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<https://doi.org/10.1016/j.ceramint.2012.04.064>

# **Aqueous sol-gel synthesis of zirconium titanate (ZrTiO<sub>4</sub>) nanoparticles using chloride precursors**

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## **Abstract**

Zirconium titanate powders were synthesized by a straightforward sol-gel method using zirconium and titanium chlorides as metal precursors, deionized water as solvent and oxygen donor, and a NaOH solution for adjusting pH to 7. According to transmission electron microscopy, amorphous particles of nearly 5 nm in size with a relatively spherical morphology were prepared. Thermogravimetry and differential scanning calorimetry analyses on the xerogel at a heating rate of 10 °C/min indicated a crystallization temperature of 690 °C, which is comparable with previous reports. Furthermore, via differential scanning calorimetry studies using the Kissinger's equation, the activation energy for ZrTiO<sub>4</sub> crystallization was determined to be 850 kJ/mol. Structural evaluations in the isothermal regime, using X-ray diffraction experiments, implied the onset of ZrTiO<sub>4</sub> crystallization at 550 °C.

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*Keywords:* Sol-gel processes (A); X-ray methods (B); Thermal properties (C)

## 1. Introduction

Zirconium titanate (ZrTiO<sub>4</sub>) is an attractive ceramic for electronic, optical, chemical, catalytic, and biomedical applications [1–6], but demanding high purity and homogeneity. One of the promising approaches to synthesize homogeneous ceramic powders is the sol-gel process. In this method, when a high level of homogeneity is achieved, the obtained amorphous materials crystallize at temperatures of several hundred degrees lower than in classical routes, such as traditional melting and powder processing [7–11]. In addition, as a result of the development of high specific surface area powders by the sol-gel process, much lower sintering temperatures are required.

The sol-gel preparation of zirconium titanate can be conducted by hydrolysis and condensation of inorganic salts [9,12,13] or alkoxides [14–16] as precursors, where the control of the hydrolysis and condensation processes is essential when establishing procedures especially for the latter. A number of non-hydrolytic sol-gel processes have been also focused on producing zirconium titanate powders [10,11]. On the other hand, nanostructured TiO<sub>2</sub>-ZrO<sub>2</sub> powders and thin films have been prepared by an aqueous particulate sol-gel route from titanium isopropoxide and zirconium acetate hydrate as precursors [17]. However, to our knowledge, no work has been reported to date on the aqueous particulate sol-gel synthesis of zirconium titanate from chloride precursors.

In this research, a straightforward aqueous particulate sol-gel method, as an environmentally friendly processing, was used to produce zirconium titanate powders. The

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process is based on the hydrolysis of metal chlorides, in which alkoxide functions are formed *in situ*. One of the advantages of this route is employing an alternative (chloride precursors) to alkoxide precursors used in polymeric sol-gel methods to develop a product at lower cost, which is a significant attribute. This paper focuses on the sol-gel synthesis and characterization of zirconium titanate starting from ZrCl<sub>4</sub> and TiCl<sub>4</sub> as the metal precursors.

## **2. Materials and methods**

### *2.1. Sample preparation*

18.2 mmol of zirconium tetrachloride (ZrCl<sub>4</sub>, Alfa Aesar, 99.5 %) was gradually added to 400 mL of deionized water and stirred for 15 min. 18.2 mmol of titanium tetrachloride (TiCl<sub>4</sub>, Alfa Aesar, 99.99 %) was added dropwise to the solution and magnetically stirred for 2 h, leading to a solution of 0.98 in pH measured using a calibrated pH meter (Sartorius Professional Meter PP-15). Subsequently, an aqueous solution of NaOH (2 M) was added dropwise under magnetic stirring conditions until the pH value approached 7, forming a white hydrogel. After aging for 30 min, in order to remove chloride ion, the hydrogel was rinsed several times with deionized water, and the absence of chloride ions was checked using AgNO<sub>3</sub>. The obtained sample was centrifuged at 6000 rpm for 10 min, washed with ethanol, again centrifuged, and finally dried at 60 °C for 2 h, developing white agglomerated powders (xerogel). The obtained powder was calcined at 500, 550, and 600 °C for 1 h with a heating rate of 10 °C/min in a nitrogen atmosphere.

### *2.2. Structural and thermal characterization*

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A transmission electron microscope (TEM, JEOL JEM-2100) operating at an acceleration voltage of 200 kV was used to evaluate the powder particle size and morphology. To do so, a small amount of the powder was dispersed in ethanol, ultrasonicated for 15 min, and then dropped on a carbon grid. To study the thermal behavior of the xerogel, simultaneous thermo-gravimetry and differential scanning calorimetry analyses (TGA/DSC, Netzsch, STA 449 F1 Jupiter<sup>®</sup>) were performed under a flowing argon atmosphere in an alumina container. The X-ray diffraction (XRD) pattern of the xerogel and calcined powders was recorded by a Bruker AXS Inco. diffractometer using Cu K $\alpha$  radiation ( $\lambda = 1.54056 \text{ \AA}$ ).

### 3. Results and discussion

The TEM observations of the xerogel powder particles (before calcination) show two typical features. As depicted in Fig. 1(a), despite dispersion in ethanol and ultrasonication, some agglomerated particles of 50-100 nm in size can be still seen, making it difficult to distinguish the real particle size and morphology. This strong tendency toward agglomeration is due to the significant surface-to-volume ratio and surface tension of nanoparticles. Nonetheless, according to Fig. 1(b), on the edge of the agglomerated particles, powder particles of approximately 5 nm in diameter with a relatively spherical morphology can be recognized. The related selected area diffraction (SAD) pattern (Fig. 1(c)) indicates homogenous featureless diffraction halos attributed to an amorphous structure. No diffraction spots or sharp rings associated with crystalline phases can be observed, suggesting the synthesis of fully amorphous nanoparticles.

The TGA and DSC profiles of the xerogel at a heating rate of 10 °C/min are provided in Fig. 2. In the TGA curve, a pronounced weight loss of 15% is observed from 100 °C to 250

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°C. A broad endothermic peak at around 120 °C and a sharp exothermic peak at 690 °C are also observed in the DSC curve. Considering both analyses simultaneously, it is evident that the first event, the endothermic one with a considerable weight loss, is due to the removal of residual water and chloride during heating. On the other hand, the second sharp event was accompanied by no weight loss and is attributed to crystallization of the amorphous phase developed by the sol-gel method, as verified by XRD and TEM studies below. In accordance with the DSC profile, the onset crystallization temperature and crystallization peak temperature are 575 and 690 °C, respectively.

To study the kinetic behavior of the crystallization event, further DSC experiments on the xerogel were carried out at different heating rates of 20, 30, 40, and 50 °C/min. Based on the data shown in Fig. 3(a), by increasing the heating rate, the crystallization peak is shifted to higher temperatures since the crystallization process from an amorphous phase is a diffusional and thermally-activated transformation controlled via nucleation and growth. The activation energy for crystallization ( $E_c$ ) can be determined by the Kissinger's equation [18]:

$$\ln\left(\frac{T_c^2}{C}\right) = \frac{E_c}{RT_c} + \text{const.} \quad (1)$$

where  $T_c$ ,  $C$ , and  $R$  are the crystallization peak temperature, heating rate, and universal gas constant respectively. Fig. 3(b) demonstrates the linear relationship between  $\ln(T_c^2/C)$  and  $(1/T_c)$ . From the slope of the curve, the value of  $E_c$  can be easily determined to be 850 kJ/mol. It has been reported that the activation energy of ZrTiO<sub>4</sub> crystallization from amorphous powders prepared by sol-gel methods is 620 kJ/mol with the crystallization peak appearing at 642 °C [19] and 139-323 kJ/mol with the peak becoming evident at 530-680 °C [2].

Compared to the results of this work, it can be concluded that the activation energy of ZrTiO<sub>4</sub> crystallization strongly depends on the preparation conditions. The activation energy of

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crystallization calculated by the Kissinger's equation is the activation energy required for growth [20]. The growth of nuclei needs an atomic rearrangement of the constituents, which is closely related to their diffusivity. Indeed, the difference of the activation energy calculated for this material synthesized by different procedures might be due to differences in the particle size, degree of hydration, homogeneity, and short-range order of the amorphous phase [2].

To assess the structural evolution in the course of heating, particularly in the case of isothermal heat treatment, the XRD analyses were conducted on the xerogel before and after calcination at 500, 550, and 600 °C, as represented in Fig. 4. The broad halo pattern of the xerogel [Fig. 4(a)] suggests a typical amorphous pattern, as confirmed by its SAD pattern [Fig. 1(c)]. As can be observed, the amorphous structure is still stable at 500 °C [Fig. 4(b)]; however, the appearance of a number of ZrTiO<sub>4</sub> reflections in the XRD pattern of the sample calcined at 550 °C, provided in Fig. 4(c), is indicative of the onset of structural ordering and crystallization. Fig. 4(d) specifies the XRD pattern of the powder calcined at 600 °C, inferring the crystalline ZrTiO<sub>4</sub> phase with a crystallite size of 13 nm, determined by the EVA software using the Scherrer equation for the XRD maxima. The instrument broadening was calibrated by using a standard sample. Note that it is reasonable to neglect the strain contribution to broadening at low Bragg angles [21]. The XRD reflections were also confirmed by the ICSD reference code of 00-034-0415 for zirconium titanate with the orthorhombic  $\alpha$ -PbO<sub>2</sub> type structure, in which Zr and Ti atoms are in octahedral oxygen coordination. No XRD reflection corresponding to TiO<sub>2</sub> and ZrO<sub>2</sub> can be detected. The direct crystallization of ZrTiO<sub>4</sub> at the low temperature is due to the high level of homogeneity of the xerogel. It has been reported that, in the case of low homogeneity, TiO<sub>2</sub> crystallization occurs before

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crystallization into ZrTiO<sub>4</sub> [22]. The TEM micrograph and SAD pattern of the sample calcined at 600 °C are illustrated in Fig. 5. Compared to the xerogel (Fig. 1), the powder particles have grown, with the driving force being a reduction in the surface energy, and have approached an average size of 50 nm with a relatively wide size range of 20 nm to 100 nm. Moreover, the high crystallinity of this powder results in distinct Debye-Scherrer diffraction spots in the SAD pattern (Fig. 5(b)).

As shown above, based on the XRD analysis, the crystallization is initiated at 550 °C (even at lower temperatures), but the crystallization peak of the DSC test is located at 690 °C. The difference of the crystallization behavior detected from the DSC and XRD evaluations is due to the difference between the thermal regimes applied. During the DSC analysis, heating is continuous at a rate of 10 °C/min; nonetheless, the XRD patterns were obtained from the samples calcined at the different temperatures for 1 h under isothermal conditions, where time for the diffusional transformation of crystallization is provided, allowing the onset of crystallization at low temperatures.

In conclusion, typical advantages of this processing route described in this paper include: i) It is a straightforward and uncomplicated procedure, ii) It allows for the successful synthesis of ZrTiO<sub>4</sub> nanoparticles with a crystallization temperature much lower than in classical ZrTiO<sub>4</sub> processing methods, and iii) It is capable of producing low cost powders compared to polymeric sol-gel methods using alkoxide precursors.

#### **4. Conclusions**

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In this work, an aqueous sol-gel procedure using zirconium and titanium chlorides was developed to synthesize ZrTiO<sub>4</sub> powder. From this study, the following conclusions could be drawn:

- According to the TEM studies, amorphous particles of almost 5 nm in size with a relatively spherical morphology were prepared.
- The crystallization peak temperature of the amorphous powder prepared by the sol-gel route was found to be nearly 690 °C, as determined by the DSC/TGA experiment at a heating rate of 10 °C/min.
- Using the Kissinger's equation, the activation energy for ZrTiO<sub>4</sub> crystallization was determined to be 850 kJ/mol.
- Calcination at 550 °C for 1 h developed ZrTiO<sub>4</sub> nanocrystals of 15 nm in size determined by XRD. In addition, high crystallinity and particle growth were confirmed by TEM studies.

### **Acknowledgements**

This report is based upon work supported by National Science Foundation (NSF) under grant no. 0933763.

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## Figures

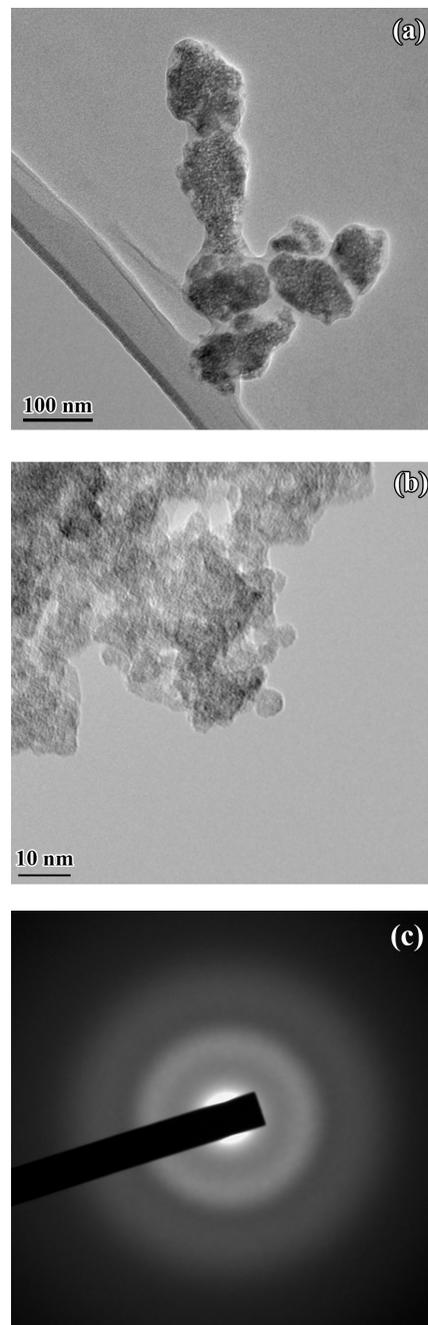


Fig. 1. TEM micrograph of the xerogel powder particles showing agglomeration (a), that showing nanoparticles of 5 nm in size (b), and the related SAD pattern suggesting an amorphous phase (c).

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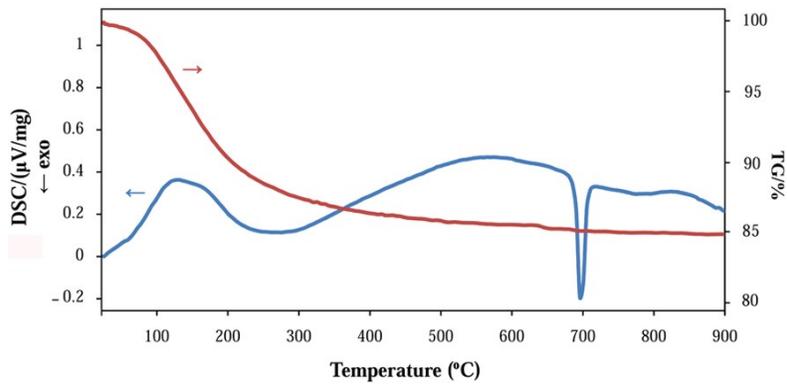


Fig. 2. TGA and DSC profiles of the xerogel at a heating rate of 10 °C/min.

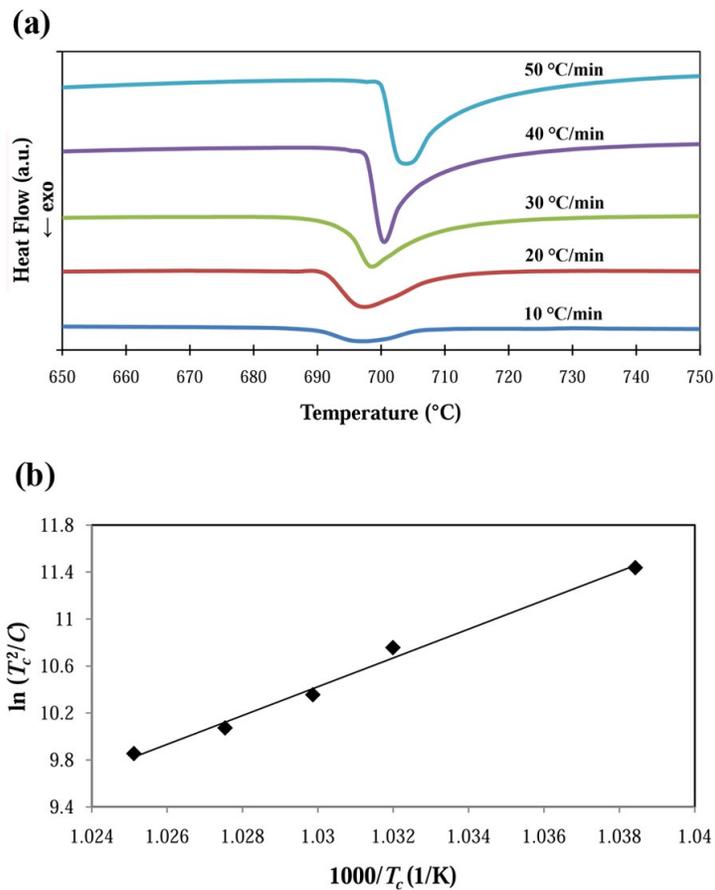


Fig. 3. DSC curves of ZrTiO<sub>4</sub> crystallization obtained at the different heating rates (a) and Kissinger plot for ZrTiO<sub>4</sub> crystallization (b).

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<https://doi.org/10.1016/j.ceramint.2012.04.064>

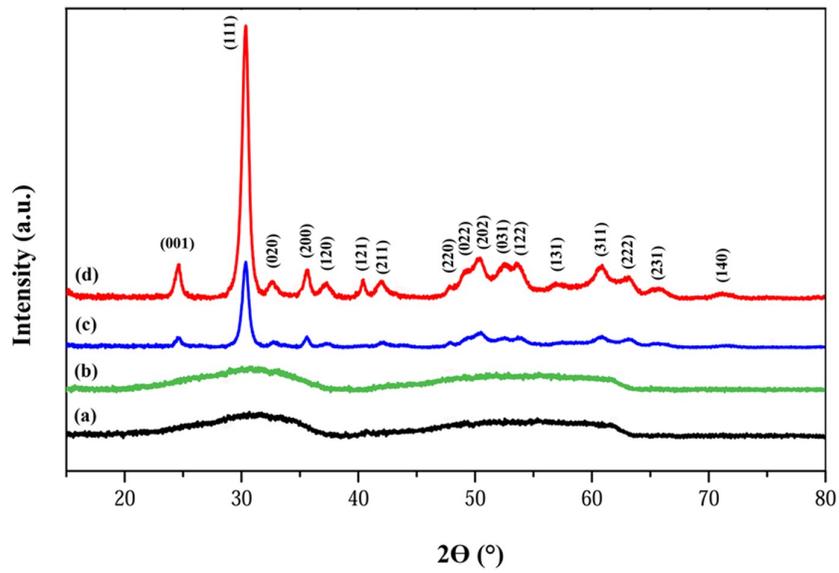


Fig. 4. XRD pattern of the xerogel (a) and the powders calcined at 500 (b), 550 (c), and 600 (d) °C.

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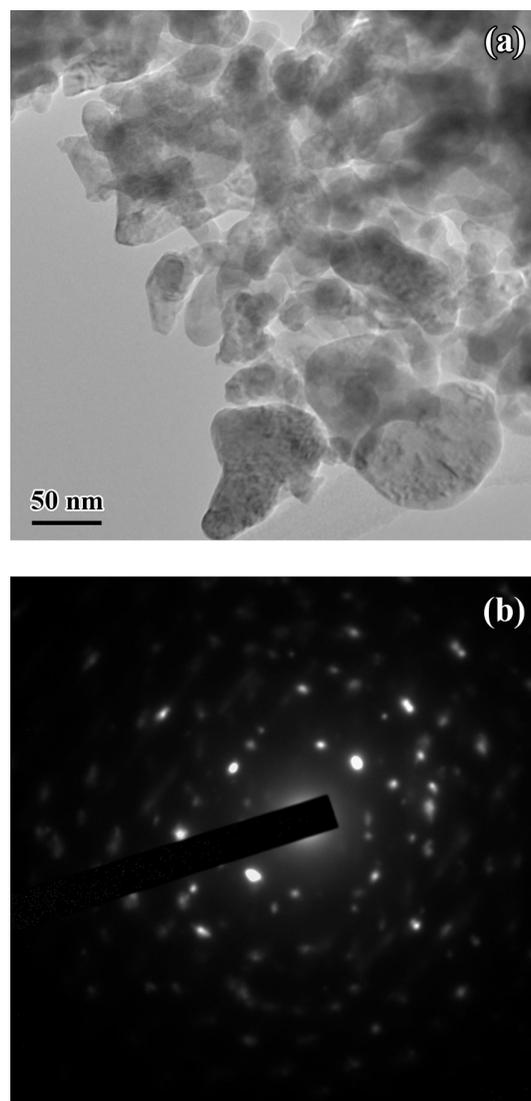


Fig. 5. TEM micrograph (a) and SAD pattern (b) of the powder specimen calcined at 600 °C.