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# Band vs. polaron: vibrational motion and chemical expansion of hydride ions as signatures for the electronic character in oxyhydride barium titanate

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The oxyhydride phase of barium titanate,  $\text{BaTiO}_{3-x}\text{H}_x$ , is a mixed hydride ion and electron conductor. The substitution of oxygen with hydrogen to form a hydride ion is accompanied by donation of an electron to the initially empty titanium 3d conduction band. It is not clear, however, whether the electron forms a delocalized state where it is shared among all titanium ions forming a bandstate, or if it localizes on a titanium ion and forms a bound electron polaron. Here, we investigate polaron formation in this material using density-functional theory (DFT) calculations, where the self-interaction error has been corrected by the DFT +  $U$  method and the HSE hybrid functional. While calculated formation energies do not provide a conclusive description of the electronic state, a comparison of the results from first-principles phonon calculations with vibrational spectra measured with inelastic neutron scattering (INS) suggests that the electrons form bandstates in bulk  $\text{BaTiO}_{3-x}\text{H}_x$ . This is further supported by comparison of the computed chemical expansion of the involved defect species with experimental data of the lattice expansion in the oxyhydride formation. The oxyhydride phase of barium titanate,  $\text{BaTiO}_{3-x}\text{H}_x$ , should thus exhibit metallic-like conductivity.

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## 1 Introduction

Oxyhydrides, where a hydride ion ( $\text{H}^-$ ) is substituted on an oxygen site, are rare in nature and the oxyhydride phases ( $\text{Ca,Sr,BaTiO}_{3-x}\text{H}_x$ ), which can form under strongly reducing conditions,<sup>1,2</sup> were the first ones discovered based on the simple perovskite structure  $\text{ABO}_3$ . Among them,  $\text{BaTiO}_{3-x}\text{H}_x$  exhibits the largest number of hydride ions with a concentration up to about  $x = 0.6$  and the material exhibits both hydride ion mobility and electronic conductivity.<sup>1,3</sup> Understanding the transport is of great importance in developing electrochemical devices but neither the ionic or the electronic transport mechanisms are fully understood.<sup>4</sup> Here we focus on the electronic part.

The substitutional hydride ion ( $\text{H}_\text{O}^\times$  in Kröger–Vink notation) acts as a donor, contributing to n-type conductivity in the initially empty titanium 3d band,<sup>5–7</sup> and measurements confirm that  $\text{BaTiO}_{3-x}\text{H}_x$  is electrically conducting.<sup>1,8,9</sup> Kobayashi *et al.*<sup>1</sup> reported a semiconducting behaviour for the bulk phase of  $\text{BaTiO}_{3-x}\text{H}_x$  (with  $x = 0.3$  and  $x = 0.6$ ). However, as the

conductivity measurement was performed on a powder sample pressed into a pellet without sintering, the reported values may not reflect the true bulk properties of the material.<sup>1</sup> More recent studies on epitaxial thin films showed that at high hydride concentrations ( $x \geq 0.2$ )  $\text{BaTiO}_{3-x}\text{H}_x$  exhibits metallic-like conductivity, while at lower concentrations ( $x \leq 0.2$ ) the conductivity is semiconductor-like.<sup>9</sup>

Based on first-principles calculations it has been suggested that electron polarons, which form localized electronic states in the band gap, are responsible for the experimentally observed semiconductor-like conductivity in bulk  $\text{BaTiO}_{3-x}\text{H}_x$ .<sup>7</sup> A polaron is a quasiparticle consisting of a charge localization in association with a local lattice distortion. In the present case this refers to a localized electron on a titanium ion that resides next to a hydride ion, thereby changing the valence of the titanium ion from +4 to +3. The concomitant distortion of the lattice influences the local atomic environment and thereby the vibrational properties of the hydride ion. In pristine  $\text{BaTiO}_3$ , (self-trapped) electron polarons have been found to be unstable,<sup>7</sup> but bound polarons have been discussed in connection to n-type doping such as  $\text{Nb}^{5+}$  substitution<sup>10,11</sup> and oxygen vacancies.<sup>12,13</sup> The question naturally arises whether the  $\text{H}^-$  substitution also can form bound polarons in  $\text{BaTiO}_3$ . In addition, electron polarons have been found in  $\text{TiO}_2$ ,<sup>14–19</sup> which, similar to  $\text{BaTiO}_3$ , has valence and conduction bands that are dominated by oxygen 2p states and titanium 3d states, respectively.

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The aim of the present study is to investigate the possibility of polaron formation in bulk  $\text{BaTiO}_{3-x}\text{H}_x$ . We use two complementary techniques: first-principles calculations based on density-functional theory (DFT) and inelastic neutron scattering (INS) experiments. We focus on: (i) the formation energy of the polaron, (ii) the chemical expansion of the lattice due to the polaronic defect, and (iii) the effect of the polaron on the hydride ion vibrational motion.

Modelling of polarons using DFT requires additional considerations due to the self-interaction error of the standard local and semi-local exchange–correlation (XC) functionals, which favour charge delocalization. The self-interaction error can be remedied by using the DFT +  $U$  method,<sup>20</sup> where the  $U$ -parameter should correct for the self-interaction error. Here we use DFT +  $U$  to study the polaronic distortion and the associated formation energy. To obtain an appropriate value of the  $U$ -parameter we determine  $U$  by restoring the piecewise linearity of the total energy as a function of fractional occupancy of the polaron level, a known property of the exact density functional.<sup>21–23</sup>

Inelastic neutron scattering (INS) is used to study the vibrational motion of the hydride ion. This experimental technique provides a straightforward way to investigate the vibrational properties of hydride ions because of the large neutron cross section of hydrogen relative to the other atomic species in  $\text{BaTiO}_{3-x}\text{H}_x$ . The vibrational spectrum is also computed and to obtain accurate frequencies we make use of the hybrid functional HSE,<sup>24,25</sup> which is known to reproduce the experimental frequencies for  $\text{BaTiO}_3$  accurately.<sup>26</sup>

We find that the electrons in bulk  $\text{BaTiO}_{3-x}\text{H}_x$  form delocalized bandstates, in contrast to ref. 7, and thus bulk  $\text{BaTiO}_{3-x}\text{H}_x$  should exhibit metallic-like conductivity.

## 2 Theoretical approach

### 2.1 Computational details

Density-functional theory (DFT) calculations were performed using the projector augmented wave (PAW) method<sup>27,28</sup> as implemented in the Vienna *ab initio* simulation package (VASP).<sup>29</sup> The semi-local functional PBE<sup>30</sup> and the hybrid functional HSE<sup>24</sup> were used to model exchange and correlation (XC) effects. For the latter we consider the conventional mixing and screening ( $\alpha = 0.25$  and  $\omega = 0.2 \text{ \AA}^{-1}$ ), which is often referred to as HSE06.<sup>25</sup> The DFT +  $U$  method was considered in its rotationally invariant form,<sup>31</sup> where the  $U$ -parameter was applied to the Ti-3d orbitals. The choice of  $U$  was based on the concept of piecewise linearity<sup>23,32</sup> and is further described in Section 2.3.

Unless otherwise stated, calculations were carried out using supercells containing  $2 \times 2 \times 2$  unit cells (40 atoms). For this supercell size the sampling of the Brillouin zone was performed using  $6 \times 6 \times 6$  Monkhorst–Pack grids for PBE and PBE +  $U$  calculations while  $4 \times 4 \times 4$  grids were used for HSE.  $k$ -point meshes for calculations with larger supercells were reduced accordingly. The rather high  $k$ -point density is necessary in order to accurately describe the delocalized electronic states. All calculations were performed spin polarized using a Gaussian

smearing of width  $\sigma = 10^{-5} \text{ eV}$  in order to properly populate the localized polaronic defect level. A plane wave cut-off energy of 500 eV was used. The calculations were converged to energies within  $10^{-7} \text{ eV}$  for the electronic structure and the ionic relaxations to forces within  $10^{-4} \text{ eV \AA}^{-1}$ .

### 2.2 Pristine $\text{BaTiO}_3$

In order to assess the validity of our calculations we begin by investigating the bulk properties of the cubic phase of pristine  $\text{BaTiO}_3$  using PBE, PBE +  $U$  ( $U = 3.3 \text{ eV}$ , see Section 2.3 for details) and HSE. The results of PBE and HSE (Table 1) agree very well with the previous calculations in ref. 26. Relative to PBE, PBE +  $U$  predicts both a slightly larger lattice constant and band gap. A more striking difference between the two methods is connected to the unstable vibrational mode, where the magnitude of the imaginary frequency is reduced substantially with PBE +  $U$ . The smaller magnitude indicates that the addition of a  $U$ -parameter increases the stability of the cubic phase, although the tetragonal phase remains favorable with the present choice of  $U$ . The real frequencies, on the other hand, are very similar for PBE and PBE +  $U$ , which shows that the addition of  $U$  does not alter the vibrational properties of pristine  $\text{BaTiO}_3$  in any significant way. PBE +  $U$  can therefore be expected to perform similarly to PBE also when defects are introduced.

When compared with experimental data<sup>35</sup> we notice that PBE and PBE +  $U$  underestimate the vibrational frequencies while HSE gives excellent results (see Table 1). The PBE error is partly due to the overestimated lattice constant but the improved performance of HSE also depends on the inclusion of exact exchange, which produces a more accurate electron density that leads to stiffening of the bonds.<sup>26,36,37</sup> The  $T_{1u}$ -modes all contain a component of stretching character in the Ti–O bonds, which make these modes pressure dependent. Wahl *et al.* showed that using a semi-local XC functional (PBEsol) with a more accurate lattice constant increases the frequencies by about 3–5% and inclusion of exact exchange (HSE) further increases the frequencies by 3–4%.<sup>26</sup> The  $T_{2u}$ -mode has more of a bending character, which makes it rather insensitive to variations of the lattice constant.<sup>38</sup> The inclusion of exact exchange, however, increases the vibrational frequencies of this mode by about 8–9%. To summarize, the use of HSE

Table 1 Lattice constant ( $\text{\AA}$ ), band gap (eV) and  $\Gamma$ -point vibrational frequencies ( $\text{cm}^{-1}$ ) for pristine  $\text{BaTiO}_3$

	This work			Wahl <i>et al.</i> <sup>26</sup>		
	PBE	PBE + $U^a$	HSE <sup>b</sup>	PBE	HSE <sup>c</sup>	Exp.
$a_0$	4.032	4.052	3.989	4.035	3.995	3.9975 (ref. 33)
$E_g$	1.71	2.12	3.10	1.70	2.92	3.22 (ref. 34)
$T_{1u}(\text{TO1})$	247i	57i	257i	239i	241i	Soft (ref. 35)
$T_{1u}(\text{TO2})$	170	167	185	169	185	181 (ref. 35)
$T_{2u}$	286	283	312	286	310	306 (ref. 35)
$T_{1u}(\text{TO3})$	452	461	478	453	480	487 (ref. 35)

<sup>a</sup>  $U = 3.3 \text{ eV}$ . <sup>b</sup>  $\alpha = 0.25$  and  $\omega = 0.2 \text{ \AA}^{-1}$ . <sup>c</sup>  $\alpha = 0.25$  and  $\omega = 0.3 \text{ \AA}^{-1}$ .



improves on the PBE description substantially as it increases the frequencies by about 5–10% and enables a very accurate description of the vibrational modes compared with experimental data (Table 1).<sup>26</sup>

### 2.3 Polaron calculations

The choice of  $U$  in the DFT +  $U$  method is of significant importance. The method is a parametric correction of standard DFT and the value of  $U$  is not transferable but will depend on the physical quantity under consideration as well as the underlying functional. Here we determine the value of  $U$  by enforcing the piecewise linearity constraint, which is a known property of the exact XC functional.<sup>21</sup> We follow the same approach as in ref. 23 and 32, and tune  $U$  so that the energy depends linearly on the charge during the population of the polaronic level.

The addition of a  $U$ -parameter on Ti introduces a pressure in the system, which increases with increasing  $U$ . Consequently, for each value of  $U$  a different equilibrium lattice constant is obtained. We chose to release this pressure by relaxing the lattice constant for each considered  $U$ -value.

The lattice distortion consistent with polaron formation cannot be known *a priori* and we chose to prepare an archetype polaron configuration by displacing the atoms randomly and performing ionic relaxations in the confinement of a cubic supercell at a large value of  $U$ . The obtained structure was confirmed to be stable through a phonon calculation, which displayed no imaginary modes. The ionic positions were then further relaxed within a cubic supercell for the respective  $U$  values at the corresponding equilibrium lattice constants.

Fig. 1 shows the deviation from a piecewise linear behaviour for four different values of  $U$  together with PBE and HSE for a polaronic  $\text{H}_2^{\times}$  configuration in a  $2 \times 2 \times 2$  supercell. As expected, PBE yields a convex behaviour consistent with delocalization, while the HSE calculation results in a slightly concave behaviour consistent with over-localization. The PBE +  $U$  method, with the appropriate  $U$ , performs reasonably well although a slight sinusoidal shape can be seen. The residual, defined as the root mean square of the deviation from piecewise linearity, was minimized at a value of  $U = 3.3$  eV, for which the lattice constant is 4.0520 Å. The value of  $U$  is not completely transferable between different supercell sizes and for a  $3 \times 3 \times 3$  supercell a smaller value of  $U$  was found to yield piecewise linearity. The residual for this system was minimized at a value of  $U = 3.1$  eV with the corresponding lattice constant of 4.0507 Å.

In a previous theoretical study of  $\text{BaTiO}_{3-x}\text{H}_x$  where polaron formation was found to be favourable, the PBE +  $U$  method with  $U = 4.49$  eV was used.<sup>7</sup> That value was determined for the LSDA functional.<sup>6</sup> However, the value of  $U$  for correcting the self-interaction error is not transferable between different XC functionals, different projection radii (PAW potentials)<sup>39</sup> and, as shown here, different supercell sizes. As indicated by Fig. 1, such a high value of  $U$  would lead to over-localization and erroneously favor polaron formation in these systems.

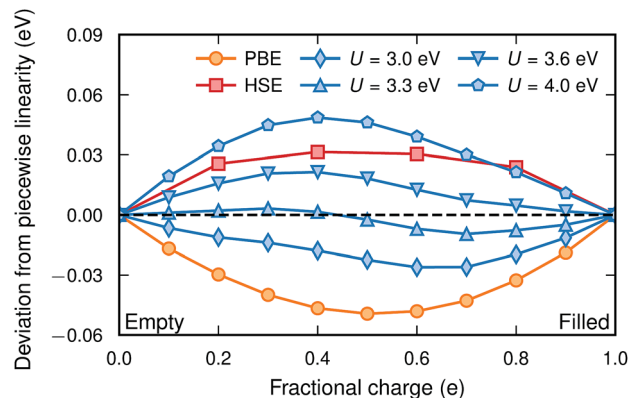


Fig. 1 Deviation from the piecewise linearity of the energy filling up the polaron level for a polaronic  $\text{H}_2^{\times}$  configuration in a  $2 \times 2 \times 2$  supercell. The yellow line shows how PBE delocalizes while the red line shows how HSE over-localizes the electron. The blue lines show how the PBE +  $U$  method can be tuned to produce a behaviour closer to linear behaviour.

### 2.4 Vibrational calculations

Vibrational calculations were conducted following two different approaches. The full phonon spectra were calculated within the harmonic approximation using the frozen phonon approach with finite displacements and phonopy<sup>40</sup> software. Forces were computed with a  $2 \times 2 \times 2$  augmented supercell consisting of 320 atoms for the defect-containing configurations, which allows for a zone folding scheme with interpolation between high symmetry points. Due to the computational cost the electronic  $k$ -point mesh was reduced to  $2 \times 2 \times 2$  for the augmented supercell. In computing the vibrational density of states, a Monkhorst-Pack  $q$ -point mesh of  $10 \times 10 \times 10$  was used.

When using the computationally more costly HSE functional, a simplified approach was used for determining the hydride ion frequencies. These oscillations are very localized (which was confirmed by the full phonon calculations, see Fig. 4) and can be well described by a 3D oscillator in a potential well provided by the relaxed positions of the surrounding atoms. The 3D potential is mapped out through displacements of the hydride ion by  $\pm 0.1$  Å and  $\pm 0.2$  Å along the three different directions, where the energy is calculated at each displacement. Second order polynomials are then used to fit the energy landscape and to obtain the vibrational frequencies. We denote this approach as the *one particle harmonic potential* (OPHP) method, and its validity is demonstrated in Section 4.4.1.

## 3 Materials and experimental technique

### 3.1 Sample preparation

The oxyhydride powder sample was prepared by mixing  $\text{CaH}_2$  powder with  $\text{BaTiO}_3$  powder and subsequently pelletizing inside an Ar filled glove box. The pellet was placed in a custom made stainless steel autoclave and heated inside a tube furnace to a temperature of 700 °C for a period of three days. Finally, the pellet was washed and subsequently dried under dynamic



vacuum at 120 °C for about 1 day. The details of the sample preparation can be found in ref. 41 and the present work makes use of the sample denoted as CA1 in ref. 41.

Powder X-ray diffraction (PXRD) measurements confirmed a simple cubic structure for the sample with lattice parameter  $a = 4.0055$  Å at ambient temperature.<sup>41</sup> Thermogravimetric analysis (TGA) of the sample showed that the total defect concentration  $x$ , as in  $\text{BaTiO}_{3-x}\text{H}_y\Box_{x-y}$ , was  $x = 0.18$ ,<sup>41</sup> where  $\Box$  denotes oxygen vacancies in the oxyhydride. The hydride ion concentration was determined to be  $y = 0.10$  using nuclear magnetic resonance (NMR) spectroscopy,<sup>41</sup> which implies an oxygen vacancy concentration of  $(x - y) = 0.08$ .

### 3.2 Inelastic neutron scattering

The INS experiment was performed on TOSCA at the ISIS Facility,<sup>42</sup> with the sample loaded into a aluminum sachet and the sachet into indium wire sealed thin-walled aluminum can. The measurement was performed at 10 K with a measuring time of approximately 6 h per sample. The INS spectra were analyzed with Mantid<sup>43</sup> and DAVE<sup>44</sup> software.

To theoretically analyze the spectrum from the multi-component poly-crystalline sample the incoherent approximation was employed.<sup>45</sup> The emission part of the one-phonon contribution to the INS law can then be written as<sup>46</sup>

$$S(Q, \omega) = \sum_v \sigma_v \frac{\hbar Q^2}{2m_v} \exp[-2W_v(Q)] \frac{g_v(\omega)}{\omega} [1 + n(\omega)], \quad (1)$$

where  $\sigma_v$  and  $m_v$  are the total neutron scattering cross section and atomic mass of the  $v$ th atom, and  $n(\omega) = [\exp(\hbar\omega/k_B T) - 1]^{-1}$  is the Bose population factor. The Debye–Waller factor,  $\exp[-2W_v(Q)]$ , is related to the mean square displacement  $\langle u_v^2 \rangle$  according to  $2W_v(Q) = (1/3)Q^2 \langle u_v^2 \rangle$ , and

$$g_v(\omega) = \frac{1}{3N} \sum_{\mathbf{q}} \left| \mathbf{e}_{\mathbf{q}j}^v \right|^2 \delta(\omega - \omega_{\mathbf{q}j}) \quad (2)$$

is the partial phonon density of states, where  $N$  is the number of atoms in the simulation cell,  $\omega_{\mathbf{q}j}$  is the eigenfrequency for phonon mode  $j$  with wavevector  $\mathbf{q}$  and  $\mathbf{e}_{\mathbf{q}j}^v$  is the corresponding eigenvector projected onto atomic species  $v$ .<sup>40</sup>

A key aspect of TOSCA<sup>42</sup> is that the momentum of the scattered neutrons is very small. The momentum transfer  $\hbar Q$  and the energy transfer  $\hbar\omega$  in the scattering event are therefore related approximately as  $(\hbar^2 Q^2)/(2m_n) = \hbar\omega$ , where  $m_n$  is the neutron mass.<sup>47</sup> The measured intensity  $S(Q, \omega)$  at low temperatures can then be written as

$$S(\omega) = \sum_v \sigma_v \frac{m_n}{m_v} g_v(\omega) \exp \left[ -\frac{2m_n \omega}{3\hbar} \langle u_v^2 \rangle \right], \quad (3)$$

where  $\langle u_v^2 \rangle$  is evaluated at  $T = 0$  K. For the total neutron scattering cross sections  $\sigma_v$  of H, Ba, Ti, and O, we use 82.0, 3.38, 4.35, and 4.23 barn, respectively.<sup>48</sup>

## 4 Results and discussion

The substitutional hydrogen on an oxygen site ( $\text{H}_\text{O}^\times$ ) acts as a donor contributing one electron to the empty Ti 3d band. Here

we investigate two possible states for this electron. The first option is the delocalized state, where the electron is shared among all titanium ions forming a *bandstate*. In the second option the electron localizes on a titanium ion that is the nearest neighbour to the hydride ion. The consequent formation of a bound polaron will be referred to as the *polaron state*.

### 4.1 Electronic and atomic structures

The valence and conduction bands of  $\text{BaTiO}_3$  consist of oxygen 2p states and titanium 3d states, respectively (see Fig. 2). In the oxyhydride phase the hydrogen level is found in the valence band with a peak at about 0.3 eV below the valence band edge. Bader analysis<sup>49,50</sup> shows an electron excess of 0.6, which confirms electron transfer to the hydrogen atom. Similar values were found by Iwazaki *et al.* for the substitutional hydrogen ion in  $\text{BaTiO}_3$  (ref. 5) as well as in  $\text{SrTiO}_3$ .<sup>51</sup>

For the bandstate the Fermi level is found inside the conduction band making the material effectively metallic [Fig. 2(a)]. The charge is delocalized and shared between all titanium atoms [see Fig. 2(c)]. The formation of a  $\text{H}^-$  defect also causes lattice distortions. The atomic displacements for the bandstate relative to pristine  $\text{BaTiO}_3$  are indicated with arrows in Fig. 2(c). The largest displacement is found for the nearest neighbour Ti, which move away from H by a distance of 0.09 Å. The nearest neighbour O moves 0.04 Å towards the hydride ion.

For the polaron state a defect level is formed in the band gap 0.37 eV below the conduction band edge [see Fig. 2(b)]. In real space the electron is localized on one of the neighbouring Ti atoms. While the bandstate maintains a principal axis of four-fold rotational symmetry, the symmetry of the polaron state is reduced to twofold rotational symmetry and the degeneracy of the perpendicular axes is removed. This can be seen in Fig. 2(d) where the electron density of the polaron is located in the  $xz$ -plane.

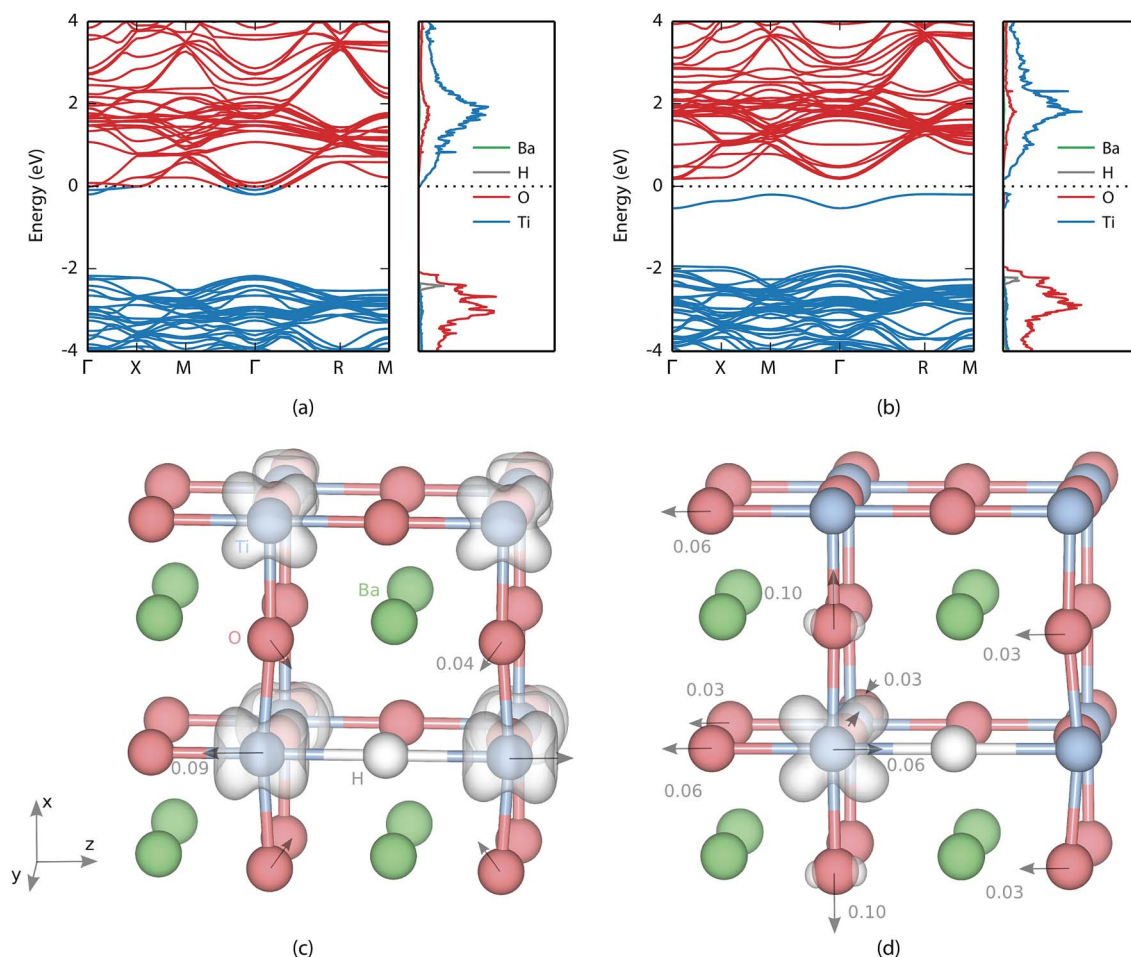
The atomic displacements due to polaron formation can be described relative to the hydride ion in the bandstate, by placing the origin on the hydrogen. Almost all major displacements are found in the vicinity of the polaron, which is not surprising since the Ti ion needs more space to accommodate the additional electron. The nearest neighbour oxygen atoms are all displaced away from the polaron along the Cartesian coordinate axes by a distance of 0.10 Å, 0.06 Å and 0.03 Å, respectively [see Fig. 2(d)]. The polaronic titanium itself is displaced by 0.06 Å towards the hydrogen, which decreases the distance between the two from 2.12 Å in the bandstate to only 2.06 Å. It is, however, still 0.04 Å larger than the oxygen-titanium distance in pristine  $\text{BaTiO}_3$  (as computed with PBE +  $U$ ). Finally, there is a shift of the entire oxygen-barium plane in the direction opposite to the titanium distortion that increases the distance between every second oxygen and titanium, which is similar to a ferroelectric transition.

### 4.2 Formation energy

In order to determine the stability of the polaron we first turn to the formation energy. For  $\text{H}_\text{O}^\times$  it has been calculated according to<sup>52</sup>







**Fig. 2** Electronic band structure and partial density of states (PDOS) for one  $\text{H}_2\text{O}$  in  $\text{BaTiO}_3$  (a) as a bandstate and (b) as a polaronic state in a  $2 \times 2 \times 2$  supercell of 40 atoms. In the band structures, blue and red colours indicate occupied and unoccupied levels, respectively. In the PDOS the hydrogen density has been magnified 10 times for clarity. The dashed black line indicates the Fermi level. The corresponding real space electronic densities for (c) the bandstate and (d) the polaron, where isosurfaces are shown at 0.1% and 1%, respectively. The arrows indicate atomic displacements (c) larger than 0.01 Å for the bandstate relative to pristine  $\text{BaTiO}_3$  and (d) larger than 0.003 Å for the polaronic state relative to the bandstate.

$$\Delta_f E_{\text{H}_2\text{O}}^{\text{X}} = E_{\text{H}_2\text{O}}^{\text{tot}} - E_{\text{pristine}}^{\text{tot}} - \frac{1}{2}\mu_{\text{H}_2} + \frac{1}{2}\mu_{\text{O}_2}, \quad (4)$$

where  $E_{\text{H}_2\text{O}}^{\text{tot}}$  and  $E_{\text{pristine}}^{\text{tot}}$  are the total energies of the oxyhydride and pristine  $\text{BaTiO}_3$ , respectively. The chemical potentials for the gas phases of  $\text{O}_2$  and  $\text{H}_2$  ( $\mu_{\text{O}_2}$  and  $\mu_{\text{H}_2}$ , respectively) were calculated as twice the total energy of the atom plus the experimental cohesive energy for the molecule, where the latter was obtained from ref. 53. Calculations were performed with  $2 \times 2 \times 2$  and  $3 \times 3 \times 3$  supercells, which correspond to hydrogen concentrations of  $x = 0.125$  and  $x = 0.037$ . PBE, PBE +  $U$  and HSE all predict rather similar formation energies that are somewhat lower in the larger supercell (see Table 2), indicating that the hydrogen concentration may influence the formation energy.

We define the polaron formation energy as the formation energy difference between the localized polaron state and the delocalized bandstate:

$$\Delta E_{\text{pol}} = \Delta_f E_{\text{H}_2\text{O},\text{pol}}^{\text{X}} - \Delta_f E_{\text{H}_2\text{O},\text{band}}^{\text{X}}. \quad (5)$$

With this definition a negative value indicates that the polaron is more stable than the bandstate. The polaron formation energy is found to depend not only on the value of  $U$  but also on the system size (see Fig. 3 and Table 2). The polaron formation energy is about 150 meV more negative for a  $2 \times 2 \times 2$  compared to a  $3 \times 3 \times 3$  supercell. Thus, in the smaller supercell a value of  $U = 3.2$  eV is sufficient to favour polaron

**Table 2** Total and polaron formation energies (eV) for  $\text{H}_2\text{O}$  in cubic  $\text{BaTiO}_3$

	$2 \times 2 \times 2$			$3 \times 3 \times 3$	
	PBE	HSE	PBE + $U^a$	PBE	PBE + $U^b$
$\Delta_f E_{\text{H}_2\text{O},\text{band}}^{\text{X}}$	4.456	4.552	4.854	4.193	4.621
$\Delta_f E_{\text{H}_2\text{O},\text{pol}}^{\text{X}}$	—	4.402	4.797	—	4.745
$\Delta E_{\text{pol}}$	—	−0.150	−0.057	—	0.124

<sup>a</sup>  $U = 3.3$  eV. <sup>b</sup>  $U = 3.1$  eV.



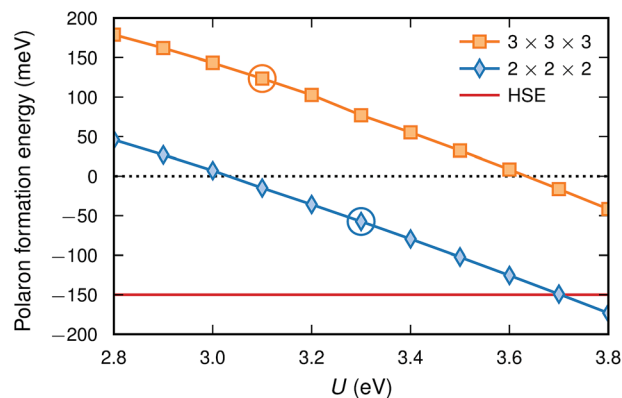


Fig. 3 Polaron formation energy [eqn (5)] calculated with PBE +  $U$  for the two supercell sizes. The circles indicate the  $U$ -values fulfilling piecewise linearity. The formation energy as calculated with HSE is also included for comparison.

formation while in the larger supercell a value of 3.7 eV is necessary. Furthermore, the  $U$ -value that fulfils piecewise linearity (see Section 2.3) is larger for a  $2 \times 2 \times 2$  supercell ( $U = 3.3$  eV) than for a  $3 \times 3 \times 3$  supercell ( $U = 3.1$  eV). This suggests that a polaron is stable ( $\Delta E_{\text{pol}} = -57$  meV) in the smaller supercell but unstable ( $\Delta E_{\text{pol}} = 124$  meV) in the larger supercell. In addition, the zero point energy is about 6 meV higher for the polaron compared to the bandstate making the polaron less stable.

Calculations with HSE suggest an even more stable polaron, which is in agreement with the trend in Fig. 1. Such an agreement between piecewise linearity and formation energy has previously been found for both electron and hole polarons in perovskites and other oxides.<sup>32,39</sup>

### 4.3 Chemical expansion

When a defect is introduced into the material it can induce an expansion (or contraction) of the material, a chemical expansion. In analogy with thermal expansion the chemical expansion coefficient  $\beta$  is defined as<sup>54</sup>

$$\beta \equiv \frac{1}{V} \frac{\partial V}{\partial x_d} \approx \frac{1}{x_d} \frac{\Delta V}{V_0}, \quad (6)$$

where  $\Delta V$  is the change of the original volume  $V_0$  when one defect is introduced into  $V_0$ . Hence, the defect concentration  $x_d$  is given by  $x_d = \Omega_c/V_0$  with  $\Omega_c$  being the volume of the primitive unit cell. For a cubic system the linear expansion coefficient is given by  $\beta_l = \beta/3$  and the relative change in the lattice constant by  $\Delta a/a_0 = x_d \beta_l = x_d \beta/3$ . If there are multiple defect species in the material the total expansion is simply given by the sum of all individual defect contributions, which in the present case, with both hydride ions and oxygen vacancies, corresponds to

$$\frac{\Delta a}{a_0} = x_H \frac{\beta_H}{3} + x_V \frac{\beta_V}{3}. \quad (7)$$

The induced strain from a single defect is generally not isotropic. It is described by the defect induced strain tensor  $\lambda$ ,

and  $\beta$  is then given by the trace of  $\lambda$ ,  $\beta = \text{Tr}(\lambda)$ .<sup>55</sup> The tensor  $\lambda$  can be computed by performing full relaxations, *i.e.*, relaxation of both ionic positions and cell parameters. The defect induced strain tensor is then obtained as<sup>55</sup>

$$\lambda = \frac{1}{x_d} (L - L_0) L_0^{-1}, \quad (8)$$

where  $L_0$  is the  $3 \times 3$  matrix that describes the cell metrics of the ideal supercell structure and  $L$  is the corresponding matrix for the defective supercell (containing a single defect) after full relaxation.

We have determined the chemical expansion for both the bandstate and the polaron state of  $\text{H}_0^\times$  using a  $2 \times 2 \times 2$  supercell. Our sample also contains oxygen vacancies (see Section 3.1). This type of cubic O-deficient oxyhydride is found to be quite common.<sup>56</sup> In  $\text{BaTiO}_3$  these vacancies act as shallow donors and we have therefore also considered the neutral vacancy ( $\text{v}_0^\times$ ).<sup>57,58</sup> We find that the lattice expands along the principal axis ( $z$ ) for all three defects and that the degeneracy of the perpendicular axes ( $x$  and  $y$ ) is lifted for the polaron (see Table 3), in agreement with the symmetries described above. The expansion of the lattice along the principal direction is largest for the neutral vacancy. For the bandstate there is a small contraction along the perpendicular axes, but the chemical expansion is still positive with  $\beta = 0.030$ . For the polaron state there is also a contraction along the  $y$  direction but an expansion along the  $x$  direction causing an overall chemical expansion of  $\beta = 0.056$ , which is nearly twice as large as that of the bandstate. For the oxygen vacancy there is also a small contraction along the perpendicular axes, and the resulting chemical expansion is  $\beta = 0.049$ , nearly as large as for the polaron state.

The magnitude of the chemical expansion for the bandstate and the polaron state is quite different and depending on which state is present the lattice expansion will be different. The measured composition is  $\text{BaTiO}_{2.82}\text{H}_{0.1}\square_{0.08}$  and the expanded lattice constant  $a = 4.0055$  Å. To obtain the change of the lattice constant we need an experimental value for  $a_0$  at 20 °C. By extrapolating the data for cubic  $\text{BaTiO}_3$  in ref. 59 to 20 °C,  $a_0 = 3.9975$  Å is obtained. This implies that experimentally  $\Delta a/a_0 = 0.0020$ . We can also compute the lattice expansion using the data in Table 3. By assuming a bandstate and polaron state, respectively, we obtain the lattice expansions  $\Delta a/a_0 = 0.0023$  and  $\Delta a/a_0 = 0.0032$  (see Table 4). Clearly, by assuming a bandstate the theoretical lattice expansion becomes much closer to the experimental value compared with assuming the polaron state.

Table 3 The defect induced strain tensor  $\lambda$  and chemical expansion coefficient  $\beta$  for the bandstate and the polaron state of  $\text{H}_0^\times$  as well as for the neutral oxygen vacancy ( $\text{v}_0^\times$ ) calculated with PBE +  $U$  ( $U = 3.3$  eV) in a  $2 \times 2 \times 2$  supercell

	$\lambda_{xx}$	$\lambda_{yy}$	$\lambda_{zz}$	$\beta$
Bandstate	−0.0045	−0.0045	0.0392	0.030
Polaron state	0.0283	−0.0179	0.0456	0.056
Oxygen vacancy	−0.0059	−0.0059	0.0608	0.049



**Table 4** Relative lattice expansion computed from the data in Table 3 according to eqn (7) assuming either the bandstate or the polaron state together with the experimentally measured lattice expansions

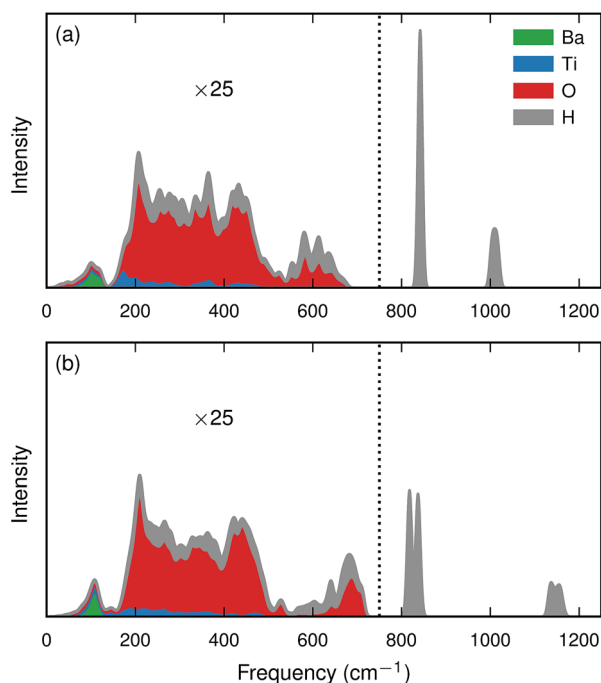
$x_{\text{H}}$	$x_{\text{V}}$	$\Delta a/a_0$		
		Bandstate	Polaron	Exp
0.10	0.08	0.0023	0.0032	0.0020 <sup>a</sup>
0.62	0.00	0.0062	0.0116	0.0065 <sup>b</sup>

<sup>a</sup> This work. <sup>b</sup> Ref. 1.

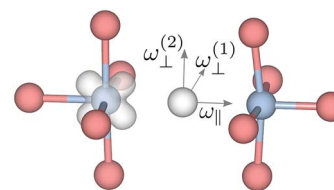
It is also interesting to compare with the experimental data by Kobayashi *et al.*<sup>1</sup> They synthesized a sample with the composition  $\text{BaTiO}_{2.38}\text{H}_{0.62}$ . It was stated that the number of oxygen vacancies was small. The expanded lattice constant was  $a = 4.0236 \text{ \AA}$ , which then corresponds to an expansion of  $\Delta a/a_0 = 0.0065$ . Again, we find that by assuming a bandstate the theoretical lattice expansion becomes much closer to the experimental value (see Table 4).

#### 4.4 Vibrational spectra

**4.4.1 Theoretical.** The vibrational spectrum for the oxyhydride phase has been computed both for the bandstate and the polaron state using the frozen phonon approach with the PBE +  $U$  functional (see Fig. 4), from which intensities are calculated according to eqn (3). The mean square displacements were found to be 1.15, 19.8, 3.45 and  $1.85 \times 10^{-3} \text{ \AA}^2$  for Ba, H, O and Ti, respectively.



**Fig. 4** Calculated partial (cumulative) intensity [eqn (3)] for a hydride ion in a  $2 \times 2 \times 2$   $\text{BaTiO}_3$  supercell for (a) the bandstate and (b) the polaron state. Intensities below  $750 \text{ cm}^{-1}$  (marked with a dotted line) have been magnified by a factor of 25.



**Fig. 5** Illustration of the vibrational modes for the substitutional hydride ion in oxyhydride  $\text{BaTiO}_3$  in the polaronic state. For the bandstate  $\omega_{\perp}^{(1)}$  and  $\omega_{\perp}^{(2)}$  are degenerate.

The hydrogen motion for the bandstate can be seen as two sharp peaks around  $843 \text{ cm}^{-1}$  and  $1018 \text{ cm}^{-1}$  [see Fig. 4(a)] and the assignment of these modes is illustrated in Fig. 5. The higher frequency mode is the stretching mode of  $A_{2u}$  symmetry along the principal axis, which is denoted as  $\omega_{\parallel}$ . The lower mode frequency is the doubly degenerate bending mode of  $E_u$  symmetry in the mirror plane perpendicular to the principal axis and is denoted as  $\omega_{\perp}$ . The vibrational frequencies of the hydride ion motion for the bandstate configuration of  $\text{H}_0^{\times}$  have been studied previously by Iwazaki *et al.*<sup>5</sup> using a method similar to our OPHP method (see Section 2.4) with the PBE functional and the results from their investigation are in good agreement with our results.

The lattice distortion associated with polaron formation breaks the mirror-plane symmetry as well as the symmetry of fourfold rotation into a two-fold rotation symmetry. The  $E_u$  bending mode is split into two modes of  $B_1$  and  $B_2$  symmetry, for which the separation is about  $20 \text{ cm}^{-1}$  [see Fig. 4(b)]. In addition, the displacement of the polaronic Ti towards the hydrogen causes a substantial upward shift of the stretching mode frequency from  $1006 \text{ cm}^{-1}$  to  $1133 \text{ cm}^{-1}$ .

Due to the small mass of the hydride ion its vibrational eigenmodes are localized and show little dispersion. Table 5 shows the maximum and minimum frequencies together with the  $\Gamma$ -point frequency for the bandstate, where it can be seen that the dispersion is small. For the stretching mode, the width of the band is only  $16 \text{ cm}^{-1}$  and the dispersion is negligible for the bending mode. These modes should therefore be well described by the OPHP method. By symmetry, the eigenmodes must lie along the lattice vectors: the principal axis and the two

**Table 5** Comparison between the vibrational frequencies ( $\text{cm}^{-1}$ ) of the  $\text{H}^-$  modes for the bandstate and the polaron state of  $\text{H}_0^{\times}$  computed using phonopy with a  $4 \times 4 \times 4$  augmented supercell ( $\Gamma$ , max, min) and the OPHP method. PBE +  $U$  calculations were performed using  $U = 3.3 \text{ eV}$ . The data that are used in the comparison with the INS spectra are highlighted in bold

	Mode	PBE + $U$			HSE
		$\Gamma$	Max	Min	
Bandstate	$\omega_{\parallel}$	1018	1018	1002	<b>1033</b>
	$\omega_{\perp}$	843	843	841	<b>899</b>
Polaron state	$\omega_{\parallel}$	1133	1168	1133	<b>1160</b>
	$\omega_{\perp}^{(1)}$	835	835	840	<b>894</b>
	$\omega_{\perp}^{(2)}$	818	815	818	<b>874</b>



perpendicular axes, which can be considered independently. The frequencies obtained with the OPHP method show good agreement with the frequencies obtained from the full phonon calculation using phonopy<sup>40</sup> software (see Table 5). This implies that the vibrational modes for the hydride ion are accurately described using the OPHP method.

As discussed in Section 2.2, the HSE functional predicts accurate frequencies for pristine BaTiO<sub>3</sub>. The stretching mode for the hydride ion is similar to the T<sub>1u</sub> modes of pristine BaTiO<sub>3</sub> and the bending mode is similar to the T<sub>2u</sub> mode. It is therefore desirable to use HSE also for the vibrational motion of the hydride ion. However, a full phonon calculation with HSE is computationally very demanding but as we have shown here that the computationally less expensive OPHP method can be used to obtain accurate frequencies. We have therefore restricted the HSE calculations to the OPHP method.

Let us first consider the bandstate. A relaxed HSE configuration at the HSE lattice constant is used to obtain the 3D potential as described in Section 2.4, from which the harmonic frequencies are determined. Compared with PBE + *U* the HSE functional shifts the stretching frequency upwards by 35 cm<sup>-1</sup>, while the shift for the bending mode is even larger, 50 cm<sup>-1</sup> (see Table 5).

Let us now consider the polaron state. The atomic relaxation with the HSE functional was conducted only for the ions along the Ti–H–Ti–O axis and its nearest neighbour oxygen ions, *i.e.*, the ions depicted in Fig. 5. The largest displacements are found along the principal axis, where the hydrogen and oxygen atoms are both displaced in the negative *z*-direction, while the non-polaronic titanium is displaced in the positive *z*-direction (see Fig. 2 for the definition of the coordinate system). The polaronic Ti itself is displaced only marginally relative to the lattice. This causes the distance between the polaronic Ti and the hydrogen to decrease by 0.04 Å, from 2.06 Å to 2.02 Å, and increase between the polaronic Ti and the oxygen by 0.02 Å. Displacements of similar magnitudes were also found for the nearest neighbour oxygen ions relative to the polaronic Ti. In the *x*-direction the distance increases by 0.02 Å and in the *y*-direction by 0.03 Å. The relaxed configuration was used to create a 3D

potential from which the harmonic frequencies are determined using the OPHP method. The HSE functional shifts the frequencies of the polaron state in a similar way to that for the bandstate. The stretching mode frequency is shifted upwards by 18 cm<sup>-1</sup> and the bending mode frequencies by 63 cm<sup>-1</sup> and 60 cm<sup>-1</sup>, respectively (see Table 5). Compared with the PBE + *U* values the frequencies are shifted upwards by about 3–4% and 7% for the stretching mode and the bending mode, respectively. These relative shifts are quite similar to the shifts for pristine BaTiO<sub>3</sub> (see Section 2.2).

**4.4.2 Experimental.** The INS spectrum at 10 K of BaTiO<sub>2.82</sub>H<sub>0.10</sub>□<sub>0.08</sub> are shown in Fig. 6(a). The most pronounced feature is the intense peak centered around 900 cm<sup>-1</sup>. At slightly higher frequencies, around 1030 cm<sup>-1</sup>, there is a second less intense peak and there is also a distinct feature around 800 cm<sup>-1</sup>. Due to the high frequencies of these modes and the large neutron cross section of hydrogen compared to the other atomic species in the material, it is safe to assume that these peaks can be assigned to hydrogen vibrational modes. Peaks at frequencies larger than 1500 cm<sup>-1</sup> are assigned to higher-order transitions of these modes. The intensities at frequencies below 700 cm<sup>-1</sup> are attributed mainly to oxygen lattice modes (*cf.* Fig. 4) but also partly to the hydrogen motion inherent in these modes. Finally, the fact that no peak is found in the range around 3500 cm<sup>-1</sup> is congruent with the fact that the hydrogen atoms are located on the oxygen sites rather than bonded as O–H species, as expected, although it should be noted that TOSCA is quite insensitive in this spectral range.<sup>59</sup>

For a more quantitative analysis we have fitted the peaks in the interval between 750 and 1250 cm<sup>-1</sup>. Four Gaussians were required to obtain a good fit. The fit parameters are listed in Table 6 and the corresponding Gaussians are plotted in Fig. 6(b).

**4.4.3 Comparison.** The two peaks at 913 and 1031 cm<sup>-1</sup> in the INS spectrum of the sample fit the calculated frequencies for the bandstate very well [see Fig. 6(b)]. These two peaks are therefore assigned to the double degenerate mode  $\omega_{\perp}$  and the mode  $\omega_{\parallel}$  with calculated frequencies 899 and 1033 cm<sup>-1</sup>,

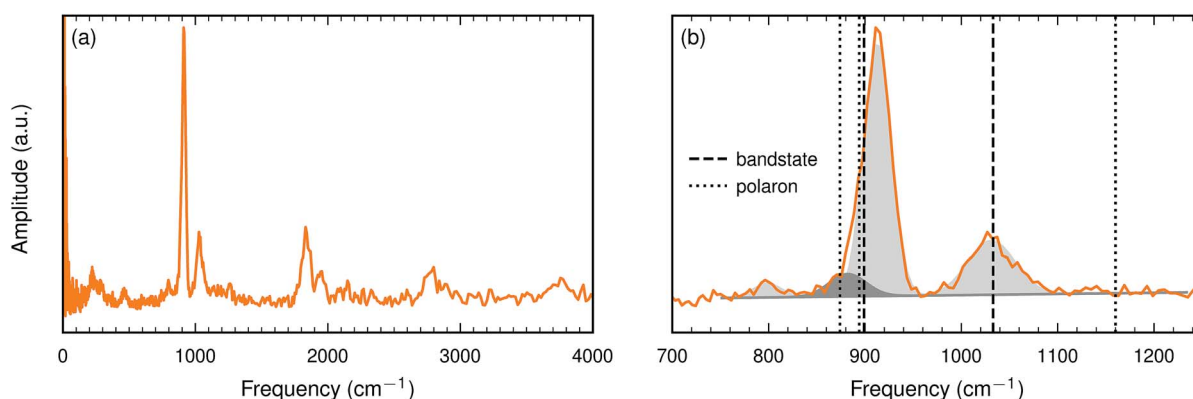


Fig. 6 (a) The INS spectrum of the BaTiO<sub>2.82</sub>H<sub>0.10</sub>□<sub>0.08</sub> sample at 10 K. (b) Close up of the spectrum in the interval 700–1250 cm<sup>-1</sup>. Frequencies calculated with HSE (see Table 5) are marked with dashed (bandstate) and dotted (polaron) lines. Gaussian fits are indicated by shaded areas (see Table 6).





**Table 6** Fitted vibrational frequencies ( $\omega$ ) of the  $H^-$  modes for the INS spectra of  $BaTiO_{2.82}H_{0.10}\square_{0.08}$  [see Fig. 6(b)] together with the corresponding fitting parameters

$\omega$ ( $cm^{-1}$ )	Area (a.u.)	FWHM ( $cm^{-1}$ )
799	0.3	33.5
882	0.7	44.8
913	6.2	33.9
1031	2.3	56.8

respectively (see Table 5). The calculated frequency for the stretch mode of the polaron state is located considerably higher, at  $1160\text{ cm}^{-1}$ . At these frequencies very little intensity is present in the measured spectrum. We also expect the bending mode for the polaron state to be broader than the measured peak around  $910\text{ cm}^{-1}$ . It follows that the presence of polarons is very unlikely.

The present oxyhydride sample contains about an equal number of hydride ions and oxygen vacancies. The interaction between these two defects is repulsive and hence we do not expect any pronounced hydride ion–oxygen vacancy association. We have computed the hydride ion frequency spectrum for a few systems with an equal number of hydride ions and oxygen vacancies, with about 10% each. We find that when the hydride ion and the oxygen vacancy are nearest neighbors the degeneracy of the bending mode is removed and one of these modes is downshifted by about  $100\text{ cm}^{-1}$ . The other two modes are essentially unaffected. It is tempting to identify the weak experimental peak around  $800\text{ cm}^{-1}$  with this defect configuration. For other defect configurations, where the hydride ion and the oxygen vacancy are not nearest neighbors, we find only small changes of the computed vibrational frequencies.

This comparison between measured and computed vibrational frequencies strongly suggests that electrons form bandstates in this O-deficient cubic oxyhydride.

## 5 Summary and conclusion

In the present contribution we have investigated the character of conduction electrons in cubic  $BaTiO_{3-x}H_x$ , where an electron can form either a delocalized bandstate or localize as a small electron polaron on titanium bound next to a hydride ion. The polaron formation energy has been determined with density-functional theory (DFT) calculations using both the DFT +  $U$  method and the hybrid functional HSE, where  $U$  in the former case was carefully chosen such that piecewise linearity was obtained and the self-interaction error was minimized. The formation energy was found to be small, on the order of thermal energy, and this result alone does not provide a conclusive description of the electronic state.

The formation of a polaron causes a local lattice distortion where the Ti–H distance is reduced, which affects the localized hydride ion vibrational modes. These vibrational frequencies have been determined using HSE and we find that the stretching mode frequency shifts up from  $1033\text{ cm}^{-1}$  to  $1160\text{ cm}^{-1}$  upon polaron formation. At the same time the degenerate bending

mode is split into two modes that are slightly downshifted by up to  $30\text{ cm}^{-1}$ . A comparison of these frequencies with data from inelastic neutron scattering (INS) experiments strongly suggests that electrons form bandstates in the present bulk material.

Furthermore, the chemical expansion has been determined, showing that the lattice expansion of the polaron state is about twice as large as that for the bandstate. By comparing the calculated data with our experimental data as well as with the experimental data for a fully hydride-ion substituted sample by Kageyama and coworkers<sup>1</sup> we conclude that the expansion due to the bandstate fits much better than the polaron state, which further supports the presence of delocalized electrons in the oxyhydride.

The delocalized nature of the electrons is in line with metallic conductivity. For thin films of  $BaTiO_{3-x}H_x$ , Kageyama and co-workers have reported metallic conductivity at high hydride concentration,<sup>8,9</sup> while at lower concentration a semi-conducting behaviour was observed.<sup>9</sup> It is possible that strain in epitaxial thin films could favour polaron formation and the small formation energies found here suggest that a rather small perturbation of the structure is sufficient for changing the relative stability of the two electronic states. Indeed, in ref. 9 it is found that the lattice structure of the thin film is not cubic but tetragonal. The semiconducting behaviour observed in bulk  $BaTiO_{3-x}H_x$  as reported in ref. 1 is, however, most likely due to grain boundaries in the powder sample.<sup>8</sup>

To conclude, we find that the electrons form delocalized bandstates in bulk  $BaTiO_{3-x}H_x$ , which implies a metallic conduction mechanism in this material. Furthermore, we have shown that first-principles calculations in combination with INS experiments are excellent tools to study the vibrational motion of the hydride ions and relate this to the character of the conduction electrons in oxyhydrides, and that chemical expansion can be used to discriminate between formation of delocalized bandstates and localized small polaron configurations in these materials.

## Conflicts of interest

There are no conflicts to declare.

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