



Cite this: *Environ. Sci.: Nano*, 2025, 12, 1896

Sorption of arsenate on cerium oxide: a simulated infrared and Raman spectroscopic identification†

Khoa Minh Ta,^a Deyontae O. Wisdom,^a Lisa J. Gillie,^a David J. Cooke,^a Runliang Zhu,^{bcd} Mário A. Gonçalves,^e Stephen C. Parker^f and Marco Molinari^{id}*^a

Ceria (CeO₂) is a candidate for arsenic removal, and characterizing its surface speciation is crucial for controlling its removal ability. Here, we focus on arsenates and exploit *ab initio* calculations to study their interaction with the three most stable surfaces of CeO₂. The adsorption of arsenate is stronger on the {100} surface followed by the {110} and {111} surfaces. We find that arsenate can potentially adsorb to CeO₂ surfaces, with a range of binding configurations. Interestingly, we discovered a 5-fold coordinated As(v) species in a trigonal bipyramidal coordination, which is stable and displays a strong interaction with the surfaces, pulling oxygen out of the surfaces, which should be a valuable model to address in As adsorption experiments such as EXAFS. We then predict the infrared (IR) and Raman spectral signatures, finding that adsorbed arsenates have a characteristic spectral fingerprint between 200 and 1200 cm⁻¹. Characteristic peaks compared with experiments gives confidence in the modelling. The 5-fold coordinated As species in particular shows potential diagnostic As–O stretching modes between 635–756 cm⁻¹ in IR spectra and 387–521 cm⁻¹ in Raman spectra. While all binding modes for arsenate adsorption on ceria provide IR active modes, interestingly this is not the case for Raman active modes. Here, we provide a set of reference spectra and binding modes for arsenates on CeO₂ that can further experimental characterization of arsenate speciation, and provide control of its impact on the removal performance of cerium dioxide.

Received 24th September 2024,
Accepted 6th January 2025

DOI: 10.1039/d4en00894d

rsc.li/es-nano

Environmental significance

Metal oxide phases are important materials for removal and control of As in natural aqueous solution because of their strong interactions with As complexes. Highly efficient adsorbents, such as cerium oxide nanoparticles, are thus worth studying for As removal in natural systems. While there is a wealth of data on As adsorption mechanisms on natural metal (hydr)oxides, less is known about Ce oxides. A new 5-fold As complex is proposed and spectroscopic data on other As surface complex species is produced aiming to assist on the characterization of adsorption mechanisms and adsorbent selective studies. These findings can equally be extended to similar metal oxide systems.

1. Introduction

Arsenic based compounds are abundant in water in several countries including China, Cambodia, India, Bangladesh, the coastal aquifers of Australia, West Africa, Western USA and Northern Chile.^{1–4} A safety guideline has been set out for the concentration of arsenic not to be greater than 10 ppb in drinking water by the World Health Organization (WHO),⁵ as the long-term exposure to the inorganic form of arsenic is reported to cause severe health problems including cancer, cardiovascular diseases and diabetes.⁶ Inorganic arsenic also relates to the suppression of cognitive development in early-age infants.^{7–9} Therefore, it is essential to monitor, detect and remove arsenic from water sources and drinking water.^{10,11}

^a Department of Physical and Life Sciences, School of Applied Sciences, University of Huddersfield, Queensgate, Huddersfield HD1 3DH, UK.

E-mail: m.molinari@hud.ac.uk

^b CAS Key Laboratory of Mineralogy and Metallogeny, Guangdong Provincial Key Laboratory of Mineral Physics and Materials, Guangzhou Institute of Geochemistry, Chinese Academy of Sciences, Guangzhou 510640, China

^c CAS Center for Excellence in Deep Earth Science, Guangzhou 510640, China

^d University of Chinese Academy of Sciences, Beijing 100049, China

^e Department of Geology and IDL, Faculty of Sciences, University of Lisbon, Campo Grande, 1749-016 Lisbon, Portugal

^f Department of Chemistry, University of Bath, Claverton Down, Bath BA2 7AY, UK

† Electronic supplementary information (ESI) available: Adsorption energies in numerical form, raw data for the IR and Raman spectra, and full details of the IR and Raman signals. See DOI: <https://doi.org/10.1039/d4en00894d>



In ground water, arsenic is found mainly to be in two ionic forms including arsenate, As(v), and arsenite, As(III). Arsenic is known to develop a strong affinity with natural metal (hydr)oxides,^{12,13} including the ability to attenuate As concentrations in mine drainage waters under proper conditions.¹⁴ Cerium oxide also shows good interaction toward both species, hence it can be considered as a potential option for arsenic removal in water.^{15–21} In Li *et al.* the “hydrous cerium oxide” produced by the authors could remove more than 100 mg g⁻¹ of both As(v) and As(III) species.¹⁶ Although depending on the synthesis conditions of nanoparticles and of the aqueous solution used, ceria nanoparticles were able to remove 95% of both arsenate and arsenite species within one hour,²⁰ while coating provides a >90% removal toward As(v).²¹ In general nanoceria exhibit a more favourable adsorption with As(III) than As(v) across the full pH range.²² Ce⁴⁺ can oxidise As(III) to As(v), while Ce³⁺ can reduce As(v) to As(III), thanks to the localised electron in the 4f orbital of Ce³⁺.²³

It is known that the surface chemistry and composition of nanoceria play an important role in the interaction with molecular species.^{24–28} However, little is known about adsorbed arsenates, As(v), on nanoceria, with some IR spectroscopy finding characteristic peaks for stretching (828 and 839 cm⁻¹)^{20,29} and bending (470 cm⁻¹)²⁹ modes. Modes of adsorption are unknown on ceria but arsenates have displayed both monodentate and bidentate complexes on other metal oxides (*e.g.* iron oxide).^{30–32} Knowing the adsorption modes has also been fundamental for the modelling and prediction of As species in macroscopic natural systems using surface complexation modelling (SCM).^{33–35} Ceria is expected to remove arsenate *via* the exchange of the surface hydroxyl with water dissolved arsenate ions.²³

Previously, we have demonstrated that the adsorption of phosphate species may occur in trigonal bipyramidal configurations wherein the oxygen atoms of the ceria surfaces are bound to the phosphorus atom.²⁵ This is speculated to be the first step towards the scavenging of CeO₂ by phosphates to form CePO₄.^{36,37} Such information is not available for arsenates, and the formation of CeAsO₄ upon adsorption of arsenates is unknown. Due to the similarity of phosphates and arsenates, with both P and As atoms in a +5 oxidation state and both species 4-fold coordinated, it is worth investigating if similar arsenate adsorption modes on ceria nanoparticles are equally likely to be formed.

This work aims to study the adsorption of arsenates on the three most important {100}, {110}, and {111} surfaces of CeO₂, employing density functional theory (DFT) calculations and lattice dynamics to simulate the surface speciation of ceria in the presence of arsenates. Furthermore, our research aims to provide computational vibrational analyses of the most stable surface bound arsenate species by simulating the IR and Raman spectra and to demonstrate that these spectroscopic techniques can be a sensitive probe for studying the interaction of arsenates with ceria. This approach and the obtained results are fundamental for the

complete characterization of As dispersion and fate in natural systems. These results obtained can help and guide the search for new forms of adsorption mechanisms and formation of structural complexes. These are fundamental steps to predict long-term As fixation and to design efficient As-removal adsorbents.

2. Methods

Calculations were performed using density-functional theory (DFT) as implemented using the Vienna *Ab initio* Simulation Package (VASP) code.^{38–40} All calculations were carried out using the generalised-gradient approximation (GGA) exchange–correlation functional of Perdew, Burke and Ernzerhof (PBE).^{41,42} A plane wave cut-off energy of 500 eV was used. The ion cores were modelled using projector augmented-wave (PAW) pseudopotentials,^{43,44} with frozen cores of [Kr] for Ce, and [He] for O and As atoms to model the valence electronic structure. The electron localisation of the Ce 4f orbitals was considered using the DFT+*U* using the Dudarev method.⁴⁵ The use of PBE +*U* with $U_{\text{eff}} = 5$ eV is standard for CeO₂.^{24,25,27,46–52}

Like in our previous study, the minimised CeO₂ bulk structure retains the *Fm* $\bar{3}$ *m* space group with a lattice constant of 5.498 Å,²⁵ which is a known overestimation compared with the experimental value of 5.411 Å.⁵³

The slab models of the {100}, {110}, and {111} surfaces were generated from the minimised bulk CeO₂ unit cell using the methodology described in our previous study²⁵ and the METADISE code.⁵⁴ The {100} and {110} surface models correspond to a $\sqrt{2} \times \sqrt{2}$ and $\sqrt{2} \times 1$ expansions of the primitive unit cell, respectively, each with 7 surface layers and 28 CeO₂ formula units, and the {111} model corresponds to a $\sqrt{2} \times \sqrt{2}$ expansion with 5 surface layers and 20 CeO₂ formula units. The surface models were optimised at constant volume, with the top and the bottom layers allowed to relax, using electronic total energy and ionic force convergence criteria of 1×10^{-5} eV and 1×10^{-2} eV Å⁻¹. A *k*-point grid with $2 \times 2 \times 1$ subdivision was used to sample the Brillouin zones.

For each surface, we attempted to minimize several models with arsenic acid adsorbed on the surface in different binding configurations. In some cases, arsenic acid dissociated following adsorption (*i.e.* its hydrogen atoms transferred to the surface oxygen atoms to form hydroxyl groups). The dissociation observed in each model is listed in Table S1.† Images of the structural models were generated using VESTA.⁵⁵

We simulated the infrared (IR) and Raman spectra of arsenates adsorbed on the surfaces of ceria using the procedure outlined in ref. 56, where harmonic lattice dynamics calculations at constant volume are employed. For each configuration, a Γ -point phonon calculation was performed employing the finite-displacement approach as implemented in the Phonopy code,⁵⁷ with a displacement length of 5×10^{-3} Å. Accurate single-point force calculations were performed using the VASP code as a minimizer, and a tight electronic convergence criterion (1×10^{-9} eV). All the



configurations examined are dynamically stable as they did not show any imaginary modes. The IR activities were computed using the Born effective-charge tensors calculated *via* the density-functional perturbation theory (DFPT)⁵⁸ as implemented in the VASP code. The Raman activities were computed from the changes in the macroscopic dielectric tensor when the atoms in the structure are displaced by $\pm 5 \times 10^{-3}$ Å along each normal-mode vector, with the tensors calculated *via* DFPT. For this approach, further details are in ref. 56 and 59. The assignments of the spectral bands were carried out using animations of the phonon modes generated *via* the Phonopy code and visualized *via* VMD.⁶⁰

3. Results and discussion

Determining the structure, energetics and spectroscopic fingerprint of adsorbed arsenates on oxide CeO₂ is key to understand and model this system, since these oxides can be effective scavengers for As. Particularly, we have focused on the 4-fold As adsorption where the arsenate species is in the common tetrahedral shape with four oxygen atoms surrounding the As atom. However, our previous study related to the adsorption of phosphate on cerium oxide has shown that a 5-fold coordinated species arises when P is adsorbed on the surfaces of CeO₂.²⁵ For this reason, we have also investigated the possibility to have such species for As when adsorbed on CeO₂. This species sees 5 oxygen atoms arranged around As in a trigonal bipyramidal configuration. Such 5-fold As species could be an intermediate state of the scavenging of As towards cerium oxide surfaces, a feature that has been reported for phosphate but not for arsenate so far.

3.1. Structure and energetics of adsorbed arsenates on CeO₂ surfaces

The three most stable {100}, {110}, and {111} surfaces of CeO₂ are adsorbed with arsenates in different binding configurations. For these configurations, we use the generic label $\{hkl\}$ - x OAs- y O_{surf}, where $\{hkl\}$ is the Miller index of the surface, x is the oxygen coordination of the As species (either 4 or 5 for tetrahedral or trigonal bipyramidal coordination), and y is the adsorption denticity of the arsenate on the surface based on the number of Ce–O bonds (denticities of 1, 2 and 3 indicate monodentate, bidentate or tridentate binding). For example, {111}-4OAs-3O_{surf} denotes arsenate adsorbed to the {111} surface as a 4-fold coordinated As species with a tridentate adsorption configuration.

Considering the wealth of EXAFS data on As adsorption, there are some general statements showing a consistency of results regarding As adsorption structure and mechanisms.³² Two of the most important refer the predominant formation of inner-sphere complexes, and the maintenance of the coordination geometries in all of the adsorption modes, which means the tetrahedral coordination. However, these authors note that such consistency may be fundamentally due to the fitting of O of the oxyanion species as one average shell, instead of considering them in separate groups. In

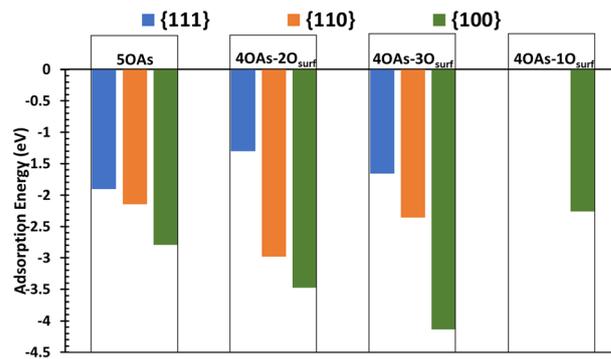


Fig. 1 Adsorption energies of arsenate species in different binding configurations on the stoichiometric {111}, {110}, and {100} surfaces of CeO₂. 5OAs denotes five-fold coordinated As species, and 4OAs-1O_{surf}, 4OAs-2O_{surf}, 4OAs-3O_{surf} denote four-fold coordinated As species adsorbed in monodentate, bidentate and tridentate configurations, respectively.

EXAFS, 4 sorption modes have been identified³² with the clear predominance of inner-sphere (IS) over outer-sphere (OS) complexes, especially the formation of the bidentate, corner-sharing, IS complex.

The strength of the interaction between adsorbed arsenates and CeO₂ surfaces is given by the adsorption energy (eqn (S1)†), which we present in Fig. 1 for all the stable configurations (numerical values are presented in Table S1†). Fig. 2 present the stable configurations, while Table S2† presents the extent of the arsenate dissociations for all configurations.

All energies are negative, implying stable adsorption configurations. The adsorption of arsenate on the {100} surface is generally more stable. The {100} surface is also the only one where a monodentate adsorption is stable ({100}-4OAs-1O_{surf}). The second most stable adsorption is on the {110} surface followed by the {111} surface. 5-Fold coordinate As species are bound to the {100} and {110} surfaces less strongly than 4-fold coordinate As species. Surprisingly this is not the case for the {111} surface where the 5OAs is more stable than any 4OAs configurations. Such trigonal bipyramidal 5-fold coordinated As has not yet been detected in adsorption experiments using either EXAFS or DFT calculations,^{61,62} but although no structure has been proposed, As models with coordination numbers between 4.6 and 4.8 have been used in the fitting of EXAFS spectra of As adsorbed onto Fe–Ce bimetal oxide.⁶³

Arsenate is very similar to phosphate, as P and As belong to the same pnictogen group of the periodic table (group 15), and thus can be compared with the available literature.²⁵ The adsorption of 5-fold arsenate species on nanoceria is more energetically stable than 5-fold phosphate species. The adsorption of 4-fold arsenate species is less energetically stable than adsorbed 4-fold phosphate species, except for {110}-4OAs-2O_{surf}. The differences in energies are presented in Table S1.†

The binding energy of arsenate is stronger than the adsorption of water on nanoceria surfaces (−0.5, −1.2, and −1.6 eV for the {111}, {110}, and {100} surfaces, respectively⁵²),



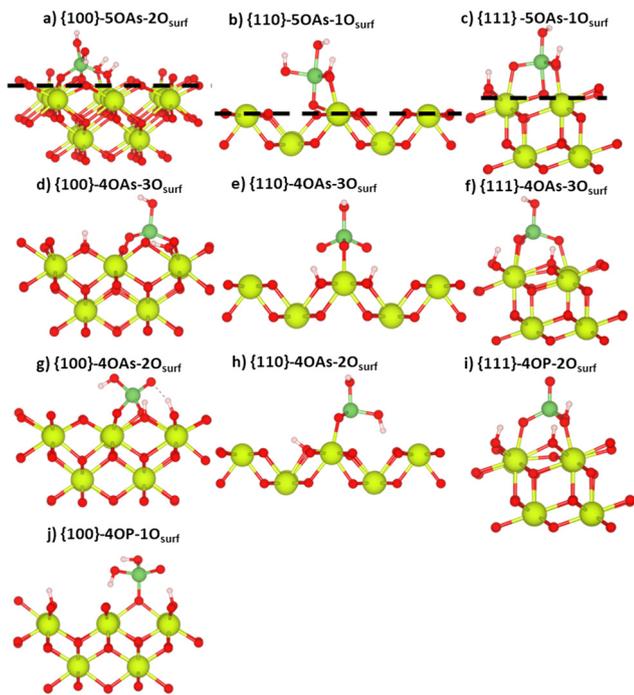


Fig. 2 Arsenate adsorption configurations on the {100}, {110}, and {111} stoichiometric surfaces of CeO_2 . (a)–(c) As in a five-fold trigonal bipyramidal configuration. (d)–(j) As in a four-fold tetrahedral configuration. The dashed blue lines in (a)–(c) show the surface plane to emphasise surface O atoms pulled out by arsenates as described in the text. The atom colours are as follows: O – red, Ce – yellow, As – green, and H – white.

implying that arsenate will adsorb as IS complexes on the surfaces of cerium oxide in an aqueous environment.

3.1.1. Five-fold coordinated As species (5OAs). When an As atom is directly bonded to a surface oxygen (*O_{surf}) it forms a 5-fold coordinated species in a trigonal bipyramidal configuration (Fig. 2a–c). The adsorption energies of these configurations are -1.90 , -2.14 and -2.79 eV for the {111}-5OAs-1O_{surf}, {110}-5OAs-1O_{surf}, and {100}-5OAs-2O_{surf} configurations, respectively (Fig. 1). The preferential surface for the formation of 5OAs species is the {100} facet, followed by the {110} and {111} facets. Like for the adsorption of phosphates, 5OAs in {111}-5OAs-1O_{surf} and {110}-5OAs-1O_{surf} shows monodentate binding, and bidentate binding in {100}-5OAs-2O_{surf}. This is possible because of the flexibility of the {100} facet, where surface oxygen ions rearrange to accommodate the adsorption of arsenates and other species.^{25,27,46,64,65} It has been shown that phosphate species may scavenge the surfaces of ceria, and that 5-fold phosphate adsorption “pulls out” surface oxygen atoms, displacing them by 0.74 Å, 0.21 Å, and 0.20 Å on the {110}, {111} and {100} surfaces, respectively.²⁵ Like phosphate, the arsenate also appears to displace a surface oxygen (*O_{surf}) above the surface plane. This is more prominent in {110}-5OAs-1O_{surf}, where the surface O atom is displaced by 0.60 Å, followed by 0.25 Å on the {100}-5OAs-2O_{surf}. The {111}-5OAs-1O_{surf} does not show any significant surface oxygen displacement. There are similarities between the adsorption of phosphate and

arsenate, with the flat {110} surface being mostly affected by the formation of 5-fold species. This could be related to the ease of formation of oxygen vacancies on the flat {110} surface, which has been reported to be also the underlying cause for the {110}/ {111} faceting of the flat {110} surface.⁴⁶ The formation of this complex does not seem to have parallel in any previous experimental studies of As adsorption onto metal oxides.^{32,33,61,62} However, it is worth noting that ref. 63, using a Fe–Ce bimetal oxide have fitted models with As–O coordination numbers between 4.6 and 4.8, in contrast with the highly consistent adsorbed As data collected by ref. 32. If this result means that dismissing the fitting to a single average shell, as done on most EXAFS spectra,³² would potentially reveal a more complex and varied set of As complex geometries, or that the presence of a Ce oxide or Ce-rich oxide substrate can actually promote the development of a 5-fold As complex, remains to be fully clarified. This 5-fold species is stable on all surfaces, and the absence of imaginary phonon modes further confirms its stability, possibly as an intermediate of the scavenging activity of arsenates towards CeO_2 like in the case of phosphates.²³

3.1.2. Four-fold coordinated As species (4OAs). The adsorption of arsenate as 4-fold coordinated species (*i.e.*, As with a tetrahedral oxygen coordination, 4OAs) on the three surfaces of CeO_2 is found to be a stable adsorption with negative adsorption energies (Fig. 1). However, the adsorption of the 4-fold As species is not as stable as the 5-fold As species on the {111} surfaces. On the {110} surface, all 4-fold As adsorbed species bind more strongly than the 5-fold As adsorbed species. The adsorption of arsenates as 4AsO on the {100} surfaces is more stable for bidentate and tridentate but not for monodentate adsorption compared to the 5AsO configuration.

We investigated three different adsorption configurations of 4OAs, *viz.* monodentate, bidentate and tridentate (Fig. 2d–j).

Both {111}-4OAs-3O_{surf} and {100}-4OAs-3O_{surf} configurations have the most negative adsorption energies compared to the monodentate and bidentate adsorptions, which indicates that 4OAs is preferably adsorbed as tridentate binding configurations on the {100} and {111} surfaces. The {110} surface expresses a different affinity towards arsenate, where the bidentate binding configuration {110}-4OAs-2O_{surf} is preferred compared to all other binding configurations.

The three arsenate oxygen atoms (O_{As}) of the {100}-4OAs-3O_{surf} configuration embed into the surface at a similar height of the surface oxygen atoms (O_{surf}) to form the tridentate binding configuration. This is due to the flexibility and mobility of the surface oxygens of the {100} surface,^{25,27,46,64,65} which allows for the surface rearrangement to accommodate the adsorption, which leads to the strongest adsorption energy (-4.14 eV) compared to all other adsorption configurations. Unlike the tridentate adsorption of 4OAs on the {100} and {111} surfaces, the {110}-4OAs-3O_{surf} configuration has three Ce– O_{As} bonds, with two of them elongated (2.36 *vs.* 2.61 Å) alongside two hydrogen bonds; however, this is not sufficient to stabilize this configuration further compared to {110}-4OAs-2O_{surf} (-2.36 eV *vs.* -2.98 eV).



The 4OAs bidentate adsorption on all three surfaces has two direct Ce–O_{As} bonds (Fig. 2). The adsorption stability follows the order {100}-4OAs-2O_{surf} > {110}-4OAs-2O_{surf} > {111}-4OAs-2O_{surf}, indicating that the strongest bidentate adsorption occurs again on the {100} surface.

We attempted to stabilize 4OAs monodentate configurations on all three surfaces of ceria, but we could only stabilize it for the {100} surface, and its adsorption energy is –2.26 eV. {100}-4OAs-1O_{surf} has a crowded hydrogen bond network, which prevents the adsorption from rearranging to bidentate or tridentate.

3.2. Vibrational signatures of adsorbed arsenate species

The simulated IR and Raman spectra of the adsorbed arsenate on nanoceria surfaces are discussed using the notation used in our previous study on phosphate species.²⁵ ν denotes stretching modes, δ bending, ρ rocking, ω wagging, sc scissoring, and t twisting modes. The subscripts s and as identify symmetric and asymmetric modes, where relevant, and we denote longitudinal (within the surface plane) and transverse (perpendicular to the surface plane) modes by L or T in parentheses. We denote bulk and surface O atoms as O_{bulk} and O_{surf}, respectively, and we further distinguish the surface O directly bonded to the As atom in the 5OP configurations as *O_{surf}. Finally, O_{As} and OH_{As} denote O and OH (hydroxyl) groups belonging to arsenate, OH_{surf} indicates a surface hydroxyl group, and H_{surf} and H_{As} denote H adsorbed on the surface and As oxygen atoms, respectively.

The Ce–O stretch for bulk CeO₂ has a single IR signal at 274 cm⁻¹,²⁵ which is a good representation of the 272 cm⁻¹ from experimental measurements.⁶⁶ Computationally, the Raman active signal is in the range of 434–437 cm⁻¹,^{25,59} which is underestimated compared to the experimental value of 464 cm⁻¹.⁶⁷ For all surfaces, both IR and Raman spectra display characteristic peaks in the region 0–500 cm⁻¹.²⁵

3.2.1. IR of adsorbed arsenate species on CeO₂ surfaces.

The main characteristics in the IR spectra of the surface models with adsorbed arsenates are summarised in Table 1 (with full details in Table S3†). The spectra display characteristic fingerprints of the interaction between the arsenate species and the surface between 200–1200 cm⁻¹ (Fig. 3). Despite the experimental and computational literature of molecular arsenate not being directly comparable to our adsorbed arsenate species, it can provide a useful comparison for the main characteristic features. The region of the IR active modes for the interaction between arsenate species and the surfaces overlaps with the main features of the IR spectrum of molecular arsenate where there are several stretching (ν) modes in the region 500–1200 cm⁻¹ (ν_s As–OH_{As}, ν_{as} As–OH_{As}, ν As=O_{As}; Table 1 and Fig. S1†). The bending modes δ As–OH_{As}, δ As–(OH_{As})₂ of the adsorbed As species are predicted to be at 428 cm⁻¹ and 1065 cm⁻¹. Experimentally the IR active bending modes δ As–OAs of the adsorbed arsenate have been measured at 470 cm⁻¹, which is the same region of the IR active modes of the surface of cerium oxide.²⁹ The IR spectra of the adsorbed arsenates on ceria surfaces also display peaks above 2000 cm⁻¹ (raw spectra in Fig. S2 and full spectra in Fig. S3†) corresponding to the stretching of the hydroxyl groups (ν O–H_{surf}/ ν O–H_{As}), which are in good agreement with the literature.^{20,30,68,69}

Fig. 3(a–c) shows the IR spectra of the 5OAs binding configurations for the {100}, {110}, and {111} stoichiometric surfaces. The displacement of the surface oxygen seen for {110}-5OAs-1O_{surf} and {100}-5OAs-2O_{surf} is also observed in the IR active mode as the ν As–*O_{surf}, where *O_{surf} is the surface oxygen atom directly bonded to As. The ν As–*O_{surf} for the {110}-5OAs-1O_{surf} and {100}-5OAs-2O_{surf} are at 493–756 cm⁻¹ and 496 cm⁻¹, respectively. Such signals are in a similar region as the 5-fold coordinated phosphate species (680–695 cm⁻¹).²⁵ The absence of displaced O_{surf} for {111}-5OAs-1O_{surf} results in no IR active mode for ν As–*O_{surf} on the {111} surface.

Table 1 Assignment of the major features (with a cut-off signal intensity of 0.1 a.u) in the simulated IR spectra of arsenate species adsorbed onto the {111}, {110}, and {100} stoichiometric surfaces of CeO₂ with different binding modes. Vibrations are labelled according to the scheme outlined in the text. The frequency ranges indicate where a given mode is seen across all binding modes on all three surfaces. Vibrational frequencies for molecular arsenate species in the gas phase and in solution are also given for comparison. Comparison with adsorbed phosphate on ceria is also provided²⁵

IR mode arsenate	Adsorbed arsenate on CeO ₂ (PBE+U, this study)	Adsorbed arsenate on CeO ₂ (Expt) ^{20,29}	Molecular arsenate (PBE+U, this study)	Molecular arsenate (DFT) ⁶⁸	Arsenate (aq) (Expt) ^{69,70}	IR mode phosphate	Adsorbed phosphate (PBE+U) ²⁵
ρ O _{As} –H _{As}	284–350, 885	—	134–317, 952–986	215–330	—	ρ O _p –H _p	547–977
ρ O _{surf} –H _{surf}	417, 778–894	—	—	—	—	—	618–870
ν O _{As} –H _{As}	2732–3694	—	3684–3699	3700–4092	—	—	—
ν O _{surf} –H _{surf}	2507–3207	—	—	—	—	—	—
t, ω O ₂ As(OH) ₂	287	—	—	250–330	—	—	—
ν , ρ , δ As–*O _{surf}	635–756	—	—	—	—	ν P–*O _{surf}	680–695
ν_s As–OH _{As}	591–979	828, 839 (ref. 20)	656–695	493–924	—	ν_s P–OH _p	796–833
ν_{as} As–OH _{As}	—	—	677	852–1008	—	ν_{as} P–OH _p	853–1007
ν_s As–O _{As}	778–848	—	—	812–924	875–878, ⁷⁰ 830–890 (ref. 69)	ν_s P–O _p	965–1077
ν_{as} As–O _{As}	783–894	—	—	852–1008	858–908, ⁷⁰ 830–890 (ref. 69)	ν_{as} P–O _p	985–1160
δ As–OH _{As} ; δ As–(OH _{As}) ₂	324–349, 975	470 (ref. 29)	278–292, 1032	1028–1157	315–385, ⁷⁰ 1170 (ref. 69)	δ P–OH _p	1043–1077
δ As–O _{As} ; δ As–(O _{As}) ₂	273–320	—	280–292	—	—	—	—



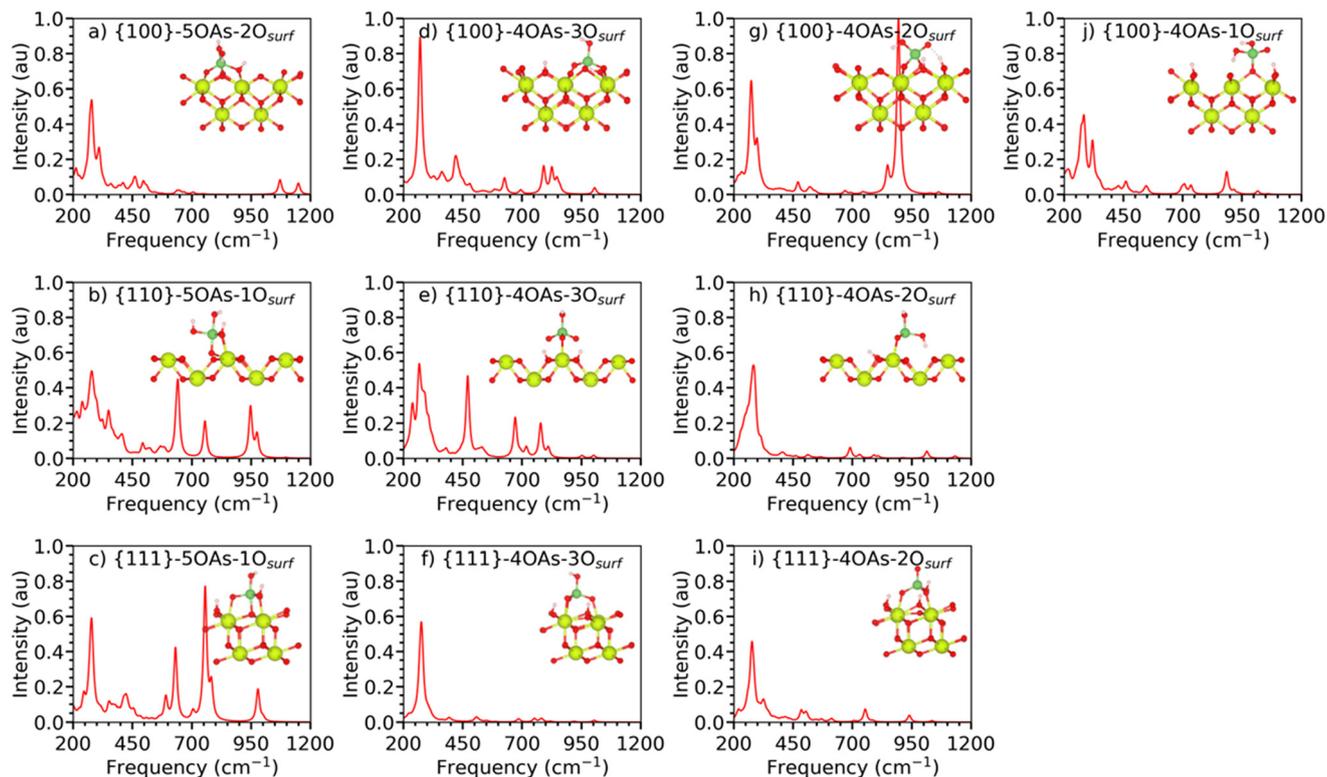


Fig. 3 Simulated IR spectra of arsenate species adsorbed onto the {111}, {110}, and {100} stoichiometric surfaces of CeO₂ with different binding modes as shown by the inset structures. The spectra are normalised relative to each other such that the highest absolute intensity across all the spectra is set to unity.

All three 5OAs binding configurations show $\nu_{\text{As-O}_{\text{As}}}$. The $\nu_{\text{As-O}_{\text{As}}}$ are at 757–783 cm⁻¹, 635–756 cm⁻¹ and 1150 cm⁻¹ for the {111}-5OAs-1O_{surf}, {110}-5OAs-1O_{surf} and {100}-5OAs-1O_{surf}, respectively. The high frequency value of $\nu_{\text{As-O}_{\text{As}}}$ on the {100}-5OAs-1O_{surf} is similar to the IR signal observed for 5-fold phosphate species adsorbed on the {100} surface (1197 cm⁻¹).²⁵

The bending modes $\delta_{\text{As-OH}_{\text{As}}}$ are not observed on {111}-5OAs-1O_{surf} and {100}-5OAs-1O_{surf}, whereas they are a strong IR feature in the region of 324–350 cm⁻¹ in {110}-5OAs-1O_{surf}, which is in the same region of the IR signal for $\delta_{\text{As-OH}_{\text{As}}}$ in molecular arsenate (278–292 cm⁻¹, 1032 cm⁻¹). The $\delta_{\text{As-O}_{\text{As}}}$ mode gives rise to IR active signals at 413 and 1072 cm⁻¹ for {111}-5OAs-1O_{surf} and {100}-5OAs-2O_{surf}. The much higher shift for the {100} surface may be due to the bidentate binding configuration of the {100}-5OAs-2O_{surf} compared to the monodentate binding configuration of the {111}-5OAs-1O_{surf}.

We predicted the IR stretching of the As=O_{As} bond ($\nu_{\text{As=O}_{\text{As}}}$) at 952 cm⁻¹ and at 1032 cm⁻¹ for molecular arsenate, Table S4.† Similar to the phosphate adsorption on ceria, the IR signal of $\nu_{\text{As=O}_{\text{As}}}$ is not observed in any of our 5OAs binding configurations, which may arise from the elongation of the As=O_{As} bond from 1.64 Å (isolated molecule) to 1.71–1.89 Å when adsorbed on ceria, potentially resulting in the band suppression in the IR spectra.^{25,71}

The IR spectra of the stable monodentate, bidentate and tridentate 4OAs adsorptions on the three surfaces of CeO₂ are

shown in Fig. 3(d–j). In the following, we discuss the spectra with respect to the denticity of the binding.

The {100}4OAs-1O_{surf} is the only monodentate configuration we could stabilize for the 4OAs adsorption (Fig. 3j). The $\nu_{\text{As-O}_{\text{As}}}$ gives rise to a strong characteristic feature at 885 cm⁻¹ in the IR spectrum. The IR bending mode ($\delta_{\text{As-O}_{\text{As}}}$) is at 320–321 cm⁻¹, which is comparable to the $\delta_{\text{As-O}_{\text{As}}}$ of molecular arsenate (280–292 cm⁻¹).

The {111}-4OAs-2O_{surf} and {100}-4OAs-2O_{surf} configurations have fingerprint characteristic signals for $\nu_{\text{As-O}_{\text{As}}}$ at 754 cm⁻¹ and 848 cm⁻¹, respectively. These signals are similar to those $\nu_{\text{As-O}_{\text{As}}}$ for molecular arsenate at 812–924 cm⁻¹ (DFT literature⁶⁸) and at 830–890 cm⁻¹ (experimental study^{69,70}). {110}-4OAs-2O_{surf} has $\nu_{\text{As-OH}_{\text{As}}}$ at 690 cm⁻¹, which is similar to the $\nu_{\text{As-OH}_{\text{As}}}$ for the isolated molecular arsenate at 656–695 cm⁻¹ (this study), and is within the range of computational studies (493–924 cm⁻¹).⁶⁸ As the IR spectra of all the configurations were standardized with the highest IR signal across all the 5OAs and 4OAs binding configurations, the {100}-4OAs-2O_{surf} has the dominant IR frequency at 894 cm⁻¹ with 1.0 a.u. intensity for $\nu_{\text{As-O}_{\text{As}}}$.

The tridentate adsorption configurations of the 4OAs arsenate species for all three surfaces have dominant IR stretching for the As–O_{As} and the As–OH_{As} bonds. The $\nu_{\text{As-O}_{\text{As}}}$ and $\nu_{\text{As-O}_{\text{As}}}$ give rise to the signals at 626–1002 cm⁻¹ and at 752–859 cm⁻¹, respectively (Table S4,† most of these peaks are weak with intensities between 0.05 and 0.1 a.u.). The $\nu_{\text{As-OH}_{\text{As}}}$ and $\nu_{\text{As-OH}_{\text{As}}}$ signals are at 626–779 cm⁻¹



Table 2 Assignment of the major features (with the cut-off signal intensity of 0.1 a.u.) in the simulated Raman spectra of arsenate species adsorbed onto the {111}, {110}, and {100} stoichiometric surfaces of CeO₂ with different binding modes. Modes are labelled according to the scheme outlined in the text. The frequency ranges indicate where a given mode is seen across all binding modes on all three surfaces. Vibrational frequencies for molecular arsenate species in the gas phase and in solution are also given for comparison. Comparison with adsorbed phosphate on ceria is also provided²⁵

Raman mode arsenate	Adsorbed arsenate on CeO ₂ (PBE+U, this study)	Isolated arsenate (PBE+U, this study)	Arsenate (aq) (experiment) ^{70,72}	Raman mode phosphate	Adsorbed phosphate (PBE+U) ²⁵
$\rho\text{O}_{\text{As}}-\text{H}_{\text{As}}$	229–786, 916–1257	280–317, 952	250 (ref. 72)	$\rho\text{O}_{\text{p}}-\text{H}_{\text{p}}$	916–1077
$\rho\text{O}_{\text{surf}}-\text{H}_{\text{surf}}$	360–786, 916–1065	—	—	$\rho\text{O}_{\text{surf}}-\text{H}_{\text{surf}}$	584–955
$\nu\text{O}_{\text{As}}-\text{H}_{\text{As}}$	2732–3701	3684–3699	—	—	—
$\nu\text{O}_{\text{surf}}-\text{H}_{\text{surf}}$	2507–3722	—	—	—	—
$\delta\text{As}-\text{O}_{\text{As}}$; $\delta\text{As}-(\text{O}_{\text{As}})_2$	229–435, 1016–1072	280	270–365, ⁷² 285–385	—	—
$\delta\text{As}-\text{OH}_{\text{As}}$; $\delta\text{As}-(\text{OH}_{\text{As}})_2$	428, 1065	—	(ref. 70)	$\delta\text{P}-\text{OH}_{\text{p}}$; $\delta\text{P}-(\text{OH}_{\text{p}})_2$	1047–1236
$\rho\text{As}-\text{O}_{\text{As}}$; $\rho\text{As}-\text{OH}_{\text{As}}$	400–484	—	—	—	—
$t, \omega\text{O}_2\text{As}(\text{OH})_2$	568–571	—	315–319 (ref. 70)	—	—
$\nu, \rho, \delta\text{As}-*\text{O}_{\text{surf}}$	387–421	—	—	$\nu\text{P}-*\text{O}_{\text{surf}}$	680–701
$\nu_{\text{s}}\text{As}-\text{OH}_{\text{As}}$	—	656	700–745 (ref. 70)	$\nu_{\text{s}}\text{P}-\text{OH}_{\text{p}}$	750–859
$\nu_{\text{as}}\text{As}-\text{OH}_{\text{As}}$	1150	—	765 (ref. 70)	$\nu_{\text{as}}\text{P}-\text{OH}_{\text{p}}$	902–1079
$\nu_{\text{s}}\text{As}-\text{O}_{\text{As}}$	561–798, 1002–1028	—	834–875 (ref. 70)	$\nu_{\text{s}}\text{P}-\text{O}_{\text{p}}$	1043–1077
$\nu_{\text{as}}\text{As}-\text{O}_{\text{As}}$	505–954	—	866–915 (ref. 70)	$\nu_{\text{as}}\text{P}-\text{O}_{\text{p}}$	1142–1160
$\text{scAs}-\text{O}_{\text{As}}$; $\text{scAs}-\text{OH}_{\text{As}}$	340–551	317	270–450 (ref. 72)	—	—
$\nu\text{As}=\text{O}_{\text{As}}$	—	952	923 (ref. 72)	—	—

and at 683–847 cm⁻¹. This wide spread of IR signals for the $\nu\text{As}-\text{O}_{\text{As}}$ is in the range of $\nu_{\text{s}}\text{As}-\text{O}_{\text{As}}$ (656–695 cm⁻¹) and $\nu\text{As}-\text{OH}_{\text{As}}$ (677 cm⁻¹) for the isolated molecular arsenate (Table 2).

Our prediction for the IR characteristic features of the adsorbed arsenates is similar to the IR signals for adsorbed phosphates on nanoceria.^{25,71} The As–OH_{As} and As–O_{As} stretching modes are generally the most intense signals (as for adsorbed phosphates on ceria), with $\nu\text{As}-\text{OH}_{\text{As}}$, $\nu_{\text{s}}\text{As}-\text{O}_{\text{As}}$, $\nu_{\text{as}}\text{As}-\text{O}_{\text{As}}$ in the regions of 591–979 cm⁻¹, 778–848 cm⁻¹, 783–894 cm⁻¹, respectively. Our predicted values are in a wider range of frequencies compared to the experimental $\nu\text{As}-\text{O}_{\text{As}}$ IR active stretching signals (828–839 cm⁻¹) for adsorbed arsenates on ceria,^{20,29} and for the IR active stretching modes of isolated aqueous arsenate (830–908 cm⁻¹),^{69,70} most likely due to the direct bonding to the surface of ceria.

Our spread of simulated $\delta\text{As}-\text{O}_{\text{As}}$ (at 273–320 cm⁻¹) and $\delta\text{As}-\text{OH}_{\text{As}}$ (at 324–349 cm⁻¹, and 975 cm⁻¹) signals is consistent with the bending modes for aqueous arsenate found at 315–385 cm⁻¹,⁷⁰ and at 1170 cm⁻¹.⁶⁹ Although we report only a few peaks for the bending modes, there are many more associated with intensities lower than 0.1 a.u. (Table S4[†]). The complex interaction between arsenate and ceria surfaces causes such a spread, where visual inspection of the vibrational modes has shown complex motions, all related to bending modes of As–O. Such a spread is, however, not seen in the experimental reported value for $\delta\text{As}-\text{O}_{\text{As}}$, which is observed at 470 cm⁻¹ for adsorbed arsenate on ceria.²⁹ This is most likely because our intensities for the bending modes are low.

3.2.2. Raman shifts of adsorbed arsenates on CeO₂ surface. Unlike the IR fingerprint, some of the adsorbed arsenate binding configurations have their Raman vibrational signature suppressed before (raw data in Fig. S4[†]) and after normalisation with respect to the highest intensity peak across all structures. Fig. 4 shows the normalised Raman

spectra of the adsorbed arsenate on the nanoceria surfaces in the range of 200–1200 cm⁻¹. The region below 600 cm⁻¹ is dominated by intense signals related to the surfaces,²⁵ as well as the bending mode (δ) of the arsenates (Table 2). The Raman active stretching modes of the hydroxyl groups ($\nu\text{O}-\text{H}_{\text{As}}$ and $\nu\text{O}-\text{H}_{\text{surf}}$) are predicted to be above 2000 cm⁻¹ and those are seen with very weak intensity in some structures (Fig. S5[†]). The common Raman active modes of the adsorbed arsenate amongst the surfaces are summarized in the Table 2.

The unique $\nu\text{As}-*\text{O}_{\text{surf}}$ stretching of the 5-fold As configurations is suppressed for the {110}-5OAs-1O_{surf} and {111}-5OAs-1O_{surf}. The {100}-5OAs-2O_{surf} has the $\nu\text{As}-*\text{O}_{\text{surf}}$ Raman active mode at 516 cm⁻¹ with a very low intensity of 0.06 a.u.

The adsorption of monodentate arsenate on the {100} surface shows a $\nu_{\text{as}}\text{O}_{\text{As}}-\text{As}-\text{OH}_{\text{As}}$ signal at 1016 cm⁻¹ (Fig. 4j), which falls close to the $\nu\text{As}=\text{O}_{\text{As}}$ signal of the isolated arsenate (952 cm⁻¹). Across all surfaces, the stretching signals are in the ranges 668–1150 cm⁻¹ for $\nu\text{As}-\text{OH}_{\text{As}}$ and 505–1028 cm⁻¹ for $\nu\text{As}-\text{O}_{\text{As}}$. These signals are also consistent with the Raman active modes for the computed isolated molecular arsenate (656 cm⁻¹ for As–OH and 952 cm⁻¹ for As–O), and in line with the reported experimental values (700–765 cm⁻¹ for As–OH and 834–915 cm⁻¹ for As–O) (Table 2).^{70,72}

The scissoring modes ($\text{scAs}-\text{O}_{\text{As}}$, $\text{scAs}-\text{OH}_{\text{As}}$) are frequent active modes (with an intensity ≥ 0.1) in the Raman spectra in the region of 340–551 cm⁻¹ (Table 2). These are in line with $\text{scAs}-\text{O}_{\text{As}}$ and $\text{scAs}-\text{OH}_{\text{As}}$ Raman active modes observed in the isolated molecular arsenate (317 cm⁻¹ with 0.08 a.u.), and the experimental values (270–450 cm⁻¹).^{69,72} Adsorbed arsenates on ceria surfaces display Raman active scissoring modes at higher frequency compared to the isolated arsenate, most likely due to the direct bonding between the arsenate and the surface.



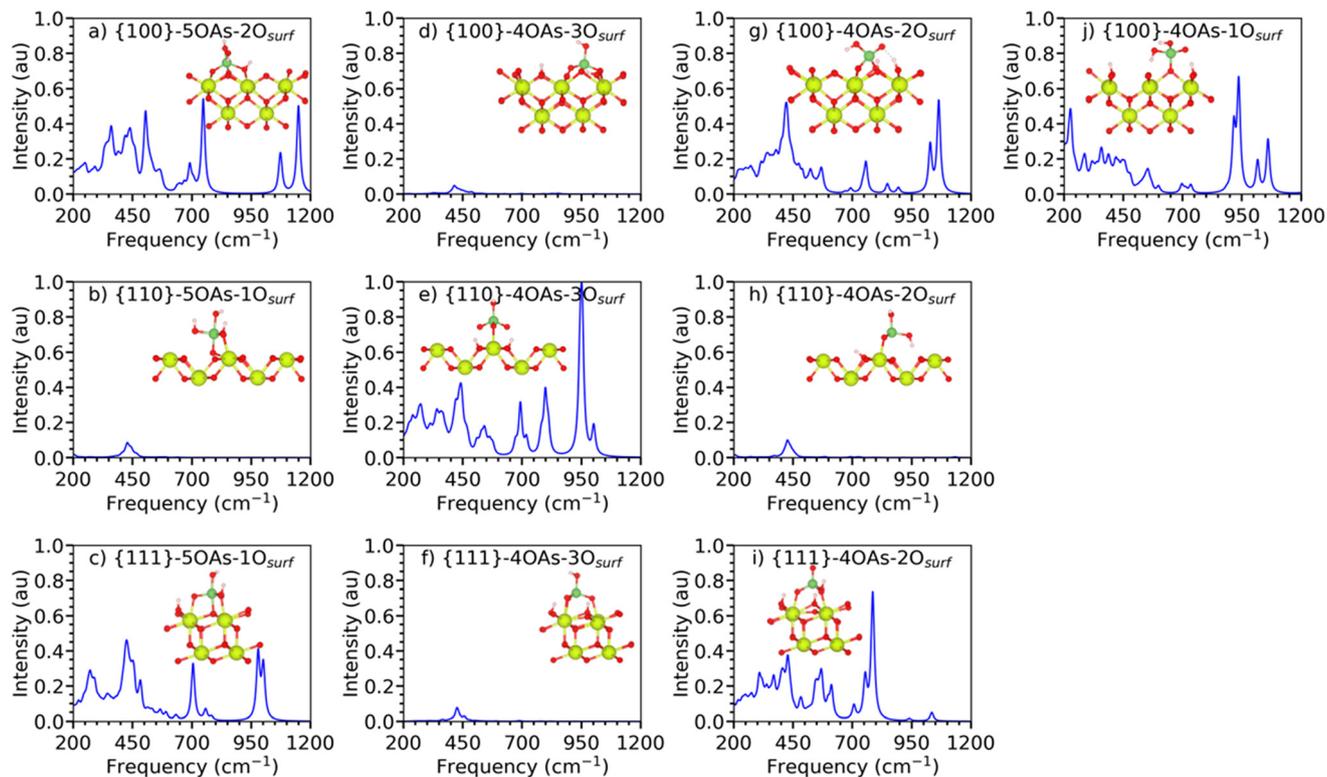


Fig. 4 Simulated Raman spectra of arsenate species adsorbed onto the {111}, {110}, and {100} stoichiometric surfaces of CeO₂ with different binding modes as shown by the inset structures. The spectra are normalised relative to each other such that the highest absolute intensity across all the spectra is set to unity.

The large range of Raman active rocking modes ($\rho_{\text{OAs-HAs}}$) are in the region 229–786 cm⁻¹, and 916–1257 cm⁻¹, which is comparable with the Raman signals for the isolated arsenate (280–317, and 952 cm⁻¹) and the rocking mode of aqueous arsenate measured at 250 cm⁻¹.⁷² The plethora of rocking modes arises from the interaction of the arsenate with the surface. However, the majority of the Raman active $\rho_{\text{OAs-HAs}}$ signals are weak, but there are some exceptions with high intensity such as {111}-5OAs-1Osurf (at 1001 cm⁻¹ with 0.37 a.u.), {100}-5OAs-2Osurf (at 1076 cm⁻¹ with 0.17 a.u.), {100}-4OAs-1Osurf (at 916 cm⁻¹, 937 cm⁻¹ and 1257 cm⁻¹ with 0.52 a.u., 0.67 a.u., and 0.45 a.u. respectively). The higher frequency shift for rocking modes in the case of adsorbed arsenate is similar to the shift observed for adsorbed phosphate (916–1077 cm⁻¹ for adsorbed phosphate species vs. 144–322 cm⁻¹ for molecular phosphate²⁵), and this again confirms the significant constraints effect due to the adsorbed arsenate interacting with the ceria surfaces.

4. Conclusions

This study provides a prediction of IR and Raman signals for adsorbed arsenates onto the three most stable Miller index {100}, {110}, and {111} surfaces of CeO₂ including different arsenate binding configurations. This study also shows the potential of IR and Raman spectroscopy to identify the

binding configurations of adsorbed arsenates onto the surfaces of ceria.

We found that arsenate can bind to the nanoceria surfaces in both trigonal bipyramidal (five-fold) and tetrahedral (four-fold) configurations. Whereas the tetrahedral binding is expected since it has been consistently identified as the main As structure bound to oxide surfaces,³² the five-fold binding is peculiar since it has never been described or suggested in experimental models, but found to be stable. The five-fold binding configurations exhibit some oxygen scavenging, pulling surface oxygen above the surface plane. Due to the nature of the binding, generally the four-fold adsorption maximised the interaction between the arsenate species with the three surfaces resulting in more thermodynamically favourable energetics, especially the tridentate adsorption which has generally the most negative binding energies.

The identification of 5-fold coordinated As has not been addressed in As adsorption experiments on oxides, using either EXAFS or DFT calculations, which in general support the formation of predominantly monodentate mononuclear and bidentate binuclear complexes.^{61,62} However, in As adsorbed onto Fe–Ce bimetal oxide, coordination numbers between 4.6 and 4.8 were used as models, meaning that our As complex structure deserves further investigation to clarify the As adsorption mechanisms.⁶³ The incorporation of this data into SCM is recognized as fundamental and may provide clues for open issues, such as the enhanced



adsorption behaviour of As into oxides with co-existing ions.⁷³

The IR and Raman spectra of the adsorbed arsenate show a stretching characteristic fingerprint in the region above 700 cm^{-1} similar to experimental findings. However, we found a plethora of stretching and bending modes (many with low intensity) in the region between 200–1200 cm^{-1} , which overlap between the IR and Raman active regions of arsenate adsorption on ceria but also with the region of IR and Raman characteristic signature of bare ceria surfaces. The five-fold arsenate configurations have unique IR fingerprint signals in the region of 635–756 cm^{-1} , and Raman fingerprint signals between 387–421 cm^{-1} for $\nu\text{As}-*\text{O}_{\text{surf}}$ due to the direct interaction between As and a surface oxygen. These characteristic signals give an indication to identify these species experimentally. We also find that some of the binding configurations have suppressed Raman spectra altogether, which would make their identification more challenging using experimental Raman measurements.

Finally, future studies should focus on more complex models including arsenite, and the effect of coverage and solvent effects are considered.

Data availability

Raw data are available from <https://doi.org/10.17632/rkbbxcmj4>.

Author contributions

Conceptualization: LJG, DJC, RZ, MAG, SCP, MM; data curation: KMT, MM; formal analysis: KMT, DOW, MM; funding acquisition: LJG, DJC, MM; investigation: KMT, DOW, MAG, MM; methodology: KMT, DOW, MM; project administration: LJG, DJC, RZ, MAG, SCP, MM; software: KMT, MM; resources: LJG, DJC, MM; supervision: LJG, DJC, MM; validation: KMT, DOW, MM; visualization: KMT, DOW, LJG, DJC, RZ, MAG, SCP, MM; writing – original draft: KMT, DOW, LJG, DJC, RZ, MAG, SCP, MM; writing – review & editing: KMT, DOW, LJG, DJC, RZ, MAG, SCP, MM.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

KMT is funded *via* the Vice Chancellor's Scholarship Scheme at the University of Huddersfield. Analysis was performed on the Orion computing facility and the Violeta HPC at the University of Huddersfield. Calculations were run on the ARCHER2 UK National Supercomputing Services *via* our membership of the UK HEC Materials Chemistry Consortium (MCC; EPSRC EP/X035859/1). MAG acknowledges funding from the project UIDB/50019/2020 to IDL, by Fundação para a Ciência e a Tecnologia, I.P./MCTES through PIDDAC National funds.

References

- 1 J. Podgorski and M. Berg, Global threat of arsenic in groundwater, *Science*, 2020, **368**, 845–850.
- 2 P. Ravenscroft, H. Brammer and K. Richards, Arsenic in Asia, in *Arsenic Pollution: A Global Synthesis*, Wiley-Blackwell, 2009, ch. 8, pp. 318–386.
- 3 P. Ravenscroft, H. Brammer and K. Richards, Arsenic in North America and Europe, in *Arsenic Pollution: A Global Synthesis*, Wiley-Blackwell, 2009, ch. 9, pp. 387–454.
- 4 P. Ravenscroft, H. Brammer and K. Richards, Arsenic in South and Central America, Africa, Australasia and Oceania, in *Arsenic Pollution: A Global Synthesis*, Wiley-Blackwell, 2009, ch. 10, pp. 455–491.
- 5 P. Sabbatini, F. Yrazu, F. Rossi, G. Thern, A. Marajofsky and M. M. Fidalgo de Cortalezzi, Fabrication and characterization of iron oxide ceramic membranes for arsenic removal, *Water Res.*, 2010, **44**, 5702–5712.
- 6 P. Ravenscroft, H. Brammer and K. Richards, Health Effects of Arsenic in Drinking Water and Food, in *Arsenic Pollution: A Global Synthesis*, Wiley-Blackwell, 2009, ch. 5, pp. 157–212.
- 7 R. Quansah, F. A. Armah, D. K. Essumang, I. Luginaah, E. Clarke, K. Marfoh, S. J. Cobbina, E. Nketiah-Amponsah, P. B. Namujju, S. Obiri and M. Dzodzomenyo, Association of arsenic with adverse pregnancy outcomes/infant mortality: a systematic review and meta-analysis, *Environ. Health Perspect.*, 2015, **123**, 412–421.
- 8 S. F. Farzan, M. R. Karagas and Y. Chen, In utero and early life arsenic exposure in relation to long-term health and disease, *Toxicol. Appl. Pharmacol.*, 2013, **272**, 384–390.
- 9 M. Tolins, M. Ruchirawat and P. Landrigan, The developmental neurotoxicity of arsenic: cognitive and behavioral consequences of early life exposure, *Ann. Glob. Health*, 2014, **80**, 303–314.
- 10 K. Nagaraj, A. N. Shetty and D. R. Trivedi, Development of multi-analyte responsive sensors: optical discrimination of arsenite and arsenate ions, ratiometric detection of arsenite, and application in food and water samples, *Sens. Diagn.*, 2024, **3**, 64–78.
- 11 P. Ravenscroft, H. Brammer and K. Richards, Removing Arsenic from Drinking Water, in *Arsenic Pollution: A Global Synthesis*, Wiley-Blackwell, 2009, ch. 7, pp. 261–317.
- 12 R. A. Root, S. Dixit, K. M. Campbell, A. D. Jew, J. G. Hering and P. A. O'Day, Arsenic sequestration by sorption processes in high-iron sediments, *Geochim. Cosmochim. Acta*, 2007, **71**, 5782–5803.
- 13 K. M. Campbell and D. K. Nordstrom, Arsenic Speciation and Sorption in Natural Environments, *Rev. Mineral. Geochem.*, 2014, **79**, 185–216.
- 14 F. Maia, C. Pinto, J. C. Waerenborgh, M. A. Gonçalves, C. Prazeres, O. Carreira and S. Sério, Metal partitioning in sediments and mineralogical controls on the acid mine drainage in Ribeira da Água Forte (Aljustrel, Iberian Pyrite Belt, Southern Portugal), *Appl. Geochem.*, 2012, **27**, 1063–1080.
- 15 Z. Li, S. Deng, G. Yu, J. Huang and V. C. Lim, As(V) and As(III) removal from water by a Ce-Ti oxide adsorbent:



- behavior and mechanism, *Chem. Eng. J.*, 2010, **161**, 106–113.
- 16 R. Li, Q. Li, S. Gao and J. K. Shang, Exceptional arsenic adsorption performance of hydrous cerium oxide nanoparticles: Part A. Adsorption capacity and mechanism, *Chem. Eng. J.*, 2012, **185–186**, 127–135.
- 17 P. P. Mon, P. P. Cho, D. Vidyasagar, P. Ghosal, G. Madras and S. Challapalli, Synergistic sorption: Enhancing arsenic (V) removal using biochar decorated with cerium oxide composite, *Mater. Today Sustain.*, 2024, **25**, 100675.
- 18 T. Basu, D. Nandi, P. Sen and U. C. Ghosh, Equilibrium modeling of As(III,V) sorption in the absence/presence of some groundwater occurring ions by iron(III)-cerium(IV) oxide nanoparticle agglomerates: A mechanistic approach of surface interaction, *Chem. Eng. J.*, 2013, **228**, 665–678.
- 19 F. Chai, R. Wang, L. Yan, G. Li, Y. Cai and C. Xi, Facile fabrication of pH-sensitive nanoparticles based on nanocellulose for fast and efficient As(V) removal, *Carbohydr. Polym.*, 2020, **245**, 116511.
- 20 R. Sawana, Y. Somasundar, V. S. Iyer and B. Baruwati, Ceria modified activated carbon: an efficient arsenic removal adsorbent for drinking water purification, *Appl. Water Sci.*, 2017, **7**, 1223–1230.
- 21 N. K. Devaraj, A. S. M. Mukter-Uz-Zaman and H. Y. Wong, Ceria-Coated Magnetite Nanoparticles for As(V) Removal from Water, presented in part at the *Recent Advances in Environmental Science from the Euro-Mediterranean and Surrounding Regions*, Cham, 4th dn, 2024.
- 22 Q. Feng, Z. Zhang, Y. Ma, X. He, Y. Zhao and Z. Chai, Adsorption and desorption characteristics of arsenic onto ceria nanoparticles, *Nanoscale Res. Lett.*, 2012, **7**, 84.
- 23 X. Bi, C. Zeng and P. Westerhoff, Adsorption of Arsenic Ions Transforms Surface Reactivity of Engineered Cerium Oxide Nanoparticles, *Environ. Sci. Technol.*, 2020, **54**, 9437–9444.
- 24 D. C. Sayle, F. Caddeo, L. M. Morgan, R. L. Neale, T. X. T. Sayle, C. Brambila, J. Nutter, U. Bhatta, K. M. Ta, J. M. Flitcroft, T. S. Sakthivel, S. Seal, G. Möbus and M. Molinari, Aging mechanisms of nanoceria and pathways for preserving optimum morphology, *Nano Today*, 2023, **51**, 101916.
- 25 K. M. Ta, D. J. Cooke, L. J. Gillie, S. C. Parker, S. Seal, P. B. Wilson, R. M. Phillips, J. M. Skelton and M. Molinari, Infrared and Raman Diagnostic Modeling of Phosphate Adsorption on Ceria Nanoparticles, *J. Phys. Chem. C*, 2023, **127**, 20183–20193.
- 26 F. Wei, C. J. Neal, T. S. Sakthivel, Y. Fu, M. Omer, A. Adhikary, S. Ward, K. M. Ta, S. Moxon, M. Molinari, J. Asiatico, M. Kinzel, S. N. Yarmolenko, V. San Cheong, N. Orlovskaya, R. Ghosh, S. Seal and M. Coathup, A novel approach for the prevention of ionizing radiation-induced bone loss using a designer multifunctional cerium oxide nanozyme, *Bioact. Mater.*, 2023, **21**, 547–565.
- 27 M. Molinari, A. R. Symington, D. C. Sayle, T. S. Sakthivel, S. Seal and S. C. Parker, Computer-Aided Design of Nanoceria Structures as Enzyme Mimetic Agents: The Role of Bodily Electrolytes on Maximizing Their Activity, *ACS Appl. Bio Mater.*, 2019, **2**, 1098–1106.
- 28 Y. Fu, C. J. Neal, E. Kolanthai, S. Munir, K. M. Ta, M. Molinari and S. Seal, Effect of acetate ions and pH on the morphology of cerium oxide nanoparticles, *Colloids Surf., A*, 2023, **679**, 132616.
- 29 Y. Zhang, M. Yang and X. Huang, Arsenic(V) removal with a Ce(IV)-doped iron oxide adsorbent, *Chemosphere*, 2003, **51**, 945–952.
- 30 S. Dudek and D. Kołodyńska, Arsenic(V) removal on the lanthanum-modified ion exchanger with quaternary ammonium groups based on iron oxide, *J. Mol. Liq.*, 2022, **347**, 117985.
- 31 Y. Wang, G. Morin, G. Ona-Nguema, F. Juillot, F. Guyot, G. Calas and G. E. Brown, Jr., Evidence for Different Surface Speciation of Arsenite and Arsenate on Green Rust: An EXAFS and XANES Study, *Environ. Sci. Technol.*, 2010, **44**, 109–115.
- 32 A. L. Foster and C. S. Kim, Arsenic Speciation in Solids Using X-ray Absorption Spectroscopy, *Rev. Mineral. Geochem.*, 2014, **79**, 257–369.
- 33 D. M. Sherman and S. R. Randall, Surface complexation of arsenic(V) to iron(III) (hydr)oxides: structural mechanism from ab initio molecular geometries and EXAFS spectroscopy, *Geochim. Cosmochim. Acta*, 2003, **67**, 4223–4230.
- 34 K. Fukushi and D. A. Sverjensky, A predictive model (ETLM) for arsenate adsorption and surface speciation on oxides consistent with spectroscopic and theoretical molecular evidence, *Geochim. Cosmochim. Acta*, 2007, **71**, 3717–3745.
- 35 R. J. Martínez, M. Villalobos, A. U. Loredó-Jasso, A. X. Cruz-Valladares, A. Mendoza-Flores, H. Salazar-Rivera and D. Cruz-Romero, Towards building a unified adsorption model for goethite based on direct measurements of crystal face compositions: I. Acidity behavior and As(V) adsorption, *Geochim. Cosmochim. Acta*, 2023, **354**, 252–262.
- 36 I. Römer, S. M. Briffa, Y. Arroyo Rojas Dasilva, D. Hapiuk, V. Trouillet, R. E. Palmer and E. Valsami-Jones, Impact of particle size, oxidation state and capping agent of different cerium dioxide nanoparticles on the phosphate-induced transformations at different pH and concentration, *PLoS One*, 2019, **14**, e0217483.
- 37 H. Liu and J. Liu, Self-limited Phosphatase-mimicking CeO₂ Nanozymes, *ChemNanoMat*, 2020, **6**, 947–952.
- 38 G. Kresse and J. Furthmüller, Efficient iterative schemes for ab initio total-energy calculations using a plane-wave basis set, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1996, **54**, 11169–11186.
- 39 G. Kresse and J. Hafner, Ab initio molecular dynamics for liquid metals, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1993, **47**, 558–561.
- 40 G. Kresse and J. Furthmüller, Efficiency of ab-initio total energy calculations for metals and semiconductors using a plane-wave basis set, *Comput. Mater. Sci.*, 1996, **6**, 15–50.
- 41 J. P. Perdew, J. A. Chevary, S. H. Vosko, K. A. Jackson, M. R. Pederson, D. J. Singh and C. Fiolhais, Erratum: Atoms, molecules, solids, and surfaces: Applications of the



- generalized gradient approximation for exchange and correlation, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1993, **48**, 4978–4978.
- 42 J. P. Perdew, K. Burke and M. Ernzerhof, Generalized Gradient Approximation Made Simple, *Phys. Rev. Lett.*, 1996, **77**, 3865–3868.
- 43 J. Heyd, G. E. Scuseria and M. Ernzerhof, Hybrid functionals based on a screened Coulomb potential, *J. Chem. Phys.*, 2003, **118**, 8207–8215.
- 44 J. Heyd, G. Scuseria and M. Ernzerhof, Erratum: “Hybrid Functionals Based on a Screened Coulomb Potential” [*J. Chem. Phys.* 118, 8207 (2003)], *J. Chem. Phys.*, 2006, **124**, 9906.
- 45 S. L. Dudarev, G. A. Botton, S. Y. Savrasov, C. J. Humphreys and A. P. Sutton, Electron-energy-loss spectra and the structural stability of nickel oxide: An LSDA+U study, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1998, **57**, 1505–1509.
- 46 U. Castanet, C. Feral-Martin, A. Demourgues, R. L. Neale, D. C. Sayle, F. Caddeo, J. M. Flitcroft, R. Caygill, B. J. Pointon, M. Molinari and J. Majimel, Controlling the {111}/ {110} Surface Ratio of Cuboidal Ceria Nanoparticles, *ACS Appl. Mater. Interfaces*, 2019, **11**, 11384–11390.
- 47 A. S. Pugazhendhi, C. J. Neal, K. M. Ta, M. Molinari, U. Kumar, F. Wei, E. Kolanthai, A. Ady, C. Drake, M. Hughes, S. Yooseph, S. Seal and M. J. Coathup, A neoteric antibacterial ceria-silver nanozyme for abiotic surfaces, *Biomaterials*, 2024, **307**, 122527.
- 48 M. Molinari, S. C. Parker, D. C. Sayle and M. S. Islam, Water Adsorption and Its Effect on the Stability of Low Index Stoichiometric and Reduced Surfaces of Ceria, *J. Phys. Chem. C*, 2012, **116**, 7073–7082.
- 49 S. Moxon, A. R. Symington, J. S. Tse, J. M. Flitcroft, J. M. Skelton, L. J. Gillie, D. J. Cooke, S. C. Parker and M. Molinari, Composition-dependent morphologies of CeO₂ nanoparticles in the presence of Co-adsorbed H₂O and CO₂: a density functional theory study, *Nanoscale*, 2024, **16**, 11232–11249.
- 50 T. Smith, S. Moxon, J. S. Tse, J. M. Skelton, D. J. Cooke, L. J. Gillie, E. L. da Silva, R. M. Harker, M. T. Storr, S. C. Parker and M. Molinari, Structural dynamics of Schottky and Frenkel defects in CeO₂: a density-functional theory study, *JPhys Energy*, 2023, **5**, 025004.
- 51 A. R. Symington, R. M. Harker, M. T. Storr, M. Molinari and S. C. Parker, Thermodynamic Evolution of Cerium Oxide Nanoparticle Morphology Using Carbon Dioxide, *J. Phys. Chem. C*, 2020, **124**, 23210–23220.
- 52 A. R. Symington, M. Molinari, S. Moxon, J. M. Flitcroft, D. C. Sayle and S. C. Parker, Strongly Bound Surface Water Affects the Shape Evolution of Cerium Oxide Nanoparticles, *J. Phys. Chem. C*, 2020, **124**, 3577–3588.
- 53 D. Prieur, W. Bonani, K. Popa, O. Walter, K. W. Kriegsman, M. H. Engelhard, X. Guo, R. Eloirdi, T. Gouder, A. Beck, T. Vitova, A. C. Scheinost, K. Kvashnina and P. Martin, Size Dependence of Lattice Parameter and Electronic Structure in CeO₂ Nanoparticles, *Inorg. Chem.*, 2020, **59**, 5760–5767.
- 54 G. W. Watson, E. T. Kelsey, N. H. de Leeuw, D. J. Harris and S. C. Parker, Atomistic simulation of dislocations, surfaces and interfaces in MgO, *J. Chem. Soc., Faraday Trans.*, 1996, **92**, 433–438.
- 55 K. Momma and F. Izumi, VESTA 3 for three-dimensional visualization of crystal, volumetric and morphology data, *J. Appl. Crystallogr.*, 2011, **44**, 1272–1276.
- 56 J. M. Skelton, L. A. Burton, A. J. Jackson, F. Oba, S. C. Parker and A. Walsh, Lattice dynamics of the tin sulphides SnS₂, SnS and Sn₂S₃: vibrational spectra and thermal transport, *Phys. Chem. Chem. Phys.*, 2017, **19**, 12452–12465.
- 57 A. Togo and I. Tanaka, First principles phonon calculations in materials science, *Scr. Mater.*, 2015, **108**, 1–5.
- 58 M. Gajdoš, K. Hummer, G. Kresse, J. Furthmüller and F. Bechstedt, Linear optical properties in the projector-augmented wave methodology, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2006, **73**, 045112.
- 59 C. Schilling, A. Hofmann, C. Hess and M. V. Ganduglia-Pirovano, Raman Spectra of Polycrystalline CeO₂: A Density Functional Theory Study, *J. Phys. Chem. C*, 2017, **121**, 20834–20849.
- 60 W. Humphrey, A. Dalke and K. Schulten, VMD: visual molecular dynamics, *J. Mol. Graphics*, 1996, **14**(33–38), 27–38.
- 61 A. C. Q. Ladeira, V. S. T. Ciminelli, H. A. Duarte, M. C. M. Alves and A. Y. Ramos, Mechanism of anion retention from EXAFS and density functional calculations: arsenic (V) adsorbed on gibbsite, *Geochim. Cosmochim. Acta*, 2001, **65**, 1211–1217.
- 62 A. Pranudta, S. Patra, W. Klysubun, P. Amonpattaratkit, T. Thanh Nguyen, V. Hong Nguyen, M. Mohamed El-Moselhy and S. Padungthon, Insight into the molecular interaction of trace As (III) and As(V) onto the hybrid anion exchanger impregnated with Fe/Mn nanoparticles (HA502P-Fe/Mn), *Chem. Eng. J.*, 2023, **454**, 139991.
- 63 X. Dou, Y. Zhang, B. Zhao, X. Wu, Z. Wu and M. Yang, Arsenate adsorption on an Fe–Ce bimetal oxide adsorbent: EXAFS study and surface complexation modeling, *Colloids Surf., A*, 2011, **379**, 109–115.
- 64 D. R. Mullins, The surface chemistry of cerium oxide, *Surf. Sci. Rep.*, 2015, **70**, 42–85.
- 65 T. X. Sayle, M. Molinari, S. Das, U. M. Bhatta, G. Möbus, S. C. Parker, S. Seal and D. C. Sayle, Environment-mediated Structure, Surface Redox Activity and Reactivity of Ceria Nanoparticles, *Nanoscale*, 2013, **5**, 6063–6073.
- 66 S. Mochizuki, Infrared optical properties of cerium dioxide, *Phys. Status Solidi B*, 1982, **114**, 189–199.
- 67 A. Filtschew, K. Hofmann and C. Hess, Ceria and Its Defect Structure: New Insights from a Combined Spectroscopic Approach, *J. Phys. Chem. C*, 2016, **120**, 6694–6703.
- 68 C. C. Pye, T. Chiome and B. L. Goodall, An ab initio study of the effect of hydration on the vibrational spectrum of hydrogen arsenate ion, *Comput. Theor. Chem.*, 2022, **1215**, 113838.
- 69 L. A. Sheludyakova, V. Afanasieva, N. V. Podberezskaya and Y. I. Mironov, Spectral and structural analysis of sodium hydrophosphates and hydroarsenates, *J. Struct. Chem.*, 1999, **40**, 869–872.
- 70 S. Goldberg and C. T. Johnston, Mechanisms of Arsenic Adsorption on Amorphous Oxides Evaluated Using



- Macroscopic Measurements, Vibrational Spectroscopy, and Surface Complexation Modeling, *J. Colloid Interface Sci.*, 2001, **234**, 204–216.
- 71 S. Bhasker-Ranganath, C. Zhao and Y. Xu, Theoretical analysis of the adsorption of phosphoric acid and model phosphate monoesters on CeO₂(111), *Surf. Sci.*, 2021, **705**, 121776.
- 72 F. K. Vansant, B. J. Van Der Veken and H. O. Desseyn, Vibrational analysis of arsenic acid and its anions: I. Description of the Raman spectra, *J. Mol. Struct.*, 1973, **15**, 425–437.
- 73 M. Kanematsu, T. M. Young, K. Fukushi, P. G. Green and J. L. Darby, Arsenic(III, V) adsorption on a goethite-based adsorbent in the presence of major co-existing ions: Modeling competitive adsorption consistent with spectroscopic and molecular evidence, *Geochim. Cosmochim. Acta*, 2013, **106**, 404–428.

