



HAL
open science

Crystallization Pathway of Size-Controlled SnO₂ Nanoparticles Synthesized via a Nonaqueous Sol–Gel Route

Alex Lemarchand, Fabien Remondiere, Jenny Jouin, Philippe Thomas, Olivier Masson

► **To cite this version:**

Alex Lemarchand, Fabien Remondiere, Jenny Jouin, Philippe Thomas, Olivier Masson. Crystallization Pathway of Size-Controlled SnO₂ Nanoparticles Synthesized via a Nonaqueous Sol–Gel Route. *Crystal Growth & Design*, 2019, 20 (2), pp.1110-1118. 10.1021/acs.cgd.9b01428 . hal-02483342

HAL Id: hal-02483342

<https://unilim.hal.science/hal-02483342v1>

Submitted on 26 Nov 2020

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

This document is confidential and is proprietary to the American Chemical Society and its authors. Do not copy or disclose without written permission. If you have received this item in error, notify the sender and delete all copies.

Crystallization pathway of size-controlled SnO₂ nanoparticles synthesized via a nonaqueous sol-gel route

Journal:	<i>Inorganic Chemistry</i>
Manuscript ID	ic-2019-03000p
Manuscript Type:	Article
Date Submitted by the Author:	11-Oct-2019
Complete List of Authors:	Lemarchand, Alex; IRCER REMONDIERE, Fabien; Universite de Limoges, Chemistry Jouin, Jenny; Universite de Limoges, Laboratoire SPCTS Thomas, Philippe; Science des Procedes Ceramiques et de Traitements de Surface, Université de Limoges Masson, Olivier; Universite de Limoges, Chemistry

SCHOLARONE™
Manuscripts

1
2
3
4
5
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24
25
26
27
28
29
30

Crystallization pathway of size-controlled SnO₂ nanoparticles synthesized *via* a nonaqueous sol-gel route

31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60

*Alex Lemarchand, Fabien Rémondière, Jenny Jouin, Philippe Thomas and Olivier Masson**

Institute of Research for Ceramics, UMR 7315 CNRS - Université de Limoges, Centre Européen
de la Céramique – 12, rue Atlantis 87068 Limoges

Abstract. This paper deals with the crystallization pathway of tin dioxide ultra-small nanoparticles synthesized *via* a nonaqueous sol-gel route based on the etherolysis of tin tetrachloride (SnCl₄) precursor. Two different ethers were used as oxygen donors: diisopropyl ether (ⁱPr₂O) and dibenzyl ether (Bn₂O). The solvothermal treatments of SnCl₄ in ⁱPr₂O at different temperatures and reaction times, revealed in the early stages of the reaction the formation of an organic polymeric phase in which crystalline nanoparticles are embedded. In the case of the SnCl₄-Bn₂O system, the formation of a brownish polymer containing crystalline oxidic nanoparticles and the production of water were evidenced. With both ethers, an unexpected intermediate disordered nano-crystalline phase is formed, which progressively transforms into rutile-type SnO₂ nanoparticles. The formation of the polymeric phase, which is suspected to act as a trap for the nanoparticles, freezes the intermediate phase, limits the particle growth and delays the restructuring into the rutile-type structure. The

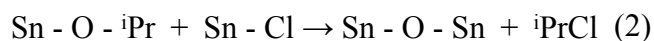
1
2
3 degradation of the polymer phase in the case of $i\text{Pr}_2\text{O}$ and/or the water action in the case of Bn_2O
4
5 are finally able to destabilize the nanoparticle-polymer assembly and to induce the restructuration
6
7 of the nanoparticles.
8
9

10
11 **Keywords.** Nonaqueous sol-gel synthesis; Tin dioxide nanoparticles; Etherolysis; Crystallization
12
13 pathway; XRD; TEM
14
15

16 17 **Introduction**

18
19 Tin (IV) dioxide with rutile-type structure (i.e. *cassiterite* mineral) [1] is a well-known
20
21 semiconductor with a band gap of 3.6 eV at 300 K widely used as solid-state gas sensor material
22
23 [2][3], oxidation catalyst [4] and transparent conductor [5]. For all these applications, a high surface
24
25 area is of great importance, which in turn emphasizes the interest of producing nano-sized particles.
26
27 Various synthesis methods are found in literature as for example flame pyrolysis [6], thermal
28
29 evaporation [7], laser-ablation techniques [8] and the common solution chemistry methods[1].
30
31 Among the latter, sol-gel aqueous [9][10][11][12][13][14] and nonaqueous routes [15][16][17] are
32
33 particularly powerful for obtaining particles of very small and controlled size. The nonaqueous
34
35 method offers the advantage of producing well-crystallized nanoparticles at moderate temperature
36
37 without the need of any additional post-thermal treatment [18][19][20]. Particularly interesting are
38
39 the solvent-controlled syntheses in which the oxygen donor solvent also controls the particle size
40
41 [18]. Among them, the so-called “benzyl alcohol route”, in which benzyl alcohol is used as solvent,
42
43 is one of the most versatile. This route has been applied in the case of tin oxide using both tin (IV)
44
45 chloride and tin (IV) *tert*-butoxide [16][17] and allowed obtaining nanoparticles of 3.5 nm and
46
47 2.2 nm respectively. However, the use of such protic solvent can lead to a water release in the
48
49 reaction medium (through solvent condensation and hydroxylation reactions), which is detrimental
50
51
52
53
54
55
56
57
58
59
60

1
2
3 for the control of the particle growth. As an alternative, the use of an aprotic oxygen donor was
4 developed by Mutin and Vioux [19][20]. The chemical pathway involved in the formation of the
5 nanoparticles is then simplified and the control over the final product is easier to achieve. This
6 method was recently applied to the synthesis of SnO₂ nanoparticles [15] and is based on the reaction
7 of SnCl₄ with a stoichiometric amount of diisopropyl ether. Organo-soluble and reactive SnO₂
8 nanoparticles exhibiting an average diameter of 4 nm with a narrow size distribution were obtained
9 with an excellent yield. The proposed synthesis mechanism is a two-step process that consists in:
10 (i) the etherolysis of the tin chloride precursor leading to the formation of tin isopropoxide groups
11 and (ii) the condensation between the isopropoxide and the chloride leading to the formation of the
12 oxidic network jointly to the release of isopropyl chloride. The corresponding chemical reactions
13 (for one ligand only) are given below:
14
15
16
17
18
19
20
21
22
23
24
25
26
27



28
29
30
31
32
33 The effective control of the nanoparticle size requires the understanding of the nanoparticle
34 formation mechanism, which is most of the time not fully understood since the classical nucleation-
35 growth theory based on the LaMer model is not pertinent in numerous cases. Recent studies
36 emphasize alternative pathways [21][22][23][24][25] (not simply based on the attachment of ions,
37 atoms or molecules to form an individual cluster) such as oriented attachment, [26][27][28][29][30]
38 mesocrystal formation [31][32] or formation of intermediate metal-organic phase. For example, Ba
39 *et al.* [33] have recently shown that the solvothermal synthesis of indium-tin oxide (ITO)
40 nanoparticles in benzyl alcohol was a two-stage process involving the formation of an
41 undetermined intermediate phase. This latter, characterized by small particles connected in an
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60

1
2
3 organic matrix, transforms with time into the final ITO nanoparticles during the solvothermal
4
5 treatment.
6

7
8 In this paper, we present a study of the early stages of crystallization of SnO₂ nanoparticles
9
10 obtained *via* a modified nonaqueous sol-gel synthesis inspired by the work of Aboulaich *et al.*[15].
11
12 In this context, two ethers were used. Firstly, ⁱPr₂O, which already showed a great ability to produce
13
14 crystalline SnO₂ nanoparticles of small size, is reinvestigated to evidence a possible unusual
15
16 crystallization sequence. Secondly, Bn₂O, which is susceptible to induce spatial hindrance because
17
18 of its aromatic ring and thus to induce a control on the particle size, is newly tested here. Moreover,
19
20 since aromatic solvents are susceptible to polymerize and to release water, the non-hydrolytic
21
22 feature of the reaction was questioned and investigated.
23
24

25 26 **Experimental section**

27
28 **Chemicals.** Tin (IV) tetrachloride (SnCl₄, 98%) was purchased from Strem Chemicals.
29
30 Diisopropyl ether (ⁱPr₂O, 99%) and dibenzyl ether (Bn₂O, 99%) were purchased from Sigma
31
32 Aldrich. Other solvents used for the synthesis and the flocculation of the nanoparticles and the
33
34 TEM grid preparation are respectively dichloromethane (99+%, Alfa Aesar), chloroform (99.8%,
35
36 Alfa Aesar), acetone (99+%, Alfa Aesar), n-heptane (99%, Fisher Scientific), 1-butanol (99%, Alfa
37
38 Aesar). All the solvents were utilized without any further purification. The anhydrous feature of
39
40 the ethers was confirmed by Karl-Fisher analysis.
41
42

43
44 **Synthesis of SnO₂ nanoparticles.** All the manipulations were carried out in a glovebox under
45
46 dry air atmosphere. The Sn⁴⁺ concentration was set to 0.5 mol.L⁻¹. This relatively high
47
48 concentration was set to minimize the economic and ecological impacts of the process.
49
50

51
52 **Syntheses with diisopropyl ether.** The typical synthesis of the tin oxide nanoparticles was
53
54 performed by adding SnCl₄ (5 mmol), ⁱPr₂O (15 mmol, [ether]/[Sn⁴⁺]=3.0) in CH₂Cl₂, to obtain a
55
56

1
2
3 Sn^{4+} concentration of 0.5 mol.L^{-1} . The mixture was then transferred in a Teflon cup of 25 mL inner
4
5 volume, placed in a steel autoclave, and heated at the targeted temperature under autogenous
6
7 pressure. To study the influence of the reaction temperature and reaction time, the heat treatment
8
9 was performed in a range of 95°C - 140°C for at least 12 h and up to 140 h. All the synthesis
10
11 parameters are summarized in Table 1.
12
13

14 ***Syntheses with dibenzyl ether.*** In this case, the synthesis of the tin oxide nanoparticles was
15
16 performed by adding SnCl_4 (7.5 mmol) in Bn_2O (72.75 mmol, $[\text{ether}]/[\text{Sn}^{4+}] = 9.7$) in order to obtain
17
18 a total volume of 15 mL. Preliminary experiments showed that the use of an ether to tin molar ratio
19
20 of 3.0 did not lead to the formation of any inorganic solid product. It was thus increased to a value
21
22 of 9.7 to ensure the formation of a powder. The reactive mixture was transferred in a sealed glass
23
24 tube and heated by a conventional oil bath. To study the influence of the reaction temperature and
25
26 reaction time, the heat treatment was performed in a range of 95°C - 140°C for at least 30 min and
27
28 up to 60 h. All the synthesis parameters are summarized in Table 1.
29
30
31

32 **Washing procedure, supernatant characterization and powder recuperation.** The brownish
33
34 suspensions obtained in the case of ${}^i\text{Pr}_2\text{O}$ were washed and centrifuged (10 minutes, 10,000 rpm)
35
36 with first a 20 mL volume mixture of heptane and acetone and finally twice with 20 mL of acetone
37
38 to eliminate the majority of the organic residues. The dark brown polymer obtained in the case of
39
40 Bn_2O was first dissolved in several portions of chloroform under a time-consuming shaking
41
42 procedure (≈ 2 -4 hours). This operation was repeated until most of the polymer was dissolved and
43
44 removed. The unsolubilized part of the sample was finally washed and centrifuged (10 minutes,
45
46 10,000 rpm) twice in 20 mL of chloroform, once with a 20 mL volume mixture of heptane and
47
48 acetone and finally twice with 20 mL of acetone to remove as much as possible of the remaining
49
50 polymer. In both cases, the as-obtained precipitate was finally collected with 15 mL of
51
52
53
54
55

dichloromethane and air dried overnight at room temperature. The final product was softly ground to powder for the chemical characterizations and kept a light-cream colour with a slight brownish tint. The characteristics of the synthesis products are summarized in Table 1. During the washing step, the first supernatant was isolated for each sample to highlight the potential presence of water during the synthesis. For that purpose, a test based on the colour change of anhydrous copper sulphate was performed. To this end, one drop of the suspension composed of the synthesis liquid and of the antisolvent mixture was deposited onto the anhydrous copper sulphate powder. The solvent used during the washing step were tested beforehand to ensure they were water-free.

Table 1. Experimental parameters for the synthesis of SnO₂ nanoparticles and corresponding reaction products.

Sample Label	[Sn ⁴⁺] (mol.L ⁻¹)	Oxygen donor	[ether]/ [Sn ⁴⁺]	Solvothermal treatment		Products
				Temperature (°C)	Reaction time (h)	
Pr-95°C-12h	0.5	ⁱ Pr ₂ O	3.0	95	12	Powder, liquid supernatant
Pr-95°C-24h					24	
Pr-95°C-60h					60	
Pr-95°C-100h					100	
Pr-95°C-140h					140	
Pr-110°C-60h				110	60	
Pr-110°C-100h					100	
Pr-110°C-140h					140	
Pr-125°C-60h				125	60	
Pr-125°C-100h					100	
Pr-140°C-60h				140	60	

Bn-95°C-60h				95	60	Polymer, powder, aqueous phase
Bn-110°C-4h				110	4	
Bn-110°C-12h					12	
Bn-110°C-24h		Bn ₂ O	9.7		24	
Bn-110°C-60h					60	
Bn-125°C-2h					125	
Bn-140°C-0.5h				140	0.5	

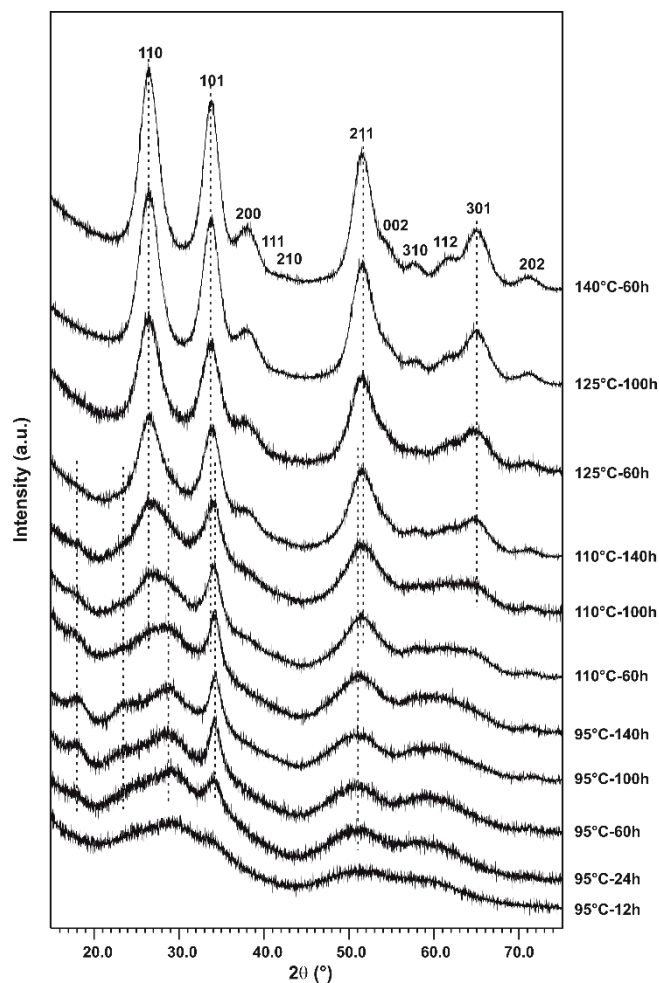
Sample characterization. The phase identification and the structural and microstructural characterizations of the samples were performed by X-ray powder Diffraction (XRD). Data were collected with the CuK α_1 radiation (1.5406 Å) from 10° to 110° (2 θ) with a step size of 0.020° (2 θ) and an equivalent counting time per step of 1567s using a Bragg-Brentano (θ -2 θ) BRUKER D8 Advance diffractometer equipped with a rapid LYNXEYE detector. The diffraction patterns were treated using the Rietveld method [34][35] implemented in the Fullprof software [36][37] and by peak by peak decomposition using the Peakoc software [38]. Particle size and micro-strains were estimated using both the Pseudo-Voigt line profile function in its Thompson-Cox-Hastings formulation [39][40] (Fullprof) and the Voigt function [41] (Peakoc). The characterizations were supplemented by High-Resolution Transmission Electron Microscopy (HRTEM) and Selected Area Electron Diffraction (SAED) using a JEOL 2100f. The samples were prepared by evaporating a drop of a diluted nanoparticle suspension onto an ultrathin carbon-A holey 400 mesh copper grid. This suspension was prepared by the dilution of one drop of the final suspension obtained at the end of the washing step in 10 mL of n-butanol. The average particle size was determined from the micrographs by the analysis of a set of 30 particles for each sample. For two samples, the 1D

1
2
3 electron powder diffraction diagrams were obtained by integrating sectors of the SAED images
4
5 using the FIT2D software [42].
6
7
8
9

10 **Results and discussion**

11
12 **Syntheses with diisopropyl ether.** The copper sulphate test conducted on the first synthesis
13
14 supernatant was negative for all the samples synthesized using $^i\text{Pr}_2\text{O}$ indicating that the formation
15
16 mechanism of the SnO_2 nanoparticles by this procedure is non-hydrolytic. The XRD patterns of
17
18 the samples prepared with $^i\text{Pr}_2\text{O}$ at 95°C , 110°C , 125°C and 140°C between 12 h and 140 h are
19
20 plotted in Figure 1. For the syntheses at 125°C and 140°C , the diagrams are typical of a single
21
22 rutile-type phase. This is confirmed by the good quality Rietveld refinements obtained for these
23
24 samples (Figure 2). The refined microstructural parameters correspond to a negligible amount of
25
26 microstrains and average particle diameters of about 3.7 nm and 3.3 nm for 140°C (60h) and 125°C
27
28 (100h) respectively. For a synthesis time reduced to 60 h at 125°C , the average particles diameter
29
30 decreases to about 2.7 nm. For lower temperatures (110°C and 95°C), the evolution of the XRD
31
32 patterns does not suggest a simple particle size decrease, but instead some important structural
33
34 changes. At 110°C , when the synthesis reaction time is reduced from 140 h to 60 h, the 110 peak
35
36 undergoes an huge asymmetric enlargement and a shift of its maximum position from $2\theta = 26.5^\circ$
37
38 to $2\theta = 27.5^\circ$ whereas the 101 peak remains almost unchanged in terms of width and position. In
39
40 addition, two peaks appear at $2\theta = 18.0^\circ$ and 23.5° . The rest of the diagram mainly undergoes a
41
42 progressive enlargement, with the exception of the 301 peak which disappears progressively. At
43
44 60h, the XRD diagram has clearly very broad lines apart from the narrow 101 peak. A similar
45
46 feature was already observed by Aboulaich *et al.* for samples synthesized following an equivalent
47
48 synthesis method but was not considered further [15]. At 95°C , for reaction times decreasing from
49
50
51
52
53
54
55
56
57
58
59
60

1
2
3 140 h to 60 h, the diagrams keep evolving similarly to finally exhibit only a narrow 101 peak and
4 two small bumps at $2\theta = 18.0^\circ$ and 23.5° . When the reaction time is further reduced to less than
5
6 two small bumps at $2\theta = 18.0^\circ$ and 23.5° . When the reaction time is further reduced to less than
7
8 24 h, these features are hardly noticeable but present.
9



41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60

Figure 1. XRD patterns of the samples obtained at different temperatures between 95°C and 140°C for different reaction times in the range of 12 h to 140 h. The pattern of Pr- 140°C -60h is indexed according to the rutile-type structure. Dotted lines to guide eye correspond to the main reflections observed on the different patterns.

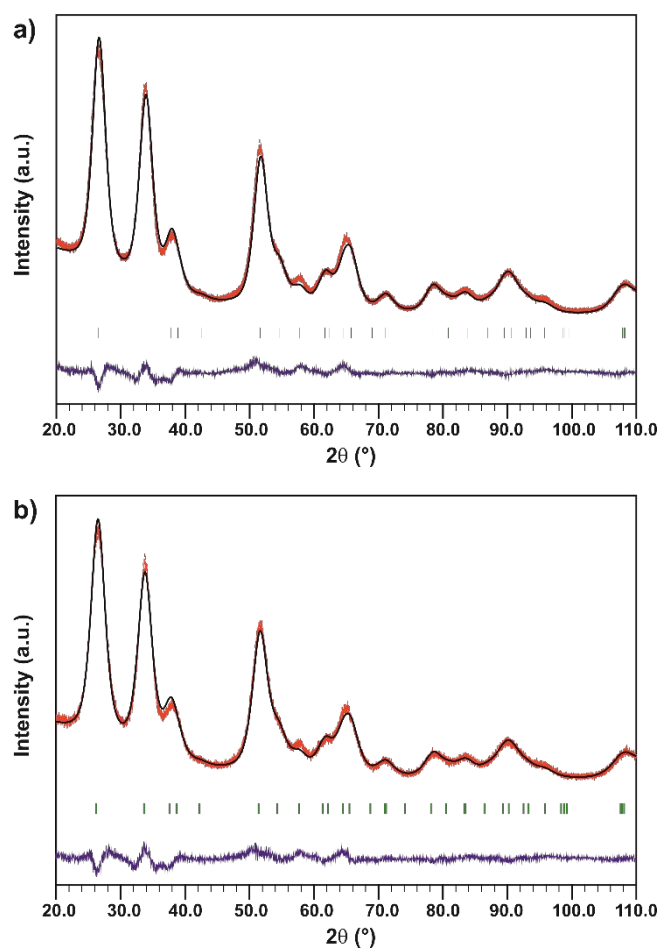
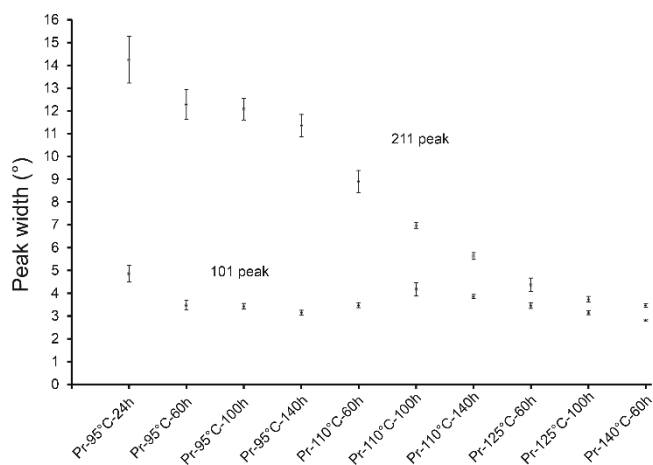


Figure 2. XRD patterns and Rietveld refinements with a rutile-type model for Pr-140°C-60h SnO₂ (a) and Pr-125°C-100h SnO₂ (b) samples.

Despite these strong changes, all XRD patterns share the same global envelop, in particular with common peaks at $2\theta \approx 34^\circ$, 51° and 71° assimilated respectively to the 101, 211 and 202 peaks of the rutile-type structure. These correlations suggest that the structure of the samples undergo a continuous evolution of an intermediate phase toward the rutile-type phase while increasing time and temperature instead of having biphasic samples, as for example a mixture of the rutile-type structure and an unknown metastable phase or a progressive conversion of an amorphous phase into the rutile-type structure.

1
2
3 The evolution of the integral widths of the 211 and 101 peaks as a function of the synthesis
4 conditions is plotted in Figure 3. The first interesting feature is that the 101 line width remains
5 almost constant, whatever the synthesis conditions. From this width, one can estimate using the
6 Scherrer equation a coherent domain size, or correlation length, perpendicular to the (101) planes.
7 We found it is about 2.7 nm. The situation is completely different for the 211 peak, which
8 undergoes large variations with the synthesis conditions. The corresponding correlation lengths
9 vary from a few angstroms (0.7 nm for 95°C-24 h) to a few nanometers (2.8 nm for 140°C-60 h).
10 These observations suggest that a precursor phase of the rutile-type structure is initially formed
11 with a structure relatively well ordered perpendicularly to the (101) lattice planes and almost no
12 order, i.e. a complete loss of coherency, in perpendicular directions. With the increase of the
13 synthesis temperature and reaction time, the structuration of the particles seems to happen gradually
14 along these directions until the final rutile-type is obtained.



31
32
33
34
35
36
37
38
39
40
41
42
43
44
45 **Figure 3.** Evolution of the peak width as a function of the synthesis parameters for the samples
46 prepared with ${}^1\text{Pr}_2\text{O}_3$.

47
48
49
50
51 In order to clarify this possible structuration of the system, TEM measurements were performed
52 on the following selected samples: (i) high temperature synthesis (125°C) for which a pure rutile-

1
2
3 type phase is obtained, (ii) low temperature synthesis (95°C) during the first stages of the
4
5 structuration when the intermediate phase appears and finally (iii) intermediate temperature
6
7 synthesis (110°C) during the restructuring of the samples from the intermediate phase to the
8
9 rutile-type one. All the TEM micrographs are presented in Figure 4.
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60

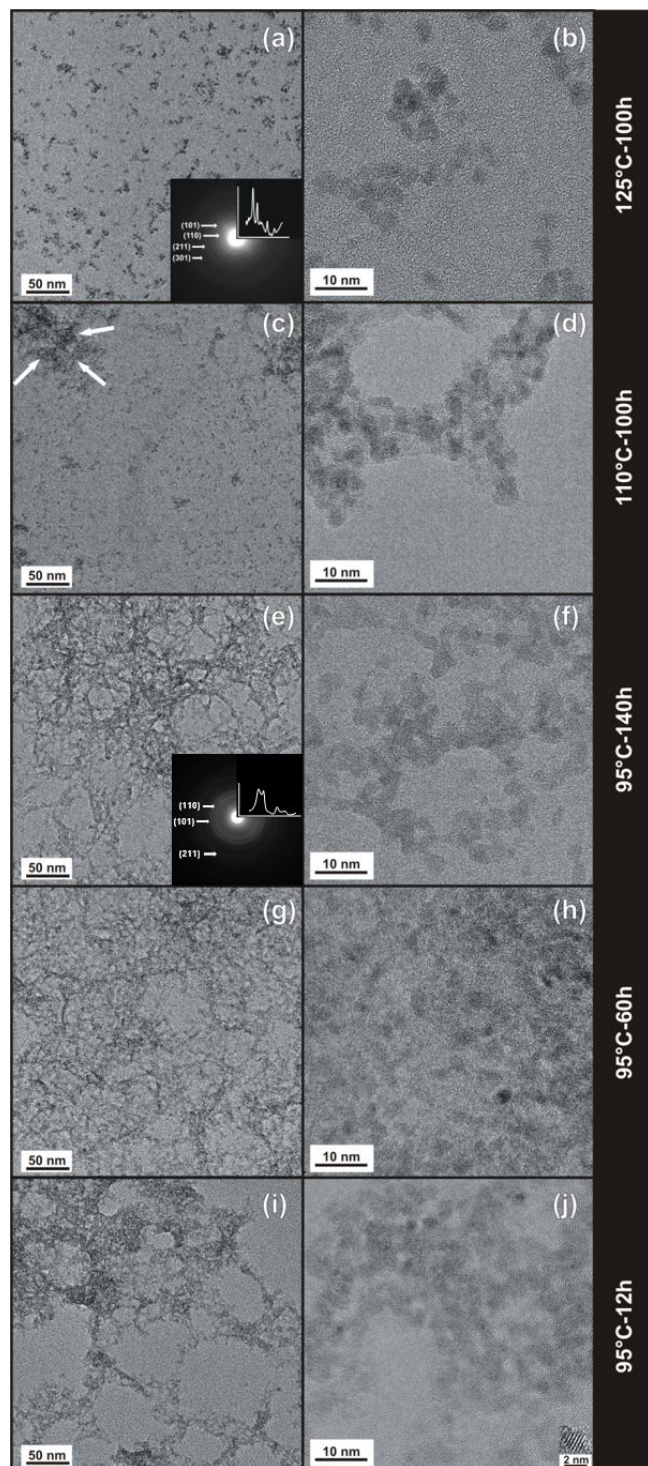


Figure 4. TEM micrographs of Pr-125°C-100h (a and b), Pr-110°C-100h (c and d) Pr-95°C-140h (e and f), Pr-95°C-60h (g and h) and Pr-95°C-12h (i and j). Arrows in (c) indicate heterogeneous

1
2
3 feature. Inset in (a) and (e): SAED pattern and 1D electron powder diffraction diagram obtained
4
5 with FIT2D.
6
7

8
9 In the case of the sample synthesized at 125°C for at least 100 h (Figure 4 a-b) isolated and
10 slightly aggregated crystallized nanoparticles are visible. The particles exhibit a spherical or
11 slightly ovoid shape and an estimated size ranging from 2.6 nm up to 4.8 nm with an average size
12 of 3.6 ± 0.3 nm, consistent with the XRD measurements. The SAED pattern shows continuous and
13 broad rings and the corresponding 1D electron powder diffraction diagram is typical of the rutile-
14 type structure. For all the samples obtained at 95°C, a special architectural feature is visible, namely
15 a network of connected crystalline nanoparticles surrounded by an amorphous matrix. The strong
16 chemical contrast between these two phases suggests that the amorphous phase is almost composed
17 of light chemical elements and thus could be organic in nature. For the Pr-95°C-12h sample
18 (Figure 4 i-j), the nanoparticles have a size ranging from 2.2 to 3.2 nm, with an average value of
19 2.8 ± 0.2 nm. The shape of the nanoparticles is hard to define since they are embedded into the
20 organic phase but it is observed that the nanoparticles are connected to each other without any
21 preferred orientation. For the Pr-95°C-60h sample (Figure 4 g-h), the typical arrangement is still
22 present and no drastic change is observed. For the Pr-95°C-140h sample (Figure 4 e-f), the lace
23 formed by the branching of the organic-nanoparticles mixture looks less dense and seems to tear
24 apart at the periphery of the large assembly shown on the micrographs. At this step, the lace is
25 almost exclusively composed of quasi-spherical nanoparticles stuck together with a very low
26 amount of organics. The Pr-110°C-100h micrographs (Figure 4 c-d) present both characteristics
27 previously exposed, namely isolated nanoparticles or partially agglomerated nanoparticles as well
28 as the specific organic-nanoparticles assembly. Nanoparticles embedded in the organic matrix have
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60

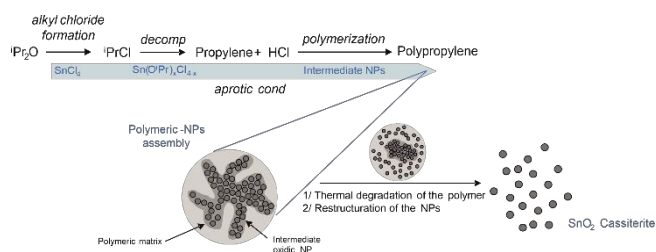
1
2
3 a size of 2.9 ± 0.2 nm while a few isolated nanoparticles, which have certainly underwent a
4 moderate growth, have a size of 3.3 ± 0.2 nm.
5
6

7
8 On all the TEM micrographs of Figure 4, particles appear crystallized as some lattice planes are
9 clearly resolved. This is also the case for the samples synthesized at low temperature (95°C) for
10 which the XRD patterns have very broad peaks (Figure 1), corresponding to small correlation
11 lengths typical of amorphous phase (Figure 3). This seemingly contradictory observations led us
12 to suspect that the microscope electronic beam could potentially induce crystallization. However,
13 the 1D electron diffraction diagram of Pr- 95°C -140h (Figure 4e) is clearly similar to that obtained
14 by XRD (Figure 2), in particular a first broad 110 peak and a second narrow 101 peak. This attests
15 that the electronic beam did not induce crystallization of the samples and that the organic-
16 nanoparticle assembly highlighted by microscopy is indeed representative of the sample, the XRD
17 diagram being more sensitive to the crystalline part of the assembly since the scattering of the
18 organic matrix is low.
19
20
21
22
23
24
25
26
27
28
29
30
31
32

33 By elevating the synthesis temperature and/or extending the reaction time, the disappearance of
34 the organic phase and the formation of rutile-type structure nanocrystals seem to concomitantly
35 happen. Here, we suggest a correlation between the two phenomena. The organic phase could act
36 as a trap for the native nanoparticles and thus limit their growth and restructuration towards the
37 rutile-type structure due to the difficulty for the precursor to diffuse or/and due to a temperature-
38 limited restructuration process.
39
40
41
42
43
44
45
46

47 The reactional mechanism leading to SnO_2 nanocrystals based on the etherolysis of tin chloride
48 thus seems complex. As it is suggested by Aboulaich et al., after an induction period, the tin
49 chloroalkoxide $\text{Sn}(\text{O}^i\text{Pr})_x\text{Cl}_{4-x}$ issued from the etherolysis of the tin (IV) chloride precursor and/or
50 the native nanoparticles play a catalytic role on the thermal decomposition of $^i\text{PrCl}$, leading to the
51
52
53
54
55
56
57
58
59
60

1
2
3 formation of propylene and HCl, and then to the polymerization of propylene [15][43][44]. These
4
5 reactions could be responsible for the formation of the organic or polymeric-nanoparticles
6
7 assembly that we evidenced for the lower synthesis temperature and shorter reaction time. Upon
8
9 heating at higher temperature, this original feature progressively disappears and only isolated
10
11 particles or aggregates are observable. At intermediate temperature (110°C), the double feature of
12
13 the sample suggests that the polymeric phase is thermally and chemically resorbing since isolated
14
15 nanoparticles are detected. Polymer degradation is known to be commonly related to thermal and
16
17 chemical effects [45]. Thereby, the increase of the synthesis temperature and the potential presence
18
19 of unreacted alkoxide groups trapped inside the polymeric matrix are thought to promote the
20
21 degradation process and the growth of the nanoparticles. However, since the particles are stuck
22
23 inside the matrix, due to the lack of mobility of the nanoparticles and to the limited diffusion of
24
25 reactants, only a slight growth is possible. Once released from the matrix, the particles are free to
26
27 grow via Ostwald ripening and to restructure into the rutile-type structure. The whole mechanism
28
29 is non-hydrolytic since no water was evidenced in the synthesis supernatant. As a conclusion, a
30
31 schematic and simplified summary of this sequence of reactions, leading first to the formation of
32
33 nanoparticles with the intermediate structure and then to their restructuration into the rutile-type
34
35 structure, is presented in Figure 5.



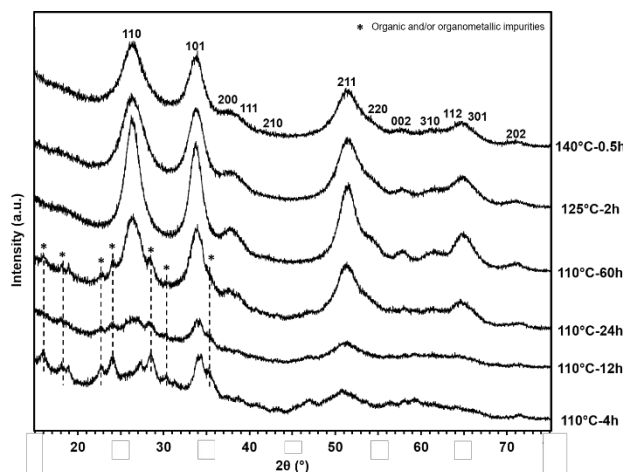
51 **Figure 5.** Illustration of the proposed crystallization pathway of SnO₂ nanoparticles synthesized
52
53 from a mixture of SnCl₄ and iPr₂O.
54
55

6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60

1
2
3 In order to investigate the impact of the solvent on the synthesis route, another ether, i.e. dibenzyl
4 ether, was used. This ether is thought to be susceptible to induce spatial hindrance through its
5 aromatic rings and thus to potentially decrease the particle size. It was also interesting to determine
6 its influence on the crystallization pathway of the nanoparticles. To the best of our knowledge, we
7 describe here the first sol-gel synthesis route of SnO₂ nanocrystals based on the use of dibenzyl
8 ether on tin chloride.
9

10
11
12 **Syntheses with dibenzyl ether.** During the washing step, the presence of a colourless liquid
13 phase which was immiscible with the organic solvents used during the procedure was evidenced
14 for all the samples. In contact to this liquid phase, the copper sulphate powder became blue thus
15 highlighting the presence of water in the system. The XRD pattern of the samples synthesized with
16 Bn₂O at 110°C, 125°C and 140°C between 30 min and 60 h are plotted in Figure 6. For 125°C and
17 140°C they are typical of a single rutile-type phase. The diffraction peaks are broad and the average
18 particle diameters are 2.8 nm and 2.5 nm for the syntheses at 125°C (2 h) and 140°C (0.5 h)
19 respectively. The sample synthesized at 110°C for 60 h has also a diagram typical of the rutile-type
20 phase but with narrower peaks corresponding to an average particle diameter of 3.7 nm. When
21 decreasing the reaction time from 24 h to 4 h, the diagrams evolve similarly to what was observed
22 with iPr₂O, namely they are characteristic of the intermediate phase evidenced previously, with in
23 particular, the presence of very broad peaks at $2\theta = 27^\circ$ and 51° and the narrower peak at 34° . This
24 evolution is however more difficult to evidence because the diagrams are polluted by numerous
25 extra peaks. As the Bn₂O samples exhibit a full organic polymerization of the reaction mixture,
26 these extra peaks are certainly the signature of an organic and/or organometallic crystalline phase
27 impurity. Even if it was not possible to perform TEM observations of the Bn₂O samples because
28 of the large remaining quantity of organic and polymeric phase, we can reasonably assume that the
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55

1
2
3 crystallization process is quite similar for both ethers. Some tests were conducted to limit the
4 production of the polymeric phase by adjusting the reaction time, the temperature, the dilution state
5 but without success. We can also notice that the ($\text{Bn}_2\text{O-SnCl}_4$) system enables the formation of
6 nanocrystalline material at shorter reaction times than with the (${}^i\text{Pr}_2\text{O-SnCl}_4$) system.
7
8
9
10
11



28 **Figure 6.** XRD patterns of the samples obtained at different temperatures between 110°C and
29 140°C for different reaction times in the range of 0.5 h to 60 h. The pattern of Bn-110°C-60h is
30 indexed according to the rutile-type structure.
31
32
33
34
35

36 We now consider the reaction sequence taking place during the nanoparticle synthesis. This is
37 related to the evolution of the initial reaction mixture composed of SnCl_4 and Bn_2O at 110°C, which
38 is quite surprising. At the beginning of the reaction, when the temperature is rising to 110°C, the
39 mixture remains brown and translucent. When the temperature reaches 110°C, the mixture becomes
40 opaque, darker, and a boil is noticeable after just a few seconds. Inside the mixture, some colourless
41 translucent drops appear and white solids form at the interface between these drops and the
42 brownish solution. Finally, the precipitate starts to accumulate on the walls of the tube. At the end
43 of the reaction, when the temperature decreases, the mixture is totally opaque, exhibits a caramel-
44 like colour and tends to harden during cooling down. These observations are evidences of three
45
46
47
48
49
50
51
52
53
54
55
56

1
2
3 concomitantly related phenomena during the synthesis: the polymerization of the reactional
4 mixture, the formation of an extra colourless liquid phase and the occurrence of a white precipitate.
5

6
7 Benzyl chloride and dibenzyl ether molecules are well-known to condense and polymerize in
8 certain conditions and in particular when certain Sn-species are present in the chemical system
9
10 [46][47][48][49]. In particular, in the case of tungsten oxide synthesis from tungsten isopropoxide
11 and benzyl alcohol, Olliges-Stadler *et al.* showed that native tungsten oxide nanowires catalyzed
12 the transformation of benzyl alcohol into dibenzyl ether and then the polymerization reaction of
13
14 dibenzyl ether into polybenzylene [50]. This reaction sequence led to a dark-blue hard and brittle
15 monolith composed of $W_{18}O_{49}$ nanowires embedded into polybenzylene polymer and also to the
16
17 release of water in the system. Based on these results, we propose that the polymeric phase we
18
19 evidenced is formed by the condensation of Bn_2O molecules and/or to the direct polymerization of
20
21 $BnCl$ issued of the etherolysis of $SnCl_4$ precursor. In both, reaction could be catalyzed by Sn
22
23 species, and lead to a polymeric matrix related to a member of the polybenzylene family. In
24
25 addition, as already stated, the colourless liquid phase recovered from the supernatant is water,
26
27 confirming the release of water during the synthesis. Since we worked with a concentrated mixture
28
29 of tin chloride and dibenzyl ether, it is not surprising that the polymerization of Bn_2O could be
30
31 activated and lead to a large release of water. Finally, the third phase, i.e. the white precipitate that
32
33 appeared at the interface of the aqueous and organic liquid phases, was recovered at the end of the
34
35 washing step and correspond to the metal oxide nanoparticles.
36
37

38
39 Considering now the global reaction pathway, we assume that the first step is strictly non-
40
41 hydrolytic and follows an alkyl halide elimination mechanism. Tin (IV) tetrachloride and/or tin
42
43 (IV) chloroalkoxide are then susceptible to catalyse dibenzyl ether polymerization and in parallel
44
45 to form the Sn-O-Sn network via aprotic condensation. The polymerization reaction leads to the
46
47

formation of polybenzylene and to the release of water, whose amount is proportional to the initial amount of Bn_2O and the reticulation rate. Water molecules released from dibenzyl ether condensation are then susceptible to switch the non-hydrolytic sol-gel process to a hydrolytic one. Doing so, they promote the precursor condensation, the particle nucleation and possibly the particle restructuring from the intermediate phase toward the final rutile-type structure, in particular in shorter time than with Pr_2O_3 at 110°C .

As a conclusion, a simplified scheme of these successive steps, involved in the formation of the SnO_2 nanoparticles *via* the etherolysis of SnCl_4 by Bn_2O , is presented Figure 7.

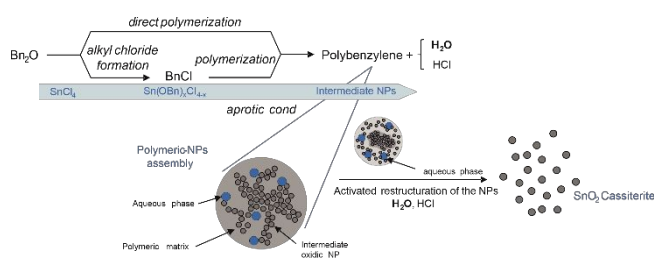


Figure 7. Illustration of the proposed crystallization pathway of SnO_2 nanoparticles synthesized from a mixture of SnCl_4 and Bn_2O .

Comparison of the two systems. In both systems, the reaction of ethers on tin (IV) tetrachloride lead to an original nanoparticle crystallization pathway involving the early formation of an intermediate precursor phase which progressively converts into the rutile-type structure. This is accompanied by the production of a polymer phase in variable quantity. We assume that the etherolysis/condensation reactions, leading to the intermediate phase formation, and the condensation/polymerization reactions, leading to the polymer formation, happen simultaneously. The polymeric phase then acts as a trap for the primary nanocrystals until the restructuring is possible.

1
2
3 In the case of diisopropyl ether, the polymerization is limited and confined near the particle
4 surface leading to the formation of the nanoparticle-polymer assembly observable on the TEM
5 micrographs (Figure 4). The restructuration depends then of the polymer degradation which
6 releases the particles. In the case of the use of dibenzyl ether, the formation of the polymeric phase
7 is more significant and the restructuration mechanism seems to happen essentially through the
8 release of water in the system. The compounds presenting aromatic rings, typically benzyl alcohol,
9 dibenzyl ether and benzyl chloride, show a great ability to polymerize, particularly in the presence
10 of tin (IV) tetrachloride, [47] and by this way are susceptible to produce hydrochloric acid and/or
11 water (for benzyl alcohol and dibenzyl ether). This ability seems to be less exacerbated in the case
12 of ethers which do not have aromatic ring, such as diisopropyl ether [44]. The difference of
13 reactivity between the two systems is probably related to the solvents properties and in particular
14 to their capacity to polymerize and to produce water since water promotes hydrolytic sol-gel
15 reactions and the restructuration of the intermediate nanoparticles into rutile-type ones at much
16 shorter reaction times.
17
18
19
20
21
22
23
24
25
26
27
28
29
30
31
32
33

34
35 For both ethers, nanoparticles with the expected rutile-type structure were obtained with similar
36 sizes of about 3 nm. Rather than the ability of the solvent to induce spatial hindrance, as we initially
37 supposed for dibenzyl ether, it is more the way it affects the crystallization pathway by
38 polymerizing, which controls the nanoparticle structure and size. Indeed, since the formation of the
39 intermediate oxide phase and its restructuration cannot be circumvented, such steps need to be
40 understood and be taken into account in order to produce small-size-controlled tin oxide
41 nanocrystals with the rutile-type structure. That is why the good knowledge of the properties and
42 the reactivity of all the reactants involved in the synthesis is necessary.
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60

Conclusion

In this study, the early stages of the synthesis of small tin oxide nanocrystals *via* the etherolysis of tin (IV) tetrachloride was investigated with diisopropyl or dibenzyl ether as oxygen donors. The results obtained from the study of the structural and morphological features obtained by XRD and TEM indicate a more complex crystallization pathway than previously reported in the literature. The SnO₂ nanocrystals synthesis begins, first, by the formation of a nanocrystalline inorganic intermediate phase embedded in a polymeric phase forming a large complex assembly. The structure of the intermediate phase is partially ordered and exhibits strong similarities with the rutile-type structure. In the case of diisopropyl ether, the degradation of the polymer matrix allows the nanoparticle restructuration into the rutile-type structure when the synthesis temperature and reaction time are sufficient. In the case of dibenzyl ether, the full polymerization of the reactional mixture is accompanied by the formation of water. The polymer, which is supposed to be a member of the polybenzylene family, is produced by the condensation of dibenzyl ether and/or benzyl chloride both catalysed by tin (IV) tetrachloride. The water release by the condensation of ether molecules seems to promote and activate the restructuration of the nanoparticles. In both cases, the occurrence of the intermediate phase cannot be circumvented and SnO₂ nanoparticles of 2-4 nm with rutile-type structure are finally obtained.

The formation of rutile-type SnO₂ nanoparticles of very small size goes through a fine control of the crystallization pathway. The formation of the polymeric phase shows that the properties of both ether-SnCl₄ couples and the formed nanocrystals has to be taken into account to control the final product. Moreover, the formation of water in a non-hydrolytic synthesis route indicates that the byproducts could also play a crucial role in the formation of the final oxide nanocrystals.

Author information

Corresponding author

*Email: olivier.masson@unilim.fr

ORCID: 0000-0003-0036-3859

Author contributions

All authors have given approval to the final version of the manuscript.

Notes

The authors declare no competing financial interest.

Acknowledgments

We acknowledge Pierre Carles for the TEM measurements and for the support in the analysis of the micrographs.

References

- [1] M. Batzill and U. Diebold, « The surface and materials science of tin oxide », *Prog. Surf. Sci.*, vol. 79, n° 2, p. 47-154, 2005.
- [2] Z. Zhang, X. Zou, L. Xu, L. Liao, W. Liu, J. Ho, X. Xiao, C. Jiang and J. Lia, « Hydrogen gas sensor based on metal oxide nanoparticles decorated graphene transistor », *Nanoscale*, vol. 7, n° 22, p. 10078-10084, 2015.
- [3] S. Das and V. Jayaraman, « SnO₂: A comprehensive review on structures and gas sensors », *Prog. Mater. Sci.*, vol. 66, p. 112-255, 2014.

1
2
3 [4] S. Wang, J. Huang, Y. Zhao, S. Wang, X. Wang, T. Zhang, S. Wu, S. Zhang and W. Huang,
4 « Preparation, characterization and catalytic behavior of SnO₂ supported Au catalysts for low-
5 temperature CO oxidation », *J. Mol. Catal. Chem.*, vol. 259, n° 1-2, p. 245-252, 2006.
6
7

8
9
10 [5] P. Rajaram, Y. C. Goswami, S. Rajagopalan and V. K. Gupta, « Optical and structural
11 properties of SnO₂ films grown by a low-cost CVD technique », *Mater. Lett.*, vol. 54, n° 2-3, p.
12 158-163, 2002.
13
14
15

16
17
18 [6] T. Sahm, L. Mädler, A. Gurlo, N. Barsan, S. E. Pratsinis and U. Weimar, « Flame spray
19 synthesis of tin dioxide nanoparticles for gas sensing », *Sens. Actuators B Chem.*, vol. 98, n° 2-3,
20 p. 148-153, 2004.
21
22
23

24
25
26 [7] S. M. Ingole, S. T. Navale, Y. H. Navale, D. K. Bandgar, F. J. Stadler, R. S. Mane, N. S.
27 Ramgir, S. K. Gupta, D. K. Aswal and V. B. Patil, « Nanostructured tin oxide films: Physical
28 synthesis, characterization, and gas sensing properties », *J. Colloid Interface Sci.*, vol. 493, p. 162-
29 170, 2017.
30
31
32
33

34
35
36 [8] M. K. Patel, J. Singh, M. K. Singh, V. V. Agrawal, S. G. Ansari and B. D. Malhotra, « Tin
37 oxide quantum dot based DNA sensor for pathogen detection », *J. Nanosci. Nanotechnol.*, vol. 13,
38 n° 3, p. 1671-1678, 2013.
39
40
41
42

43
44 [9] E. R. Leite, E. J. H. Lee, T. R. Giraldi, F. M. Pontes and E. Longo, « A simple and novel
45 method to synthesize doped and undoped SnO₂ nanocrystals at room temperature », *J. Nanosci.*
46 *Nanotechnol.*, vol. 4, n° 7, p. 774-778, 2004.
47
48
49
50
51
52
53
54
55
56

1
2
3 [10] N. M. Shaalan, D. Hamad, A. Y. Abdel-Latief and M. A. Abdel-Rahim, « Preparation of
4 quantum size of tin oxide: Structural and physical characterization », *Prog. Nat. Sci. Mater. Int.*,
5 vol. 26, n° 2, p. 145-151, 2016.
6
7

8
9
10 [11] M. Epifani, E. Pellicer, J. Arbiol and J. R. Morante, « Metal oxide nanocrystals from the
11 injection of metal oxide sols in a coordinating environment: Principles, applicability, and
12 investigation of the synthesis variables in the case study of CeO₂ and SnO₂ », *Chem. Mater.*, vol.
13 21, n° 5, p. 862-870, 2009.
14
15
16
17

18
19
20 [12] J. S. Chen and X. W. Lou, « SnO₂-based nanomaterials: Synthesis and application in
21 lithium-ion batteries », *Small*, vol. 9, n° 11, p. 1877-1893, 2013.
22
23
24

25
26 [13] H. Zhu, D. Yang, G. Yu, H. Zhang and K. Yao, « A simple hydrothermal route for
27 synthesizing SnO₂ quantum dots », *Nanotechnology*, vol. 17, n° 9, p. 2386-2389, 2006.
28
29

30
31 [14] S. Gnanam and V. Rajendran, « Influence of ethylene glycol on the nanostructured pure
32 and V-doped SnO₂ nanoparticles via sol-gel process and application in photocatalysts », *J.*
33 *Optoelectron. Adv. Mater.*, vol. 12, n° 11, p. 2199-2207, 2010.
34
35
36
37

38
39 [15] A. Aboulaich, B. Boury and P. H. Mutin, « Reactive and organosoluble SnO₂ nanoparticles
40 by a surfactant-free non-hydrolytic sol-gel route », *Eur. J. Inorg. Chem.*, n° 24, p. 3644-3649, 2011.
41
42
43

44 [16] J. Ba, J. Polleux, M. Antonietti and M. Niederberger, « Non-aqueous synthesis of tin oxide
45 nanocrystals and their assembly into ordered porous mesostructures », *Adv. Mater.*, vol. 17, n° 20,
46 p. 2509-2512, 2005.
47
48
49
50
51
52
53
54
55
56
57
58
59
60

1
2
3 [17] N. Pinna, G. Neri, M. Antonietti and M. Niederberger, « Nonaqueous synthesis of
4 nanocrystalline semiconducting metal oxides for gas sensing », *Angew. Chem. - Int. Ed.*, vol. 43,
5 n° 33, p. 4345-4349, 2004.
6
7

8
9
10 [18] M. Niederberger and N. Pinna, « Metal Oxide Nanoparticles in Organic Solvents:
11 Synthesis, Formation, Assembly and Application », *London: Springer-Verlag*, 2009.
12
13

14
15 [19] P. H. Mutin and A. Vioux, « Nonhydrolytic Processing of Oxide-Based Materials: Simple
16 Routes to Control Homogeneity, Morphology, and Nanostructure », *Chem. Mater.*, vol. 21, n° 4,
17 p. 582-596, 2009.
18
19
20

21
22 [20] A. Vioux, « Nonhydrolytic Sol–Gel Routes to Oxides », *Chem. Mater.*, vol. 9, n° 11, p.
23 2292-2299, 1997.
24
25

26
27 [21] M. Niederberger and H. Cölfen, « Oriented attachment and mesocrystals: Non-classical
28 crystallization mechanisms based on nanoparticle assembly », *Phys. Chem. Chem. Phys.*, vol. 8, n°
29 28, p. 3271-3287, 2006.
30
31
32

33
34 [22] J. F. Banfield, S. A. Welch, H. Zhang, T. T. Ebert and R. L. Penn, « Aggregation-based
35 crystal growth and microstructure development in natural iron oxyhydroxide biomineralization
36 products », *Science*, vol. 289, n° 5480, p. 751-754, 2000.
37
38
39

40
41 [23] S. Sun, D. Gebauer and H. Cölfen, « Alignment of Amorphous Iron Oxide Clusters: A Non-
42 Classical Mechanism for Magnetite Formation », *Angew. Chem. - Int. Ed.*, vol. 56, n° 14, p. 4042-
43 4046, 2017.
44
45

46
47 [24] B. Ludi and M. Niederberger, « Zinc oxide nanoparticles: Chemical mechanisms and
48 classical and non-classical crystallization », *Dalton Trans.*, vol. 42, n° 35, p. 12554-12568, 2013.
49
50

1
2
3 [25] D. Gebauer, A. Völkel and H. Cölfen, « Stable prenucleation calcium carbonate clusters »,
4
5 *Science*, vol. 322, n° 5909, p. 1819-1822, 2008.
6

7
8 [26] Z. Zhuang, J. Zhang, F. Huang, Y. Wang and Z. Lin, « Pure multistep oriented attachment
9
10 growth kinetics of surfactant-free SnO₂ nanocrystals », *Phys. Chem. Chem. Phys.*, vol. 11, n° 38,
11
12 p. 8516-8521, 2009.
13

14
15 [27] D. G. Stroppa, L. A. Montoro, A. Beltrán, T. G. Conti, R. O. da Silva, J. Andres, E. R. Leite
16
17 and A. J. Ramirez, « Anomalous oriented attachment growth behavior on SnO₂ nanocrystals »,
18
19 *Chem. Commun.*, vol. 47, n° 11, p. 3117-3119, 2011.
20
21

22
23 [28] R. L. Penn and J. F. Banfield, « Imperfect oriented attachment: Dislocation generation in
24
25 defect-free nanocrystals », *Science*, vol. 281, n° 5379, p. 969-971, 1998.
26
27

28
29 [29] E. J. H. Lee, C. Ribeiro, E. Longo and E. R. Leite, « Oriented attachment: An effective
30
31 mechanism in the formation of anisotropic nanocrystals », *J. Phys. Chem. B*, vol. 109, n° 44, p.
32
33 20842-20846, 2005.
34
35

36
37 [30] D. Zitoun, N. Pinna, N. Frolet and C. Belin, « Single crystal manganese oxide multipods by
38
39 oriented attachment », *J. Am. Chem. Soc.*, vol. 127, n° 43, p. 15034-15035, 2005.
40
41

42
43 [31] H. Cölfen and M. Antonietti, « Mesocrystals: Inorganic superstructures made by highly
44
45 parallel crystallization and controlled alignment », *Angew. Chem. - Int. Ed.*, vol. 44, n° 35, p. 5576-
46
47 5591, 2005.
48
49

50
51 [32] R.-Q. Song and H. Cölfen, « Mesocrystals - Ordered nanoparticle superstructures », *Adv.*
52
53 *Mater.*, vol. 22, n° 12, p. 1301-1330, 2010.
54
55

1
2
3 [33] J. Ba, A. Feldhoff, D. F. Rohlfing, M. Wark, M. Antonietti and M. Niederberger,
4 « Crystallization of indium tin oxide nanoparticles: From cooperative behavior to individuality »,
5
6 *Small*, vol. 3, n° 2, p. 310-317, 2007.
7

8
9
10 [34] H. M. Rietveld, « A profile refinement method for nuclear and magnetic structures », *J.*
11
12 *Appl. Crystallogr.*, vol. 2, n° 2, p. 65-71, 1969.
13

14
15 [35] R. A. Young, « The Rietveld Method », Oxford University Press, 1995.
16

17
18 [36] J. Rodríguez-Carvajal, « Recent advances in magnetic structure determination by neutron
19
20 powder diffraction », *Phys. B Condens. Matter*, vol. 192, n° 1, p. 55-69, oct. 1993.
21

22
23 [37] T. Roisnel and J. Rodríguez-Carvajal, « WinPLOTR: A windows tool for powder
24
25 diffraction pattern analysis », *Mater. Sci. Forum*, vol. 378-381, n° I, p. 118-123, 2001.
26

27
28 [38] O. Masson, « Peakoc ». <http://www.esrf.eu/computing/scientific/PEAKOC/MAIN.htm>.
29

30
31 [39] P. Thompson, D. E. Cox and J. B. Hastings, « Rietveld refinement of Debye–Scherrer
32
33 synchrotron X-ray data from Al₂O₃ », *J. Appl. Crystallogr.*, vol. 20, n° 2, p. 79-83, 1987.
34

35
36 [40] J. I. Langford, « A rapid method for analysing the breadths of diffraction and spectral lines
37
38 using the Voigt function », *J. Appl. Crystallogr.*, vol. 11, n° 1, p. 10-14, 1978.
39

40
41 [41] P. Scherrer, « Bestimmung der Größe und der inneren Struktur von Kolloidteilchen mittels
42
43 Röntgenstrahlen », *Nachrichten Von Ges. Wiss. Zu Gött. Math.-Phys. Kl.*, vol. 1918, p. 98-100,
44
45 1918.
46

47
48 [42] A. P. Hammersley, « FIT2D: a multi-purpose data reduction, analysis and visualization
49
50 program », *J. Appl. Crystallogr.*, vol. 49, n° 2, p. 646-652, 2016.
51

1
2
3 [43] T. Matsumoto, T. Nakano, Y. Yokoyama and T. Makiguchi, « Process for producing
4 isopropyl chloride », US20040215041A1, 2004.
5
6

7
8 [44] F. H. Gayer, « The Catalytic Polymerization of Propylene », *Ind. Eng. Chem.*, vol. 25, n°
9 10, p. 1122-1127, 1933.
10
11

12
13 [45] L. Shibryaeva, « Thermal Oxidation of Polypropylene and Modified Polypropylene -
14 Structure Effects ». INTECH Open Access Publisher, 2012.
15
16

17
18 [46] C. P. Tsonis, « Homogeneous catalytic polymerization of benzyl chloride leading to linear
19 high molecular weight polymers: an elusive goal », *J. Mol. Catal.*, vol. 57, n° 3, p. 313-323, 1990.
20
21

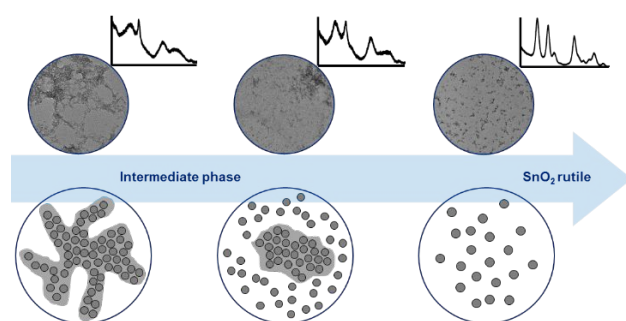
22
23 [47] T. M'Hiri, C. Catusse, R. Catusse and J. L. J. Dubry, « Polymerization of benzyl alcohol
24 and its derived compounds with an ion exchange resin », *React. Kinet. Catal. Lett.*, vol. 22, n° 3-4,
25 p. 425-428, 1983.
26
27

28
29 [48] R. A. Jacobson, « Polymers from benzyl chloride and related compounds », *J. Am. Chem.*
30 *Soc.*, vol. 54, n° 4, p. 1513-1518, 1932.
31
32

33
34 [49] O. C. Dermer and E. Hooper, « Catalysts for the Polymerization of Benzyl Chloride », *J.*
35 *Am. Chem. Soc.*, vol. 63, n° 12, p. 3525-3526, 1941.
36
37

38
39 [50] I. Olliges-Stadler, M. D. Rossell and M. Niederberger, « Co-operative formation of
40 monolithic tungsten oxide-polybenzylene hybrids via polymerization of benzyl alcohol and study
41 of the catalytic activity of the tungsten oxide nanoparticles », *Small Weinh. Bergstr. Ger.*, vol. 6,
42 n° 8, p. 960-966, 2010.
43
44
45
46
47
48
49
50
51
52
53
54
55
56

TOC graphic and synopsis



For table of content only

Synopsis. The crystallization pathways of tin dioxide nanoparticles synthesized by a nonaqueous sol-gel method based on the etherolysis of a tin (IV) tetrachloride precursor was investigated. Two ethers were used as oxygen donor, namely diisopropyl ether and dibenzyl ether. In both cases, the formation of ultra-small nanoparticles proceed via an original and complex pathway involving the progressive transformation of an intermediate phase as well as the formation of a polymeric phase.