


 Cite this: *Nanoscale*, 2024, **16**, 9436

Optical study of Te₈ ring clusters: comparison with density functional theory and a step towards materials design using nanoporous zeolite space

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The Te₈ ring molecule (cluster) is poorly investigated due to the lack of experimental data. Here, we report an experimental and theoretical study of a regular array of oriented Te₈ rings formed in the ~1.14 nm diameter cavities of zeolite LTA, which are arranged in a cubic lattice with a spacing of ~1.2 nm. Single crystals of LTA with encapsulated tellurium (LTA-Te) were studied using Raman spectroscopy (RS) and optical absorption spectroscopy (OAS). The experimental LTA-Te spectra were found to be in agreement with those calculated using density functional theory (PBE0 hybrid functional and def2-TZVP basis sets) for the crown-shaped Te₈ ring molecule with *D*_{4d} symmetry. Using polarization–orientation RS, we show that the Te₈ rings are oriented by their major axes along the 4-fold axes of cubic LTA. We also show that the site symmetry of Te₈ in LTA-Te is lower than *D*_{4d}. Te₈ bond-bending modes are well described in the harmonic approximation, while bond-stretching modes are mixed due to the reduced ring symmetry and, probably, anharmonicity. Importantly, OAS data of LTA-Te display dependence on the Te₈ concentration, implying the interaction of the rings from neighbouring LTA cavities with the generation of the valence and conduction electron bands of such a cluster crystal.

 Received 9th January 2024,
 Accepted 12th April 2024

DOI: 10.1039/d4nr00114a

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Introduction

Tellurium (Te) belongs to the same group of the periodic table as oxygen (O), sulphur (S) and selenium (Se). Since its discovery at the end of the 18th century, Te was widely used in metal alloys. Nowadays, Te and its compounds have become a real frontier of science and technology. They have found important applications in electronics, optoelectronics, photonics, phase-change materials, topological insulators, Li-Te batteries, thermoelectrics and solar cells.^{1–6} However, elemental Te is mainly known in its chain form analogous to crystalline trigonal tellurium (t-Te) while it is nearly unknown in the ring form.

Zeolites provide a unique opportunity to form and accommodate uniform guest species in their cavities/channels where the species are oriented due to the crystalline nature of zeolites.^{7–13} For example, only zeolites allowed obtaining polarized Raman spectra (RS) and optical absorption spectra (OAS) of isolated Te chains and rings.^{7–11} Moreover, fabrication of such regular high-density Te species arrays can be considered as an important direction in the development of new functional materials, so-called cluster crystals.^{14–16}

Here, we study Te₈ rings regularly arranged in the large cavities of zeolite LTA. Importantly, we are using the polarization–orientation RS examination of zeolite single crystals. On one hand, this method can be considered as an alternative to X-ray diffraction (XRD) for the structural study of zeolite-confined clusters. On the other hand, it provides very useful information about the cluster properties and their interaction with zeolites. We also study the OAS spectra of Te₈ and demonstrate the effect of the interaction between Te₈ rings in neighboring cavities, which is important for the fabrication of real cluster crystals.

The idea of Te₈ ring formation in LTA cavities was proposed in ref. 17 where the only observed RS feature at ~168 cm⁻¹ was assigned to the ring with an argument that its frequency is different from the dominant trigonal tellurium (t-Te) band frequency at ~120.5 cm⁻¹.^{18,19} However, this observation was not confirmed later. In ref. 20, the RS of LTA-Te microcrystalline powder was re-examined. A strong band at ~182 cm⁻¹ and a weaker one at ~45 cm⁻¹ were observed and attributed rather to Te₁₂ than to Te₈ since Te₈ requires a relatively strong symmetric bond-bending mode band at around 60 cm⁻¹, which was not clearly detected due to laser-induced band broadening, as was found later. The LTA-Te band at ~62 cm⁻¹ accompanied by ~182 cm⁻¹ and ~45 cm⁻¹ bands was observed in ref. 21. This was a rather strong argument in favor of Te₈ ring formation in LTA cavities. The observed bands at ~45,

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~ 62 and ~ 182 cm^{-1} look similar to the bending E_2 and A_1 mode bands and stretching A_1 mode bands of S_8 (~ 153 , ~ 222 , ~ 480 cm^{-1}) and Se_8 (~ 76 , 111 , 267 cm^{-1}) rings.²¹ The RS of another zeolite AFI with encapsulated tellurium (AFI-Te) showed similar bands at 41 – 47 cm^{-1} , 65 cm^{-1} and 184 cm^{-1} .⁷ The bands were attributed to the Te_8 ring. However, no polarized RS study of LTA-Te single crystals has been performed yet, which could provide crucial evidence for Te_8 ring formation. Here, we present this evidence. Importantly, no other species but Te_8 was found in LTA-Te. This is beneficial for realizing the idea of cluster crystal formation.

Experimental and theoretical methods

Synthetic zeolite LTA with the composition $Na_{12}Al_{12}Si_{12}O_{48}$ was used in this work. The crystal growth procedure for LTA is described in ref. 22. The sizes of the cubic LTA crystals were ~ 20 μm along the edge of the cube. LTA has nearly spherical-shaped large cavities with a diameter of ~ 1.14 nm, which are connected through narrow windows of ~ 0.42 nm diameter (Fig. 1). The LTA cavities are arranged in a simple cubic lattice. The zeolite structures are accessible at the International Zeolite Association website https://www.iza-structure.org/IZA-SC/ftc_table.php.

Dehydration of zeolites in a vacuum was performed in Pyrex ampoules for several hours at $t \sim 550$ $^\circ\text{C}$. The design of the ampoules made it possible to add crystalline Te powder to the zeolite after dehydration without exposing the dehydrated samples to air. The subsequent adsorption of the Te vapour was carried out at $t \sim 550$ $^\circ\text{C}$ for a few days until the brownish colour of the sample was saturated.

RS spectra of the LTA-Te single crystals were studied using a Renishaw micro-Raman spectrometer equipped with Semrock edge-filters and an Olympus micro-objective lens allowing an ~ 1 μm focused laser probe size. A 633 nm wavelength line of the He-Ne laser and a 785 nm line of the light-emitting diode

laser were used for RS excitation. Weak absorption of Te_8 at these wavelengths is beneficial for Te_8 compared to the non-absorbing LTA matrix for obtaining the cluster Raman signal with nearly no contribution of LTA. An additional measurement with 561 nm laser diode line excitation was performed using a Nanofinder-30 Raman/AFM system (Tokyo Instruments Inc.) specially designed for the maximal RS detection sensitivity around this wavelength, which allowed LTA-Te Raman measurement at relatively high Te_8 absorption and very low laser excitation power in order not to strongly heat or destroy the clusters. Moreover, a Nanofinder-30 (Tokyo Instruments Inc.) was equipped with a set of three volume Bragg grating notch filters (OptiGrate Co), allowing the detection of the Raman signal with frequencies down to ~ 6 cm^{-1} . UV-visible OAS was performed using a Carl Zeiss micro-optical spectrometer with a light probe size of ~ 5 μm . The samples were intentionally broken to minimize their optical density in the ultraviolet (UV) spectral range and placed in glycerol between two cover glasses to minimize surface light scattering.

The Te_8 RS, infrared (IR) spectra and OAS were simulated using density functional theory (DFT).^{23–26} The structure and vibrational frequencies of Te_8 were computed using the hybrid PBE0 exchange–correlation functional²⁶ and the resolution-of-identity approximation for the Coulomb part (RI-J).²⁷ The def2-TZVP basis sets²⁸ together with the corresponding auxiliary basis sets²⁹ and small-core effective core potentials (ECPs)³⁰ were employed for Te. RS intensities were obtained in the double harmonic approximation using analytical derivatives of static electronic polarizabilities³¹ and analytical energy Hessians.^{32,33} Calculations of static polarizability derivatives used the PBE0 functional and augmented def2-TZVPD basis sets.³⁴ IR spectra were computed in the double harmonic approximation from the analytical dipole and energy Hessians.^{32,33} The vibrational frequencies were not scaled in this work. OAS was performed with time-dependent DFT (TDDFT)³⁵ using the PBE0 functional with the RI-J approximation³⁶ and def2-TZVPD basis sets. All calculations were performed with the Turbomole program package, version 7.5.^{23,37}

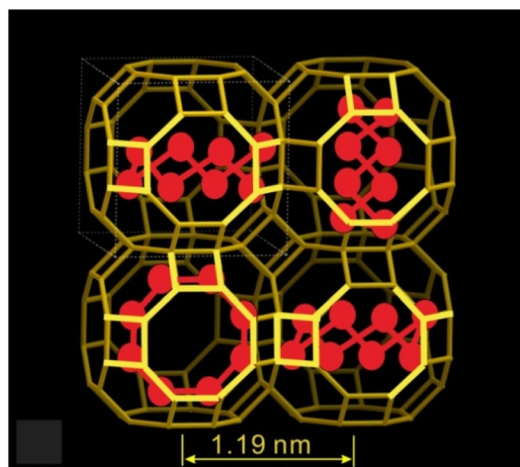


Fig. 1 Schematic view of the LTA framework with Te_8 rings in large cavities.

Structure and Raman and absorption spectra of single Te_8 rings via DFT calculations

The calculated Te_8 (D_{4d} point group symmetry) ring structure is shown in Fig. 2(a). The bond length of Te_8 is $r = 0.271$ nm and the second neighbour distance is $R = 0.432$ nm, while its bond and dihedral angles are 105.8° and 101.2° , respectively. The obtained structure parameters are in agreement with previous DFT calculations.^{38–42} The ring Z axis is directed along its 4-fold axis, while the X and Y axes lie in the molecule plane normal to the Z axis (Fig. 2(a)).

Theoretical point group analysis for the vibration modes of Te_8 is similar to that made for the S_8 molecule with a similar crown shape.⁴³ The Te_8 vibration irreducible representations



Fig. 2 Te_8 ring with the bond length r and angle ϕ (a); calculated Raman spectra of a single free-standing Te_8 ring for XX , ZZ , XY and XZ polarization configurations (b); Te_8 atomic displacements for A_1 bond-bending and A_1 bond-stretching modes with corresponding Raman tensors (c); Te_8 MO diagram with dominant MO contributions to the strongest optical transitions shown in red, all for $E//X$ (d); and calculated Te_8 OAS including transitions with photon energies of <5.5 eV (e).

(symmetries) with calculated frequencies and IR/Raman activities are shown in Table 1. The E_1 symmetry corresponds to the IR activity for the light polarized in the ring plane ($E//X$ or $E//Y$), while the B_2 symmetry corresponds to the $E//Z$ IR activity. Calculated Te_8 Raman intensities for different polarization configurations are equal to their squared Raman tensor components (XX , ZZ , XY , XZ) for each mode, which are determined by the corresponding atomic displacements and related molecular polarizability derivatives. The corresponding values in Table 1 are given in atomic units of $10^{-2}e^2a_0/E_h$, where e , a_0 and E_h are the electron charge, Bohr radius and Hartree energy, respectively.

The calculated RS spectra of Te_8 for different polarization configurations, corresponding to the Raman tensor components from Table 1, are shown in Fig. 2(b). Raman bands are represented as Lorentz curves with the amplitudes corresponding to the calculated Raman activities and half-widths of 5 cm^{-1} for all bands.

The dominant band in the calculated RS spectra of Te_8 , displaying comparable XX and ZZ activities, corresponds to the A_1 symmetric bond-stretching mode at a frequency of 186.9 cm^{-1} . In contrast, the A_1 bond-bending mode at 65.4 cm^{-1} displays high XX activity and nearly negligible ZZ activity. Fig. 2(c) shows atomic displacements and Raman tensors of these two

Table 1 Te_8 normal vibrations: symmetries, computed frequencies, IR intensities and Raman activities

Symmetry	Frequency (cm^{-1})	IR intensity (km mol^{-1})	Raman activity XX	Raman activity ZZ	Raman activity XY	Raman activity XZ
E_2	21.7	0	-0.49	0	-2.08	0
	21.7	0	-2.08	0	0.49	0
E_2	43.8	0	-4.38	0	-3.36	0
	43.8	0	3.36	0	-4.38	0
E_1	56.9	0.524	0	0	0	0
	56.9	0.524	0	0	0	0
A_1	65.4	0	-7.04	0.56	0	0
B_2	68.6	0.128	0	0	0	0
E_3	72.7	0	0	0	0	-1.86
	72.7	0	0	0	0	-0.86
B_1	179.9	0	0	0	0	0
E_3	185.3	0	0	0	0	-3.54
	185.3	0	0	0	0	-0.17
A_1	186.9	0	-8.48	-6.95	0	0
E_1	188.1	1.29	0	0	0	0
	188.1	1.29	0	0	0	0
E_2	192.1	0	1.21	0	-1.05	0
	192.1	0	-1.05	0	-1.21	0

A_1 modes, which explain the difference in their ZZ activities. One can consider the A_1 bond-stretching as a breathing mode since both in-plane and off-plane Te_8 dimensions expand and contract the in-phase, while the A_1 bond bending mode is an important instrument for Te_8 ring orientation determination.

The E_2 bond-bending mode at 43.8 cm^{-1} is rather active in the XX and XY configurations, while inactive in the ZZ configuration. Similar but weaker activity is displayed by the E_2 torsion and stretching modes at 21.7 cm^{-1} and 192.1 cm^{-1} , respectively. The E_3 bond-bending mode at 72.7 cm^{-1} and the bond-stretching mode at 185.3 cm^{-1} dominate in the XZ configuration spectrum.

In addition to the Te_8 Raman-active modes, there are several inactive modes: E_1 bond-bending at 56.9 cm^{-1} , B_2 bond-bending at 68.6 cm^{-1} , B_1 bond stretching at 179.9 cm^{-1} and E_1 bond stretching at 188.1 cm^{-1} , which can become active when the ring D_{4d} symmetry is reduced. As we mentioned above, the E_1 and B_2 modes are IR-active.

The calculated partial molecular orbital (MO) diagram of Te_8 including the highest occupied MOs and the lowest unoccupied MOs is presented in Fig. 2(d). These MOs consist of combinations of $Te\ 5p$ orbitals, as determined by Mulliken population analysis. The occupied orbitals correspond to Te lone pairs, while the unoccupied orbitals have antibonding character with respect to Te - Te bonds. The largest MO contributions to the allowed electronic transitions with photon energies $<4\text{ eV}$ and the largest oscillator strengths are shown in red. It should be noted that the electron transition energies take into account excitonic effects. Therefore, the transition energies are less than the orbital energy differences of the dominant MO contributions. All these four transitions have E_1 symmetry, implying the polarization of light $E//X$.

There are also $E//Z$ -allowed transitions of B_2 symmetry. However, their oscillator strengths appeared to be much lower than those of the strongest $E//X$ transitions (Table 2). The calculated OAS spectra of Te_8 for polarizations of light $E//X$ and $E//Z$ are shown in Fig. 2(e). Allowed electronic transitions (E_1

for $E//X$ and B_2 for $E//Z$) with photon energies of up to 5.5 eV were computed. Due to the difference in the oscillator strengths, the $E//X$ absorption is stronger than the $E//Z$ one. The $E//X$ absorption is mainly represented by transitions at photon energies of ~ 2.9 , ~ 3.1 , ~ 4.2 , ~ 4.6 and $\sim 5.4\text{ eV}$, while the $E//Z$ one is determined by transitions at photon energies of ~ 3 , ~ 3.6 , ~ 4 and $\sim 5.2\text{ eV}$. The corresponding absorption bands are approximated by the Lorentz curves with half-widths of $\sim 0.6\text{ eV}$.

Experimental polarized Raman spectra of LTA-Te single crystals

Using Te vapor adsorption, we obtained LTA-Te samples with Te loading densities of ~ 3 and ~ 8 atoms per cavity (at. per cav.). Unlike the RS of LTA-Se with different loading densities showing changes in the Se_8/Se_{12} intensity ratio,⁴⁴ LTA-Te RS spectra did not show any qualitative dependence on the Te loading density except an increase in the Raman signal of the same bands with an increase in the Te density. This is a strong argument in favour of the formation of only one stable Te cluster in the LTA cavities.

RS spectra of LTA-Te with a loading density of ~ 8 at. per cav. at an excitation wavelength of 785 nm are shown in Fig. 3(a). We performed measurements in four different polarization configurations: aa , cc , ab and cd (see the inset in Fig. 3(a)) similar to the polarization-orientation Raman study of LTA with sulphur LTA-S¹² and LTA with selenium LTA-Se.^{12,44,45} Experimentally, we rotated the LTA-Te crystals in the ab plane with incident and scattered light polarizations (1) parallel and (2) perpendicular to each other. The procedure was described in detail earlier.¹² Theoretical RS spectra of LTA-Te for the aa , cc , ab and cd polarization configurations were obtained *via* summation of the Raman responses of three Te_8 rings in their three possible orientations in LTA crystals [Fig. 3(b)].

The experimental LTA-Te RS spectra display rather good correspondence with the calculated ones. Indeed, the bands at 162 cm^{-1} and 183 cm^{-1} can be assigned to the $Te_8\ E_3$ and A_1 bond-stretching modes similar to the 442 cm^{-1} and 480 cm^{-1} bands of S_8 in LTA-S [Fig. 3(c)]. The correspondence between the experiment and theory looks especially good for the A_1 mode frequency (experimental: $\sim 183\text{ cm}^{-1}$ vs. theoretical: 186.9 cm^{-1}) and intensity (the strongest band in the spectrum). At first glance, the polarization dependence of its intensity also looks reasonable: the band is strong in aa and cc polarization configurations, while it is weak in ab and cd configurations as expected for the A_1 modes. Actually, as we have shown below, the situation is more complicated for the bond stretching modes due to a possible mode mixing effect but, in a rough approximation, the assignment of the 183 cm^{-1} band to the A_1 bond-stretching mode looks reasonable.

Polarization dependencies of the LTA-Te bands in the Te_8 bond-bending mode region appear to be qualitatively similar to those of the bond-bending modes of S_8 ¹² [Fig. 3(c)] and

Table 2 Computed dipole-allowed electron transitions of Te_8

Symmetry	Photon energy (eV)	Oscillator strength
E_1	2.90	0.104
B_2	3.03	0.016
E_1	3.09	0.053
B_2	3.45	0.003
E_1	3.53	0.001
B_2	3.61	0.003
E_1	3.89	0.012
B_2	3.96	0.004
E_1	4.11	0.006
E_1	4.21	0.357
E_1	4.35	0.002
E_1	4.57	0.525
B_2	5.21	0.071
E_1	5.42	0.333
B_2	5.77	0.008
E_1	5.82	0.229
B_2	5.94	0.002



Fig. 3 LTA-Te Raman spectra of *aa*, *cc*, *ab* and *cd* polarization configurations at room temperature and 785 nm wavelength excitation with the inset schematically showing LTA-Te cubic crystals and the *a*, *b*, *c* and *d* axes (a); summation of the calculated Raman spectra of three Te_8 orientations in LTA (see the inset) for *aa*, *cc*, *ab* and *cd* polarization configurations (b); LTA-S Raman spectra of *aa*, *cc*, *ab* and *cd* polarization configurations (c); LTA-Te Raman spectra of *aa*, *cc*, *ab* and *cd* polarization configurations at $T \sim 77$ K and 633 nm wavelength excitation (d); LTA-Te Raman spectra of *aa*, *cc*, *ab* and *cd* polarization configurations at room temperature and 633 nm wavelength excitation (e); and LTA-Te Raman spectra of the *cc* polarization configuration at different excitation wavelengths with the unpolarized spectrum of bare LTA (black curve) (f).

Se_8 ^{12,44,45} rings in LTA. Indeed, the 46–50 cm^{-1} band polarization-orientation dependence is similar to that of the E_2 band at 74–76 cm^{-1} or 77–79 cm^{-1} of Se_8 ⁴⁵ and the E_2 band at 153–155 cm^{-1} of S_8 [Fig. 3(c)]. The calculated Te_8 E_2 bond-bending mode frequency of 43.8 cm^{-1} is in good agreement with the observed 46–50 cm^{-1} band frequencies of LTA-Te. The splitting of the degenerate E_2 mode indicates structural distortion of Te_8 in LTA and therefore a reduced symmetry compared to the initial D_{4d} symmetry of the ring. For example, a slight reduction of the Te_8 symmetry from D_{4d} to C_{4v} would split the E_2 mode into B_1 and B_2 modes, active in *XX* and *XY* polarization configurations, respectively. This is exactly what we experimentally observed. Interestingly, the splitting of the Te_8 E_2 mode in LTA is very strong, $\sim 8.3\%$, in contrast to the weaker splitting of the corresponding E_2 modes of S_8 , $\sim 1.3\%$, and Se_8 , $\sim 2.6\%$.

The LTA-Te 64 cm^{-1} band displays polarization dependence similar to the A_1 symmetric bond-bending mode band of S_8 at ~ 221 cm^{-1} [Fig. 3(c)] and the analogous band of Se_8 at ~ 112 cm^{-1} .^{12,44,45} The frequency of 64 cm^{-1} is very close to the calculated frequency of 65.4 cm^{-1} of the Te_8 A_1 bond-bending mode [Fig. 2(b); 3(b)]. The experimental 64 cm^{-1} band is rather strong in the *aa* and *cc* polarization configurations and it is still not weak in the *cd* configuration, while the band completely disappears in the *ab* configuration. This is a consequence of the 2-dimensional character of the Raman tensor of this mode with only two strong diagonal tensor components in contrast to the rather 3-dimensional tensor of the A_1 bond-stretching mode [Fig. 2(c)]. Correspondingly, the single free-

standing Te_8 A_1 bond-bending mode band is strong in the *XX* polarization configuration and negligibly weak in the *ZZ* configuration with the zero *XY* and *XZ* activities [Fig. 2(b)]. Theoretical calculation of this mode activity in the RS spectra of LTA-Te [Fig. 3(b)] is also in very good agreement with the experiment, namely high *aa* activity and no *ab* activity. This is a clear indication of the orientation of the Te_8 4-fold axis along the 4-fold axis of LTA.

The experimental LTA-Te ~ 83 cm^{-1} band looks similar to the 248 cm^{-1} E_3 bond-bending mode band of S_8 [Fig. 3(c)]. It can be attributed to the E_3 bond-bending mode of Te_8 with a calculated frequency of 72.7 cm^{-1} . The ~ 29 cm^{-1} band of LTA-Te can be attributed to the E_2 torsional mode of Te_8 with a calculated frequency of 21.7 cm^{-1} . A broad ~ 25 cm^{-1} band, probably, originates from the superposition of the second component of the E_2 torsional mode and the ring librations in the LTA cavity.

As we mentioned above, the bands of LTA-Te in the bond-stretching mode region require more detailed consideration. Fig. 3(d) shows LTA-Te RS in this region taken at temperature $T \sim 77$ K and an excitation wavelength of 633 nm with better resolved bands due to their temperature-induced narrowing. Fig. 3(e) shows room temperature LTA-Te RS at the same excitation wavelength in a wider spectral range. Four bands at ~ 163 cm^{-1} , ~ 173 cm^{-1} , and ~ 183 cm^{-1} and its shoulder at ~ 186 cm^{-1} can be observed in Fig. 3(d). The shoulder is, probably, associated with the Te_8 E_2 bond-stretching mode (theoretical frequency of ~ 192 cm^{-1}). Taking into account our preliminary assignment of the 182–183 cm^{-1} and 162–163 cm^{-1}

bands to the Te_8 A_1 and E_3 modes, respectively, we have to attribute the 171–173 cm^{-1} band to the forbidden B_1 (theoretical frequency of $\sim 180 \text{ cm}^{-1}$) mode which can be activated due to the reduced symmetry of Te_8 in the LTA cavity, and we found that the bands in the bond-stretching mode region can be mixed.

The band at 182–183 cm^{-1} shows slightly stronger activity in the cc polarization configuration than that in the aa configuration, which is clearly seen in the room temperature RS of LTA-Te obtained by excitation with 633 nm light, as shown in Fig. 3(e). This can be associated with the mode mixing due to the reduced symmetry of the Te_8 rings. Interestingly, the 2nd order Raman band of Te_8 can be recognized in the cc configuration spectrum at $\sim 360 \text{ cm}^{-1}$ [Fig. 3(e)]. An even stronger 2nd order Raman band can be clearly seen at an excitation wavelength of 561 nm [Fig. 3(f), green curve] due to the resonant Raman enhancement. The band is very broad covering the spectral range from $\sim 300 \text{ cm}^{-1}$ to $\sim 375 \text{ cm}^{-1}$. Importantly, LTA bands do not contribute to the 2nd order Raman band of Te_8 since even the strongest LTA band at $\sim 490 \text{ cm}^{-1}$ [Fig. 3(f), black curve] does not contribute to LTA-Te RS obtained by excitation with 561 nm light.

Summarizing this section, the polarization dependence of the A_1 bond-bending mode Raman band of Te_8 indicates the orientation of the ring by its 4-fold axis along the LTA 4-fold axis, while that of the E_2 bond-bending mode suggests a noticeable symmetry reduction of the ring from the ideal D_{4d} point group symmetry. Te_8 bond-stretching modes are mixed due to the ring symmetry reduction. The Raman bands at 162–163 cm^{-1} and 171–173 cm^{-1} , probably, originate from the E_3 mode and formerly inactive B_1 mode while the Raman bands at 182–183 cm^{-1} and 186 cm^{-1} originate from the A_1 and E_2 modes of Te_8 . The originally inactive E_1 bond-stretching mode may also contribute to the observed Raman bands.

Optical absorption spectra of LTA-Te single crystals

Fig. 4(a and b) show the OAS of LTA-Te with the loading densities of ~ 3 atoms per cavity [LTA-Te(3)] and ~ 8 atoms per cavity [LTA-Te(8)]. Absorption bands at ~ 2.8 eV, 3.2 eV, 4.3 eV and 4.6 eV are observed in the OAS of LTA-Te(3), while absorption bands at ~ 2.6 eV, 3.2 eV and 4.4 eV are observed in the OAS of LTA-Te(8). The spectra indicate that the energy band gap decreases when the Te loading density increases. Moreover, LTA-Te(8) bands look slightly broader than those of LTA-Te(3). In particular, the absorption band doublet at ~ 4.3 eV and 4.6 eV, resolved in LTA-Te(3), looks like a single band at ~ 4.4 eV.

First, the observed experimental OAS spectra of LTA-Te [Fig. 4(a and b)] appeared to be in nearly perfect agreement with the theoretical OAS spectra of Te_8 [Fig. 2(e)]. The agreement with theoretical OAS for a single Te_8 ring is especially good for LTA-Te(3) with approximately two rings per five cavities. We have to note that surface light scattering, which

increases with the photon energy, contributes to the experimental spectra. In contrast, theoretical spectra do not include this effect and do not include high-energy transitions (>5.5 eV in Fig. 2(e)). This may produce an impression that the experimental and theoretical spectra are different. However, actually, they are very similar.

Secondly, differences between the OAS of LTA-Te(3) and LTA-Te(8) indicate a noticeable interaction between the Te_8 rings in neighbouring LTA cavities, in particular, for LTA-Te(8) with one ring per cavity. In Fig. 4(b), we show the spectral positions of the Raman excitation laser lines, clearly indicating rather strong absorption of LTA-Te(8) at a wavelength of 561 nm corresponding to the resonant Raman effect [Fig. 3(f)]. To avoid laser-induced heating/destruction of Te_8 , we used a very low 561 nm laser power of ~ 0.07 mW. At weaker LTA-Te absorption corresponding to wavelengths of 785 nm and 633 nm [Fig. 4(b)], we were able to use higher laser powers of ~ 1 mW and ~ 0.3 mW, respectively [Fig. 3(f)].

In order to examine a possible effect of the ring interaction on the OAS spectrum of LTA-Te, we theoretically considered three possible Te_8 ring dimers with a separation of ~ 1.2 nm between the ring centres [Fig. 4(c)]. The corresponding calculated dimer OAS spectra are shown in Fig. 4(d). The effect of the ring interaction is indicated by the absorption enhancement with an increase in the ring interaction, which is the weakest in the “face-to-face” configuration and the strongest in the “side-to-side” configuration. Due to the inter-ring interaction, the unoccupied MOs of the Te_8 rings organize conduction bands of the LTA-Te cluster crystals, while the highest occupied MOs organize its valence band. The conduction and valence bands should be rather narrow. Therefore, a variety of non-linear electric conductance effects predicted for narrow semiconductor superlattice mini-bands⁴⁶ can be expected in the LTA-Te electric properties. A negative differential electric resistance of LTA-Te observed in ref. 47 confirms this.

Te_8 interaction with LTA and its librations vs. temperature

According to the calculated Te_8 structure [Fig. 2(a)], the diameter of the ring, including the van der Waals radius of ~ 0.2 nm of the Te atom, is ~ 1 nm, which is slightly smaller than the LTA large cavity diameter. Therefore, one can expect that the Te_8 ring is slightly shifted from the cavity center along the 4-fold LTA axis similar to the S_8 and Se_8 rings^{12,44,45} but not as much as in the cases of S_8 and Se_8 . This should cause Te_8 symmetry reduction from D_{4d} at least to C_{4v} [Fig. 5(a)]. We imposed the bond angle changes in order to simulate such structural distortion. The angles of type ϕ_1 were enlarged while those of type ϕ_2 were reduced. The second nearest neighbor distances $R = 0.418$ nm and 0.446 nm in Fig. 5(a) correspond to $\Delta\phi = \phi_1 - \phi_2 = 5^\circ$ ($R = 0.432$ nm for undisturbed Te_8).

The experimentally observed splitting of the E_2 bond-bending mode [Fig. 3(a, d and e)] is, definitely, a sign of D_{4d}

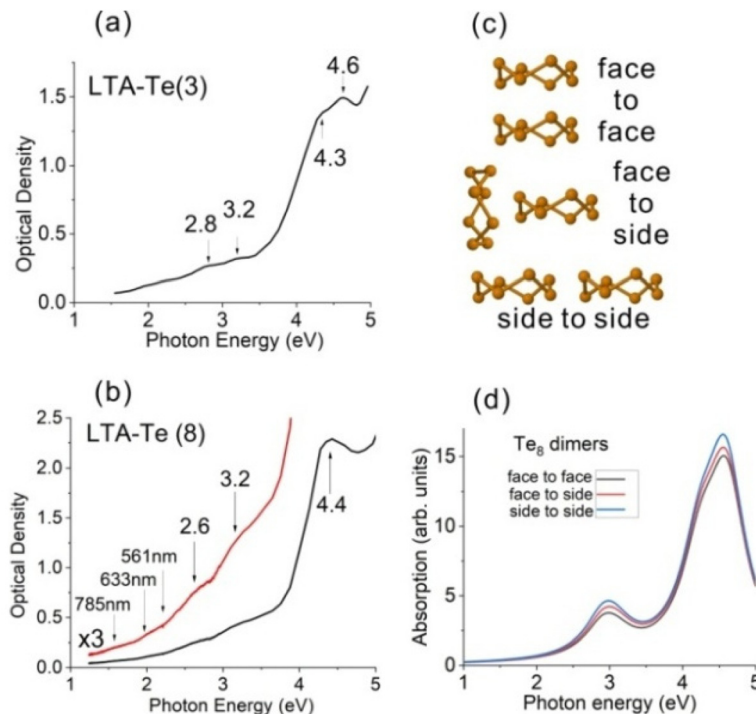


Fig. 4 Experimental absorption spectra of LTA-Te(3) (a) and LTA-Te(8) (b); three Te_8 dimers considered for the calculation of the effect of ring interaction on the absorption spectrum (c); and calculated absorption spectra of the considered Te_8 dimers including transitions with photon energies of <5 eV (d).

symmetry reduction. It is important to verify whether the observed E_2 mode splitting fits the ring structure distortion shown in Fig. 5(a) or not. Due to the Te_8 symmetry reduction from D_{4d} to C_{4v} , the E_2 bond-bending mode splits into B_1 and B_2 modes, displaying XX and XY Raman activities, respectively. We performed calculations of the B_1 and B_2 mode frequencies and intensities as a function of the distortion $\Delta\phi$. The results are shown in Fig. 5(b). The B_1 mode frequency increases while the B_2 mode frequency decreases with $\Delta\phi$. This corresponds well to our experimental observation for the E_2 bond-bending mode with the LTA-Te *aa*-active component frequency higher than that of the *ab*-active component. Thus, we conclude that the Te_8 structural distortion shown in Fig. 5(a) reasonably describes the experimentally observed splitting of the E_2 bond-bending mode of Te_8 . However, even stronger symmetry reduction of Te_8 is possible.

As we mentioned in the Introduction, Raman polarization-orientation spectroscopy in application to the zeolite-confined cluster structure study can be considered in some sense as an alternative to XRD. Indeed, using Raman polarization-orientation spectroscopy, we can determine a type of cluster (molecule) and its orientation in a zeolite. Then we can suggest its possible location in the zeolite when we know its size. However, we can only guess what sort of interaction occurs between the Te atoms and the zeolite. For understanding this point, the XRD data would be very useful. However, a structural analysis of LTA-Te by XRD is rather difficult because Te is distributed statistically inside the cavities. The resulting scatter-

ing power of Te sites with partial occupancy is comparable to those of Na^+ cations and water molecules filling spaces not occupied by Te.

The ~ 20 micron size of LTA-Te is insufficient for single-crystal XRD, while powder XRD of zeolites with clusters is, definitely, not effective for the examination of the cluster structure. It shows that the zeolite filling with clusters or molecules causes just changes in the relative intensities of the XRD peaks like, for example, in ref. 48 and its ESI, where a number of different types of zeolites, including LTA, with/without sulphur clusters were examined using powder XRD. In contrast, XRD analysis of ~ 65 micron size LTA single crystals containing sulphur clearly showed double S_8 ring cluster formation in each large cavity,⁴⁹ which is in agreement with the published RS results^{12,50,51} and this work [Fig. 3(c)].

An XRD study of ~ 60 micron size LTA-Te single crystals was performed in ref. 52. The XRD analysis suggested the interaction of Te atoms with the zeolite oxygen atoms and cations, while it showed low occupancy and high temperature factors for Te atomic sites. Therefore, the authors could not make any univocal conclusion about the Te cluster structure and they proposed several options including Te_8 rings.

Using the RS of LTA-Te, the interaction between the Te_8 ring and the zeolite can be probed *via* observation of the Te ring librations *vs.* temperature. Taking into account the S_8 libration frequency of ~ 37 cm^{-1} [Fig. 3(c)] and assuming approximately the same van der Waals interaction force with LTA for both S_8 and Te_8 , we can roughly estimate the expected

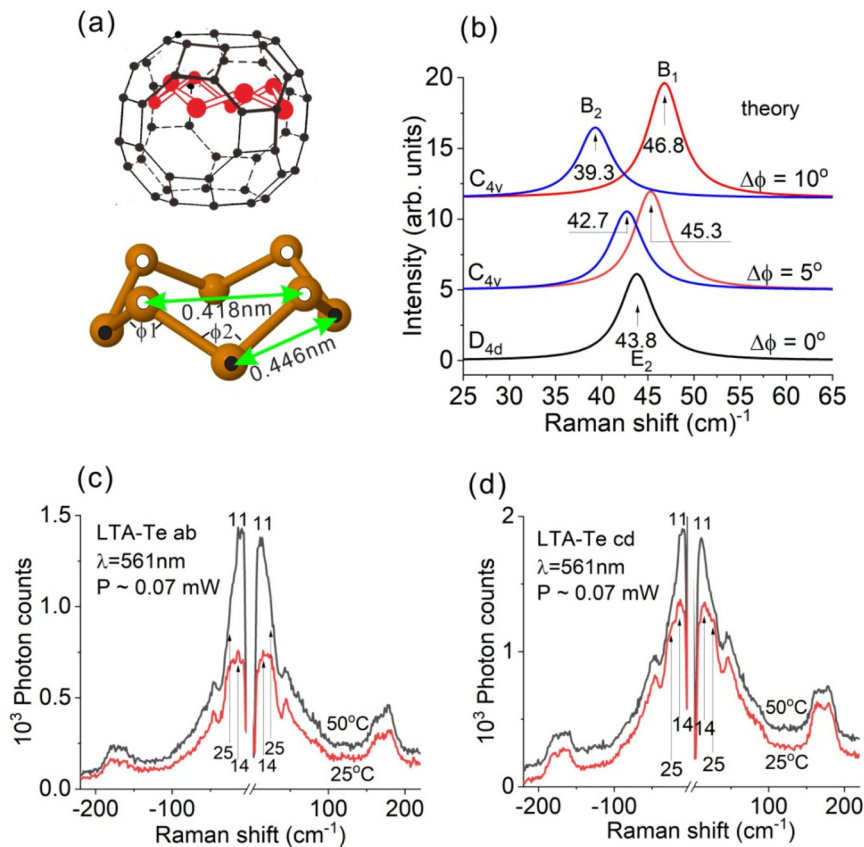


Fig. 5 Te_8 ring shifted up from the centre of the LTA large cavity (top) with the C_{4v} structural distortion corresponding to the bond angle difference $\Delta\phi = \phi_1 - \phi_2 = 5^\circ$ (bottom), the white and black spots show Te atoms in the top and bottom layers, respectively (a); calculated RS spectra of the Te_8 (D_{4d}) E_2 bond-bending mode (black) and Te_8 (C_{4v}) B_1 (red) and B_2 (blue) bond-bending modes (b); and experimental low-frequency RS of LTA-Te at stage temperatures of $\sim 25^\circ\text{C}$ and $\sim 50^\circ\text{C}$ for *ab* (c) and *cd* (d) polarization configurations.

Te_8 room temperature libration frequency as $\sim 18.5\text{ cm}^{-1}$ using the square root of the Te and S mass ratio, which is ~ 2 .

In the LTA-Te RS spectra obtained by excitation with 785 nm light, Te_8 librations, probably, contribute to the broad $\sim 22\text{--}25\text{ cm}^{-1}$ band [Fig. 3(a)]. Instrumental limitations did not allow RS detection at frequencies $< 20\text{ cm}^{-1}$ at this excitation wavelength. In contrast, such measurements were available at 561 nm light excitation. A weak point of this wavelength is that the corresponding Te_8 absorption and light-induced heating are not negligible.

Fig. 5(c and d) show the Stokes and anti-Stokes low-frequency RS of LTA-Te in *ab* and *cd* polarization configurations at $\sim 25^\circ\text{C}$ and $\sim 50^\circ\text{C}$ temperatures of the heat-controlling table. We roughly estimated the laser-induced heating from the Stokes/anti-Stokes intensity ratio as $\sim 20\text{--}30^\circ\text{C}$, the value that should be added to the heating table temperature. In the RS taken at $\sim 25^\circ\text{C}$ table temperature, one can see a $\sim 25\text{ cm}^{-1}$ band in combination with a $\sim 14\text{ cm}^{-1}$ band for both configurations. The last band, definitely, originates from the Te_8 librations. With the increase of the table temperature to $\sim 50^\circ\text{C}$, it shifts to a lower frequency of $\sim 11\text{ cm}^{-1}$ and displays significant enhancement. This corresponds to the intensification of librations and the reduction of the ring interaction with LTA.

The position of the $\sim 25\text{ cm}^{-1}$ feature remains unchanged, confirming its origin from the internal torsional E_2 mode of Te_8 .

Thus, the Te_8 libration amplitudes strongly increase with a moderate increase in the LTA temperature. This corresponds to a rather weak van der Waals interaction of Te_8 with LTA. At room temperature, the libration amplitudes are, probably, quite strong as well. This explains high temperature factors observed for Te sites,⁵² which complicates the XRD analysis of LTA-Te. In contrast, this effect does not cause any trouble for polarization-orientation Raman spectroscopy, which appears to be a very fruitful method of zeolite-confined cluster study providing information on the cluster structure with distortions, orientation and interaction with zeolites.

Conclusions

In summary, the electronic and vibrational properties of Te_8 rings were studied experimentally using Raman and absorption optical spectroscopy and theoretically using density functional theory. A good agreement is obtained between (1) the theoretical optical absorption and Raman spectra of isolated Te_8 and (2) those of Te_8 formed in the large cavities of the

zeolite LTA. Owing to zeolite utilization, this is a big step in the experimental study of Te₈, which is limited by the fabrication of exotic compounds like Cs₃Te₂₂⁵³ or cluster registration in molecular beams.⁵⁴ Polarization–orientation Raman spectra of LTA-Te show that the rings are oriented by their major axes along the 4-fold axes of LTA. The observed splitting of the Te₈ bond-bending modes suggests the reduction of the ring *D*_{4d} symmetry in the LTA cavity. The bond-stretching modes demonstrate mixing due to the reduced symmetry of the ring and, probably, the essential anharmonicity of the vibrations. Even moderate heating of LTA-Te causes significant intensification of Te₈ librations in the LTA cavities, which corresponds to a rather weak van der Waals interaction between the ring and the zeolite. Experimental optical absorption spectra of LTA-Te display dependence on the Te loading density due to the electronic interaction of Te₈ from neighbouring cavities, which increases with an increase in the loading density. The effect is confirmed using DFT calculations of the absorption spectra of Te₈ dimers. The observed Te₈ ring–ring interaction is beneficial for the formation of cluster crystals with narrow electron bands and predicted non-linear electric properties. On the other hand, our results can be interesting in the sense of relationships with other inorganic 8-membered ring clusters such as Pd₈.⁵⁵

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

The authors thank V. P. Petranovskii for LTA crystal synthesis and A. V. Fokin for the Te vapour adsorption procedure.

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