Design of nanoscaled heterojunctions in precursor-derived $t$-$\text{ZrO}_2$/SiOC(N) nanocomposites: Transgressing the boundaries of catalytic activity from UV to visible light

Shakthipriya Bhaskar$^1$, Eranezhuth Wasan Awin$^1$, K. C. Hari Kumar$^1$, Abhijeet Lale$^2$, Samuel Bernard$^2$ & Ravi Kumar$^1$*

In this work, nanocomposites made of nanosized zirconia crystallized in situ in an amorphous silicon oxycarbo(nitride) (SiOC(N)) matrix have been designed through a precursor route for visible light photocatalytic applications. The relative volume fraction of the starting precursors and the pyrolysis temperatures not only influences the phase fraction of zirconia crystallites but also stabilizes the tetragonal crystal structure of zirconia ($t$-$\text{ZrO}_2$) at room temperature. The presence of carbon in interstitial sites of zirconia and oxygen vacancy defects led to drastic reduction in the band gap (2.2 eV) of the nanocomposite. Apart from being a perfect host avoiding sintering of the active phase and providing mechanical stability, the amorphous matrix also reduces the recombination rate by forming heterojunctions with $t$-$\text{ZrO}_2$. The reduction in band gap as well as the formation of heterojunctions aids in harnessing the visible light for photocatalytic activity.

Large scale industrialization and urbanization has led to extensive pollution and contamination of rivers and water bodies, posing a threat to future generations and sustainable development$^1$. Chemical industries, leather factories, paint manufactures, dyeing and plating units release harmful untreated waste directly into our water bodies$^2$–$^4$. These are non-biodegradable and toxic to humans as well as flora and fauna. Filtration with activated carbon, coagulation using chemical agents, ozonization, flocculation and reverse osmosis are some of the techniques that are frequently used to remove these harmful substances$^5$–$^8$. However, these techniques are not effective in complete removal and merely result in the transfer of chemicals from one source to another. The photocatalytic phenomenon is capable of converting these toxic substances (EDTA, Cr(VI), nitrite) to harmless counterparts such as hydrogen, Cr(III) and nitrates$^9$–$^{11}$. This is achieved via a catalyst that upon subsequent excitation produces electrons and holes. These electrons and holes produce superoxide and hydroxyl radicals that are responsible for the oxidation and reduction processes$^{12}$–$^{15}$. But the efficiency of photocatalyst is largely dependent on the recombination rate of these electrons and holes. This drives the need for developing suitable catalysts which could be eventually scaled up for a large-scale clean-up drive$^{16}$–$^{19}$.

The wide band gap (3.25–5.1 eV) combined with a negative conduction band potential ($\approx$ 1.0 V vs. NHE at pH 0)$^{11}$ has recently spurred a lot of research on zirconia as a photocatalyst$^{20}$–$^{23}$. It is important to note that the band gap value depends on the crystal structure (3.84 eV for cubic, 4.11 eV for tetragonal and 4.51 eV for monoclinic)$^{24}$, defects and processing route of the ceramics$^{25}$. From the available limited literature, it is known that the presence of both monoclinic and tetragonal phases of zirconia$^{26}$–$^{28}$ is beneficial for photocatalysis. The high temperature tetragonal phase of zirconia namely $t$-$\text{ZrO}_2$ can be stabilized at room temperature by using suitable...
dopants like CaO, MgO and Y2O3 or by reducing the crystallite size. It is well established that nanostructured materials with large surface area greatly enhance the absorption of the irradiated light during photocatalysis and hence it appears advantageous to synthesize zirconia with crystallite size restricted to nanoscale dimensions for photocatalytic applications.

Anodization and combustion method are some of the common routes to synthesize nano-zirconia. However, the tedious, time-consuming processing steps, agglomeration and non-uniform size distribution, instability of the metastable tetragonal phase and its subsequent transformation to a pure monoclinic structure are some of the major drawbacks of these synthesis routes. Furthermore, the majority of research has been concentrated on UV light degradation of zirconia (bandgap values in the range of 4.71–5.0 eV and 3.1–3.54 eV have been reported).

Recently, black ZrO2 particles with very low bandgap (~1.5 eV) were reported to be photocatalytically active in the visible range and this was attributed to the presence of surface defects. Doping of zirconia with carbon and nitrogen is also seen to reduce the band gap and increase the absorption of visible light. Researchers have reported band gap values of 2.37 and 3.8 eV for carbon and nitrogen doped zirconia respectively. It is clear that the photocatalytic response of ZrO2 is governed to a large extent by the crystal structure, crystallite size, specific surface area, surface adsorbed species, surface defects and chemical composition of zirconia. This necessitates the need for developing a simplified processing approach wherein control of a large number of factors which contribute to the photocatalytic effect in the visible region is possible.

Dire et al. have synthesized zirconia crystallites in an amorphous silicon oxycarbide (SiOC) matrix through a sol-gel route using diethoxydimethylsilane as SiOC precursor and zirconium n-propoxide as ZrO2 precursor. Ionescu et al. have prepared ZrO2/SiOC ceramic nanocomposites through both in-situ addition of zirconium tetra(n-propoxide) as precursor and ex-situ addition of zirconia nano particles to polymethylsilsesquioxane as a matrix precursor (= preceramic polymer). Ceramics synthesized through the ex-situ addition of zirconia showed better crystallization resistance of matrix. This could be attributed to the higher zirconia content obtained through the ex-situ process. Also, the residual free carbon content was lower in ex-situ process than in-situ. Both of them clearly indicate that with the increase in the zirconia content there is great enhancement in the properties of matrix. In both cases, the photocatalytic behavior of nanocomposites was not investigated.

In this work we report the photocatalytic behavior of t-ZrO2/SiOCN nanocomposites with in-situ formation of crystallized zirconia stabilized with a tetragonal crystal structure in a structurally stable SiOCN matrix via the concept of "Nanocomposites Through Chemistry of Single-Source Precursors". The basis for this approach comes from the design of a suitable synthetic precursor in which uniform chemical composition is established at molecular scale. Then, precursors are converted in a first pyrolysis step into single-phase amorphous ceramics in which considerable weight loss occur. These are subsequently heat-treated at higher temperature under nitrogen to initiate the crystallization of the nanophase and provide the material with tuned phase composition and nano-/microstructure organization. The proposed new nanocomposites provide new insights not only into the material design by taking advantage of the weight loss occurring during the pyrolysis to form foams but also into the understanding of the photocatalytic behavior not explored before.

Results and discussion

Material design. We chose the Z10 sample to discuss the synthesis process. A representative FT-IR spectrum of Z10 cross-linked at 200 °C (Fig. 1) was recorded to identify the mechanisms occurring during reaction.

The spectrum of the Z10 sample (Fig. 1) shows peaks at 2965, 2173, 1263 cm⁻¹ corresponding to vibrations from νas(C-H), Si-H and νs(Si-CH3) stretching respectively. The peaks at 1037 and 1093 cm⁻¹ correspond to D units of Si-O vibrations. The peaks at 755 and 804 cm⁻¹ are associated with Si-C and νas(Si-CH3) vibrations.
spectrum indicates the modification of PHMS by ZB. ZB seems to crosslink the siloxane chains through the formation of Si-O-Zr bridges. The cross-linking reaction between ZB and PHMS is exemplified in Fig. 2.

The cross-linked polymer later decomposes to form amorphous SiZrOC. The thermogravimetric analysis of the Z10 sample (Fig. S1) confirms the completion of ceramization at around 858 °C. The ceramic sample obtained at 1400 °C (for 5 h in nitrogen atmosphere) labeled Z10-1400 has been fully characterized in parallel to other selected samples obtained with different Zr contents.

The FT-IR spectra of Z10-1400, Z30-1400 and Z50-1400 samples are exemplified in Fig. 3. The band observed at 1094 cm\(^{-1}\) corresponds to Si-O-Si stretching vibrations while the last two bands at 787 and 580 cm\(^{-1}\) could be associated with the ordered zirconia phase\(^{35}\). The overlapping of Si-O-Si and Si-O-Zr peaks results in the broadening of the band at 1094 cm\(^{-1}\). As expected, the intensity of Zr-O vibrations becomes sharper with increase in volume percentage of ZB. The band at 1625 cm\(^{-1}\) is a result of C\(=\)C stretching vibrations. It is also seen that the Si-C band at 755 cm\(^{-1}\) observed in the polymer has completely disappeared. Also, this could indicate possible cleavage of Si-C bond and presence of free carbon in the matrix. The broad band at 3400 cm\(^{-1}\) observed in all

Figure 2. Cross-linking reaction between ZB and PHMS.

Figure 3. FTIR spectra for different volume fractions of ZB pyrolyzed at 1400 °C.
samples comes from O-H functional groups. The intensity of this band increases with increase in the volume fraction of ZB. Earlier reports also suggest an increase of hydroxyl groups with increase in concentration of zirconia. This suggests that zirconia surface has greater ability to adsorb hydroxyl groups. It is believed that these surface hydroxyl groups interact with the holes generated during photocatalysis.

X-ray diffractograms (XRD) of the same samples are shown in Fig. 4. In addition, the XRD patterns of Z20-1400 and Z40-1400 are reported. The XRD diffractograms are composed of peaks at 30.2, 34.7, 50.2 and 59.7° which correspond to the (011), (110), (112) and (121) planes of tetragonal zirconia, respectively. The corresponding d-spacings are 2.94, 2.55, 1.81 and 1.54 Å. A broad hump observed at around 21° in Z10-1400 and Z20-1400 can be assigned to silica and the broadness of the hump indicates the amorphous nature of the phase. This amorphous hump disappears with increase in volume fraction of ZB. The broad amorphous hump weakened for Z40-1400 and completely disappeared for Z50-1400. The decrease/disappearance of SiO2 hump could be either due to suppression of the crystallization of silica by ZrO2 or the increase in other composition. For further understanding of the effect of pyrolysis temperature on the crystallization behavior, only samples with 10, 30 and 50 volume fractions of ZB were investigated and presented in the manuscript.

Figure 5 shows the XRDs patterns of (a) Z10, (b) Z30 and (c) Z50 samples pyrolyzed at temperatures ranging from 1100 °C to 1400 °C for 5 h in nitrogen atmosphere. Crystallization of zirconia appears to begin at 1300 °C for Z10 whereas for Z30, the onset of crystallization is further lowered by 100 °C. In the case of Z50, well defined peaks corresponding to t-ZrO2 are observed even in the samples pyrolyzed at 1100 °C. Hence, with the increase in the volume fraction of ZB the onset of crystallization is shifted to lower temperatures as observed by Dire et al.

The crystallite size of t-ZrO2 was calculated using the Scherrer’s equation as given below.

\[ d = \frac{k\lambda}{B \cos \theta} \]  

where d stands for the volume-averaged particle/crystallite size, K for the shape factor, \( \lambda \) represents the wavelength of CuKα, B refers to the peak broadening at FWHM obtained after correcting for the instrumental broadening and 2θ corresponds to the Bragg angle in degrees. The calculated crystallite size of Z10-1400, Z30-1400 and Z50-1400 were found to be 4.5 and 7 nm respectively.

Srinivasan et al. have reported phase separation of SiOC ceramics at temperatures exceeding 1300 °C. SiOC decomposes into silicon dioxide and silicon carbide. Silicon dioxide further reacts with the excess free carbon and forms silicon carbide. From the XRD (Fig. 5), the matrix was found to be stable till 1400 °C. The retention of the matrix in its amorphous state at 1400 °C is attributed to the presence of zirconia. Ionescu et al. and Dire et al. have shown that presence of zirconia strongly retards the carbothermal decomposition of SiOC. Increase in the zirconia content promotes the cleavage of Si-C bond discouraging SiC crystallization. This is due to the formation of strong interconnected network between the zirconia crystals and D units of silicon in the matrix. Some of the zirconia is still retained as Si-O-Zr units in the matrix as shown in the FT-IR analysis. While the structural and photocatalytic investigations were carried out on all the samples produced, for the sake of clarity and the higher fraction of crystallinity exemplified by Z50-1400, the discussion in the manuscript is henceforth restricted to the aforementioned sample.

**Oxygen defects and interstitial carbon.** In order further confirm the tetragonal structure of ZrO2, Raman spectra of Z50-1400 was analyzed (Fig. 6). There are six vibrational modes (one A_{1g}, three E_g and two B_{1g}) of tetragonal zirconia that are Raman active and these modes represent the different symmetries of point groups. The bands observed at 148, 266, 474 and 637 cm^{-1} (Fig. 6a) correspond to the characteristic bands of tetragonal zirconia. The broadness in the bands observed is primarily due to the presence of disordered oxygen sub-lattice. According to Cai et al., disordered lattice activates the Raman scattering from Brillouin zone points. It is this first order process that results in the broadening of the spectrum.
The D (1358 cm\(^{-1}\)) and G (1596 cm\(^{-1}\)) bands indicate the presence of free carbon in the matrix (Fig. 6b). The intensity ratio of the D band to G band gives an estimate about the defect densities\(^{45}\). The I\(_D/I_G\) ratio of 1.61 is obtained signifying the presence of large number of defects in carbon which in turn could be attributed to the formation of Zr-O-C bond. As mentioned earlier, increase in the zirconia content promotes the cleavage of Si-C bond resulting in the segregation of free carbon in the matrix. The presence of carbon is shown to increase the absorption of visible light, increase the adsorbing tendency of pollutants, prevent recombination and help in the better transport of electrons from the conduction band to the adsorbed species\(^{33,46}\). In order to understand the surface elemental composition as well as the chemical states of zirconium, X-ray photoelectron spectroscopy (XPS) was performed. Figure 7 exhibit the XPS survey of Z50-1400 which clearly reveals the presence of Si, Zr, O, C and N. In Fig. 8a, the Zr (3d) spectra shows a doublet with Zr (3d\(_{5/2}\)) and Zr (3d\(_{3/2}\)) at 182.9 and 185.3 eV.

Figure 5. X-ray diffractograms of (a) Z10, (b) Z30 and (c) Z50 as a function of pyrolysis temperature revealing the phase evolution and nanocrystalline nature of stabilized t-ZrO\(_2\).

Figure 6. Raman spectra of Z50-1400 indicating the presence of (a) tetragonal zirconia and (b) free carbon.
respectively which confirms the Zr$^{4+}$ oxidation state in the composite. The undoped ZrO$_2$ is reported to exhibit binding energies at 182.2 and 184.5 eV for Zr (3d$^{5/2}$) and Zr (3d$^{3/2}$) respectively which clearly indicates that there is an increase of 0.7 and 0.8 eV from the respective doublets in this study. The shift in doublets could be attributed to lattice distortions. Interestingly, the deconvolution of C (1s) peak resulted in three peaks at 282.1, 284.4 and 286.7 eV which can be attributed to the presence of Zr-C, C-C and C-O/C=O. The peak at 284.4 eV refers to the sp$^2$ hybridized carbon. The presence of Zr-C indicates that some of the oxygen atoms in the ZrO$_2$ lattice are replaced by carbon atoms (O-Zr-C) creating oxygen vacancies. The presence of zirconia in an amorphous

Figure 7. Full survey XPS spectra of Z50-1400 confirming the presence of silicon, oxygen, carbon, nitrogen and zirconia in the nanocomposite.

Figure 8. XPS spectra of Z50-1400 (a) Zr 3d (b) C 1s (c) N 1s and (d) O 1s.
carbon based matrix could also be responsible for the shift in Zr (3d) peaks
d. A scan explicitly done in the region from 390–405 eV shows an intensity at a binding energy of 399.5 from N 1 s peak (Fig. 8c). The absence of Zr-N peak at 396.8 eV excludes the possibility of nitrogen doping in ZrO$_2$ lattice. Hence the peak at 399.5 eV could be assigned to the bonding of nitrogen to SiOC matrix giving the SiOC(N) label. The deconvolution of O (1 s) region resulted in two peaks at 532.3 and 530.6 eV (Fig. 8d). The peak at 530 eV corresponds to lattice oxygen in ZrO$_2$ whereas the peak at 532 eV signifies surface lattice oxygen vacancy. The presence of oxygen vacancies in zirconia nanostructures (as indicated in Raman) is confirmed through XPS analysis. The area under the peak gives a relative oxygen vacancy of 52% (assuming negligible amount of OH adsorption) on the surface of ZrO$_2$. The presence of oxygen vacancies is also confirmed through EPR analysis (g value of 1.992) (Fig. S6). These oxygen vacancies serve as electron traps and prevent recombination of electron and holes during photocatalysis. The peak at 532 eV could also refer to the oxygen in surface adsorbed O-H groups. The ability of nanocrystalline zirconia to adsorb hydroxyl groups is highly beneficial for photocatalysis.

**Microstructural and nanostructural features.** The SEM images as shown in Fig. 9 show porosity at different length scales with pore sizes ranging from 120 μm down to less than 10 nm, resulting predominantly due to the evolution of oligomers (in the low temperature regime of the pyrolysis) and gases during the pyrolysis. The BET analysis (Fig. S4) revealed the pore diameter to be varying between 4 nm to 150 nm. The porous structure allows adequate contact facilitating the mass transfer between the pollutant fluid and the catalyst, increasing the scattering of light aiding in the photocatalytic process. The pores obtained for Z10-1400 and Z30-1400 were non-uniform in nature (Fig. S3).

Figure 9a,b illustrate bright field TEM image of Z50-1400 nanocomposite. TEM investigation confirms the nanocomposite structural organization of the materials: an uniform distribution of fine spherical zirconia nanocrystals in an amorphous matrix is clearly observed. On an average, around 4–5 nanocrystals of zirconia are present for every 1000 nm$^2$ area of the matrix. This means that four active sites are available for photocatalysis per 1000 nm$^2$.

The nanocrystallite size distribution obtained from the TEM micrographs is shown in Fig. 10a inset. The histogram was generated from five different micrographs of different volume fractions of ZB. The crystals are of different sizes ranging from 4–20 nm. Maximum number of crystals falls in the range of 6–14 nm. The crystallite sizes obtained from TEM are in agreement with the calculated values using Scherrer’s equation from XRD. It was also observed that the crystallite size of zirconia increases with the volume fraction of ZB, Fig. S5, which substantiate the XRD results.
The SAED pattern recorded for aggregates of crystals in Z50-1400 is shown in Fig. 10a. The radius of the rings was measured and ratios between them correspond well to the ratios of the d-spacing of (011), (002), (112) and (121) planes of t-ZrO2 (ICDD-157618). The diffraction from (112) plane is the characteristic feature that distinguishes the tetragonal from cubic phase. The d spacing of 0.18 nm calculated from the lattice fringe spacing (Fig. 10b) corresponds to the (112) plane of the tetragonal structure. It is also evident from the diffraction pattern that matrix is entirely amorphous and there is no crystallization of silica.

Stabilization of the tetragonal phase. Monoclinic polymorph of zirconia is the most stable crystal structure at room temperature and the tetragonal transformation occurs around 1170 °C. It has been shown that a single strain free zirconia nanocrystal can exist in tetragonal form at room temperature with a critical size less than 10 nm as the surface energy difference overcomes the volume free energy difference53. In case of a cluster of crystallites, interfacial energy comes into play increasing the critical size to 33 nm53. As shown in Table 1, it is seen that for all types of interfaces the interfacial energy for tetragonal structure is lower than the monoclinic structure54.

A recent study by Jayakumar et al.55 for pure ZrO2 established crystallite size values around 7–20 nm for stable tetragonal phase at room temperature. It is generally observed that at high annealing temperatures (~800–1000 °C), there is considerable size growth resulting in spontaneous transformation to monoclinic structure46,48. In the current work, extremely fine crystallite sizes (6–12 nm) are observed even at pyrolysis temperature as high as 1400 °C for extended durations of time most probably because of the presence of a predominantly covalently-bonded matrix surrounding nanocrystals. It is also seen that with the increase in the ZB volume fraction or temperature, the coarsening of t-ZrO2 is rather minimal, possibly owing to the matrix confinement effect. However, Shukla et al.56 have shown that with increase in oxygen vacancy concentration, the activation energy for grain growth decreases considerably. Hence, oxygen vacancies seem to contribute in the stabilization of the tetragonal structure of zirconia to maintain its 7-fold coordination. Since pyrolysis of all samples was carried in nitrogen atmosphere at low oxygen partial pressure in the current work, it is expected that the pyrolyzed samples are oxygen deficient which is also confirmed from EPR data (Fig. S6). Also, Xian et al.48 demonstrated stabilization of tetragonal structure for crystallite sizes as high as ~47 nm and attributed this effect to the presence of carbon. The presence of carbon in the creation of oxygen vacancies can either occur by occupying the interstitial positions of zirconia lattice (as observed in XPS, Fig. 8) or by the reaction between the residual carbon in the matrix and oxygen in the zirconia lattice. The strain energy associated with the SiOC(N) matrix can also strongly retard the transformation to monoclinic structure55. Shukla et al.56 have shown that hydrostatic strain energy values could be as high as 2.1 × 109 J m−3 pushing the critical size to a value as high as 200 nm. Hence, the authors believe that the synergistic effect of nanoscale dimensions of ZrO2, oxygen vacancies, presence of interstitials such as carbon and nitrogen and the confinement effect of SiOC(N) matrix assist in the retention of tetragonal crystal structure of zirconia at room temperature.

**Bandgap and role of interstitial carbon.** The DRS spectrum of Z50-1400 is shown in Fig. 11. The band gap (Eg) was determined by using Tauc Eq. (2)29.
$\alpha h\nu = K(E - E_g)^n$

(2)

where, $\alpha$ is the absorption coefficient, $h\nu$ is the photon energy, $K$ is a constant, and $n$ is either 1/2 for a direct transition or 2 for an indirect transition.

The bang gap was calculated by extrapolating the linear portion of the plot $(\alpha h\nu)^2$ vs $h\nu$ (assuming ZrO$_2$ as a direct semiconductor). The Tauc plot of Z50-1400 exhibits a band gap of 2.2 eV respectively (Fig. 11).

In general, reported values of band gap for tetragonal structure are around 4-5 eV$^{9,25,30}$. However, the band gap values as low as 3 eV has also been reported by Navio et al. for the tetragonal ZrO$_2$.$^{57}$ The drastic reduction in band gap (2.2 eV) could be attributed to oxygen vacancies and presence of carbon as revealed from XPS (Fig. 7b,d). It is highly probable that carbon enters the interstitial sites of zirconia (O-Zr-C bond) and forms a mid-band gap state between the conduction band and valence band.$^{32}$ The presence of oxygen vacancies might also create new energy states between the valence band and conduction band thereby reducing the bang gap and enhancing the visible light absorption$^{31}$.

**Photocatalytic studies.** The degradation of methylene blue dye after exposure to simulated solar radiation in the presence of catalysts was analyzed using UV-vis spectroscopy. The absorbance pattern of the samples collected from photocatalytic experiments were recorded in the wavelength range of 200–900 nm. The decrease in the concentration of methylene blue dye with time after the adsorption-desorption equilibrium for Z50-1400 is given in Fig. 12a. The absorbance value reached a maximum at a wavelength of 663 nm. With the increase in the exposure time absorption peak value at 663 nm decreases, showing the decolorization of the solution in the presence of the catalyst and thus indicating the degradation of the dye. The degradation percentage of methylene blue was calculated according to the following equation$^{32}$.

$$D = \left(\frac{C_0 - C}{C_0}\right) \times 100$$

(3)

where $C_0$ and $C$ are the initial and final concentration in mg/L.

According to the Langmuir-Hinshelwood model$^{28}$ the photocatalytic reaction rate after adsorption-desorption equilibrium is

$$\ln\left(\frac{C}{C_0}\right) = -kt$$

(4)

where $C_0$ and $C$ are the concentration at time $t = 0$ and $t = t$ minutes respectively and $k$ is the reaction rate constant.

It was observed that after 4 h of visible light exposure, 72% of methylene blue was degraded for Z50-1400 nanocomposite. The self-degradation of methylene blue without the presence of catalyst was taken into account and the corrected photodegradation rate constant $k$ ($\times 10^{-3}$) was found to be 3.2 min$^{-1}$ (Fig. 12b).

**Photocatalysis mechanism.** The absence of long range ordered structure in amorphous silicon oxycarbonitride results in the formation of localized band tails in the valence and conduction bands. The presence of dangling bonds often results in the formation of gap states.$^{58}$ The amorphous material is hence expected to possess multiple electron states. The presence of band tails as well as the mid band gap states results in the reduction of band gap in amorphous materials. The formation of heterojunction between an amorphous and crystalline material is a viable approach to improve the photocatalytic efficiency. The interfacial lattice mismatch at the heterojunction between crystalline domains could hinder the transport of the carriers. However, the existence of
heterojunctions between amorphous and crystalline material provides an efficient way to separate the photogenerated holes and electrons. This separation combined with the reduced band gap helps in harnessing visible light for photocatalysis. Albeit the detrimental contribution of t-ZrO₂ towards photocatalytic activity, the formation of heterojunctions as well as the retention of nanodomains of t-ZrO₂ with an average size of 10 nm in an amorphous matrix is believed to enhance the catalytic activity. The degradation mechanism and the heterojunction formation are exemplified in Figs. 13 and 14 respectively. When the nanocomposite is exposed to solar radiation, the electrons are excited from the valence band to the conduction band leaving behind holes. Concomitantly, the photogenerated electrons from the conduction band of amorphous SiOC(N) are transferred to the conduction band of t-ZrO₂. On similar grounds, holes are injected from the valence bands in the opposite direction (i.e., from t-ZrO₂ to amorphous SiOC(N)). This transfer mechanism results in effective charge separation and prevents charge recombination thereby increasing the catalytic degradation efficiency. Further, in order to elucidate the above proposed mechanism, reactive species trapping experiments and EPR measurements were performed (Figs. S7 and S8). The valence and conduction band edges of a-SiOC(N) (Fig. S9) and t-ZrO₂ has been calculated and included in the supplementary data.

The reaction between surface hydroxyl groups and holes produces hydroxyl radicals which are powerful oxidizers.

$$\text{OH}^- + \text{h}^+ \rightarrow \text{OH}^* \hspace{1cm} (6)$$

Subsequently, the captured electrons then react with dissolved oxygen in the polluted water and generate superoxide anions.

$$\text{O}_2 + \text{e}^- \rightarrow \text{O}_2^- \hspace{1cm} (7)$$
These superoxide anions further react with water molecules generating large number of hydroxyl radicals. These hydroxyl radicals are finally responsible for efficient dye degradation. The decomposition of methylene blue in the presence of hydroxyl radicals is given by Soltani et al.\textsuperscript{29}

\[
\text{MB} + \text{OH} \cdot \rightarrow \text{CO}_2 + \text{H}_2\text{O} + \text{NH}_4^+ + \text{NO}_3^- + \text{SO}_4^{2-} + \text{Cl}^-
\] (8)

This is the phenomenon responsible for the decrease in the concentration and decolorization of methylene blue and on a large scale reduces the chemical oxygen demand (COD) of textile industry effluents. The synergistic effect of nano t-ZrO\textsubscript{2} and amorphous SiOC(N) matrix were confirmed by X-ray diffraction and electron microscopy. The structural characterization depicted the hierarchical porous nature of the nanocomposites. The presence of carbon as well as oxygen vacancies in zirconia lattice confirmed via spectroscopy techniques has resulted in drastic reduction of band gap (2.2 eV). The retention of t-ZrO\textsubscript{2} confined in a matrix (t-ZrO\textsubscript{2}/SiOC(N) nanocomposite) exhibited an unusually high photodegradation efficiency of 72\% possibly because of the formation of heterojunctions reducing the recombination rate. The synergistic effect of nanoscaled heterojunctions and the presence of carbon and oxygen defects resulted in a paradigm shift pushing the limits of photocatalytic activity zirconia-based nanocomposites from UV to visible region with high degradation efficiency.

Conclusions

The crystallization and stabilization of extremely fine nanocrystals (less than 15 nm) of t-ZrO\textsubscript{2} in an amorphous SiOC(N) matrix were confirmed by X-ray diffraction and electron microscopy. The structural characterization depicted the hierarchical porous nature of the nanocomposites. The presence of carbon as well as oxygen vacancies in zirconia lattice confirmed via spectroscopy techniques has resulted in drastic reduction of band gap (2.2 eV). The retention of t-ZrO\textsubscript{2} confined in a matrix (t-ZrO\textsubscript{2}/SiOC(N) nanocomposite) exhibited an unusually high photodegradation efficiency of 72\% possibly because of the formation of heterojunctions reducing the recombination rate. The synergistic effect of nanoscaled heterojunctions and the presence of carbon and oxygen defects resulted in a paradigm shift pushing the limits of photocatalytic activity zirconia-based nanocomposites from UV to visible region with high degradation efficiency.

Methods

Zirconium (IV) butoxide (ZB) was mixed with polymethylhydrosiloxane (PHMS) (Sigma Aldrich, Bangalore, India) in varying volume fractions (10–50\%) and the mixture was stirred in a magnetic stirrer for about 30 min. In order to facilitate the cross-linking of polymers the stirred mixture was held at 200 °C for 4 h in nitrogen atmosphere. The cross-linked polymers were pyrolyzed in nitrogen atmosphere at temperatures ranging from 1100–1400 °C for 5 h. Nitrogen gas was purged at a flow rate of 2.5l/min and a constant heating rate of 5 °C/min was maintained during pyrolysis. The samples were cooled down to room temperature and a foamy structure with well-defined pores was obtained. The pyrolyzed samples, hereafter will be referred to as ZX−Y throughout the manuscript, where X represents the volume fraction of ZB in PHMS and Y the pyrolysis temperature. The general process for the synthesis of t-ZrO\textsubscript{2}/SiOCN nanocomposite is shown in Fig. S1.

The Fourier transform infrared spectra (FT-IR) of ceramic powders were recorded to understand the binding characteristics (Perkin Elmer Spectrum, USA). The powders were mixed with KBr in the ratio 1:9 and the analysis was done in the transmission mode. X-ray diffractograms (X’PertPRO PANalytical X-ray diffractometer, Netherlands) for all the ceramic samples were obtained in the 2θ range of 10–90° with a step size of 0.008 using Cu K\textalpha\, radiation. The Raman spectroscopy (Labram HR 800 Horiba, USA) measurements were done at an excitation wavelength of 480 nm. The foamy structure of ceramics was imaged using FEI Quanta 200, USA scanning electron microscope (SEM). In order to minimize the charging effect, the samples were sputter coated with gold for 120 s before analysis. The morphology and the lattice fringes of nanocrystals were imaged using Philips CM12 and JEOL 3010 high resolution transmission electron microscopes (HRTEM) respectively at an accelerating voltage of 200kV. The powder samples were ultrasonicated in ethanol solution and then a few drops were placed on the carbon-coated copper grids. X-ray photoelectron spectroscopy (XPS) was done using SPECS with PHIBOS100 energy analyzer (SPECS GmbH, Germany) to understand the surface elemental composition. Two anodes, Mg and Al act as the X-ray source. The binding energies of the different elements present were analyzed with respect to elemental carbon (284.4 eV) as the standard.

Photocatalytic measurements were carried out in a photoreactor (Heber Scientific, India) with a 500 W tungsten lamp as the light source. In all the measurements the catalyst concentration was kept constant at 50 mg for 100 ml of methylene blue solution of 0.03 mM concentration. The light source was continuously cooled by water circulation through an outer jacket for dissipation of the excessive heat generated. Prior to the exposure, the solution was maintained in dark for two hours under aerated condition to ensure adsorption-desorption equilibrium. The experiments were carried out in visible light for 4 h. At the end of every hour, few ml of solution was collected and centrifuged to remove any solid catalyst particles. The collected samples were further analyzed in UV-vis spectrophotometer (ThermoFischer Scientific, Evolution 220, USA) to study their adsorption characteristics and decolorization behavior. The diffuse reflectance spectra (DRS) were measured with BaSO\textsubscript{4} as a reference.

Received: 13 February 2019; Accepted: 27 December 2019;
Published online: 16 January 2020

References

1. Connor, R., Uhlenbrook, S., Koncaogül, E. & Ortigara, A. R. C. The United Nations World Water Development Report (2017).
2. Babu, B. R., Parande, A. K., Raghu, S. & Kumar, T. P. Cotton Textile Processing: Waste Generation and Effluent Treatment. J. Cotton Sci. 153, 141–153 (2007).
3. Kant, R. Textile dyeing industry an environmental hazard. Nat. Sci. 04, 22–26 (2012).
4. Singaree, P. & Dhabardeb, S. Toxic metals pollution due to industrial effluents released along Dombivali Industrial Belt of Mumbai, India. Eur. J. Env. Saf. Sci. 2, 5–11 (2014).
5. Faria, P. C. C., Órfão, J. M. & Pereira, M. F. R. Activated carbon and ceria catalysts applied to the catalytic ozonation of dyes and textile effluents. *Appl. Catal. B Environ.* **88**, 341–350 (2009).

6. Hu, E. et al. Catalytic ozonation of simulated textile dyeing wastewater using mesoporous carbon aerogel supported copper oxide catalyst. *J. Clean. Prod.* **112**, 4710–4718 (2016).

7. Pelaiz-Cid, A.-A., Herrera-González, A.-M., Salazar-Villanueva, M. & Bautista-Hernández, A. Elimination of textile dyes using activated carbons prepared from vegetable residues and their characterization. *J. Environ. Manage.* **181**, 269–278 (2016).

8. Harinclas, F. E., Aziz, A., Vaconah, A., Benhammou, A. & Poss, M. N. Treatment of textile dye effluents using coagulation-flocculation coupled with membrane processes or adsorption on powdered activated carbon. *Desalination* **235**, 330–339 (2009).

9. Suresh, P., Vijaya, J. J. & Kennedy, L. J. Photocatalytic degradation of textile-dyeing wastewater by using a microwave combustion-synthesized zirconium oxide supported activated carbon. *Mater. Sci. Semicond. Process.* **27**, 482–493 (2014).

10. Labaki, M., Siffert, S., Lamonier, J., Zhilinskaya, E. A. & Abouka, A. Total oxidation of propane and toluene in the presence of zirconia doped by copper and yttrium Role of anionic vacancies. *Appl. Catal. B Environ.* **43**, 261–271 (2003).

11. Botta, S. G., Nav, J. A., Hidalgo, C., Restrepo, G. M. & Litter, M. I. Photocatalytic properties of ZrO2 and Fe/ZrO2 semiconductors prepared by a sol – gel technique. *J. Photochem. Photobiol. A Chem.* **129**, 89–99 (1999).

12. Chen, X. et al. Increased photocatalytic activity of tube-brush-like ZnO nanostructures fabricated by using PVP nanofibers as templates. *Appl. Surf. Sci.* **319**, 216–220 (2014).

13. Kapridaki, C., Pinho, L., Mosquera, M. J. & Maravelaki-Kalaitzaki, P. Producing photoactive, transparent and hydrophobic SiO2-crystalline TiO2 nanocomposites at ambient conditions with application as self-cleaning coatings. *Appl. Catal. B Environ.* **156–157**, 416–427 (2014).

14. Hojamberdiev, M. et al. Template-free synthesis of polymer-derived mesoporous SiOxCr2O3 powders. *Ceram. Int.* **39**, 1183–1190 (2013).

15. Ou, C.-C., Yang, C.-S. & Lin, S.-H. Selective photo-degradation of Rhodamine B over zirconia incorporated titania nanoparticles: a quantitative approach. *Catal. Sci. Technol.* **1**, 295 (2011).

16. Chen, F. et al. Thickness-Dependent Facet Junction Control of Layered BioIO3 Single Crystals for Highly Efficient CO2 Photoreduction. *Adv. Funct. Mater.* **28**, 1–11 (2018).

17. Chen, F., Huang, H., Guo, L., Zhang, Y. & Ma, T. The role of polarization in photocatalysis. *Angew. Chemie Int. Ed.* (2019).

18. Huang, H. et al. Macroscopic Polarization Enhancement Promoting Photo- and Piezoelectric-Induced Charge Separation and Molecular Oxygen Activation. *Angew. Chemie - Int. Ed.* **56**, 11860–11864 (2017).

19. Huang, H. et al. Anionic Group Self-Doping as a Promising Strategy: Band-Gap Engineering and Multi-Functional Applications of High-Performance CO 3 2– Doped Bi 2 O 3 CO 3. *ACS Catal.* **5**, 4094–4103 (2015).

20. Stojadindrović, S., Vasilč, R., Radič, N. & Grbić, B. Zirconia films formed by plasma electrolytic oxidation: Photoluminescence and photocatalytic properties. *Opt. Mater. (Amst.)* **40**, 20–25 (2015).

21. Yin, L., Niu, J., Shen, Z., Bao, Y. & Ding, S. Preparation and photocatalytic activity of nanoporous zirconia electrospun fiber mats. *Mater. Lett.* **65**, 3131–3133 (2011).

22. Moafi, H. F., Shojaie, A. F. & Zanjanchi, M. A. The Comparative Study of Photocatalytic Self-Cleaning Properties of Synthesized Nanoscale Titanium and Zirconia onto Polyacrylonitrile Fibers. *J. Appl. Polym. Sci.* (2010).

23. Vignesh, K., Suganthi, A., Min, B. & Kang, M. Applied Surface Science Fabrication of meso-porous BiO1 nano-sensitized zirconia nanoparticles with enhanced photocatalytic activity under simulated solar light irradiation. *Appl. Surf. Sci.* **324**, 652–661 (2015).

24. Bechepeche, A. P., Treu, O., Longo, E., Paiva-Santos, C. O. & Varela, J. A. Experimental and theoretical aspects of the stabilization of zirconia. *J. Mater. Sci.* **34**, 2751–2756 (1999).

25. Basahel, S. N., Ali, T. T., Mokhtar, M. & Narasimharao, K. Influence of crystal structure of nanosized ZrO2 on photocatalytic degradation of methyl orange. *Nanoscale Res. Lett.* **73**, 1–13 (2015).

26. Jiang, W. et al. Preparation and photocatalytic performance of ZrO2 nanotubes fabricated with anodization process. *Appl. Surf. Sci.* **307**, 407–413 (2014).

27. Wang, D. et al. Photoluminescence properties and photocatalytic activities of zirconia nano-tube arrays fabricated by anodization. *Opt. Mater. (Amst.)* **35**, 1461–1466 (2013).

28. Ashkarran, A. A., Afshar, S. A. A., Afshar, S. M. & Kavianipour, M. Photocatalytic activity of ZrO2 nanoparticles with enhanced photocatalytic activity under simulated solar light irradiation. *Appl. Surf. Sci.* **93**, 121–125 (2016).

29. Matsui, H., Ohkura, N., Karuppuchamy, S. & Yoshihara, M. The effect of surface area on the photo-catalytic behavior of ZrO2/ carbon clusters composite materials. *Ceram. Int.* **39**, 5827–5831 (2013).

30. Sudrajat, H., Babel, S., Sakai, H. & Takizawa, S. Rapid enhanced photocatalytic degradation of dyes using novel N-doped ZrO2. *J. Environ. Manage.* **165**, 224–234 (2016).

31. Oestreich, F., Ceccato, R., Gialanella, S. & Babonneau, F. Thermal evolution and crystallisation of polydimethylsiloxane–zirconia nanocomposites prepared by the sol–gel method. *J. Eur. Ceram. Soc.* **19**, 2849–2858 (1999).

32. Ionescu, E. et al. Polymer-Derived SiOxCr2O3 Ceramic Nanocomposites with Excellent High-Temperature Stability. *J. Am. Ceram. Soc.* **93**, 241–250 (2010).

33. Bechelany, M. C. et al. Nanocomposites through the Chemistry of Single-Source Precursors: Understanding the Role of Chemistry behind the Design of Monolith-Type Nanostructured Titanium Nitride/Silicon Nitride. *Chem. - A Eur. J.* **23**, 832–845 (2017).

34. Mera, G., Galiet, M., Bernard, S. & Ionescu, E. Ceramic Nanocomposites from Tailor-Made Preceramic Polymers. *Nanomaterials* **5**, 468–540 (2015).

35. Bechelany, M. C. et al. In Situ Controlled Growth of Titanium Nitride in Amorphous Silicon Nitride: A General Route Toward Bulk Nitride Nanocomposites with Very High Hardness. *Adv. Mater.* **26**, 6548–6553 (2014).

36. Drie, S., Cazzato, P. R., Ceccato, R. & Porrino, F. Ryolitische Chemistry of Sol – Gel-Derived Poly (dimethylsiloxane) – Zirconia Nanocomposites. Influence of Zirconium on Polymer-to-Ceramic Conversion. *Chem. Mater.* **24**, 268–278 (1999).

37. Nedunchezhian, S., Sujith, R. & Kumar, R. Processing and characterization of polymer precursor derived silicon oxy-carbide ceramic foams and composites. *J. Adv. Ceram.* **2**, 318–324 (2013).

38. Naumenko, A. P., Berezovska, N. I., Bily, M. M. & Shevchenko, O. V. Vibrational Analysis and Raman Spectra of Tetragonal Zirconia. *Phys. Chem. Solid State* **1**, 121–125 (2008).

39. Tan, D. et al. Synthesis of nanocrystalline cubic zirconia using femtosecond laser ablation. *J Nanoparticle Res.* **13**, 1183–1190 (2011).

40. Cai, J., Raptis, C., Raptis, Y. S. & Anastassakis, E. Temperature dependence of Raman scattering in stabilized cubic zirconia. *Phys. Rev. B* **51** (1995).
57. Navio, J. A., Macias, M. & Co, G. Surface characterization of ZrO2-SiO2 systems prepared by a sol-gel method.

58. Yang, X.

59. Huang, H.

60. Garvie, R. C. Stabilization of the tetragonal structure in zirconia microcrystals.

61. Shukla, S. & Seal, S. Mechanisms of room temperature metastable tetragonal phase stabilisation in zirconia.

62. Yu, H.

63. Jayakumar, S.

64. Shukla, S., Seal, S., Vij, R. & Bandyopadhyay, S. Reduced activation energy for grain growth in nanocrystalline yttria-stabilized zirconia.

65. Milošev, I., Strehblow, H.-H., Gaberšček, M. & Navinšek, B. Electrochemical Oxidation of ZrN Hard (PVD) Coatings Studied by XPS. Surf. Interface Anal. 24, 448–458 (1996).

66. Teeparthi, S. R., Awin, E. W. & Kumar, R. Dominating role of crystal structure over defect chemistry in black and white zirconia on visible light photocatalytic activity. Sci. Rep. 8, 1–11 (2018).

67. Yu, H. et al. Three-in-One Oxygen Vacancies: Whole Visible-Spectrum Absorption, Efficient Charge Separation, and Surface Site Activation for Robust CO2 Photoreduction. Angew. Chemie - Int. Ed. (2019).

68. Shukla, S. & Seal, S. Mechanisms of room temperature metastable tetragonal phase stabilisation in zirconia. Int. Mater. Rev. 50, 45–64 (2005).

69. Garvie, R. C. Stabilization of the tetragonal structure in zirconia microcrystals. J. Phys. Chem. 82, 218–224 (1978).

70. Jayakumar, S. et al. Nanosize stabilization of cubic and tetragonal phases in reactive plasma synthesized zirconia powders. Mater. Chem. Phys. 140, 176–182 (2013).

71. Shukla, S., Seal, S., Vij, R. & Bandyopadhyay, S. Reduced activation energy for grain growth in nanocrystalline yttria-stabilized zirconia. Nano Lett. 3, 397–401 (2003).

72. Navio, J. A., Macias, M. & Co, G. Surface characterization of ZrO2-SiO2 systems prepared by a sol-gel method. Appl. Surf. Sci. 71, 226–229 (1993).

73. Yang, X. et al. Amorphous molybdenum selenide as highly efficient photocatalyst for the photodegradation of organic dyes under visible light. Appl. Surf. Sci. 457, 214–220 (2018).

74. Huang, H. et al. Fabrication of multiple heterojunctions with tunable visible-light-active photocatalytic reactivity in BiOBr-BiOI full-range composites based on microstructure modulation and band structures. ACS Appl. Mater. Interfaces 7, 482–492 (2015).

75. Du, X. et al. In situ assembly of BiOBr@Bi 12 O 17 Cl 2 p – n junction: charge induced unique front-lateral surfaces coupling heterostructure with high exposure of BiOI [001] active facets for robust and nonselective photoredox catalysis. Appl. Catal. B Environ. 199, 73–86 (2016).

Acknowledgements

We would like to thank the funding received from Indo-French Centre for the Promotion of Advanced Research (IFCPAR/CEFIPRA), a bilateral Indo-French project (Project No. 5108-1) to carry out this work. We would like to thank Sophisticated Instrument Analytical Facility (SAIF, IIT Madras) for the assistance provided to carry out FTIR and EPR measurements. We would also like to thank the DST Unit of Nanoscience, Department of Chemistry, IIT Madras and Central Electron Microscopy Facility (CEMF), IIT Madras for providing access to HR-TEM and SEM respectively. We would also like to thank Prof. Umit B Demirci, University of Montpellier, for his scientific inputs.

Author contributions

S.B.1. and R.K. conceived the idea and designed the experiments. E.W.A. and R.K. supervised all the experiments and analysis. S.B.1. prepared materials, performed characterizations, photocatalytic measurements and analyzed the results. K.C.H., A.L. and S.B.2. commented on the manuscript writing. S.B.1. wrote the manuscript and all authors discussed the results and commented on the manuscript of the work.

Competing interests

The authors declare no competing interests.

Additional information

Supplementary information is available for this paper at https://doi.org/10.1038/s41598-019-57394-8.

Correspondence and requests for materials should be addressed to R.K.

Reprints and permissions information is available at www.nature.com/reprints.

Publisher’s note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made. The images or other third party material in this article are included in the article’s Creative Commons license, unless indicated otherwise in a credit line to the material. If material is not included in the article’s Creative Commons license and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this license, visit http://creativecommons.org/licenses/by/4.0/.

© The Author(s) 2020