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# Local coordination geometry within cobalt spinel oxides mediates photoinduced polaron formation†

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Understanding the photophysics of transition metal oxides is crucial for these materials to realize their considerable potential in applications such as photocatalysis and optoelectronics. Recent studies suggest that formation of localized excited states consisting of polarons (quasi-particles comprising a charge carrier strongly coupled to a proximal lattice distortion) plays a crucial role in the photophysics of these materials. Cobalt-containing spinel oxides ( $\text{Co}_3\text{O}_4$  and  $\text{ZnCo}_2\text{O}_4$ ) offer a unique opportunity to investigate the influence of local geometry, and cation inversion on photoinduced polaron formation. Here, we use Hubbard-corrected density functional theory (DFT +  $U$ ) paired with resonance Raman and temperature-dependent optical spectroscopies to demonstrate that low-energy transitions observed in  $\text{Co}_3\text{O}_4$  are associated with d–d transitions involving cobalt ions occupying tetrahedral sites within the spinel lattice. These low-energy optical transitions exhibit strong coupling to phonon modes associated with tetrahedral sites. Replacing most tetrahedral cobalt ions with zinc produces the slightly inverted ternary spinel material,  $\text{ZnCo}_2\text{O}_4$ , in which we observe a phonon-coupled optical transition that occurs at the same energy as observed in  $\text{Co}_3\text{O}_4$ . We propose that these phonon-coupled optical transitions enable direct access to a polaronic state upon photoexcitation; however, the intensity of this optical transition depends on temperature in  $\text{Co}_3\text{O}_4$ , whereas no significant temperature dependence is observed in  $\text{ZnCo}_2\text{O}_4$ . We therefore hypothesize that in  $\text{Co}_3\text{O}_4$  the mechanism of polaron formation is coupling of the optical transition to dynamic, thermally-gated lattice distortions, whereas, in  $\text{ZnCo}_2\text{O}_4$ , the transition couples to static lattice defects that arise from the presence of a small population of tetrahedrally-coordinated cobalt ions.

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## Introduction

Transition metal oxides have promising characteristics for solar energy conversion technologies because of their visible band gap energies, stability, abundance, cost-effectiveness and low toxicity.<sup>1,2</sup> However, these materials contain weakly dispersive bands originating from the metal 3d orbitals that lead to low charge-carrier mobility,<sup>3,4</sup> rapid charge recombination,<sup>5,6</sup> and formation of localized, “self-trapped” states near the band-edges.<sup>7</sup> These states, which comprise one or more localized charge carriers coupled to proximal lattice distortions arising from one or more phonons, are known as polarons.<sup>7,8</sup> Small polarons, first described by Holstein,<sup>9,10</sup> are described by

a short-range carrier–phonon interaction ( $\text{radius}_{\text{polaron}} \sim \text{lattice parameter}$ ) and carrier mobility that increases with increased temperature.<sup>11</sup> Conversely, large polarons have long-range carrier–phonon interactions ( $\text{radius}_{\text{polaron}} \gg \text{lattice parameter}$ ) with carrier mobility that decreases with increased temperature.<sup>12,13</sup> Many transition metal oxides are reported to host small polarons,<sup>7,14–19</sup> and in many cases the performance of these materials for electrocatalytic, photoelectrocatalytic, and optoelectronic applications is impacted by small polaron formation.<sup>20–25</sup>

Although conductivity studies have contributed to a thorough understanding of polaron transport mechanisms in transition metal oxides,<sup>11,15,17,26,27</sup> there is still a limited understanding of the mechanisms by which polarons form in photoexcited states. In hematite ( $\alpha\text{-Fe}_2\text{O}_3$ ), for example, one proposed mechanism involves indirect population of polaron states *via* relaxation from an initially excited, higher-energy non-polaronic state.<sup>20,28</sup> Our group recently identified an additional mechanism involving the direct population of polaronic excited states at room temperature upon band-edge excitation in  $\alpha\text{-Fe}_2\text{O}_3$ .<sup>29,30</sup> Fig. 1 depicts the difference between indirect and direct population of polaronic states *via* photoexcitation.

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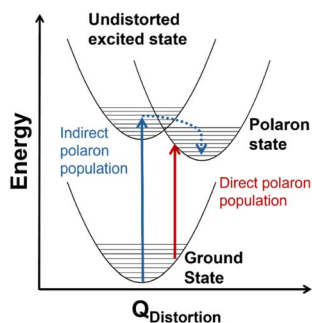


Fig. 1 Conceptual configuration coordinate diagram depicting mechanisms of photoexcited polaron formation. The red arrow illustrates direct photoexcitation from a thermally distorted ground state into a polaronic state and the blue arrow shows relaxation into a polaronic state *via* an undistorted ground state.

Importantly, optical population of polaronic states in  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub> is proposed to occur even in a pristine, defect-free lattice: the optical transitions couple to intrinsic, thermally-activated phonon distortions within the crystal.<sup>29,30</sup> There is also evidence of polaron formation arising from charge carriers coupling to intrinsic lattice distortions in rutile TiO<sub>2</sub> and LiNbO<sub>4</sub>.<sup>16,31</sup> These self-trapping mechanisms differ from other descriptions of small polarons forming *via* charge carriers coupling to dopants or lattice defects.<sup>32–35</sup> Distinguishing the mechanism of photoinduced polaron formation (mediated by dynamic thermally activated lattice distortions or static lattice defects) is fundamentally important to the development of

strategies for engineering metal oxide materials for photoapplications.

Spinel oxides (AB<sub>2</sub>O<sub>4</sub>) offer a unique opportunity to understand how the mechanism of photoinduced polaron formation depends on orbital composition of the band edge, coordination geometry of metal centers, and the presence of substitutional defects. These materials are mixed-valent with *Fd3m* symmetry and two different site symmetries for metals: tetrahedral (T<sub>d</sub>) and octahedral (O<sub>h</sub>). Ternary spinel oxides, in which A and B are different metals, are described by an inversion parameter  $x$  ( $0.0 < x < 1.0$ ) that quantifies the percentage of the A cations that occupy octahedral sites. When  $x = 0.0$ , meaning all of the A cations are in tetrahedral sites, the spinel is considered “normal,” whereas when  $x = 1.0$ , meaning all of the A cations are in octahedral sites, the spinel is fully inverted. Intermediate values of  $x$  correspond to population of A cations in both tetrahedral and octahedral sites. Spinel oxides containing cobalt are of particular interest because of the difference in crystal field splittings of Co<sup>2+</sup> T<sub>d</sub> and Co<sup>3+</sup> O<sub>h</sub> (Fig. 2A). Transitions between the  $e$  and  $t_2$  states within the T<sub>d</sub> sites are allowed by the Laporte selection rule whereas this rule forbids transitions between the  $t_{2g}$  and  $e_g$  states within the O<sub>h</sub> sites. Normal Co<sub>3</sub>O<sub>4</sub>, which contains both Co<sup>2+</sup> T<sub>d</sub> and Co<sup>3+</sup> O<sub>h</sub>, exhibits optical transitions associated with localized d–d transitions at 0.8, 0.9 and 1.6 eV, in addition to an optical transition at 2.5 eV associated with a ligand-to-metal-charge transfer (LMCT) transition.<sup>36–40</sup> Normal ZnCo<sub>2</sub>O<sub>4</sub>, with only Co<sup>3+</sup> O<sub>h</sub>, is reported to have only the LMCT-type transition at 2.5 eV.<sup>27</sup> The coordination geometry of cobalt in spinel oxides thus impacts their

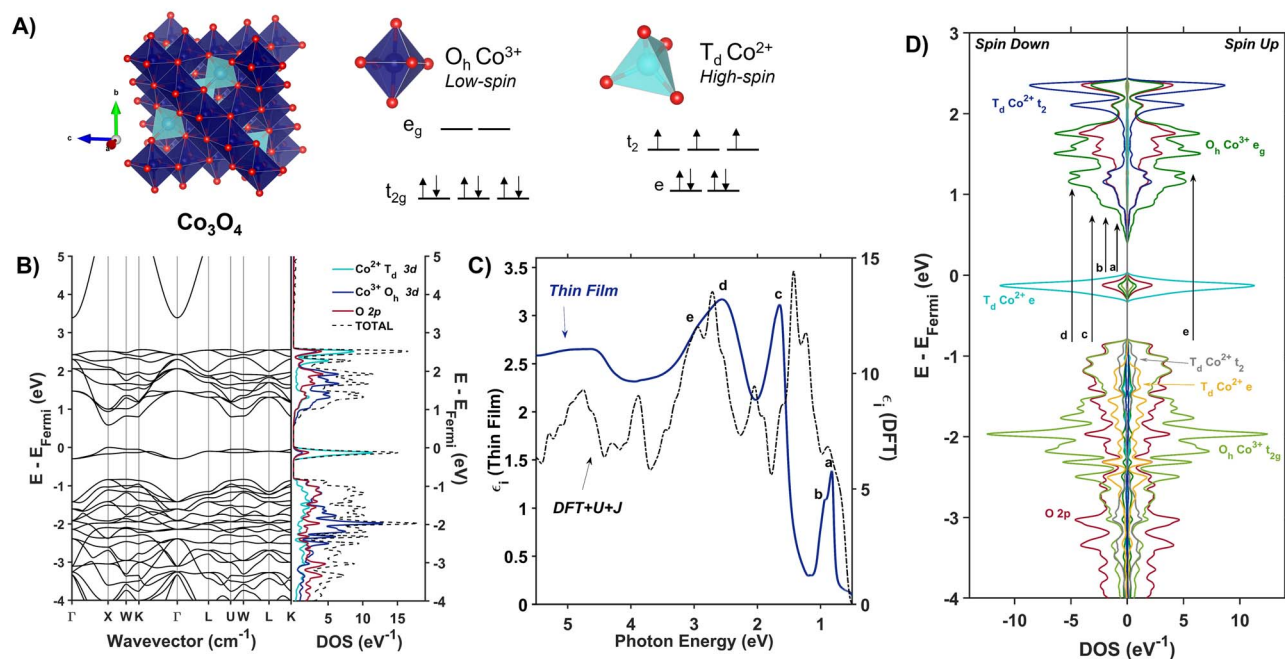


Fig. 2 (A) The Co<sub>3</sub>O<sub>4</sub> unit cell (*Fd3m*) with associated d-splitting diagrams for Co<sup>3+</sup> (O<sub>h</sub>) and Co<sup>2+</sup> (T<sub>d</sub>). (B) Electronic band structure and projected density of states of Co<sub>3</sub>O<sub>4</sub> calculated with Hubbard-corrected DFT. (C) Plot of experimentally (solid blue line) and computationally (dashed black line) determined imaginary dielectric spectra of Co<sub>3</sub>O<sub>4</sub>. Labels a–e correlate with arrows in (D), the spin-symmetrized projected density of states arising from the primitive cell of Co<sub>3</sub>O<sub>4</sub>.



optical spectra; we aim to understand how this coordination geometry impacts photoinduced polaron formation.

The formation of small polarons in Co spinel oxides ( $\text{Co}_3\text{O}_4$  and  $\text{ZnCo}_2\text{O}_4$ ) has been inferred from the observation of thermally activated charge transport in these materials;<sup>26,27</sup> however, as with other transition metal oxides, descriptions of photoinduced polarons in these materials are scarce. Transient absorption (TA) studies of  $\text{Co}_3\text{O}_4$  have reported that the strongest TA signal is induced by thermal effects,<sup>41,42</sup> which mirrors the behavior of hematite, a material known to undergo photoinduced polaron formation.<sup>29,30,43,44</sup> In other work using extreme ultraviolet (XUV) spectroscopy, Zhang, *et al.* describe indirect formation of small polarons from self-trapped photocarriers in  $\text{Co}_3\text{O}_4$ .<sup>45</sup> Using a combined approach of resonance Raman spectroscopy, temperature-dependent optical spectroscopy and Hubbard-corrected density functional theory, we investigate the influence of cobalt coordination geometry on photoinduced polaron formation in  $\text{Co}_3\text{O}_4$  and  $\text{ZnCo}_2\text{O}_4$ . We report evidence that  $T_d$  Co in  $\text{Co}_3\text{O}_4$  mediates resonance Raman enhancement of specific phonon modes, indicating the presence of phonon-coupled optical transitions that lead to polaron formation. Parallel studies of partially inverted  $\text{ZnCo}_2\text{O}_4$ , in which the majority of the  $T_d$  Co ions are replaced with Zn, confirm the involvement of  $T_d$  Co in phonon-coupled transitions; however, in  $\text{ZnCo}_2\text{O}_4$  these transitions do not depend on temperature. This observation suggests that photoinduced polarons in  $\text{ZnCo}_2\text{O}_4$  form at static defects rather than dynamic lattice distortions. The fundamental understanding of photophysical properties as a function of coordination geometry presented here is crucial to designing transition metal oxides for optical applications.

## Results and discussion

### Assignment of the optical spectra of $\text{Co}_3\text{O}_4$

$\text{Co}_3\text{O}_4$  adopts a spinel crystal structure (space group  $Fd\bar{3}m$ ) with two different metal sites:  $\text{Co}^{3+}$  ions occupy octahedral sites and  $\text{Co}^{2+}$  ions occupy tetrahedral sites (Fig. 2A). The two coordination sites give rise to two distinct crystal field splittings of the 3d orbitals as shown in Fig. 2A. Transitions between the  $e$  and  $t_2$  states within the  $T_d$  sites are allowed by the Laporte selection rule whereas this rule forbids transitions between the  $t_{2g}$  and  $e_g$  states within the  $O_h$  sites. The unpaired electrons of adjacent  $T_d$   $\text{Co}^{2+}$  atoms in  $\text{Co}_3\text{O}_4$  are antiferromagnetically coupled.<sup>46</sup> Density Functional Theory with Hubbard and Hund corrections (DFT +  $U$  +  $J$ ) was used to calculate the ground-state electronic structure of  $\text{Co}_3\text{O}_4$  within the Born–Oppenheimer approximation, in which nuclear motion is neglected. Hubbard (electron correlation correction)<sup>47</sup> and Hund (local magnetization correction)<sup>48</sup> corrections are used in highly correlated materials such as  $\text{Co}_3\text{O}_4$  to mitigate self-interaction errors.<sup>30</sup> Using a linear response method,<sup>30,47</sup>  $U$  and  $J$  parameters were calculated for  $\text{Co}_3\text{O}_4$  from first principles *via* perturbation of the local environments of open-shell ions ( $T_d$   $\text{Co}^{2+}$  and  $O_h$   $\text{Co}^{3+}$  in the case of  $\text{Co}_3\text{O}_4$ , see ESI† for more computational details). The DFT +  $U$  +  $J$ -computed band structure and projected density of states (pDOS) shown in Fig. 2B indicate that 3d orbitals

associated with  $\text{Co}^{2+}$   $T_d$  atoms are the primary contributors to an energetically isolated band at the valence band maximum (VBM), whereas both  $\text{Co}^{2+}$   $T_d$  and  $\text{Co}^{3+}$   $O_h$  3d orbitals contribute to the conduction band minimum (CBM). These computed results suggest isolated valence bands with  $\text{Co}^{2+}$   $T_d$  character participate in the low-energy transitions observed in the experimental imaginary dielectric spectrum at 0.82 and 0.92 eV (labeled a, b in Fig. 2C).

Fig. 2C plots the DFT +  $U$  +  $J$ -computed dielectric spectrum overlaid with the experimental dielectric spectrum extracted from transmission and reflection spectra measured from a 53.7-nm thick  $\text{Co}_3\text{O}_4$  film (see ESI† for details of dielectric spectrum determination and powder X-ray diffraction pattern). We applied a rigid shift of +0.3 eV to all conduction band eigenvalues to bring the computed dielectric function into alignment with the measured spectrum. Herein, this shift is applied to all electronic band diagrams and electronic density of states plots of  $\text{Co}_3\text{O}_4$ . In order to fulfill the  $f$ -sum rule governing total oscillator strength, all computed optical spectra are subsequently renormalized by a factor of  $(1 - (0.3 \text{ eV}/\hbar\omega))$ .<sup>49,50</sup> The computed single-particle dielectric spectrum shows good agreement with the measured spectrum, confirming the features at 0.82, 0.92 and 1.64 eV involve  $\text{Co}^{2+}$   $T_d$  bands (a–c). From the spin-resolved density of states separated into band contributions from  $t_{2g}$  and  $e_g$  ( $O_h$ ),  $e$  and  $t_2$  ( $T_d$ ) and O 2p (Fig. 2D), optical transitions are assigned while also considering spatial wavefunction overlap. The onsets of the conduction bands derived from  $O_h$   $\text{Co}^{3+}$   $e_g$ ,  $T_d$   $\text{Co}^{2+}$   $t_2$ , and O 2p orbitals occur at the same energy; however, the largest projected density of states comes from  $O_h$   $\text{Co}^{3+}$   $e_g$ . Thus, when considering transitions from  $T_d$   $\text{Co}^{2+}$   $e$  to the conduction band, these DFT +  $U$  +  $J$  computations exhibit no energetic difference among transitions to  $O_h$   $\text{Co}^{3+}$   $e_g$ ,  $T_d$   $\text{Co}^{2+}$   $t_2$ , and O 2p. The oscillator strength of the experimental dielectric peak at 0.82 eV is higher than the 0.92 shoulder, indicating the higher likelihood of the 0.82-eV transition; however, the shoulder is not resolved in the imaginary dielectric computed by DFT +  $U$  +  $J$ . Therefore, to assign this shoulder, we qualitatively assessed the spatial overlap of orbital wavefunctions by mapping the contributions of specific atoms to specific projected densities of states (Fig. S10†). This approach reveals that the most probable transition contributing to the 0.82 eV feature is an intra-atomic transition of  $\text{Co}^{2+}$   $T_d$  that satisfies the spin transition selection rule (Fig. S10A and B†). The spatial overlap of the orbitals involved in an intra-atomic transition is greater than the overlap of orbitals between a  $T_d$  Co and an  $O_h$  Co (inter-sublattice charge transfer), further supporting the assignment of the 0.82-eV optical transition to an intra-atomic d-to-d transition in  $T_d$   $\text{Co}^{2+}$  ( $e \rightarrow t_2$ ). From the spin-resolved density of states combined with our spatial overlap analysis, the feature at 0.92 eV is best described as a charge transfer transition from the tetrahedral sublattice to the octahedral sublattice ( $T_d$   $\text{Co}^{2+}$   $e \rightarrow O_h$   $\text{Co}^{3+}$   $e_g$ ). Similarly, the transition at 1.62 eV can be described as an inter-sublattice charge transfer from  $O_h$   $\text{Co}^{3+}$   $t_{2g} \rightarrow T_d$   $\text{Co}^{2+}$   $t_2$ . The covalency between  $T_d$   $\text{Co}^{2+}$  and O 2p, as well as  $O_h$   $\text{Co}^{3+}$  and O 2p, allows these metal-to-metal (inter-sublattice charge transfer) transitions to occur. The feature at 2.55 eV



Table 1 Assignment of optical transitions in  $\text{Co}_3\text{O}_4$ 

Peak center (eV)	Label	Transition	Description
0.82	a	$\text{T}_d \text{Co}^{2+} \text{e} \rightarrow \text{t}_2$	Intra-atomic transition
0.92	b	$\text{T}_d \text{Co}^{2+} \text{e} \rightarrow \text{O}_h \text{Co}^{3+} \text{e}_g$	Inter-sublattice charge transition
1.64	c	$\text{O}_h \text{Co}^{3+} \text{t}_{2g} \rightarrow \text{T}_d \text{Co}^{2+} \text{t}_2$	Inter-sublattice charge transition
2.55	d	$\text{O } 2\text{p} \rightarrow \text{T}_d \text{Co}^{2+} \text{t}_2$	Ligand-to-metal charge transition
2.90	e	$\text{O } 2\text{p} \rightarrow \text{O}_h \text{Co}^{3+} \text{e}_g$	Ligand-to-metal charge transition

and its shoulder at 2.90 eV arise from ligand-to-metal charge transfer (LMCT) type transitions. The assignment of these two LMCT transfers is resolved from comparing the experimental spectrum of  $\text{Co}_3\text{O}_4$  with  $\text{ZnCo}_2\text{O}_4$ , which has majority  $\text{O}_h \text{Co}^{3+}$  (Fig. S12†). It becomes apparent that the  $\text{O } 2\text{p} \rightarrow \text{O}_h \text{Co}^{3+} \text{e}_g$  transition is slightly higher in energy than the  $\text{O } 2\text{p} \rightarrow \text{T}_d \text{Co}^{2+} \text{t}_2$  transition, leading to the following assignments:  $\text{O } 2\text{p} \rightarrow \text{T}_d \text{Co}^{2+} \text{t}_2$  (2.55 eV) and  $\text{O } 2\text{p} \rightarrow \text{O}_h \text{Co}^{3+} \text{e}_g$  (2.90 eV). All transitions are spin-conserved (Fig. 2D and Table 1).

The electronic density of states and band structure of  $\text{Co}_3\text{O}_4$  have been previously calculated using many approaches including, but not limited to GGA +  $U$ , hybrid functional PBE0, range-separated exchange–correlation functional HSE06, and many-body Green's function GW approximation.<sup>27,51,52</sup> Commonly, the density of states shows some degree of Co 3d and O 2p wavefunction overlap at the band edges; however, depending on the computational approach, the band gap varies from 0.78 to 1.6 eV.<sup>27,51,52</sup> Singh, *et al.* explored many different DFT approaches to calculating the electronic structure of  $\text{Co}_3\text{O}_4$ , including PBE, PBE +  $U_{\text{eff}}$ , HSE06, and many-body Green's function using the GW approximation (Sc-GW0).<sup>52</sup> Of these, the Sc-GW0 method produces the most accurate representation of the electronic structure of  $\text{Co}_3\text{O}_4$  based on computed electronic bands, density of states, and absorption spectra.<sup>52</sup> Although the absorption spectrum computed with Sc-GW0 exhibits a high oscillator strength between ~0.5 and 2.0 eV,<sup>52</sup> unlike our DFT +  $U$  +  $J$  approach, it cannot resolve the two distinct transitions at 0.8 and 1.6 eV reported in experimental optical spectra.<sup>27,36,37,41,45</sup>

The experimental presence of the 0.8-eV optical transition and discrepancies in the band gap energies computed with various approaches has led to debate over defining the optical band gap of  $\text{Co}_3\text{O}_4$ : some report it as 0.8 eV,<sup>27,39,53</sup> while others define it to be 1.6 eV.<sup>41,54</sup> The incongruity in reported band gap energies is scrutinized by Smart, *et al.* in their work modelling  $\text{Co}_3\text{O}_4$  optical transitions with DFT +  $U$  and a hybrid functional that includes a fraction of Hartree–Fock exchange.<sup>55</sup> They propose the 0.8-eV optical transition arises from photoinduced formation of a small hole polaron, and that this transition becomes most apparent upon application of uniaxial lattice strain.<sup>55</sup> Our calculation of the ground-state electronic structure of  $\text{Co}_3\text{O}_4$  using a Hubbard- and Hund- corrected plane-wave pseudopotential approach demonstrates that the low-energy transition at 0.8 eV (and its 0.9-eV shoulder) originate from localized  $\text{Co}^{2+} \text{T}_d$  valence bands (e). Additionally, the corresponding empty  $\text{Co}^{2+} \text{T}_d$  conduction bands ( $\text{t}_2$ ) participate in the observed transition at 1.64 eV, indicating the importance of  $\text{T}_d \text{Co}^{2+}$  electronic character to this transition as

well. Although the computational approach used here is a single-particle method that neglects many-body perturbations and nuclear motion, its ability to capture all the features observed in the experimental dielectric spectrum supports its accuracy in describing the nature of the bands that contribute to optical transitions in  $\text{Co}_3\text{O}_4$ . Because the dielectric spectrum reported here is calculated within the Born–Oppenheimer approximation, transitions originating from ground states containing nuclear displacements are not captured, contrasting the description of the 0.8 eV transition as involving a lattice strain-induced small hole polaron by Smart, *et al.*<sup>55</sup>

### Optical phonon enhancement in $\text{Co}_3\text{O}_4$

We assess phonon coupling to various optical transitions in  $\text{Co}_3\text{O}_4$  using Stokes resonance Raman spectroscopy. Fig. 3A plots a series of resonance Raman spectra collected for a 428-nm thick  $\text{Co}_3\text{O}_4$  film deposited on a sapphire substrate using a variety of excitation lasers with photon energies ranging from 1.49 to 3.06 eV, which spans the  $\text{Co}_3\text{O}_4$  absorption spectrum (Fig. 3B). These Raman spectra of  $\text{Co}_3\text{O}_4$  each contain five phonon modes, consistent with previous reports.<sup>56,57</sup> The phonon mode at 86 meV (oxygen breathing about  $\text{T}_d \text{Co}^{2+}$ , Fig. 4D) is the most intense at all excitation energies, except  $h\nu_{\text{exc}} = 1.88$  eV, which corresponds to the inter-sublattice charge transfer transition ( $\text{O}_h \text{Co}^{3+} \text{t}_{2g} \rightarrow \text{T}_d \text{Co}^{2+} \text{t}_2$ ). When  $\text{Co}_3\text{O}_4$  is excited with a photon energy of 1.58 eV, which correspond to the lower-energy edge of this inter-sublattice charge transfer peak, the 86 meV phonon mode becomes most intense again. Fig. 3C plots the excitation spectrum for each phonon mode corrected for scattering cross section and sample absorption, which enables the comparison of phonon mode intensities across different excitation energies. This quantitative analysis of relative intensities reveals that there is amplified resonance enhancement of all modes upon excitation at 1.49 eV, indicating strong phonon coupling to this optical transition. This analysis was repeated on a thin film of  $\text{Co}_3\text{O}_4$  deposited on quartz, and the same trends are apparent (Fig. S7†). Given that the 86-meV phonon mode is most intense at an excitation energy of 1.49 eV, we propose it is this phonon mode that most strongly couples to the optical transition at 1.49 eV.

DFT +  $U$  +  $J$  was used to calculate the displacement vectors of all 39 optical phonon modes in  $\text{Co}_3\text{O}_4$  at  $k$ -point  $\Gamma$  (Table S2†). White and DeAngelis determined through evaluating the Raman selection rules that there are five Raman active modes in normal spinel oxides.<sup>58</sup> The symmetry of these modes was compared with the displacement vectors computed with DFT +





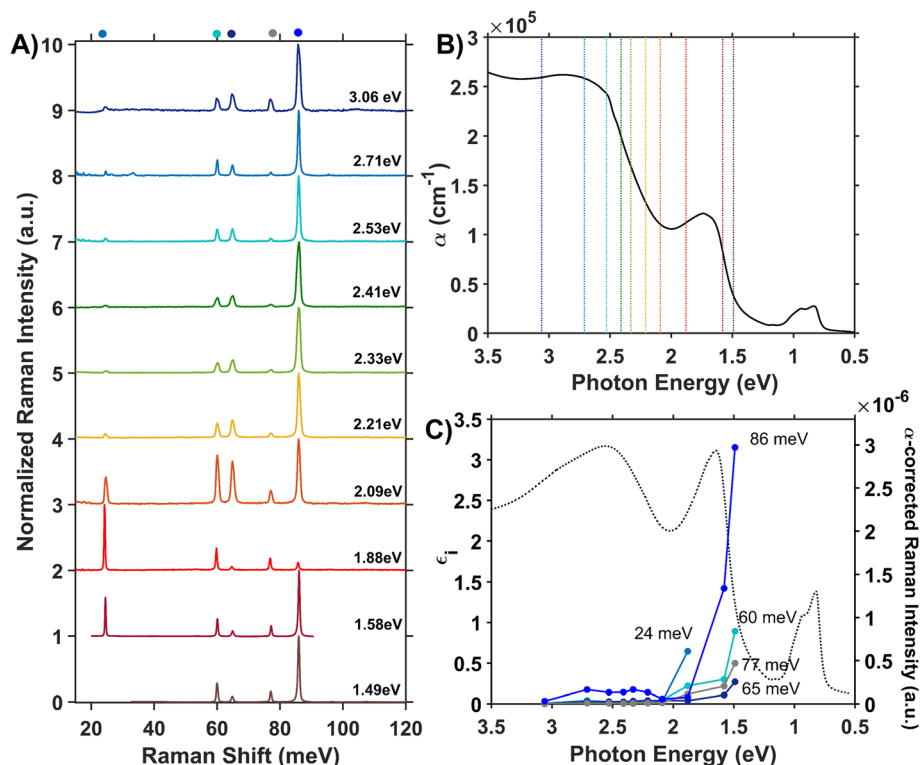


Fig. 3 (A) Internally normalized Stokes Raman spectra of  $\text{Co}_3\text{O}_4$  collected with various excitation energies that span the absorption spectrum plotted in (B). Raman spectra are vertically offset for clarity. Because the spectra are internally normalized, only relative changes in phonon mode intensity can be assessed as a function of excitation energy. Note that the spectrum excited with 1.49 eV begins at a Raman shift of 30 meV due to instrumental constraints, primarily the bandwidth of the filter used to remove Rayleigh scattering. (C) Plot of the intensities of the various Raman modes shown in part A corrected for scattering cross section and sample absorption superimposed on the imaginary dielectric spectrum. Note the corresponding color point of each phonon mode indicated above the relevant Raman peak in A. For the full correction, see ESI Fig. S5.†

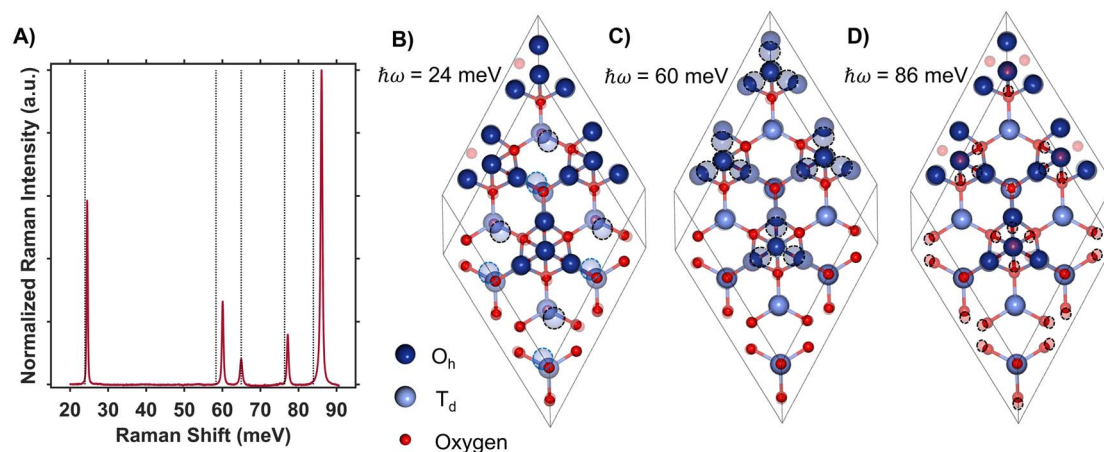


Fig. 4 (A) Raman spectrum of  $\text{Co}_3\text{O}_4$  collected with a excitation photon energy of 1.58 eV overlaid with the energies of DFT +  $U$  +  $J$ -computed Raman-active phonon modes. The atomic displacements corresponding to the modes at 24 meV (B), 60 meV (C) and 86 meV (D) are shown. The circled atoms demonstrate the vector displacements of the phonon modes in a  $2 \times 2 \times 1$  super cell, where  $T_d$  sites are most shifted at 24 meV,  $O_h$  sites at 60 meV and oxygens surrounding  $T_d$  sites at 86 meV. Note that the extra oxygen atoms in the 24 meV and 86 meV vector-displacement images appear from the displacement of the neighboring super-cell.

$U + J$  to identify the Raman active modes, the energies of which are overlaid with the Raman spectrum in Fig. 4A. From the resonance Raman profile (Fig. 3A and C), the phonon modes

with energies of 24, 60, and 86 meV exhibit the most significant resonance enhancement upon excitation at 1.49 eV, corresponding to the onset of the absorption feature centered at

1.64 eV. These modes correspond to motion of tetrahedral  $\text{Co}^{2+}$  (24 meV), octahedral  $\text{Co}^{3+}$  (60 meV), and oxygen stretching symmetrically about  $\text{Co}^{2+}$   $\text{T}_d$  atoms (86 meV, Fig. 4B–D).

The observed enhancement of the 24-meV ( $\text{T}_d$   $\text{Co}^{2+}$  motion) and 60-meV ( $\text{O}_h$   $\text{Co}^{3+}$  motion) phonons at a Raman excitation energy of 1.49 eV is related to the contribution of both  $\text{T}_d$   $\text{Co}^{2+}$  and  $\text{O}_h$   $\text{Co}^{3+}$  electronic character to this optical transition, which corresponds to population of empty  $\text{T}_d$   $\text{Co}^{2+}$   $t_2$  conduction band states from  $\text{O}_h$   $\text{Co}^{3+}$   $t_{2g}$  valence band states. Although both  $\text{T}_d$   $\text{Co}^{2+}$  and  $\text{O}_h$   $\text{Co}^{3+}$  phonon modes exhibit resonance enhancement because of associated electronic character in the optical transition centered at 1.64 eV, it is the oxygen breathing mode about  $\text{Co}^{2+}$   $\text{T}_d$  ions at 86 meV that exhibits the most enhancement. The degree of energetic overlap between the O 2p and  $\text{Co}^{2+}$   $\text{T}_d$  projected density of states in the conduction band at the energy of the  $\text{O}_h$   $\text{Co}^{3+}$   $t_{2g} \rightarrow \text{T}_d$   $\text{Co}^{2+}$   $t_2$  transition is significant (Fig. 2B and D) and supports the observed enhancement of the 86-meV phonon, which corresponds to motion of oxygen atoms along their bond axes to  $\text{T}_d$   $\text{Co}^{2+}$ .

### Temperature dependence of optical transitions

To further understand the role of phonon coupling in the optical transitions of  $\text{Co}_3\text{O}_4$ , we measured the dependence of its dielectric spectrum on temperature between 82 and 470 K. Fig. 5A and B plot the resulting thermal difference spectra (TDS) calculated according to eqn (1).

$$\Delta\epsilon_i(T) = \epsilon_{i,T} - \epsilon_{i,294\text{K}} \quad (1)$$

We assessed the impact of temperature on the dielectric spectrum by integrating the intensity of the thermal difference spectra. Fig. 5C (blue) plots the absolute value of the thermal difference spectra integrated from 0.68 to 1.86 eV and normalized to the integrated intensity obtained from the spectrum collected at 82 K. To account for sign changes, the integrands of spectra collected below room temperature are shown as negative, and those collected above room temperature are positive. For phonon-coupled optical transitions, we expect the change in intensity with changing temperature to be proportional to the change in population of the coupled phonons. The thermal population of phonon modes is dictated by the Bose–Einstein distribution shown in eqn (2), where  $\hbar\Omega$  is the phonon energy. Normalization of the differential spectra to an arbitrary temperature, in this case 82 K, leads to eqn (3), where we compare the change in intensity of the dielectric spectrum ( $\Delta\epsilon$  defined in eqn (1)) to the change in population of the Raman active phonon modes at various temperatures.

$$N(\hbar\Omega, T) = \left( \frac{\hbar\Omega}{e^{kT} - 1} \right)^{-1} \times \rho(\hbar\Omega) \quad (2)$$

$$\frac{\Delta\epsilon(T)}{|\Delta\epsilon(82\text{ K})|} = \frac{N(\hbar\Omega, T) - N(\hbar\Omega, 294\text{ K})}{|N(\hbar\Omega, 82\text{ K}) - N(\hbar\Omega, 294\text{ K})|} \quad (3)$$

The temperature dependence of the TDS intensity overlays well with the change in population of the 24-meV Raman-active phonon with temperature predicted by the Bose–Einstein distribution (eqn (3)). This agreement suggests that the low energy optical transitions (1.64 eV, 0.96 eV and 0.82 eV) are coupled strongly to the thermal population of this phonon. Interestingly, the 24-meV phonon mode is not the most enhanced in resonance Raman spectra collected with an excitation photon energy of 1.49 eV; however, both the 24-meV and 86-meV phonon modes are described by displacement vectors primarily of or around  $\text{T}_d$   $\text{Co}^{2+}$  ions (Fig. 4B and D). Evidence of thermally activated optical transitions combined with strong phonon coupling to optical transitions observed in resonance Raman spectra (*vide supra*) suggests an optically accessed polaronic state related to  $\text{T}_d$   $\text{Co}^{2+}$ . Compared to resonance Raman measurements, thermal difference spectra are not as precise in determining the energies of the specific phonon modes coupling to the optical transition, as there may be multiple phonon modes contributing to thermal activation. However, the Bose–Einstein distribution corresponding to a phonon energy of 24 meV overlays the temperature-dependent TDS intensities much better than the distributions corresponding to other Raman-active phonon modes (Fig. 5C). We therefore interpret the threshold phonon energy of 24 meV to be where thermal activation occurs. Similar temperature-dependent behavior is observed in  $\alpha\text{-Fe}_2\text{O}_3$ : at the excitation energy where maximal Raman enhancement is observed ( $\sim 2.2$  eV), the strongest temperature-dependence in the optical spectrum is also present.<sup>29</sup> Thermal activation of optical transitions in  $\alpha\text{-Fe}_2\text{O}_3$  is the proposed mechanism by which direct excitation into intrinsic polaronic states occurs.<sup>29,30</sup> The similarities in the trends of Raman enhancement and the temperature-dependence of the dielectric spectrum observed for  $\text{Co}_3\text{O}_4$  and  $\alpha\text{-Fe}_2\text{O}_3$  support the presence of an optically accessible polaronic state in  $\text{Co}_3\text{O}_4$  arising from coupling to intrinsic, dynamic lattice distortions (phonons) (Fig. 1, red arrow).

### $\text{Co}_3\text{O}_4$ compared to $\text{ZnCo}_2\text{O}_4$ : assessing the role of $\text{T}_d$ Co in spinel oxides

The resonance Raman and thermal difference spectra of  $\text{Co}_3\text{O}_4$  suggest that  $\text{Co}^{2+}$  ions occupying  $\text{T}_d$  sites are involved in the phonon-coupled optical transitions. To further understand the role of  $\text{T}_d$   $\text{Co}^{2+}$  in the optical spectra of  $\text{Co}_3\text{O}_4$ , we assessed the optical properties of  $\text{ZnCo}_2\text{O}_4$ , where  $\text{Zn}^{2+}$  replaces  $\text{T}_d$   $\text{Co}^{2+}$  (Fig. 6A). We computed the imaginary dielectric spectrum for normal  $\text{ZnCo}_2\text{O}_4$  with DFT +  $U$  (see ESI for details<sup>†</sup>). Within normal  $\text{ZnCo}_2\text{O}_4$ , the  $\text{T}_d$   $\text{Zn}^{2+}$  3d orbitals are completely filled, and the  $\text{O}_h$   $\text{Co}^{3+}$  3d orbitals have a low-spin configuration, termed “quasi close-shelled”.<sup>59,60</sup> With no unpaired electrons present, the Hund parameter,  $J$ , is not implemented in these calculations. The DFT +  $U$  calculations of normal  $\text{ZnCo}_2\text{O}_4$  reveal two absorption bands at and above  $\sim 2.5$  eV, while the experimental spectrum only has one transition in this region. Literature reports assign the experimentally observed feature at 3.0 eV to an LMCT-type transition,<sup>27,38</sup> which matches the lower energy



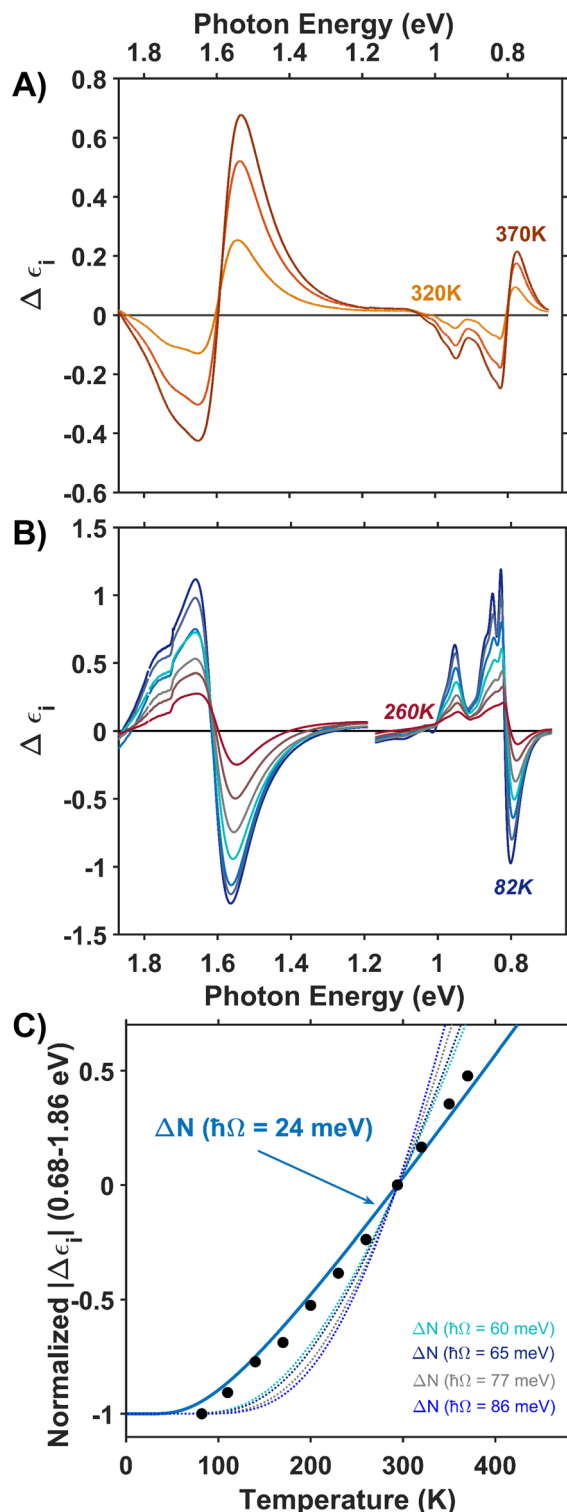


Fig. 5 Thermal difference imaginary dielectric spectra of  $\text{Co}_3\text{O}_4$  collected at temperatures above (A) and below (B) room temperature (294 K). (C) Absolute value of the thermal difference spectra integrated from 0.68 to 1.86 eV and normalized to the integrated intensity of the spectrum collected at 82 K plotted versus temperature (black circles). The solid blue line plots the temperature-dependent change in Bose–Einstein population of a phonon with an energy of 24 meV. Bose–Einstein distributions corresponding to the energies of the other Raman active phonon modes are plotted as dotted lines.

peak (2.5 eV) observed in the DFT +  $U$  dielectric spectrum. The 3.5 eV peak in the DFT +  $U$  dielectric spectrum computed here is associated with an intra-sublattice charge transfer ( $\text{O}_h \text{Co}^{3+} t_{2g} \rightarrow \text{O}_h \text{Co}^{3+} e_g$ ). Because this is a charge transfer between two different lattice sites, the Laporte selection rule does not apply. Due to the hybridized nature of the Co 3d and O 2p orbitals in both the valence and conduction bands, we assign the experimental dielectric peak at 3.0 eV to a combination of LMCT and intra-sublattice charge transfer transitions (see Fig. S11 in the ESI†). Interestingly, when comparing the experimental peak shape of the dielectric transition at 3.0 eV in  $\text{ZnCo}_2\text{O}_4$  (Fig. 6B) to the analogous transition in  $\text{Co}_3\text{O}_4$  (Fig. 2C and S12†), it becomes apparent that the shoulder in  $\text{Co}_3\text{O}_4$  (labeled e in Fig. 2C) is at the same energy as the peak center of the experimental  $\text{ZnCo}_2\text{O}_4$  transition. This energetic alignment in the experimental dielectric spectra, used to assign the 2.90-eV shoulder in  $\text{Co}_3\text{O}_4$  as excitation into  $\text{O}_h \text{Co}^{3+} e_g$  bands, has been observed previously in a Zn dopant study of  $\text{Co}_3\text{O}_4$ .<sup>38</sup>

The lower-energy region of the experimental dielectric spectrum of  $\text{ZnCo}_2\text{O}_4$  contains broad and weak features at 1.79, 0.98 and 0.82 eV, which coincide with where  $\text{Co}_3\text{O}_4$  has transitions associated with  $\text{Co}^{2+} T_d$ . X-ray fluorescence analysis indicates the stoichiometry of the  $\text{ZnCo}_2\text{O}_4$  film used to produce the dielectric spectrum is  $1.94 \pm 0.04 \text{ Co} : \text{Zn}$ , and the powder X-ray diffraction pattern indicates the film is phase-pure spinel (Fig. S3†). We therefore suspected that our  $\text{ZnCo}_2\text{O}_4$  film may be slightly inverted and contain a small population of cobalt in tetrahedral sites, consistent with previous reports.<sup>61</sup> In an effort to control cation distribution in  $\text{ZnCo}_2\text{O}_4$ , we changed the temperature at which  $\text{ZnCo}_2\text{O}_4$  films were annealed following spin-coating. We observe that annealing above 600 °C induces phase separation as features associated with ZnO become apparent in the X-ray diffraction pattern and Raman spectra (Fig. S13 and S14†). Additionally, as annealing temperature increases, the low energy peaks observed in the dielectric spectrum increase in intensity (Fig. S15†). We therefore suspect that the films containing ZnO also contain  $\text{Co}_3\text{O}_4$ , which accounts for the presence of the intense low-energy peaks in the dielectric spectrum (Fig. S15†). With no ZnO peaks present in X-ray diffraction or ZnO phonon modes observed in Raman for the films annealed at 600 °C, we hypothesize that the weak, low-intensity features observed in the dielectric spectra of these films arise from cation inversion, whereby a fraction of the Co ions occupy  $T_d$  sites and a fraction of the Zn ions occupy  $\text{O}_h$  sites. We describe the Co ions in  $T_d$  sites as substitutional lattice defects.

To investigate the impact of cation inversion on the optical spectra of  $\text{ZnCo}_2\text{O}_4$ , we computed the electronic structure of inverted  $\text{ZnCo}_2\text{O}_4$  using DFT +  $U$  +  $J$  (see ESI for details†). Inverted  $\text{ZnCo}_2\text{O}_4$  (i- $\text{ZnCo}_2\text{O}_4$ ) was modeled by switching two  $\text{O}_h \text{Co}^{3+}$  ions with two  $T_d \text{Zn}^{2+}$  ions within a  $2 \times 2 \times 2$  supercell of normal  $\text{ZnCo}_2\text{O}_4$  to produce an inversion factor of  $x = 0.125$  while maintaining charge balance and stoichiometry. The resulting two  $T_d \text{Co}^{3+}$  ions were modeled to be antiferromagnetically coupled to maintain net-zero magnetization. Compared to normal  $\text{ZnCo}_2\text{O}_4$  (n- $\text{ZnCo}_2\text{O}_4$ ), i- $\text{ZnCo}_2\text{O}_4$  contains an isolated state at the valence band-edge with primarily  $T_d \text{Co}$

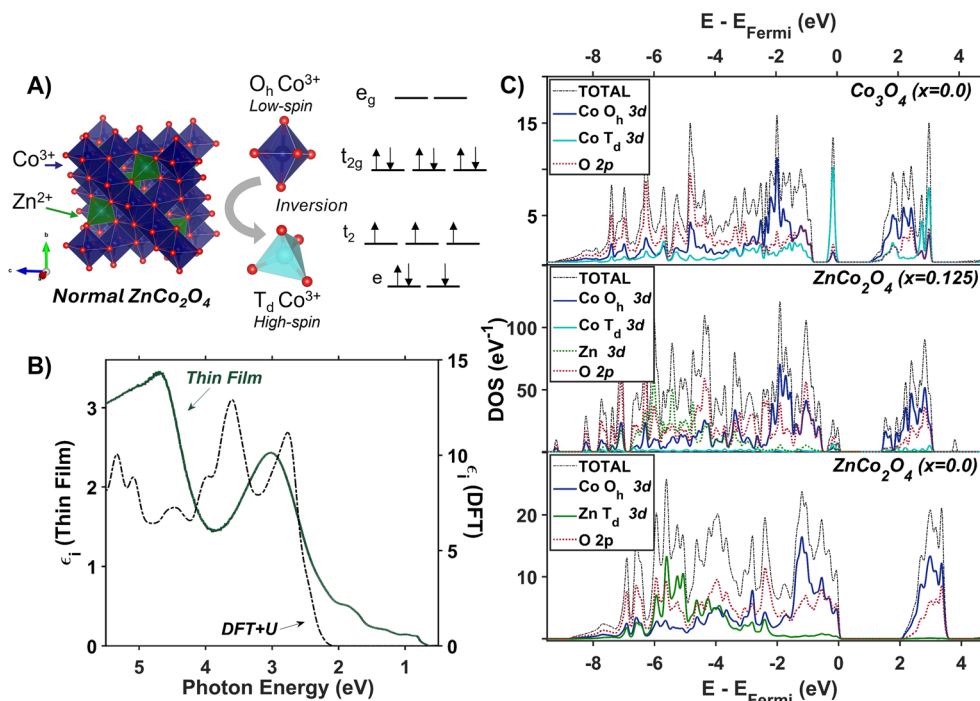


Fig. 6 (A) Unit cell of normal ZnCo<sub>2</sub>O<sub>4</sub> with associated crystal field splitting diagrams of Co<sup>3+</sup> in octahedral and tetrahedral coordination. (B) Plot of experimentally (solid green line) and computationally (dashed black line) determined imaginary dielectric spectra of ZnCo<sub>2</sub>O<sub>4</sub>. The computed dielectric spectrum was calculated for normal ZnCo<sub>2</sub>O<sub>4</sub>. (C) The electronic density of states calculated with Hubbard-corrected DFT for normal Co<sub>3</sub>O<sub>4</sub> (top), ZnCo<sub>2</sub>O<sub>4</sub> with an inversion factor of 0.125 (middle), and normal ZnCo<sub>2</sub>O<sub>4</sub> (bottom).

and O 2p character (Fig. 6C, middle and bottom). The conduction band edge in i-ZnCo<sub>2</sub>O<sub>4</sub> appears at a lower energy than that in n-ZnCo<sub>2</sub>O<sub>4</sub> (similar to Co<sub>3</sub>O<sub>4</sub>), and i-ZnCo<sub>2</sub>O<sub>4</sub> contains an isolated region at the conduction band-edge of primarily O<sub>h</sub> Co and O 2p character (Fig. 6C, middle and top). The band-edge character of i-ZnCo<sub>2</sub>O<sub>4</sub>, despite the 3+ oxidation state of the cobalt ion in T<sub>d</sub> sites, has more similarities to that of Co<sub>3</sub>O<sub>4</sub> than n-ZnCo<sub>2</sub>O<sub>4</sub> (Fig. 6C). Thus, the presence of Co (T<sub>d</sub>) in spinel oxides results in low-energy d-to-d transitions, regardless of its oxidation state (+2 or +3). Because there is no evidence of phase separation (*i.e.* presence of Co<sub>3</sub>O<sub>4</sub>) to describe the low-energy features in the dielectric, we conclude the synthesized ZnCo<sub>2</sub>O<sub>4</sub> contains at least a small degree of inversion.

The Raman spectrum of ZnCo<sub>2</sub>O<sub>4</sub> has five distinct modes (Fig. 7A and B) consistent with previous reports.<sup>62,63</sup> The phonon mode frequencies are similar to those observed in Co<sub>3</sub>O<sub>4</sub>, which is expected, as both materials adopt the spinel crystal structure. Additionally, the 23-meV, 61-meV, and 89-meV phonon modes in ZnCo<sub>2</sub>O<sub>4</sub> are described by the same phonon motions as in Co<sub>3</sub>O<sub>4</sub>: motion of T<sub>d</sub> metal center motion of O<sub>h</sub> metal center, and oxygen stretching about T<sub>d</sub> sites respectively (Fig. 4B–D). Fig. 7B (top) plots the experimental Raman spectrum of ZnCo<sub>2</sub>O<sub>4</sub> overlaid with the computed phonon modes of normal ZnCo<sub>2</sub>O<sub>4</sub>. While the five prominent phonon modes are accounted for by DFT + *U* phonon calculations, there are two broad and weak features at 26 meV and 85 meV that do not appear in calculations. Both phonons have similar energies to corresponding modes observed in Co<sub>3</sub>O<sub>4</sub>, which are both related predominantly to T<sub>d</sub> sites (Fig. 7B, bottom). The

discrepancy between the energies of experimentally observed Raman-active optical phonon modes and the computed modes of normal ZnCo<sub>2</sub>O<sub>4</sub> can be explained by the sample crystallizing with a degree of inversion. With a small percentage of tetrahedral sites occupied by Co<sup>3+</sup>, and the rest by Zn<sup>2+</sup>, the phonon modes dominated by tetrahedral motion (~24 meV and ~89 meV) split into two distinct distributions. This phenomenon has been observed in other inverted spinel oxides<sup>64</sup> and is further evidence of the ZnCo<sub>2</sub>O<sub>4</sub> films crystallizing with a small percent occupation of cobalt in tetrahedral sites (Fig. 7B).

Unlike Co<sub>3</sub>O<sub>4</sub>, the resonance Raman profile of ZnCo<sub>2</sub>O<sub>4</sub> does not exhibit increased resonance enhancement of the phonons associated with T<sub>d</sub> motion (23 and 89 meV) with decreasing excitation energy. In contrast, the 61-meV phonon, related to Co O<sub>h</sub> motion (described by Fig. 4B), exhibits enhanced intensity in ZnCo<sub>2</sub>O<sub>4</sub> relative to all the other modes as excitation photon energy decreases (Fig. 7A). Interestingly, when the Raman profile is corrected for scattering cross section and sample absorption, the 61-meV mode is most enhanced at an excitation photon energy of 1.49 eV – the same excitation photon energy at which the most significant resonance enhancement is observed in Co<sub>3</sub>O<sub>4</sub> (Fig. 3C and Fig. 7C). This analysis was repeated on a thin film of ZnCo<sub>2</sub>O<sub>4</sub> deposited on quartz, and the same trends are apparent (Fig. S8†). The similarity in excitation photon energy where resonance enhancement is observed in ZnCo<sub>2</sub>O<sub>4</sub> and Co<sub>3</sub>O<sub>4</sub> indicates that the onset of the optical transition centered at ~1.6 eV is significant for both materials.

Based on the calculated density of states of inverted ZnCo<sub>2</sub>O<sub>4</sub> (Fig. 6C), the experimental optical transition observed at 1.6 eV



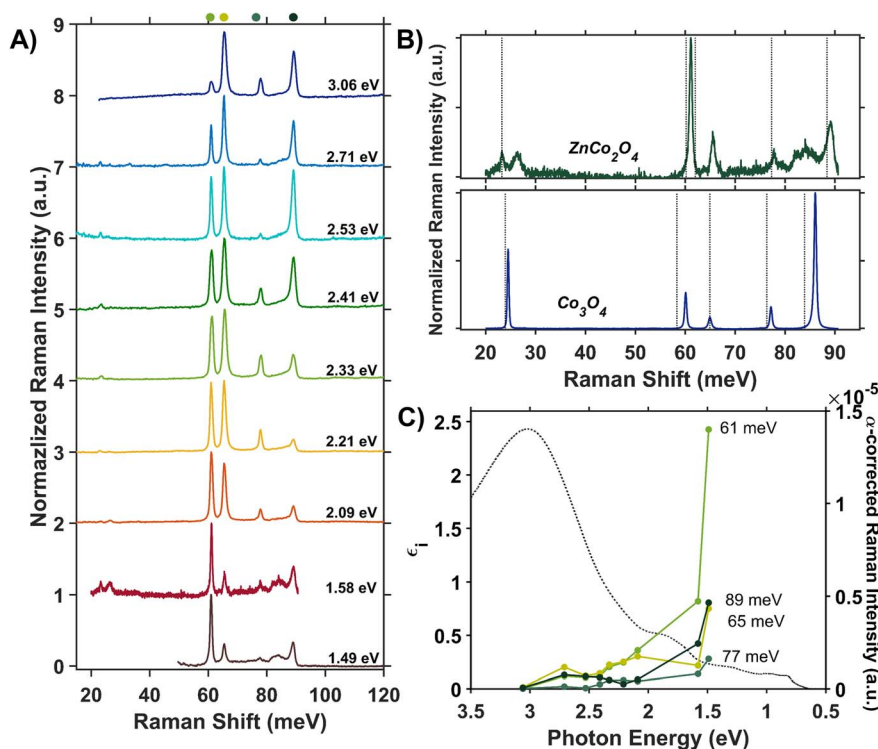


Fig. 7 (A) Internally normalized Raman spectra of a  $\text{ZnCo}_2\text{O}_4$  thin film deposited on a sapphire substrate collected with excitation photon energies that span its absorption spectrum. Because the spectra are internally normalized, only relative changes in phonon mode intensity can be assessed as a function of excitation energy (see ESI† for full work-up of Raman data). (B) Raman spectra of  $\text{ZnCo}_2\text{O}_4$  (top) and  $\text{Co}_3\text{O}_4$  (bottom) excited at a photon energy of 1.58 eV are overlaid with the Raman-active phonon modes computed for normal  $\text{ZnCo}_2\text{O}_4$  and  $\text{Co}_3\text{O}_4$ , respectively. (C) Plot of the intensities of the various Raman modes shown in part A corrected for scattering cross section and sample absorption superimposed on the imaginary dielectric spectrum of  $\text{ZnCo}_2\text{O}_4$ . Note the corresponding color point of each phonon mode indicated above the relevant Raman peak in A.

is assigned to metal-to-metal charge transfer (MMCT) from bands with  $T_d \text{ Co}^{3+}$  character (arising from the  $T_d \text{ Co}$  substitutional defects) to bands with  $O_h \text{ Co}^{3+}$  character. The most enhanced Raman spectrum ( $h\nu_{\text{exc}} = 1.49 \text{ eV}$ ) occurs upon excitation at the onset of the 1.6-eV transition, implying that the presence of the  $\text{Co}^{3+} T_d$  substitutional defect is crucial for the observed enhancement of the  $O_h$  phonon in  $\text{ZnCo}_2\text{O}_4$ . The conduction band character associated with the 1.6-eV transition arises from 3d orbitals associated with  $\text{Co}^{3+} O_h$ , the same site associated with the main vibrational motion of the 61-meV phonon (Fig. 4C). Although the strong resonance enhancement observed for the 61-meV phonon in  $\text{ZnCo}_2\text{O}_4$  is evidence of strong coupling between this mode and the optical transition at 1.6 eV, we do not observe significant temperature dependence of this transition in thermal difference spectra when compared to  $\text{Co}_3\text{O}_4$  (see Fig. S16 in ESI†). In fact, unlike  $\text{Co}_3\text{O}_4$ , the thermal difference spectra collected for  $\text{ZnCo}_2\text{O}_4$  at elevated temperatures do not exhibit any well-defined features corresponding to spectral features observed in the dielectric spectrum at room temperature.

The spectral and computational results reported here for  $\text{ZnCo}_2\text{O}_4$  and  $\text{Co}_3\text{O}_4$  indicate that Co 3d character at the band edge mediates phonon-coupled optical transitions (Fig. 3C, 4C, D and 7C). However, the contrast in temperature-dependence of the optical transition where both materials exhibit the strongest

phonon coupling (1.6 eV) suggests fundamentally different processes dictating the observed enhancement. We interpret our findings in  $\text{Co}_3\text{O}_4$  as direct population of an intrinsic polaronic state *via* photoexcitation, which is strongly influenced by the contributions of  $\text{Co}^{2+} T_d$  ions to band-edge states. In  $\text{ZnCo}_2\text{O}_4$ , we interpret the experimental  $\text{ZnCo}_2\text{O}_4$  sample to have a small percent occupation of tetrahedral sites by Co based on the presence of low-energy optical transitions associated with  $T_d \text{ Co}$ . Although the occupation of  $\text{Co}^{3+}$  in tetrahedral sites enables the optical transition and phonon enhancement at 1.6 eV, the lack of significant temperature dependence of this optical transition indicates that the phonon-coupled optical transitions leading to Raman enhancement in  $\text{ZnCo}_2\text{O}_4$  are fundamentally different from those observed in  $\text{Co}_3\text{O}_4$ .

We suspect that the differences observed in the thermal difference spectra of  $\text{Co}_3\text{O}_4$  and  $\text{ZnCo}_2\text{O}_4$  are related to the disruption in the translational symmetry of the lattice induced by cation inversion in  $\text{ZnCo}_2\text{O}_4$ . From assignment of optical transitions in  $\text{Co}_3\text{O}_4$ , the low-energy optical transitions are either highly localized transitions between neighboring Co atoms or intra-atomic transitions in  $T_d \text{ Co}$  (Table 1). With fewer Co atoms in  $T_d$  sites, these optical transitions become suppressed, as observed in  $\text{ZnCo}_2\text{O}_4$  (Fig. 6B); however, the localized transitions still exhibit phonon coupling, leading to the observed resonance Raman enhancements (Fig. 7C). The lack of



temperature dependence of these transitions in  $\text{ZnCo}_2\text{O}_4$  suggests that, although these transitions access localized phonon-coupled states, the localization is related to the static lattice defect of tetrahedral cobalt rather than thermally induced dynamic lattice displacements within a pristine, translationally symmetric lattice. The lack of thermal dependence in  $\text{ZnCo}_2\text{O}_4$ , but presence of resonance Raman enhancement of  $\text{O}_h$  phonon modes upon excitation of a transition from  $\text{T}_d$  Co to  $\text{O}_h$  Co, is strong evidence of an optically accessed polaronic state. We therefore conclude that in both  $\text{Co}_3\text{O}_4$  and  $\text{ZnCo}_2\text{O}_4$  there is evidence of an optically accessible polaronic state. However, in  $\text{Co}_3\text{O}_4$  formation of the polaronic state is due to intrinsic, dynamic lattice deformations caused by thermally activated phonons (exactly analogous to the mechanism we observe in hematite), whereas in  $\text{ZnCo}_2\text{O}_4$ , photoinduced polaron formation is mediated by static lattice defects arising from cation inversion.

## Conclusions

The optical characterization of  $\text{Co}_3\text{O}_4$  and  $\text{ZnCo}_2\text{O}_4$  reported here shows that low energy transitions at 0.8, 0.9 eV and 1.6 eV arise from tetrahedrally coordinated cobalt ions, regardless of oxidation state ( $\text{Co}^{2+}$ ,  $\text{Co}^{3+}$ ). The d-d transition observed at 1.6 eV in both  $\text{Co}_3\text{O}_4$  and  $\text{ZnCo}_2\text{O}_4$  is a phonon-coupled optical transition. In  $\text{Co}_3\text{O}_4$ , the combination of temperature-dependent intensity and resonance Raman enhancement at the onset of the  $\text{O}_h$   $\text{Co}^{3+} \rightarrow \text{T}_d$   $\text{Co}^{2+}$  optical transition (1.64 eV) is evidence that this transition directly populates an intrinsic polaron state coupled to thermally activated phonons. Although the resonance Raman profile of  $\text{ZnCo}_2\text{O}_4$  exhibits a similar Raman enhancement upon excitation of the 1.6-eV transition, the observed inversion and lack of temperature dependence suggests that this photoexcited polaron forms due to the presence of tetrahedral cobalt substitutional defects within the lattice. We conclude that  $\text{T}_d$ -coordinated cobalt is a significant factor in phonon-coupled transitions for cobalt-containing spinel oxides. The contrast in the mechanism of polaron formation (intrinsic or self-trapped polaron vs. defect-mediated polaron) observed for these similar materials is an important insight that we anticipate will help uncover mechanisms of photoinduced polaron formation in other oxide materials. Although both mechanisms of polaron formation observed here have been previously reported, understanding the spectral signatures that distinguish them is crucial for further development of oxide materials for photo-applications.

## Data availability

Data supporting this article have been included as part of the ESI.† Primary data are available from the authors upon reasonable request.

## Author contributions

E. P. C., J. L. S., and K. E. K. contributed to conceptualization of the project and interpretation of the data; K. E. K. supervised

the project. E. P. C. performed most experiments and computations. J. L. S. contributed to some computations for  $\text{Co}_3\text{O}_4$ . M. T. R. collected low-frequency Raman spectra. E. P. C. wrote the manuscript. All authors reviewed the manuscript.

## Conflicts of interest

There are no conflicts to declare.

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