *J. Inorg. Nucl. Chem.*, 1967, **29**, 2701–2714.
- 68 F. Tielens, M. Calatayud, R. Franco, J. M. Recio, J. Pérez-Ramírez and C. Minot, *J. Phys. Chem. B*, 2006, **110**, 988–995.
- 69 J. D. Dunitz and L. E. Orgel, in *Advances in Inorganic Chemistry and Radiochemistry*, ed. H. J. Emeleus and A. G. Sharpe, Academic Press, 1960, vol. 2, pp. 1–60.
- 70 L. Yuan, C. Han, M.-Q. Yang and Y.-J. Xu, *Int. Rev. Phys. Chem.*, 2016, **35**, 1–36.
- 71 N. Guijarro, P. Borno, M. Prévot, X. Yu, X. Zhu, M. Johnson, X. Jeanbourquin, F. Le Formal and K. Sivula, *Sustain. Energy Fuels*, 2018, **2**, 103–117.



- 72 M. Sundararajan, L. John Kennedy, P. Nithya, J. Judith Vijaya and M. Bououdina, *J. Phys. Chem. Solids*, 2017, **108**, 61–75.
- 73 Q. Liang, G. Brocks and A. Bieberle-Hütter, *J. Phys.: Energy*, 2021, **3**, 026001.
- 74 H. Elnaggar, R. Wang, M. Ghiasi, M. Yañez, M. U. Delgado-Jaime, M. H. Hamed, A. Juhin, S. S. Dhesi and F. de Groot, *Phys. Rev. Mater.*, 2020, **4**, 026001.
- 75 B. Liu, C. Piamonteze, M. U. Delgado-Jaime, R.-P. Wang, J. Heidler, J. Dreiser, R. Chopdekar, F. Nolting and F. M. F. de Groot, *Phys. Rev. B*, 2017, **96**, 026001.
- 76 R. A. D. Pattrick, G. Van Der Laan, C. M. B. Henderson, P. Kuiper, E. Dudzik and D. J. Vaughan, *Eur. J. Mineral.*, 2002, **14**, 1095–1102.
- 77 V. Blanco-Gutierrez, E. Climent-Pascual, M. J. Torralvo-Fernandez, R. Saez-Puche and M. T. Fernandez-Diaz, *J. Solid State Chem.*, 2011, **184**, 1608–1613.
- 78 M. Á. Cobos, P. de la Presa, I. Puente-Orench, I. Llorente, I. Morales, A. García-Escorial, A. Hernando and J. A. Jiménez, *Ceram. Int.*, 2022, **48**, 12048–12055.

