

Investigation of strong shock wave interactions with CeO₂ ceramic

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Abstract: Strong shock wave interactions with ceramic material ceria (CeO₂) in presence of O₂ and N₂ gases were investigated using free piston driven shock tube (FPST). FPST is used to heat the test gas to very high temperature of about 6800–7700 K (estimated) at pressure of about 6.8–7.2 MPa for short duration (2–4 ms) behind the reflected shock wave. Ceria is subjected to super heating and cooling at the rate of about 10⁶ K/s. Characterization of CeO₂ sample was done before and after exposure to shock heated test gases (O₂ and N₂). The surface composition, crystal structure, electronic structure and surface morphology of CeO₂ ceramic were examined using X-ray photoelectron spectroscopy (XPS), X-ray diffraction (XRD), Fourier transform infrared (FTIR) spectrometry, scanning electron microscopy (SEM) and high resolution transmission electron microscopy (HRTEM). Results obtained from the experimental investigations show that CeO₂ can withstand high pressure accompanied by thermal shock without changing its crystal structure. Reducible CeO₂ releases lattice oxygen making it possible to shift between reduced and oxidized states upon the interaction with shock heated gas. Due to such reaction mechanism, CeO₂ ceramic undergoes nitrogen doping with decrease in lattice parameter. Investigations reveal that CeO₂ retains its crystal structure during strong shock interaction, even at elevated pressure.

Keywords: shock tube; shock interaction; high enthalpy gas; ceria; nitrogen-doped; surface reaction

1 Introduction

Ceria based materials have been explored for several applications such as heterogeneous catalysis [1,2], gas sensors [3], fuel cells [4], bio-materials [5,6], bio-photonics [7], etc. In many applications, knowledge of the cerium oxidation state is necessary for understanding the properties of the ceria compounds. These ceria based catalysts are very useful as high temperature materials for space applications

and also for three-way catalysis. Among rare earth oxides, CeO₂ ceramic material draws special attention due to its role as a promoter for the control of toxic emission from automobile exhaust for many years. Ceria is usually treated as an active support for precious metal as well as base metal catalyst in heterogeneous catalysis. Reducible CeO₂ can release lattice oxygen making it possible to shift between reduced and oxidized states (Ce⁴⁺(4f⁰) ↔ Ce³⁺(4f¹)) indicating high endurance of ceria towards fluctuating oxygen vacancies under oxygen excess and deficient conditions. The extent of lattice oxygen that can be released and restored is known as “oxygen storage capacity (OSC)” [8]. This unique property of ceria has

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made it a special component for three-way catalyst since it can store oxygen during oxygen-lean conditions and release oxygen during oxygen-rich conditions. Under oxygen-rich condition, cerium ion is in 4+ oxidation state, and reduces partially to 3+ oxidation state under oxygen-lean condition. In addition, ceria keeps the same cubic crystal structure even during the alternate storage and release of oxygen and its volume change is observed to be very small.

Stoichiometric CeO₂ has a cubic fluorite lattice (space group *Fm3m*) with four cerium and eight oxygen atoms per unit cell. The Ce⁴⁺ cations occupy the face centered cubic lattice sites, and the oxygen anions (O²⁻) are located at the eight tetrahedral sites. From structural considerations, the addition or removal of oxygen atoms should involve a minimal reorganization of the skeleton arrangement of the cerium atoms [9,10]. This structural property should definitely facilitate the excellent reversibility of the reduction–oxidation process. The unit cell is constructed out of eight unit cells of CeO₂ with 25% of the oxygen vacancies created in a particular arrangement and the cations are six coordinated [11]. High pressure study of bulk CeO₂ at room temperature shows phase transformation from cubic fluorite to orthorhombic phase on the application of 35 GPa pressure. Study at higher temperature (1473 K) at 14.5 GPa shows similar phase transformation, but at room temperature on decompression it comes back to cubic fluorite structure [12].

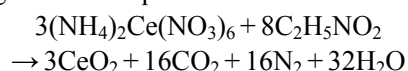
Free piston driven shock tube (FPST) which is a part of hypersonic shock tunnel 3 (HST3) has been used to investigate the effect of strong shock compression on several nano materials [13–16]. Thermal protection system (TPS) materials are generally tested in presence of air for space applications. In general, shock heated O₂ and N₂ (instead of air) gases are used to study aerothermodynamic reactions during re-entry of space vehicles. Studies of oxidation and nitridation reactions are essential to understand the catalytic activity of high temperature ceramic materials. The novel methods of investigating the interaction of real gases with the materials for short duration using shock tubes are not reported in the literature. In this paper we report one such study where we present the experimental results obtained on the interaction of CeO₂ with strong shock heated O₂ and N₂ gases for short duration (2–4 ms) using FPST.

2 Experimental methods

Ceria ceramic nanopowders were synthesized by the solution combustion method to study their behavior on shock loading. Experiments were performed on CeO₂ using shock heated O₂ and N₂ at high temperature and pressure. High enthalpy thermodynamic conditions generated in FPST were utilized to investigate the shock interaction behavior of CeO₂. The characterization of CeO₂ was carried out before and after the interaction with shock heated test gases using different experimental methods.

2.1 Solution combustion synthesis of nanopowders of ceramic material CeO₂

CeO₂ was synthesized by the solution combustion method [17]. The chemical combustion reaction led to the formation of solid oxide by using 5.4826 g (0.01 mol) of ammonium cerium (IV) nitrate (NH₄)₂Ce(NO₃)₆·6H₂O and 1.98 g (0.01 mol) of glycine (C₂H₅NO₂) dissolved in distilled water, taken in a crystallizing dish. The compounds were dissolved into clear solution. The solution was rapidly heated in a furnace at 723 K. After dehydration, the solution was ignited into a flame, rising the temperature to about 1273 K for 1–2 min, leaving behind pale yellow color nanocrystalline ceramic powders of CeO₂ as per the following chemical equation:



Nanocrystalline ceramic powders were then made into pellets using hydraulic press unit by applying 75 kN load before exposing to shock heated test gases. Pellets were made by using 0.2 g CeO₂. Synthesized CeO₂ was exposed to strong shock heated oxygen and nitrogen gases using FPST.

2.2 Experimental facilities and conditions of test gases

FPST consists of a high pressure gas reservoir, compression tube filled with driver gas (helium for the present study) and shock tube with a provision to mount the test sample on the end flange. Operation of the FPST involves placing the piston at mounting location of the gas reservoir and holding it rigidly by vacuum after evacuation. Aluminum diaphragm is placed in between the compression tube filled with helium gas at 1 atm and the shock tube. After evacuating the shock tube, gas handling system is used

to fill the test gas at a required pressure. Ultra-high pure (UHP) O₂ or N₂ is purged many times. The shock tube is initially evacuated using vacuum pump and then filled with test gas (O₂ or N₂) at required pressure. Sudden supply of the high pressure gas by opening valves behind the piston sets its motion in the compression tube and as a result, the piston gets maximum acceleration. Motion of 20 kg piston in compression tube adiabatically compresses the helium gas and thereby increases the pressure and temperature. This high pressure and high temperature helium gas bursts the primary diaphragm. It produces a normal primary shock wave, traveling into the driven section

of a shock tube filled with test gas, which is reflected finally at the end of the shock tube to generate higher stagnation temperature and pressure. The reflected shock wave can be used to produce a stagnation enthalpy of about 10 MJ/kg. A simple working principle of FPST was described in earlier references [18,19]. Detail experimental procedure, working condition and development of the FPST were described elsewhere [20,21]. The schematic diagram of FPST is shown in Fig. 1. The panoramic view of the completely assembled 21 m long FPST (HST3) is shown in Fig. 2.

The temperature behind the reflected shock wave (T_5)

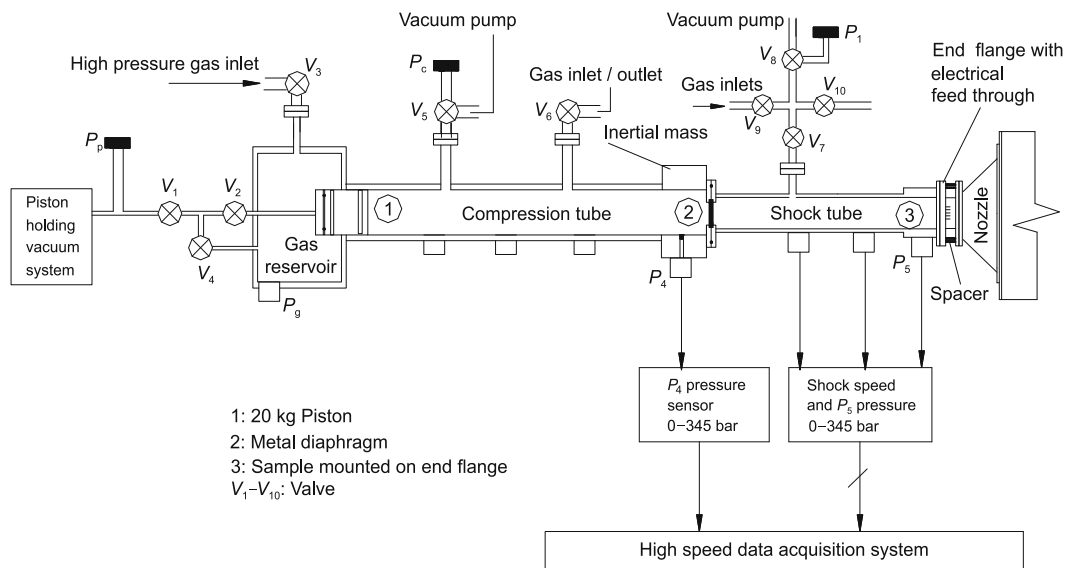


Fig. 1 Schematic diagram of FPST.

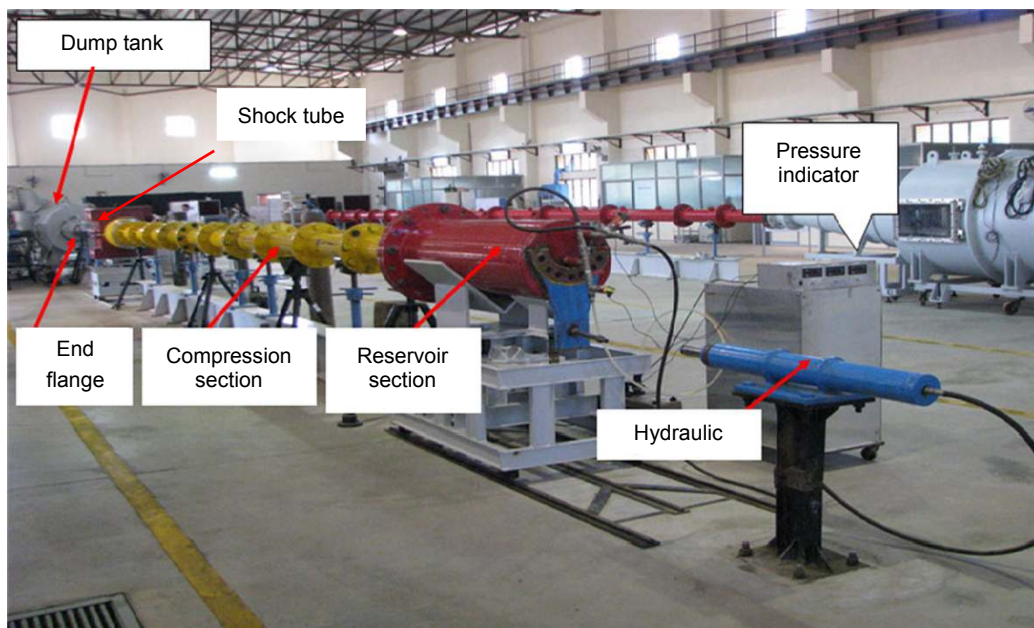


Fig. 2 Panoramic view of completely assembled 21 m long FPST.

and the corresponding specific enthalpy of the test gas at the end of the shock tube are estimated using normal shock relation [22]:

$$\frac{T_5}{T_1} = \frac{[2(\gamma - 1)M_s^2 + (3 - \gamma)][(3\gamma - 1)M_s^2 - 2(\gamma - 1)]}{(\gamma + 1)^2 M_s^2}$$

where γ is the specific heat ratio of the test gas; M_s is the measured shock Mach number; and T_1 is the initial temperature of the test gas in the shock tube.

Experiments were performed by filling helium gas at 0.1 MPa in the compression tube while shock tube was filled with test gas at initial pressure of 0.01 MPa. The shock heated O₂ test gas was made to react for 2–4 ms with the CeO₂ pellets mounted on the end flange of the shock tube. Typical experimentally measured and recorded parameters are shock speed 2440 m/s corresponding to shock Mach number of 6.9 and reflected shock pressure ~6.8 MPa. Hence in this experiment, the sample was subjected to the reflected shock pressure of 6.8 MPa measured using pressure sensor mounted at the end of the shock tube and estimated reflected shock temperature of about 6400 K for about 2–4 ms. Similar experimental conditions were set for nitrogen test gas with corresponding values of 2630 m/s shock speed, 7.4 shock Mach number and reflected shock pressure 7.2 MPa, at estimated reflected shock temperature of about 7700 K. The typical shock speed and reflected shock pressure signals have been recorded by using piezoelectric pressure transducer, as shown in Figs. 3(a) and 3(b), respectively. The ceramic test compound was mounted at the end of the shock tube in the form of pellets for interaction with the shock heated test gases.

2.3 Characterizations

Different experimental techniques were used to

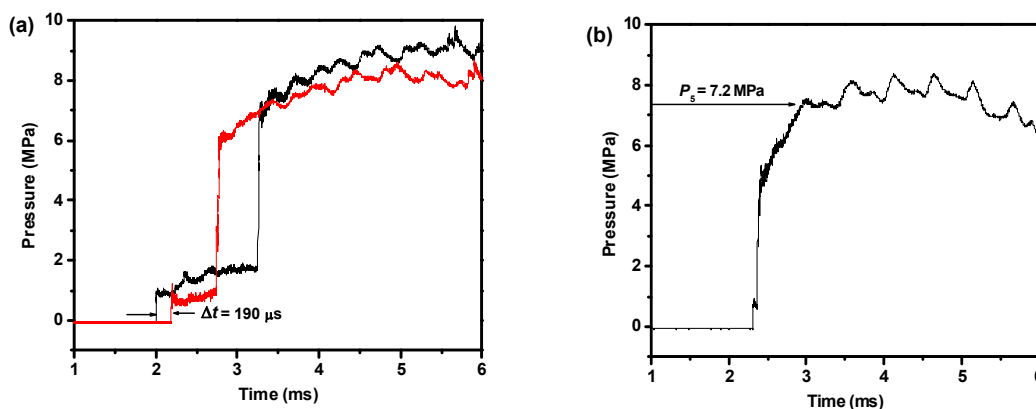


Fig. 3 Typical pressure signals acquired from pressure transducers: (a) time history data for the shock to travel 0.5 m distance; (b) reflected shock pressure (P_s) data.

characterize the sample before and after exposure to shock heated test gases. Electronic structure of the ceramic was characterized by X-ray photoelectron spectroscopy (XPS) using Al K α radiation (1486.6 eV) where C(1s) at 284.5 eV was taken as reference and the accuracy of the binding energy (BE) reported here was ± 0.1 eV (MultiLab 2000, Thermo Fisher Scientific). X-ray diffraction (XRD) of the compound was recorded (Phillips X'Pert diffractometer) using Cu K α radiation at scan rate of 0.25 ($^\circ$)/min with 0.01 $^\circ$ step size in the 2θ range between 20 $^\circ$ and 80 $^\circ$. Fourier transform infrared (FTIR) spectra were recorded using FTIR spectrometer (Perkin Elmer, SPECTRUM-1000) in the 400–4000 cm^{-1} range by preparing KBr pellets. Scanning electron microscopy (SEM, Philips SIRION) and high resolution transmission electron microscopy (HRTEM, Technai T-20) operated at 200 kV were used to study the morphology and nano crystalline structure of CeO₂ before and after shock treatment.

3 Results and discussion

XPS spectra of Ce(3d) and N(1s) core levels of as-prepared and shock exposed CeO₂ are shown in Fig. 4. Ce(3d_{5/2}) peak at 881.7 eV along with satellite peaks at 6.4 eV and 16 eV below the main peak is characteristic of Ce⁴⁺ in CeO₂ [23]. Characteristic satellite peak observed at 916.8 eV as shown in Fig. 4(i)(a) confirms that Ce is in 4+ state in this compound. After exposure to shock heated test gases like O₂ or N₂, BE of the main and satellite peaks of Ce(3d) remain the same as shown in Figs. 4(i)(b) and 4(i)(c) indicating that Ce is in 4+ state before and after exposure to shock heated gases. Thus, XPS results confirm no change in the electronic structure of CeO₂

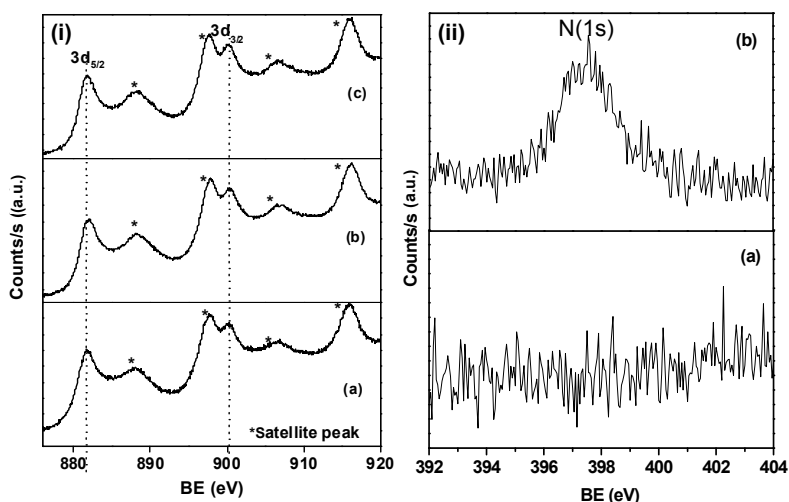


Fig. 4 (i) XPS spectra of Ce(3d): (a) as-prepared ceria ceramic, (b) after exposure to shock heated O₂ and (c) after exposure to shock heated N₂. (ii) XPS spectra of N(1s): (a) as-prepared sample and (b) after exposure to shock heated N₂.

in presence of shock heated O₂. Core level N(1s) spectra in CeO₂ before and after exposed to N₂ are shown in Fig. 4(ii). N(1s) peak is not observed in as-prepared CeO₂ sample as shown in Fig. 4(ii)(a). On the other hand, core level N(1s) peak at 397.5 eV after exposure to nitrogen is attributed to the formation of Ce–N bond due to the doping of nitrogen during shock treatment as shown in Fig. 4(ii)(b). XPS of N(1s) in nitrogen doped cerium in presence of NH₃ was reported in the literature [24]. The formation of CeO_{2-x}N_x (nitrogen doped CeO₂) under strong shock compression is the signature of three-body non-catalytic surface reactions.

Powder XRD patterns of the nanoceramic CeO₂ before and after exposure to shock heated test gases are shown in Fig. 5. All the peaks are indexed into fluorite structure according to JCPDS No. 81-0792, and no impurity peak is detected in the XRD pattern of as-prepared CeO₂ given in Fig. 5(a). Thus, the compound explicitly crystallizes in fluorite structure (space group *Fm*3*m*, lattice parameter *a*=5.411 Å). The diffraction lines are sharper and more intense in comparison with that of as-prepared ceramic due to melting and re-crystallization with increase in the size of crystallite. The characteristic diffraction lines are assigned with *hkl* values for CeO₂. XRD of CeO₂ ceramic remains cubic with increase in the intensity of diffraction lines after exposure to shock heated O₂ and N₂ as shown in Figs. 5(b) and 5(c). Also there is a clear shift of 0.3° towards the higher Bragg angle with the shock heated N₂ gas. With shock heated O₂ gas, the oxidation state of CeO₂ remains with Ce⁴⁺, and Ce³⁺

can go to Ce⁴⁺ state. However there is no shift in the diffraction lines due to this process as shown in Fig. 5(b). During the interaction of CeO₂ with shock heated N₂, the oxide ion vacancies in the CeO₂ lattice due to Ce³⁺ ions are replaced by nitrogen atoms and form CeO_{2-x}N_x. Doping of nitrogen atoms to CeO₂ during shock compression reduces the lattice parameter by 0.033 Å with a clear shift of 0.3° towards higher angle in the diffraction lines as shown in Fig. 5(c). The study shows that there is no phase transformation of CeO₂ after subjecting CeO₂ to shock compression in presence of O₂ and N₂ gases.

FTIR spectra of as-prepared CeO₂ and the samples

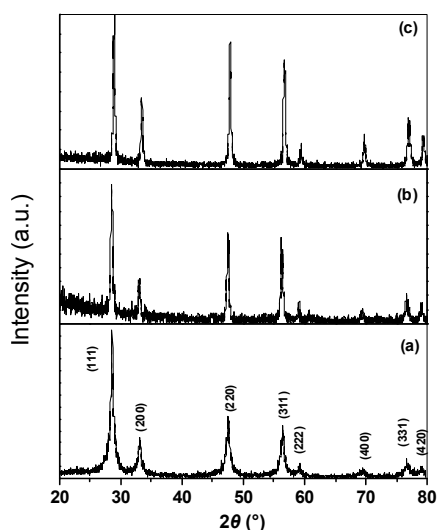


Fig. 5 XRD of CeO₂: (a) before exposure to shock, (b) after shock exposure to O₂ and (c) after shock exposure to N₂.

after shock exposure to O₂ and N₂ gases are shown in Figs. 6(a), 6(b) and 6(c), respectively. Small peaks at 2960 cm⁻¹ and 2850 cm⁻¹ are due to asymmetric stretching vibrations of -CH₂ bonds of glycine. The wide peak at 3000–3600 cm⁻¹ range corresponds to -OH vibration band of H₂O present in the compound. The peaks around 3393 cm⁻¹ and 1623 cm⁻¹ are attributed to the O-H stretching and bending vibrations, respectively, and are in comparison with the literature [25,26]. There are shifts in O-H frequencies at 1623 cm⁻¹ and 3393 cm⁻¹ towards higher wavelength region after interaction with O₂. Peak around 3700 cm⁻¹ as shown in Fig. 6(b) is due to bi- or tri-coordinated O-H species formed in presence of huge O₂ pressure during shock wave treatment [27]. The results demonstrate that as-prepared sample has absorbed H₂O molecules on the surface of CeO₂. The band at 1385 cm⁻¹ is due to CO₂ present on the surface after exposure to shock heated O₂ gas. The absorption peaks of Ce-N and Ce-O bonding located at 900 cm⁻¹ and 1150 cm⁻¹ respectively indicate the presence of distinct nitrogen doped CeO₂ (CeO_{2-x}N_x) and CeO₂ phase as shown in Fig. 6(c). The main peak at 850–450 cm⁻¹ is due to the presence of Ce-O bond.

SEM micrographs of ceramic CeO₂ samples before and after exposure to shock heated test gases are shown in Fig. 7 with 10000× magnification. Micrograph of CeO₂ before exposure to shock is shown in Fig. 7(a). When normal shock wave impinges on the sample for a short duration of 2–4 ms, the re-crystallization phenomenon spreads throughout the

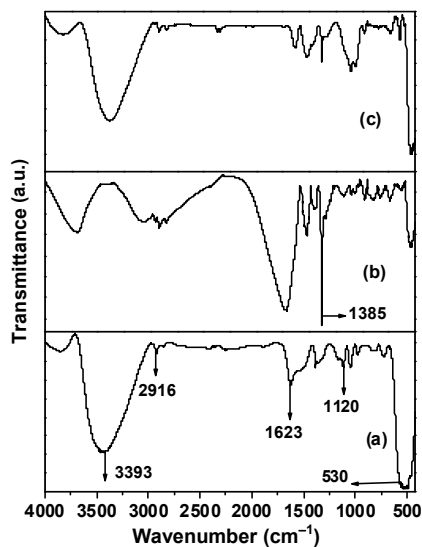


Fig. 6 FTIR spectra of CeO₂: (a) as-prepared sample, (b) after exposure to shock heated O₂ and (c) after exposure to shock heated N₂.

surface of the CeO₂ pellets. After exposure to shock heated O₂ and N₂ gases, the compound melts and the surface morphology of re-crystallized sample are shown in Figs. 7(b) and 7(c), respectively. During the exchange of lattice oxygen and formation of CeO_{2-x}N_x, pores and cracks of different sizes are developed on the surface of CeO₂ (Figs. 7(b) and 7(c)).

Bright field image, HRTEM image and electron diffraction pattern of as-prepared CeO₂ are shown in Fig. 8. Bright field image clearly shows that the sizes of ceramic crystallites are in the range of 10–15 nm as shown in Fig. 8(a). Ring type selected area electron diffraction pattern is indexed to fluorite structure which is shown in the inset of Fig. 8(a). Bright field image, HRTEM image and electron diffraction pattern of CeO₂ after exposure to shock heated O₂ gas are shown in Fig. 9. Crystallites are in the range of

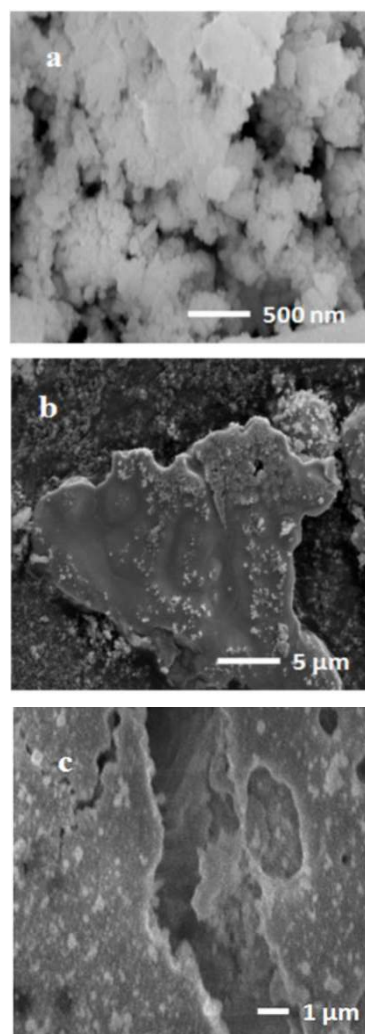


Fig. 7 SEM micrographs of CeO₂: (a) before shock, (b) after exposure to shock heated O₂ and (c) after exposure to shock heated N₂.

20–60 nm as evident from bright field image as shown in Fig. 9(a). This indicates that crystallite sizes are increased due to shock interaction. Electron diffraction pattern is indexed to fluorite structure and more intense bright spots are seen. Lattice fringe spacing of 3.11 Å is shown in Fig. 9(b), which corresponds to d_{111} plane of fluorite CeO_2 lattice (as shown in the inset of Fig. 9(a)). Bright field image, HRTEM image and electron diffraction pattern of CeO_2 after exposure to shock

heated N_2 gas are shown in Fig. 10. Bright field image shows that crystallites are in the range of 20–80 nm, which indicates an increase in crystallite size. Electron diffraction shows clear rings with more intense bright spots, which are indexed to different crystal lattice. Lattice fringe spacing in HRTEM image is again 3.11 Å as shown in Fig. 10(b), which corresponds to d_{111} plane of fluorite CeO_2 lattice (as shown in the inset of Fig. 10(a)).

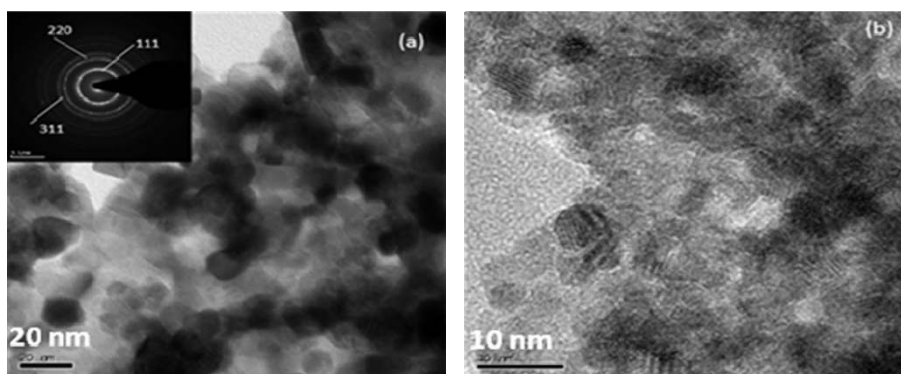


Fig. 8 As-prepared CeO_2 ceramic: (a) bright field image (inset: electron diffraction pattern) and (b) HRTEM image.

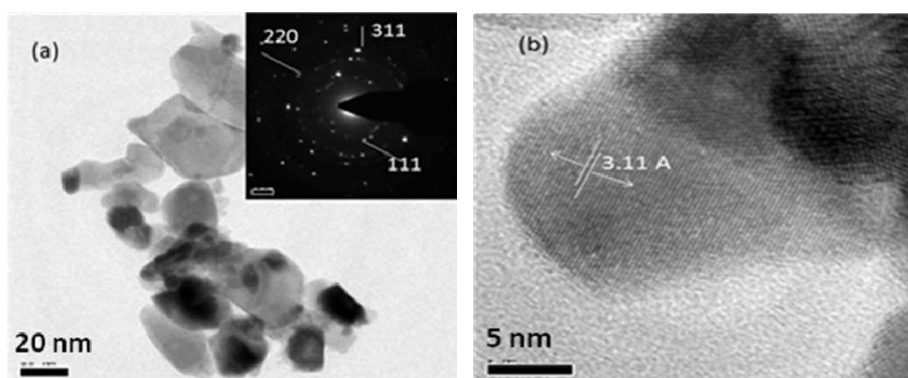


Fig. 9 CeO_2 sample after shock exposure to O_2 : (a) bright field image (inset: electron diffraction pattern) and (b) HRTEM image.

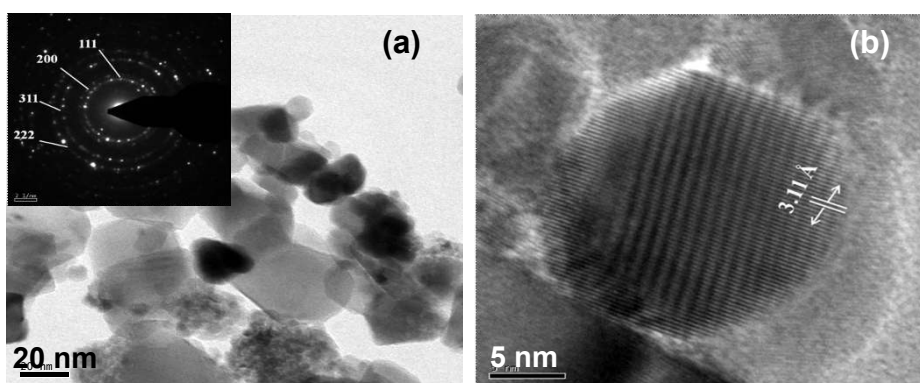
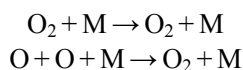


Fig. 10 CeO_2 sample after shock exposure to N_2 : (a) bright field image (inset: electron diffraction pattern) and (b) HRTEM image.

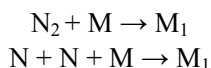
CeO₂ undergoes three-body surface recombination reaction with non-dissociated and dissociated gas after shock treatment as follows.

(1) Reaction with non-dissociated and dissociated O₂ gas:



In both the cases, the third body M (CeO₂) retains the same chemical composition and electronic structure. The crystal structure also remains as fluorite on shock loading. In practical applications, the extent of CeO₂ reduction occurring in the catalysis process is very small compared to complete reduction to Ce₂O₃ (or CeO_{1.5}). On shock heating CeO₂ → CeO_{2-δ} + δ/2O₂ transformation takes place, where only partial oxygen vacancies are created by releasing lattice oxygen from CeO₂ by retaining its skeleton with fluorite structure.

(2) Reaction with non-dissociated and dissociated N₂ gas:



In these cases, the third body M (CeO₂) will change to M₁ (CeO_{2-x}N_x) when oxygen vacancies are occupied by nitrogen atom. The chemical composition and electronic structure of new CeO_{2-x}N_x were characterized by XRD, FTIR and XPS studies.

4 Conclusions

Shock-tube techniques are very useful to produce variable thermal shock for short duration at elevated pressure. Shock tubes are also used to study three-body catalytic and non-catalytic surface reactions with high enthalpy test gases. Three-body reactions were studied on ceramic CeO₂ sample with shock heated O₂ and N₂ test gases. The results obtained from different experimental techniques like XPS, XRD, FTIR, SEM and HRTEM show no change in the phase of CeO₂ before and after exposure to shock wave in the presence of O₂ gas. Further due to strong shock compression in the presence of N₂, the peak positions of diffraction lines are observed to be shifted equally towards the higher Bragg angle and XPS confirms that nitrogen is incorporated into the cubic fluorite structure. Nitrogen doped CeO₂ is also confirmed by FTIR studies through the characteristic Ce–N bond. SEM and HRTEM studies show increase in the crystallization of CeO₂ after shock treatment. The experimental results show the effect of strong shock heated gas interactions with CeO₂.

Acknowledgements

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