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Published on: 17 Aug 2011 - Langmuir (American Chemical Society)

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O. Nikonova, Jean-Marie Nedelec, V.G. Kessler, G. Seisenbaeva. Precursor-Directed Assembly of Complex Oxide Nanobeads: The Role of Strongly Coordinated Inorganic Anions. *Langmuir*, American Chemical Society, 2011, 27 (18), pp.11622. 10.1021/la2028719 . hal-00673070

HAL Id: hal-00673070

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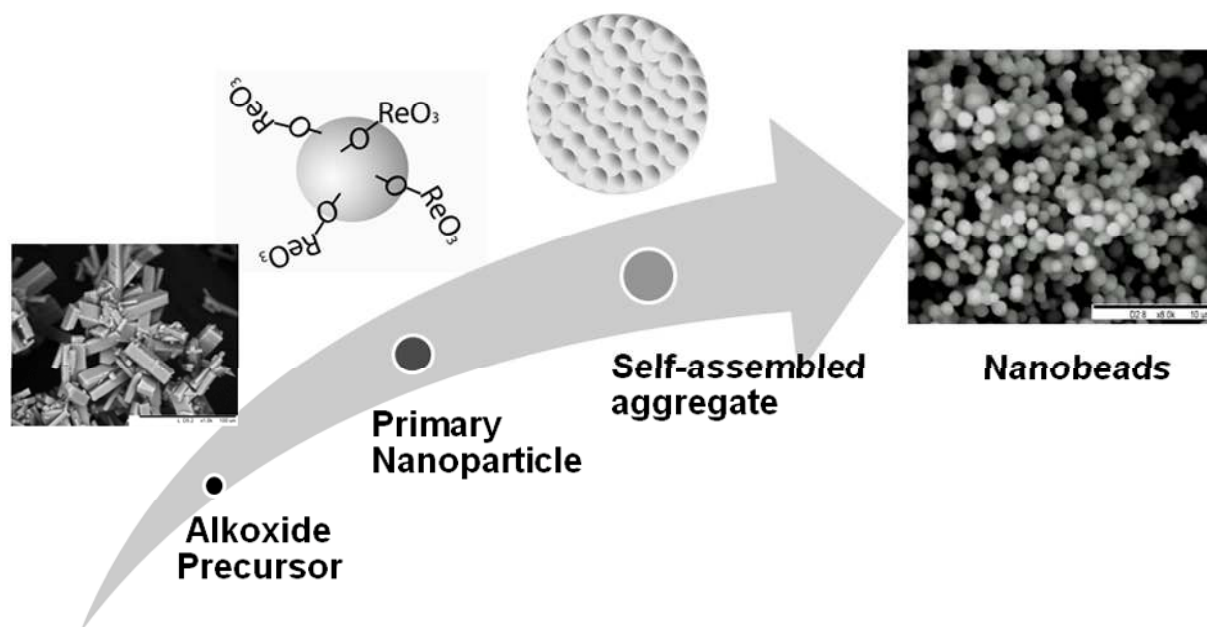
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Journal:	<i>Langmuir</i>
Manuscript ID:	la-2011-028719
Manuscript Type:	Article
Date Submitted by the Author:	25-Jul-2011
Complete List of Authors:	Nikonova, Olesya; SLU, Chemistry Nedelec, Jean-Marie; Clermont université, Chemistry Kessler, Vadim; SLU, Department of Chemistry Seisenbaeva, Gulaim; SLU, Chemistry

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Precursor-directed assembly of complex oxide nanobeads. **The role of strongly coordinated inorganic anions**

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ABSTRACT

Use of an inorganic perrhenate ligand in the structure of early transition metal alkoxide precursors permits to achieve uniform self-assembly of the primary nanoparticles produced by their hydrolysis. The latter has been carried out in hydrocarbon reaction medium by addition of water on vigorous stirring, either in the pure form or in solutions in parent alcohols. **The self-assembly is guided by the surface charge enhanced by the presence of strongly coordinated anions as determined by zeta-potential measurements. The aggregation process has been followed in real time by Nanoparticle Tracking Analysis (NanoSight technique).** The reaction products are spherical aggregates with the size that can be efficiently controlled through polarity of the reaction medium. The produced nanobeads have been characterized by TEM, SEM-EDS, **DLS, nitrogen adsorption** and FTIR. **The coordination of metal centers has been investigated using EXAFS-spectroscopy.** The aggregates remain amorphous on thermal treatment up to 700°C (24h treatment), but crystallize, when treated at 1000°C. This latter process is associated with the total loss of rhenium content and offers early transition metal oxides as products.

Introduction

Formation of 3D-aggregates of self-assembled nanoparticles has recently attracted much attention^{1,2,3,4,5} in the view of a broad spectrum of applications foreseen for these constructions, ranging from optical^{1,2} and magnetic³ lenses to high-stability electrode materials for lithium-ion batteries⁴ and dye-sensitized solar cells⁵. Among the approaches proposed to control the self-assembly major focus has been on either application of surfactants^{3,5}, including graft-copolymers⁶. Even in the absence of surfactants, the action of weak inter-particle forces, such as Van-der-Waals or electrostatic interactions have been proposed to explain uniform self-assembly in solution⁷, observed generally for metal oxide nanoparticles^{4,8,9,10}. The aggregation of metal oxide nanoparticles has been proved to be facilitated even by hydro- or solvothermal synthesis^{4,11}.

A special attention has been recently directed towards oxide nanobeads involving one or several early transition elements as perspective materials for Li-battery electrodes and for oxidation or acidic catalysis^{12,13}. The applied structure-directing tools have in this case been either application of sol-gel technology in complex solvent mixtures¹² or using external templates¹³. Formation of spherical particles on spontaneous hydrolysis of alkoxide precursors on liquid-air interface in contact with moist air has been reported long ago, but no means for control of particle size and aggregation have been developed yet¹⁴. The advantage of use of metal alkoxides as precursors lies in the formation via their hydrolysis of uniform oxometallate primary nanoparticles with the size below 5 nm – products of coordination equilibrium.^{10,15} A lot of effort has been put recently on the investigation of the role of different factors to control formation and aggregation of nanoparticles in aqueous media, envisaging, in particular, the role of coordinated anions.¹⁶ In the present work we have successfully exploited introduction of an inorganic perhenate anion in the structure of metal-organic precursors to achieve the uniform self-assembly of the initial particles. The size of the produced spherical aggregates could be efficiently controlled through the polarity of the medium (introduction of alcohols into hydrocarbon reaction medium).

Experimental

All the preparative procedures were carried out in dry nitrogen atmosphere using a dry box. Toluene was dehydrated by refluxing over LiAlH_4 with subsequent distillation. Methanol was purified by distillation over magnesium methoxide. Ethanol was dried by refluxing over Ba ethoxide, and n-propanol – by refluxing over aluminium n-propoxide with subsequent distillation.

The alkoxides, used in this work as starting materials for the synthesis of bi- and trimetallic complexes, $\text{Nb}_2(\text{OMe})_{10}$ and $\text{Ta}_2(\text{OMe})_{10}$, were prepared by anodic oxidation of the corresponding metals in methanol and purified according to conventional techniques¹⁹. The $\text{Ta}(\text{OCH}_2\text{CH}_3)_5$ 99+% (L10288) and $\text{Nb}(\text{OC}_3\text{H}_7)_5$ 99% (36571) have been purchased from Alfa Aesar. Preparation of the precursor complexes $(\text{Nb}_x\text{Ta}_{1-x})_4\text{O}_2(\text{OEt})_{14}(\text{ReO}_4)_2$ (I), $(\text{Nb}_x\text{Ta}_{1-x})_4\text{O}_2(\text{OMe})_{14}(\text{ReO}_4)_2$ (II), $\text{Nb}_4\text{O}_2(\text{O}^n\text{Pr})_{14}(\text{ReO}_4)_2$ (III) $\text{Nb}_4\text{O}_2(\text{OEt})_{14}(\text{ReO}_4)_2$ (IV), $\text{Nb}_4\text{O}_2(\text{OMe})_{14}(\text{ReO}_4)_2$ (V), $\text{Ta}_4\text{O}_2(\text{OEt})_{14}(\text{ReO}_4)_2$ (VI) and $\text{Ta}_4\text{O}_2(\text{OMe})_{14}(\text{ReO}_4)_2$ (VII), followed procedures described earlier for compounds II and IV in^{20,21} and are described in detail in the Supplementary.

IR spectra of tablets pressed with dry KBr were obtained with a Perkin Elmer FT-IR spectrometer Spectrum-100. Mass-spectra were recorded using JEOL JMS-SX/SX-102A mass-spectrometer applying electron beam ionization ($U = 70$ eV) with direct probe introduction. ^1H NMR spectra were collected for the precursors on Bruker DRX 400 spectrometer. The details of spectroscopic and crystallographic characterization of precursor compounds can be obtained from the Supplementary (Tables TS1-TS3).

SEM micrographs of the samples were obtained in scanning electron microscope HITACHI TM-1000 μDex , equipped with EDAX analysis. The RSD for all elements have been determined by repeating the EDS analysis 10 times from the same area on the highest magnification. TEM images were obtained with Philips TEM/STEM 400 instrument. Specific surface areas (SSA) were measured at 77 K on a Quantachrome Autosorb 1 apparatus. SSA was determined from at

1
2 least 5 points in the 0.05-0.3 P/P₀ range. Hydrodynamic size and ζ potential were measured using
3
4 the dynamic light scattering (DLS) technique on Malvern Instruments Zetasizer Nano-ZS
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6 instrument operating under a He-Ne 633 nm laser and equipped with the Dispersion Technology
7
8 Software for data collection and analysis. The samples were sonicated in 15 min before each
9
10 measurement. All measurements have been carried out in duplicate and revealed high
11
12 reproducibility.
13
14

15
16 EXAFS - *Data collection*. Tantalum and Rhenium K edge X-ray absorption spectra were
17
18 recorded at the wiggler beam line I811 at the MAXLab, Lund University, Lund; Sweden. The
19
20 EXAFS station was equipped with a Si [111] double crystal monochromator. The data collection
21
22 was performed in transmission mode. Higher order harmonics were reduced by detuning the
23
24 second monochromator to 30 % of maximum intensity at the end of the scans. The energy scale
25
26 of the X-ray absorption spectra were calibrated by assigning the first inflection point of the K
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28 edge of a tantalum foil to 9881 eV¹⁷. The EXAFSPAK program package¹⁸ was used for the data
29
30 treatment.
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34 *Preparation of nanobeads*

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36 Two types of hydrolysis: natural and forced have been applied to obtain spherical aggregates
37
38 denoted below as nanobeads. Certain masses of the precursor crystals were taken from the
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40 mother liquor and dissolved in different volumes of toluene, depending on the solubility of the
41
42 alkoxides. Natural hydrolysis was carried out by slow diffusion of the ambient atmosphere into a
43
44 flask with solution of alkoxide precursor connected with air through a syringe needle piercing
45
46 the serum cup. The forced hydrolysis was carried out by addition of liquid water in two different
47
48 protocols: (1) 0.01 ml of H₂O was quickly added by syringe on shaking to 0.5 ml of toluene
49
50 solution of the precursor, or (2) 0.05 ml of the respective alcohol (depending from precursor) and
51
52 0.01 ml of H₂O were added consequently by syringe on shaking to 0.5 ml of toluene solution of
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54 the precursor. Directly after addition of water a milky colloid solution was obtained with
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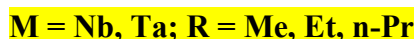
1 following formation of nanobeads in both cases. The particles were separated by centrifugation
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4 with subsequent removal of solvent by decantation and dried in vacuum.
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6 Starting solutions for preparation of nanobead samples were obtained by dissolution of 1.017 g
7
8 of (I) in 7 ml toluene, for the synthesis of **NBI**, **NBI*** and **NBI**** respectively (the first sample
9
10 originating from natural hydrolysis, the one marked with one star – from **forced** hydrolysis with
11
12 pure water and the one with two stars – from **forced** hydrolysis by water in parent alcohol); 0.121
13
14 g of (III) in 3 ml toluene were used for **NBIII**, **NBIII*** and **NBIII****; 0.294 g of (IV) in 4.8 ml
15
16 toluene were used for **NBIV**, **NBIV*** and **NBIV****; and 0.719 g of (VI) in 2 ml toluene were
17
18 used for the synthesis of **NBVI** and **NBVI***. Poor solubility of precursors (II) and (V) hinder
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20 preparation of the corresponding nanobead materials from them.
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28 **Results and discussion**

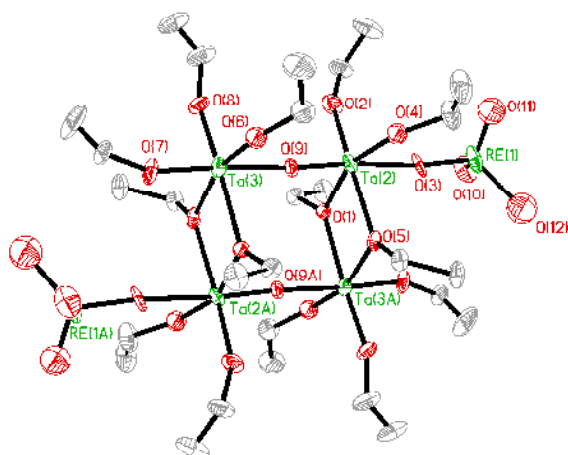
29 **Precursor compounds**

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35 Introduction of perrhenate anions into the structures of alkoxide precursors was achieved by a
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37 simple procedure involving refluxing of the commercially available metal alkoxides with
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39 rhenium(VII) oxide, Re_2O_7 , taken in stoichiometric amounts. The reaction follows practically
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41 quantitatively the general scheme:
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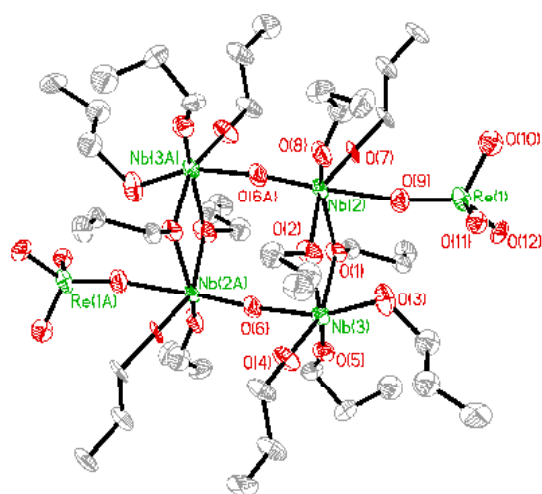


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54 The structures of all synthesized complexes (see Fig. 1) are following the same structure type
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56 based on a tetranuclear $\text{M}_4(\mu\text{-O})_2(\mu\text{-OR})_4\text{X}_{12}$ core, $\text{X} = \text{OR, ReO}_4$, with rectangular planar M_4 -
57
58 arrangement, composed of two pairs of edge-sharing octahedra, joined via two almost linear oxo-
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60 bridges. The perrhenate ligands are attached in a centrosymmetric manner to the two opposite

corners of the rectangle and are thus incorporated in the coordination spheres of the transition metal (Nb, Ta) atoms of the alkoxide precursor.



a



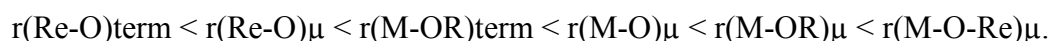
b

Fig. 1 Ortep view of the molecular structures of a - **(I)** and b - **(III)**, representative for the precursor complexes.

It should be mentioned that the structures of trimetallic complexes **(I)** and **(III)** have non uniform distribution of niobium and tantalum between the two symmetrically independent positions. As it was described previously²⁰ for the methoxide complex **(II)** the niobium atoms (70 %) preferably

1
2 are situated in the oxo-position which is connected with perrhenate group in contrast to tantalum
3
4 ones (30%) which are more abundant in the alkoxo-position where 70% of tantalum atoms and
5
6 30% of niobium ones are present. The same tendency can be seen in the new ethoxocomplex (**I**),
7
8 but the distribution of niobium and tantalum atoms is even more preferential: in oxo-position **M1**
9
10 is situated 85% of niobium and 15 % of tantalum atoms while in alkoxo-position **M2** 35% of Nb
11
12 and 65 % of Ta are present. According to the structure refinement statistics the ratio between Nb
13
14 and Ta in the $(\text{Nb}_x\text{Ta}_{1-x})_4\text{O}_2(\text{OEt})_{14}(\text{ReO}_4)_2$ complex is 60:40, while for the trimetallic
15
16 methoxide complex (**II**) this ratio is 50:50. To confirm the ratios of M:Re (M = Nb, Nb+Ta) and
17
18 Nb:Ta in the bulk samples the EDS analysis has been performed (See table TS2).
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20

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22 The increasing size of the alkoxide ligand influences also the M – O bond energy, which is
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24 confirmed by analysis of the IR spectra (see Table TS3). They show that the peaks shift toward
25
26 smaller wavenumbers, which indicated that the interaction energy in case of trimetallic (**I**) and
27
28 tantalum-rhenium bimetallic ethoxides (**VI**) is lower than in methoxo complexes (**II**) and (**VII**)
29
30 respectively. In case of niobium-rhenium bimetallic complexes the same trend can be observed
31
32 for the difference between the normal propoxide and ethoxide and methoxide respectively. In
33
34 agreement with the earlier observations^{21a,21b}, the bond lengths increase in the series:



44 45 **Nanobeads**

46
47 Following the hypothesis that application of perrhenate **anions** will improve stability of primary
48
49 particles, originating from hydrolysis of metal alkoxides, and permit their more uniform
50
51 aggregation, the solutions of perrhenate alkoxides were first subjected to slowly diffusing
52
53 moisture via the procedure of natural hydrolysis.
54
55

56
57 For all the tested precursors it produced mostly spherical beads with different degree of
58
59 aggregation. Relatively small fraction of bigger poorly shaped aggregates was also present. The
60

1 size of the spheres was mostly uniform and showed no trend in relation to either nature of the
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8 The sample **NBI** reveal almost the same size of well-shaped spheres approximately 500–700 nm
9
10 in average (see Fig 2a and 2b), mostly not aggregated. In case of spherical particles from the
11
12 bimetallic precursors we can see that Niobium-containing ones distinctly are not uniform and
13
14 have different size starting from 1 μm and up to more than 40 μm . The Tantalum-containing
15
16 sample **NBVI** shows well-formed, practically uniform particles 500 nm to 2 μm in size, even
17
18 when the solution concentration is quite high. It should be noticed that all nanobeads have dense
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20 oxide surface based on Nb or/and Ta metal oxides, resulting in very low active surface as
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22 measured by nitrogen sorption.
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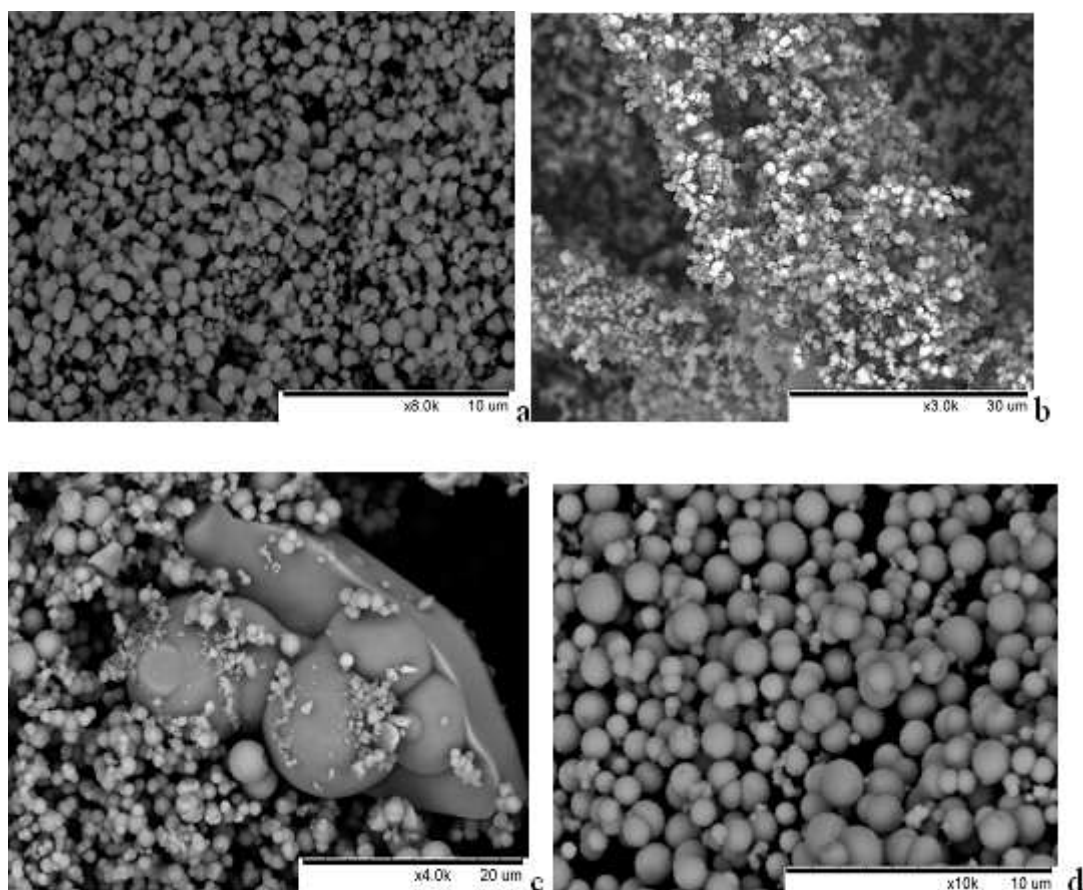


Fig. 2 SEM images of nanobeads obtained by natural hydrolysis: a – **NBI**, b – **NBIII**, c – **NBIV**,
d – **NBVI**

The perrhenate ligand was quantitatively retained through the hydrolysis procedure according to the EDS analysis (see Tab. 1), which indicated the absence of hydrolysis for the M-O-Re bond and potential for their use as structure-directing units.

Table 1 Results of the EDS analysis for produced nanobeads

Nanobeads	Atomic %			Ratio Nb:Ta	Ratio M:Re M=Nb or/and Ta
	Nb	Ta	Re		
NBI	46.42	23.67	29.91	0.66:0.34	2.3:1
NBI*	42.49	27.80	29.54	0.60:0.40	2.4:1
NBI**	47.85	23.20	28.95	0.67:0.33	2.5:1
NBIII	69.52		30.48		2.3:1
NBIII*	67.55		32.45		2.1:1
NBIII**	65.59		34.41		1.9:1
NBIV	70.19		29.80		2.4:1
NBIV*	71.03		28.97		2.5:1
NBIV**	70.62		29.37		2.4:1
NBVI		61.99	38.01		1.6:1
NBVI*		63.21	36.79		1.7:1
NBVI**		63.93	36.07		1.8:1

Retention of perrhenate ligands is clearly observed from the FTIR spectra, where the M-O-Re frequencies are situated at practically same wavelength at 700 – 800 cm^{-1} as in the initial precursors and have quite strong intensity, while the frequencies, corresponding to the organic ligands, strongly decrease (see Tab. 2). In the IR spectra the characteristic bands at 400 – 650 cm^{-1} could be assigned to Nb(Ta)-O-Nb(Ta) vibrations, corresponding to the bridging bonds

1
2 analogous to those in the spectra of precursor complexes (see table TS 2). The characteristic set
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4
5 Re=O bands indicate that Re is present in the beads as ReO_4^- -ligands.

6
7 The coordination of the metal centers does not change during the hydrolysis as indicated by the
8
9 EXAFS spectra (Fig. S7-S10), remaining octahedral for Ta(Nb) and tetrahedral (conservation of
10
11 the perrhenate ligand structure) for Re.
12

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16 Table 2 Comparison of the FTIR spectra* of the nanobeads obtained by natural hydrolysis

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18 M = Nb or/and Ta
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	NBI	NBIII	NBIV	NBVI
v(M–O)			410 w	
v(M–O–M)			422 w	
		495 w br	465 w br	
	516 sh		502 m br	513 w br
		587 s	570 s br	
	638 s			640 s br
v(M–O– Re)		719 w	730 w br	
	813 m br	820 m br	802 w br	813 m br
			839 w br	
	909 w	909 s	911 w	909 w
v(ReO_3)	931 sh	930 m br	931 w br	
belongs to ReO_4^-	939 s br	946 sh	947 w	942 s br

* s – strong, w – weak, br – broad, sh – shoulder

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7 The forced hydrolysis is resulting in the formation of spherical particles with size and degree of
8 aggregation strongly dependent on the polarity of the solvent (see Fig. 3). The kinetics of
9 aggregation of the initially formed primary particles was followed by laser reflection microscopy
10 using NanoSight (NTA Version 2.1) instrument (see real-time video VS1 in the Supplementary).
11
12 The initial particles with the size below 10 nm aggregated uniformly to spherical particles of
13 several hundred nm in size within minutes. The aggregates can be split into initial small particles
14 again by sonication in a standard ultrasound bath in 5 min and the re-aggregation can then be
15 followed to occur reproducibly with formation of again same type spherical aggregates in the
16 same time scale. This is an important evidence for the formation of the observed oxide particles
17 through so called Micelle Templated by Self-Assembly of Ligands (MTSAL) mechanism^{10,15}.
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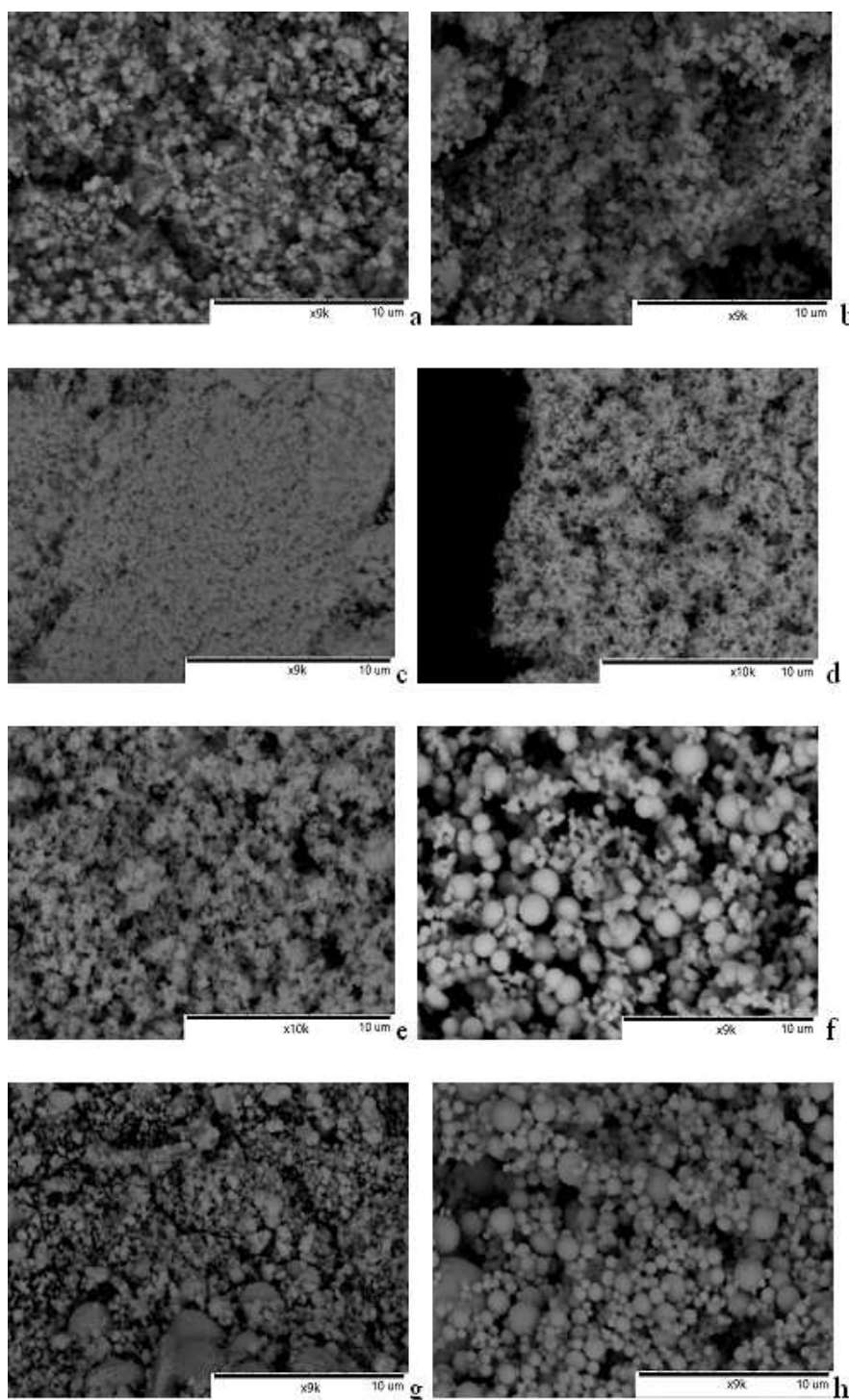


Fig. 3 SEM images of nanobeads obtained by forced hydrolysis: a – NBI*, b – NBII**, c – NBIII*, d – NBIII**, e – NBIV*, f – NBIV**, g – NBVI*, h – NBVI**

As it can be seen from the figures 2b, 2d, 2f and 2h the addition of the parent alcohol to the system leads to formation of better formed nanobeads, while the hydrolysis only with water results in the particles with the smaller size and less developed shape. The bigger aggregates are

absent, because the coalescence of the primary particle occurs more uniformly as the distribution of water in the volume is more uniform due to agitation. Even in this case the retention of Re is complete and occurs in the form of ReO_4 -ligands according to EDS and FTIR.

In spite of well-defined geometry of the metal centers, the produced nanobeads remain amorphous on thermal treatment up to 700 °C even after 24h. Increasing the temperature to 1000°C leads to complete loss of Re (in the form of volatile Re_2O_7) with formation of crystalline nanobeads based on only Nb or/and Ta oxides (see Fig. 4). The surface area increases on this treatment considerably from less than 1 m^2/g to about 1.2 – 4 m^2/g according to the nitrogen sorption data (see FS9-FS10). The low value of the surface area observed indicate that the pores produced by removal of perrhenate anions in the form of volatile anhydride, Re_2O_7 , are mostly cured at this high temperature, increasing only the surface roughness as observed in SEM (Fig. 4).

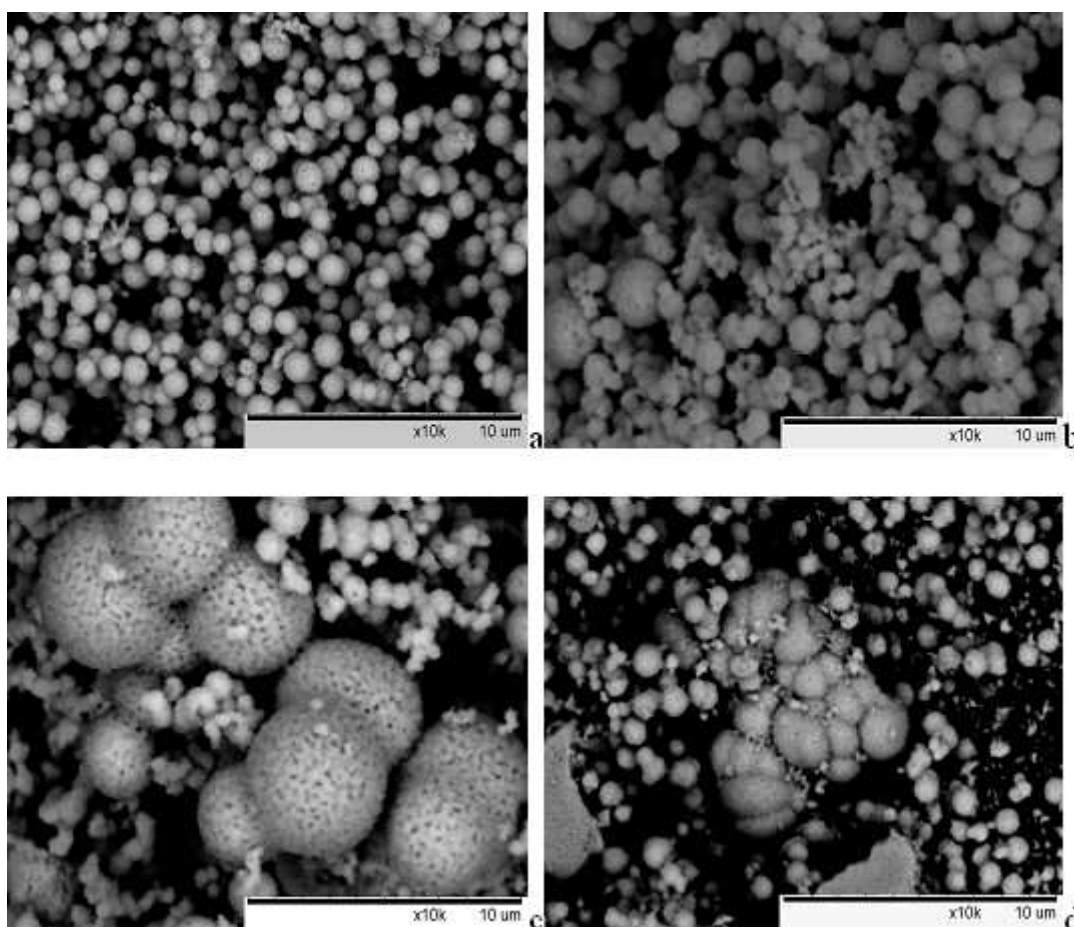
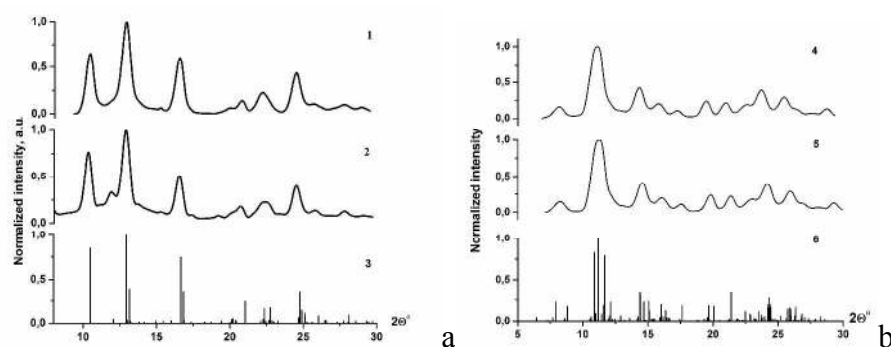


Fig. 4 SEM images of heat-treated nanobeads a – NBI, b – NBIII, c – NBIV, d – NBVI

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4 The non-heat-treated beads are thus possessing fully dense surface (completely closed porosity)
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6 formed via coalescence of the original nanoparticles as it is often observed for the niobium and
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8 tantalum oxides produced by sol-gel.²³ The products of thermal treatment were investigated by
9
10 X-ray powder diffraction analysis (see Fig. 5). The XPD results have shown that **NBIII** and
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12 **NBIV** based on the monoclinic phase of α - Nb_2O_5 oxide, for the samples of **NBI** and **NBVI** are
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14 resulting in formation of the orthorhombic modification of γ - Ta_2O_5 oxide.
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31 Fig. 5 XPD of the samples heat-treated at 1000°C a: 1 – **NBI**, 2 – **NBVI**, 3 – γ - Ta_2O_5 ICDD-
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33 JCPDS 25-0922; b: 4 – **NBIII**, 5 – **NBIV**, 6 – α - Nb_2O_5 ICDD- JCPDS 37-1468
34
35

36 Even if no Rietveld refinement of XPD patterns is performed we could add that the broad
37
38 diffraction peaks observed are in good agreement with nanocrystalline domains.
39

40 The EDS analysis indicated that the ratio between Nb and Ta in thermal product of **NBI** remains
41
42 the same as in the initial sample NBI 65.98 and 34.02 at% respectively.
43
44

45 In order to understand the mechanism guiding the self-assembly process, we have carried out
46
47 DLS and ζ potential measurements on both the samples produced by forced hydrolysis by water
48
49 and the products of their thermal treatment at 1000°C. In all the cases the samples were dried in
50
51 vacuum and re-dispersed in water by sonication. Because of rapid re-agglomeration of particles
52
53 only aggregates size could be measured by DLS but all samples have been strictly measured in the
54
55 same conditions (sonification conditions, time and delay after sonification). The results are
56
57 summarized in Tab. 3
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59
60

Table 3. ζ potential and size of aggregates as observed by DLS measurements. The T-marking refers to the samples heat-treated at 1000°C.

Sample	ζ (mv)	Peak 1 r(nm)	Peak 2 r(nm)	Integral intensity of the main peak
NBVI*	34,6	548,1	0	0,594
NBVI*T	-2,78	376,8	87,11	0,379
NBIII*	-9,31	364,8	66,6	0,569
NBIII*T	-23	309,1	2619	0,328
NBIV*	15,8	392,6	2746	0,318
NBIV*T	-27,2	333,6	2515	0,285
NBI*	16,8	335,9	2780	0,600
NBI*T	-29,4	296,9	2633	0,388

The observed sizes of the aggregates are in good agreement with the SEM data. The ζ potentials reveal a clear and consistent insight in the role of the introduced anion as a source for stabilization of surface charge, related also in turn to the acidity of the transition metal oxide. Generally, the presence of an anion in the inner coordination sphere of the atoms on the surface facilitates their ability to bear a positive charge. Ta₂O₅ is much less acidic (more basic) compound compared to Nb₂O₅, which explains much higher positive potential for NBVI* derived from Ta₄O₂(OEt)₁₄(ReO₄)₂ (VI) compared to all other samples. When the anions are removed by thermal treatment (in NBVI*T) the surface charge practically disappears and the aggregation and de-aggregation will only be due to hydrogen bonding on the surface. For the analogous ethoxide-derived samples NBIV* and NBI* (produced from Nb₄O₂(OEt)₁₄(ReO₄)₂ (IV) and (Nb_xTa_{1-x})₄O₂(OEt)₁₄(ReO₄)₂ (I) respectively) the positive charge is lower due, apparently to higher acidity (lower basicity) of niobium oxide. When the ligands are removed in NBIV*T and NBI*T, the particles gain distinct negative charge due to the same reason. The same trend is even visible for NBIII*, obtained from an n-propoxide complex, but the observed potential here is negative already for the non-heat-treated sample. The explanation for this deviation can be sought in the relative hydrophobicity of the residual n-propoxide ligands on the surface of the particles, influencing formation of the electric double layer. The thermal treatment

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2 removes all the surface ligands and offers distinctly negative charged particles as it should be
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4 observed for the appreciably acidic niobium oxide.
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8 9 **Conclusions**

10 Application of perrhenate as structure-directing ligand permits successful and reproducible
11 preparation of spherical particles conserving all of its Re content from small primary particles
12 originating on hydrolysis of the alkoxide precursor. The self-assembly process is guided by a
13 relative positive charge of the particles stabilized via inclusion of negatively charged perrhenate
14 ligands into the coordination sphere of transition metal atoms on the surface. The aggregation
15 can be reversed in solution by sonication, but the primary particles assemble again into spherical
16 beads. The particles remain amorphous and retain Re until rather high temperatures (700°C),
17 most probably due to formation of dense oxide layer on their surface. The Re content is lost at
18 1000°C with resulting increase in surface roughness as observed by SEM.
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35 **Supplementary materials available:** Details of synthesis, spectroscopic and crystallographic
36 characterization for molecular precursors applied, results of FTIR, EXAFS, TEM, nitrogen
37 adsorption studies, a real-time movie demonstrating the aggregation process, and statistical
38 description of the NanoSight measurements on the aggregation process for the produced
39 nanobeads. This material is available free of charge via the Internet at <http://pubs.acs.org>.
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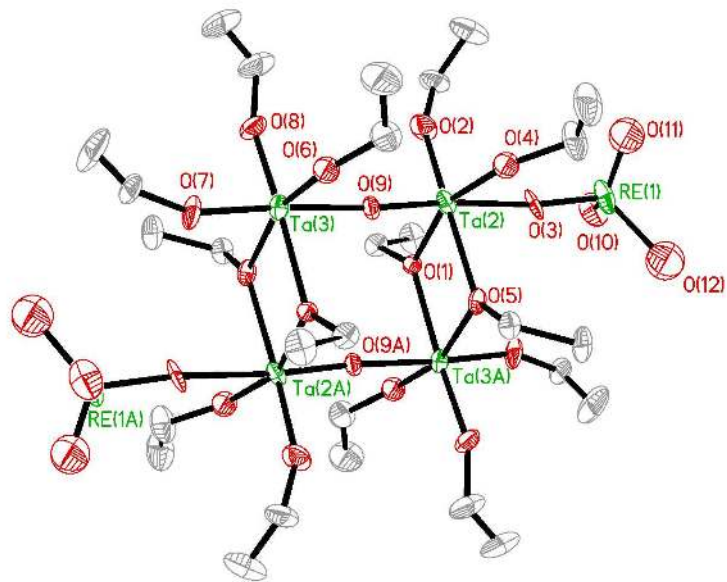
49 **Acknowledgements**

50 Authors would like express their gratitude to Corine Sandström for performing the NMR
51 experiment, to Suresh Gohil from Department of Chemistry, SLU, to Jonathan Smith from
52 NanoSight Ltd for assistance with size distribution data collection on NBI, NBVI and to MAX-
53 lab for the giving time to perform the EXAFS data collection from Lund.
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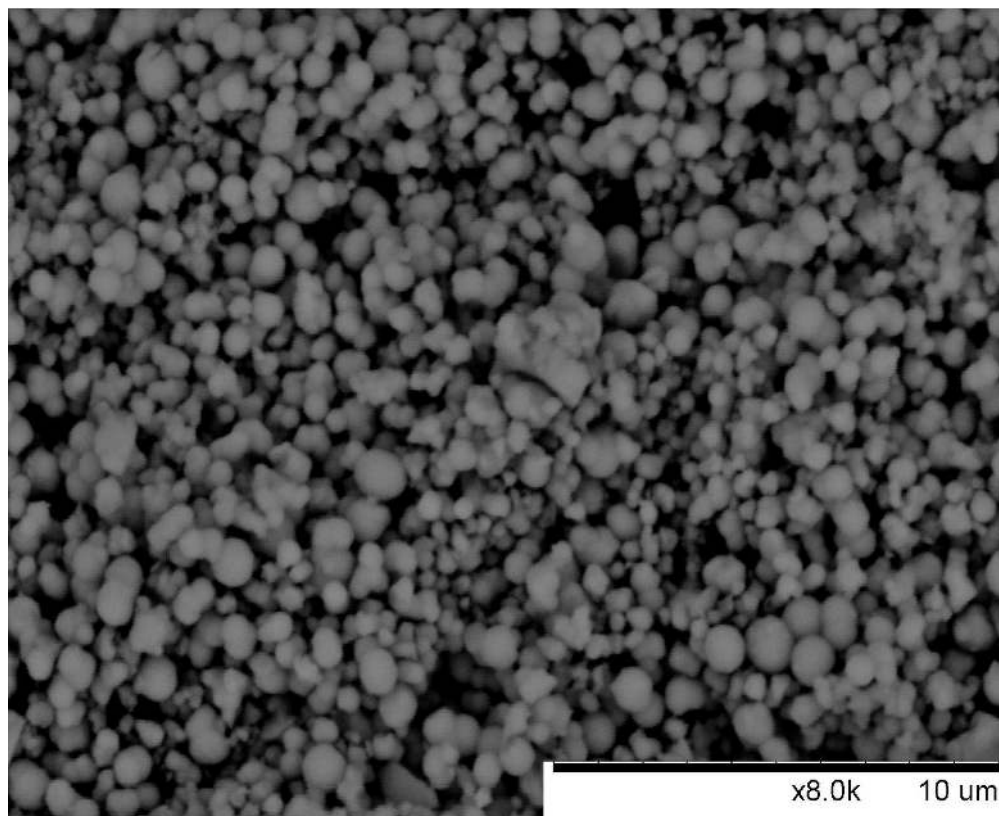
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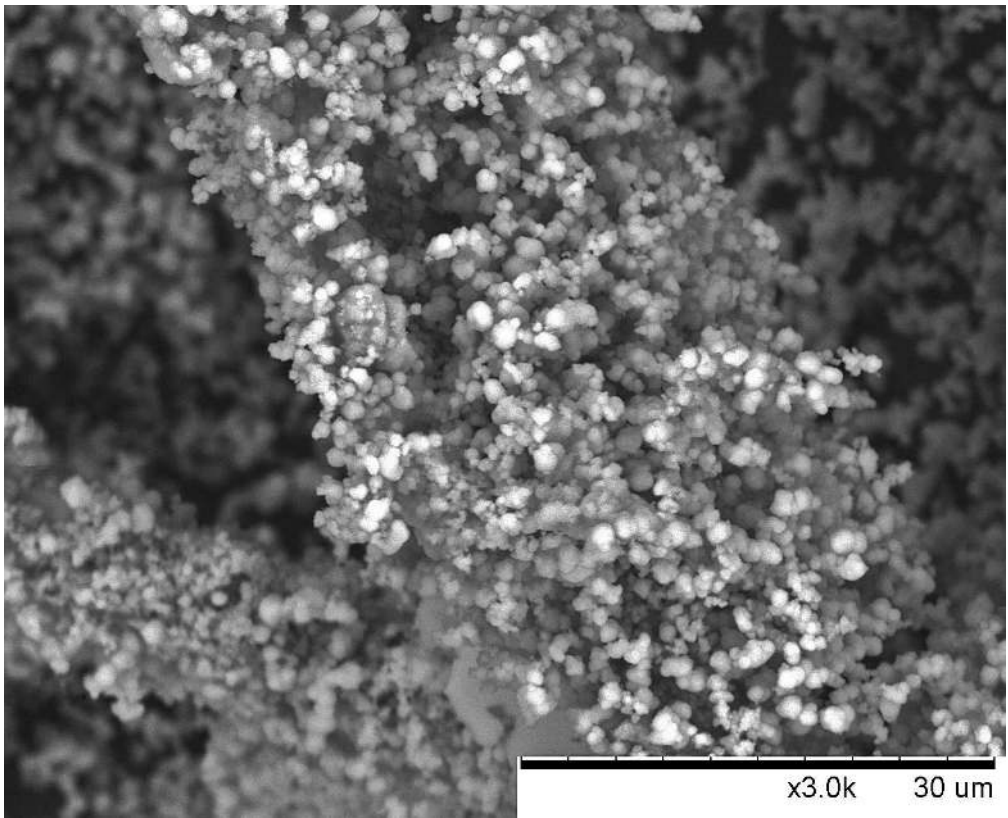
Ortep view...
279x215mm (300 x 300 DPI)



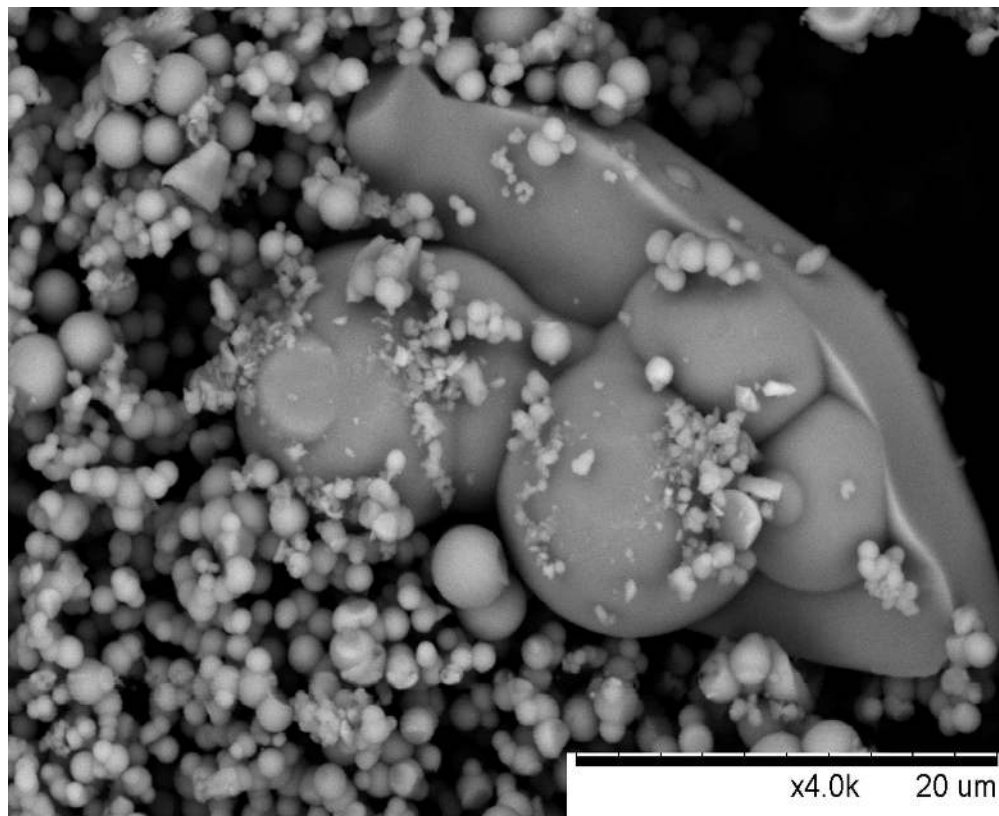
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Obtained by natural hydrolysis from NBI
179x145mm (300 x 300 DPI)

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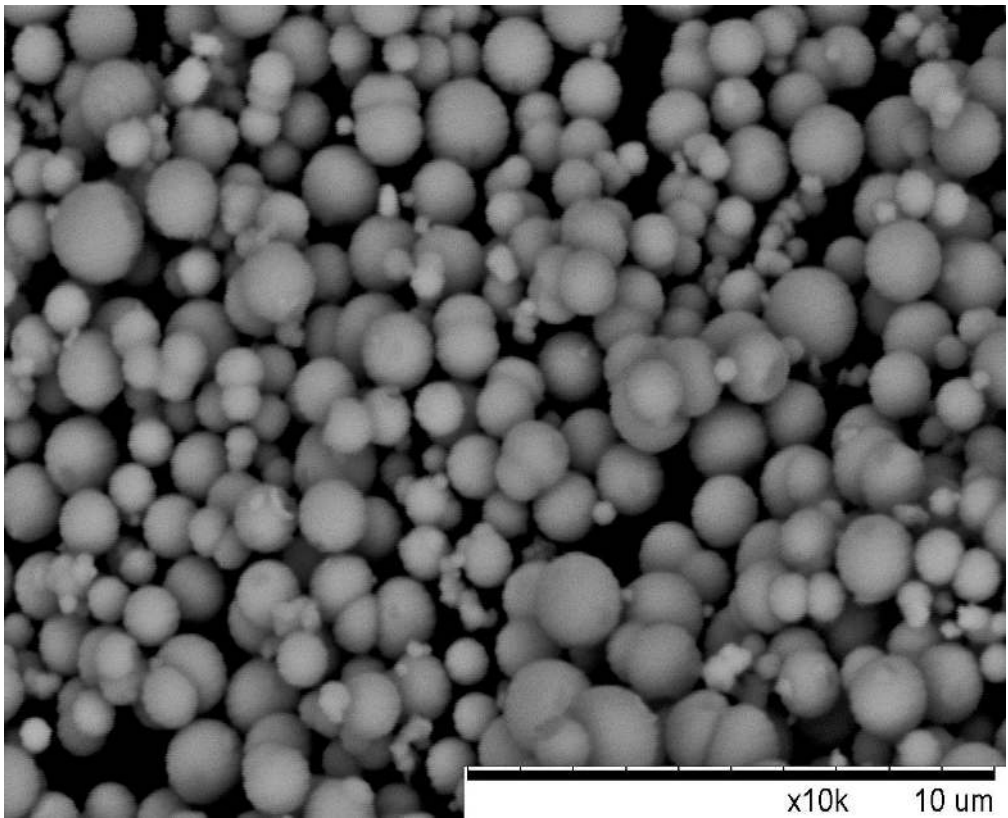
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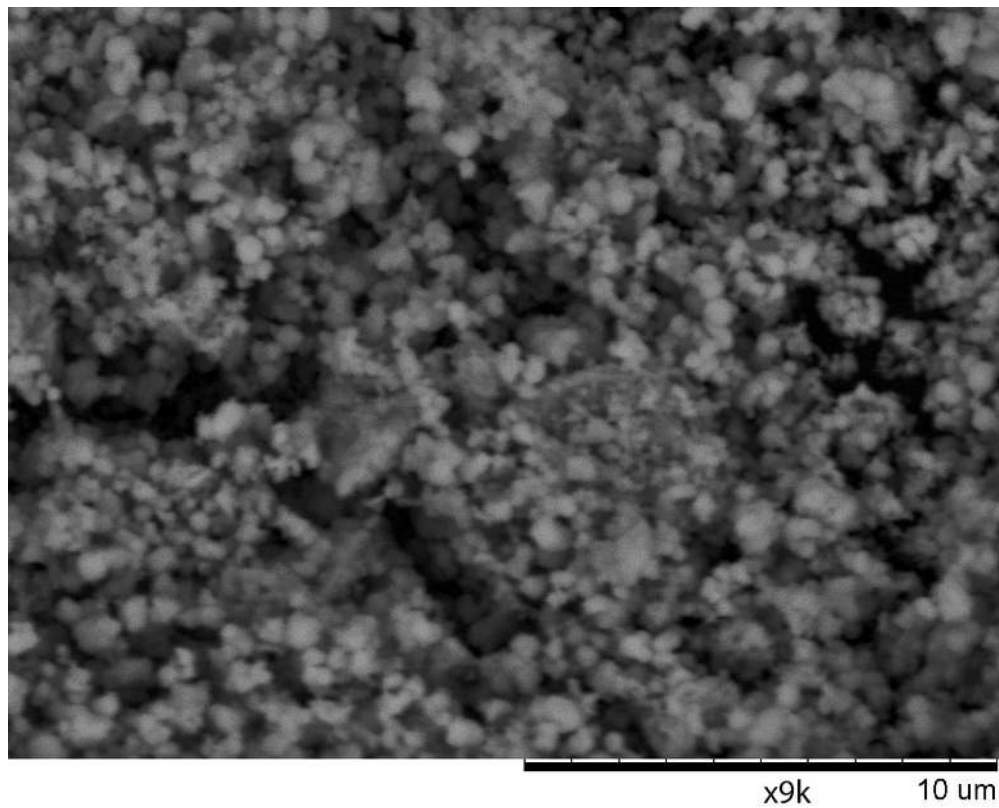
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Obtained by natural hydrolysis from NBIV
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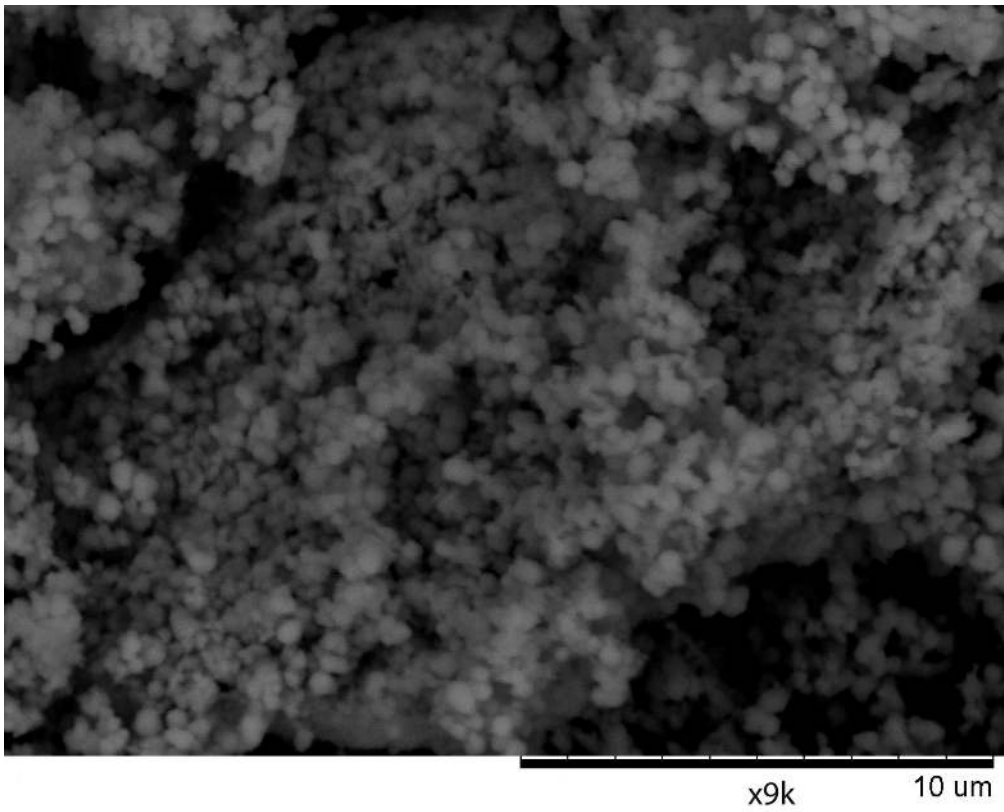


Obtained by natural hydrolysis of NBVI
190x154mm (171 x 171 DPI)

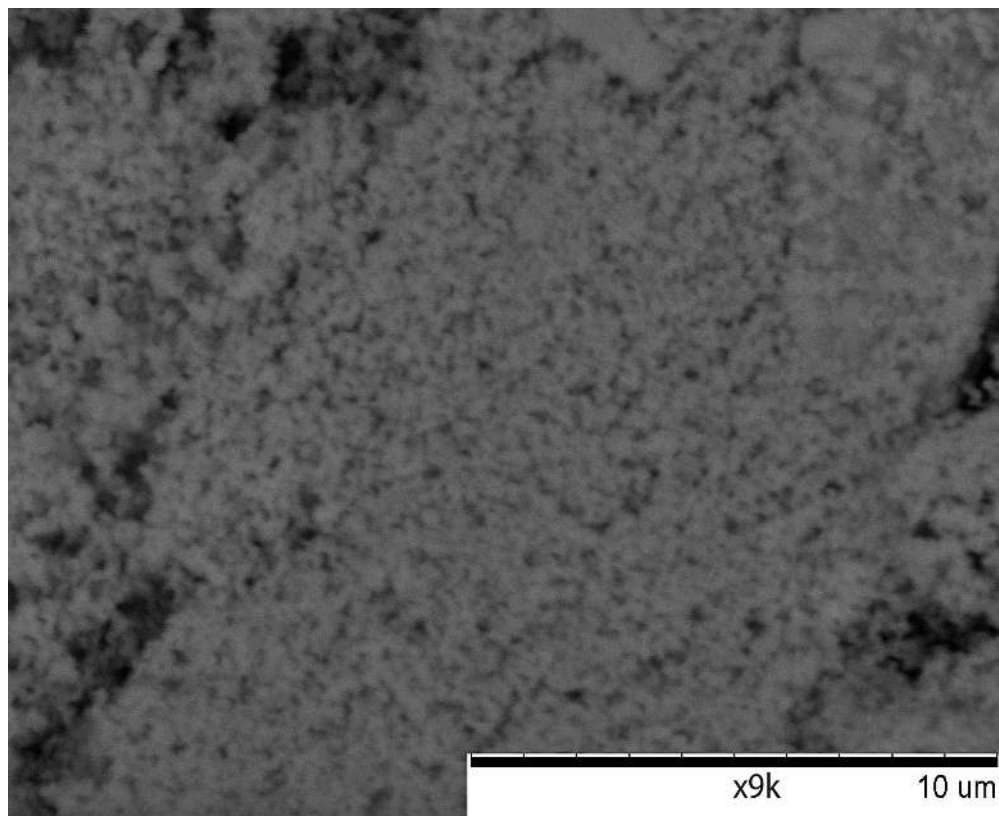


Obtained by forced hydrolysis of NBI*
190x154mm (300 x 300 DPI)

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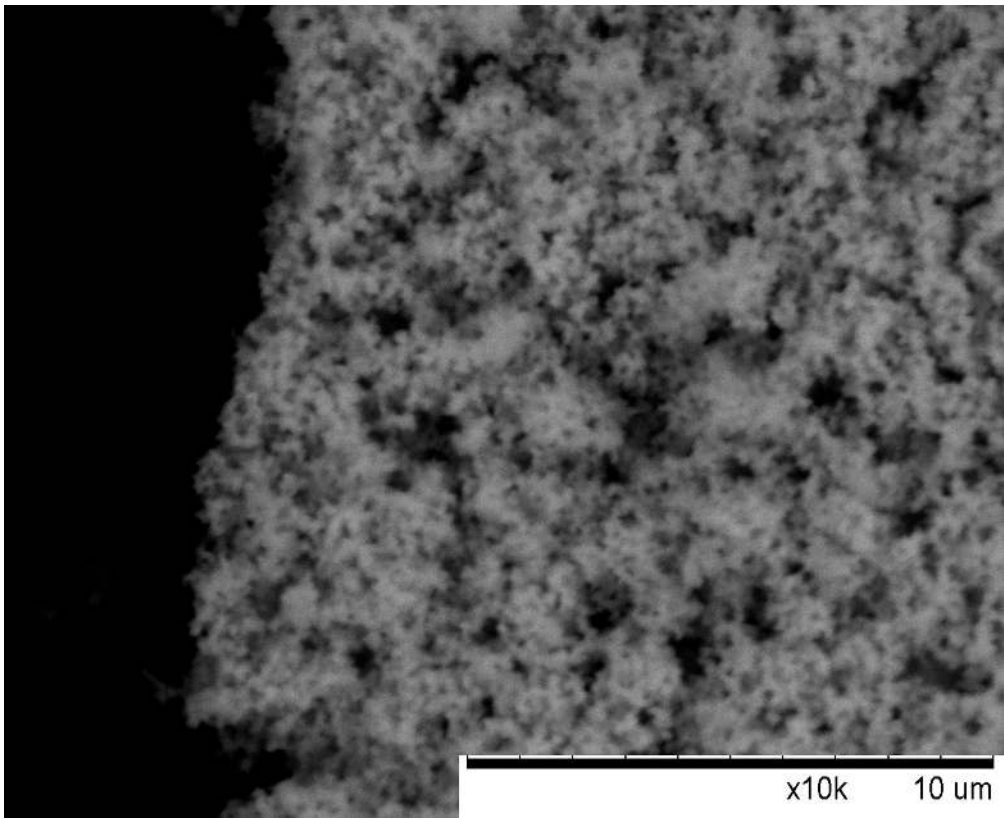
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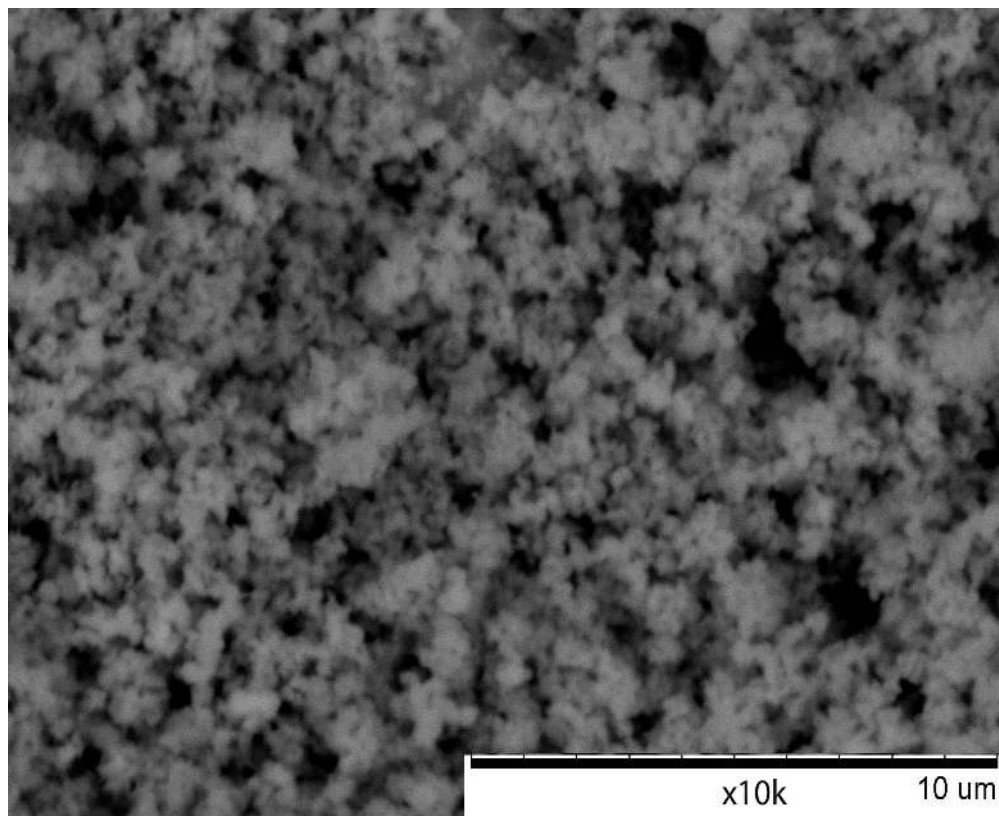
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Obtained by forced hydrolysis of NBIII*
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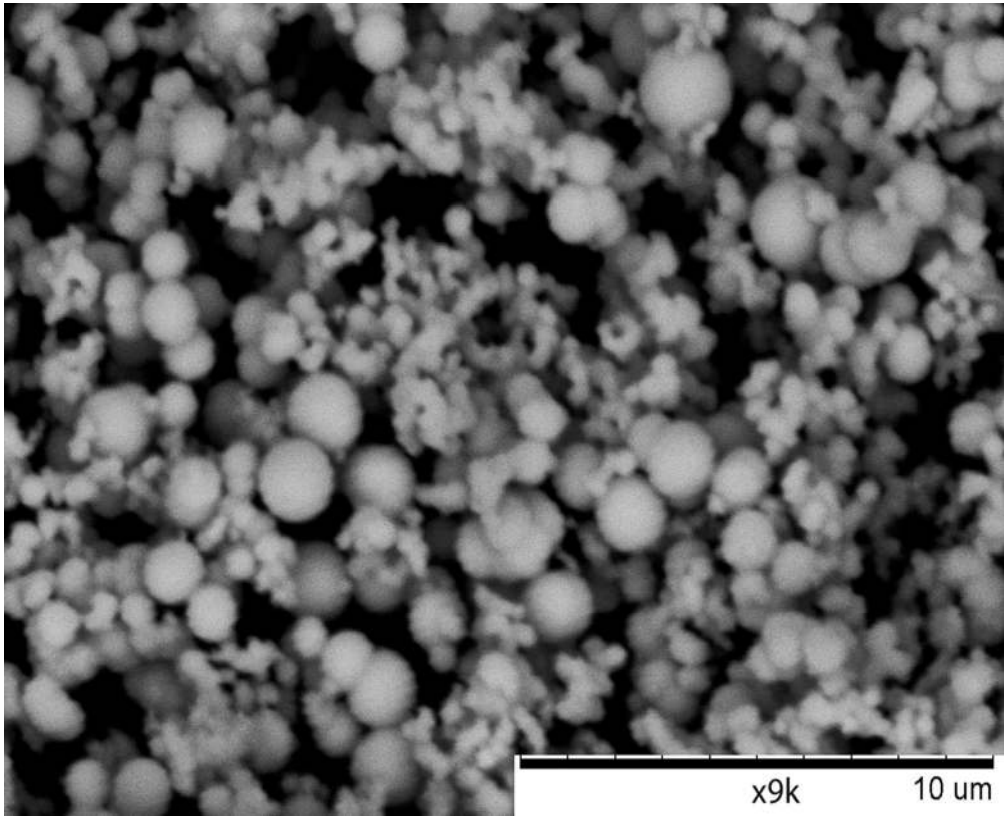
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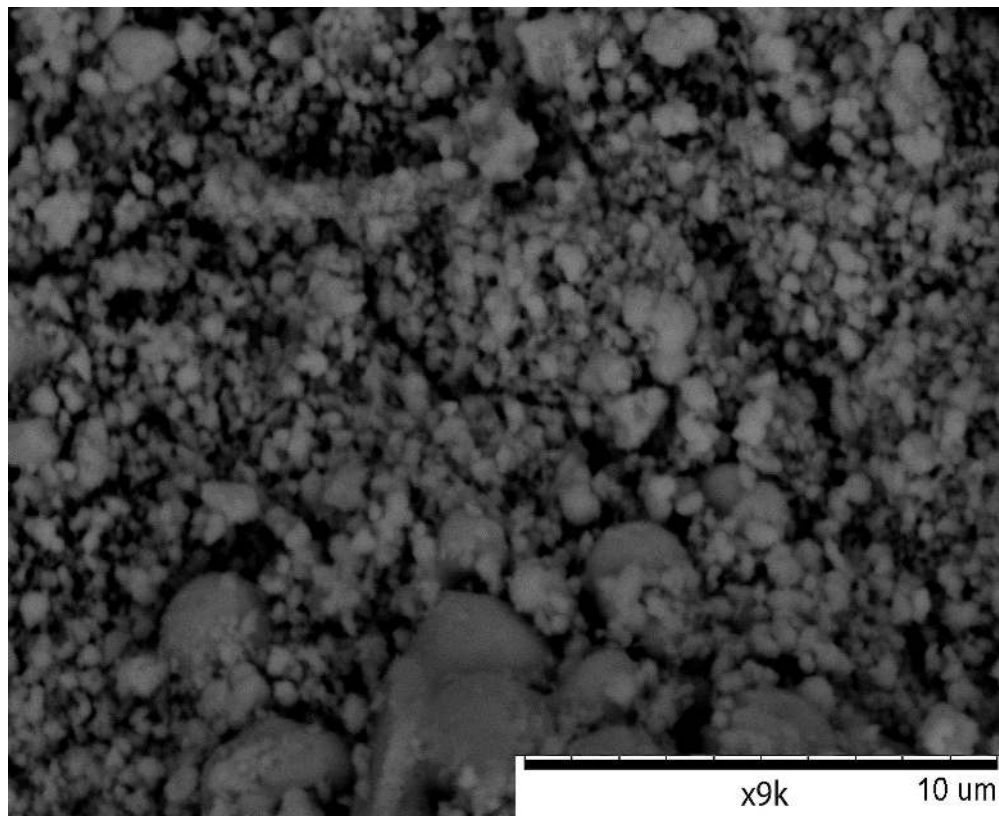
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Obtained by forced hydrolysis of NBIV*
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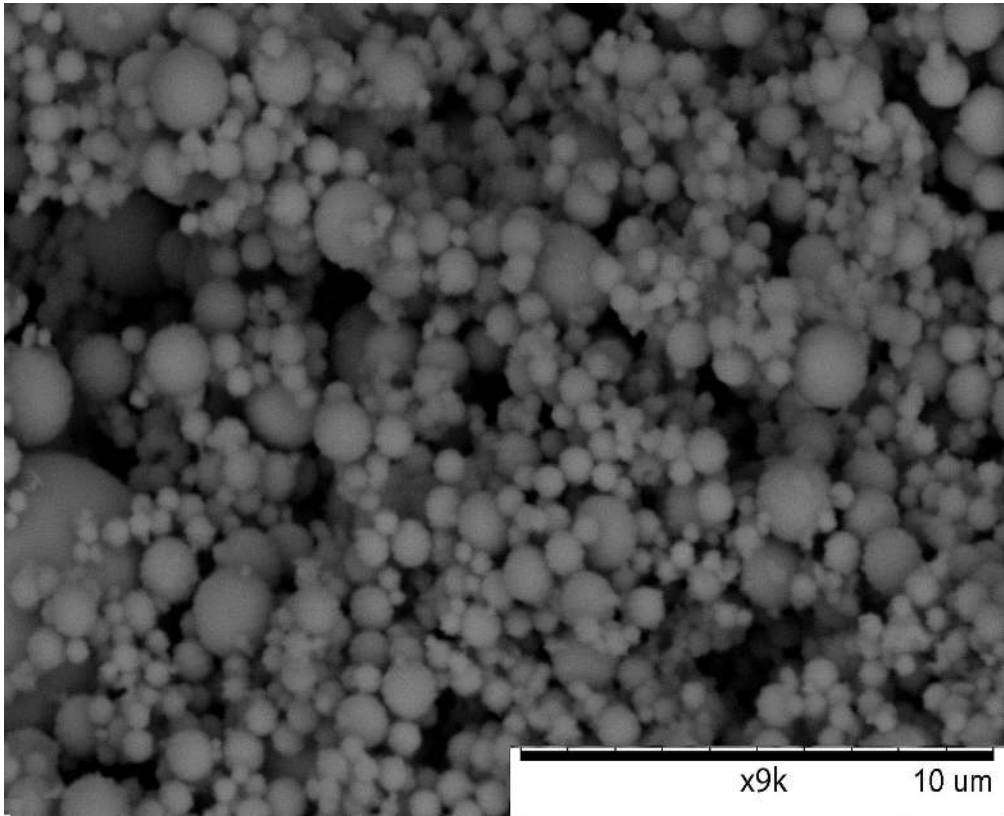
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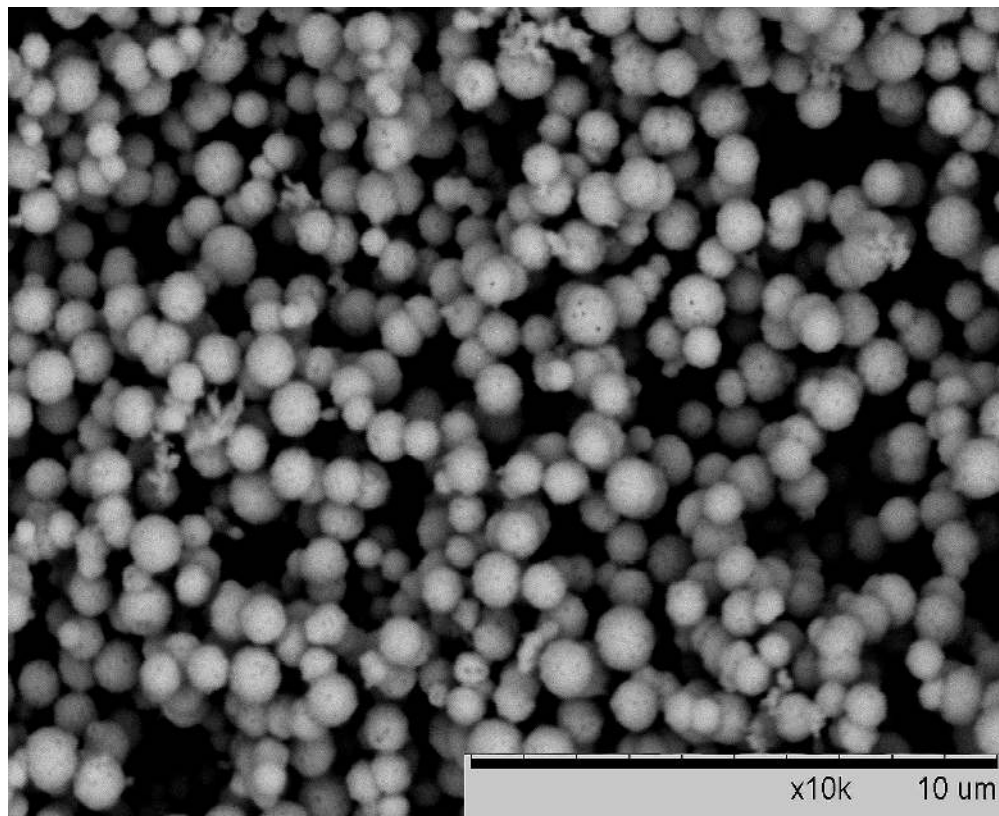
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Obtained by forced hydrolysis of NBVI*
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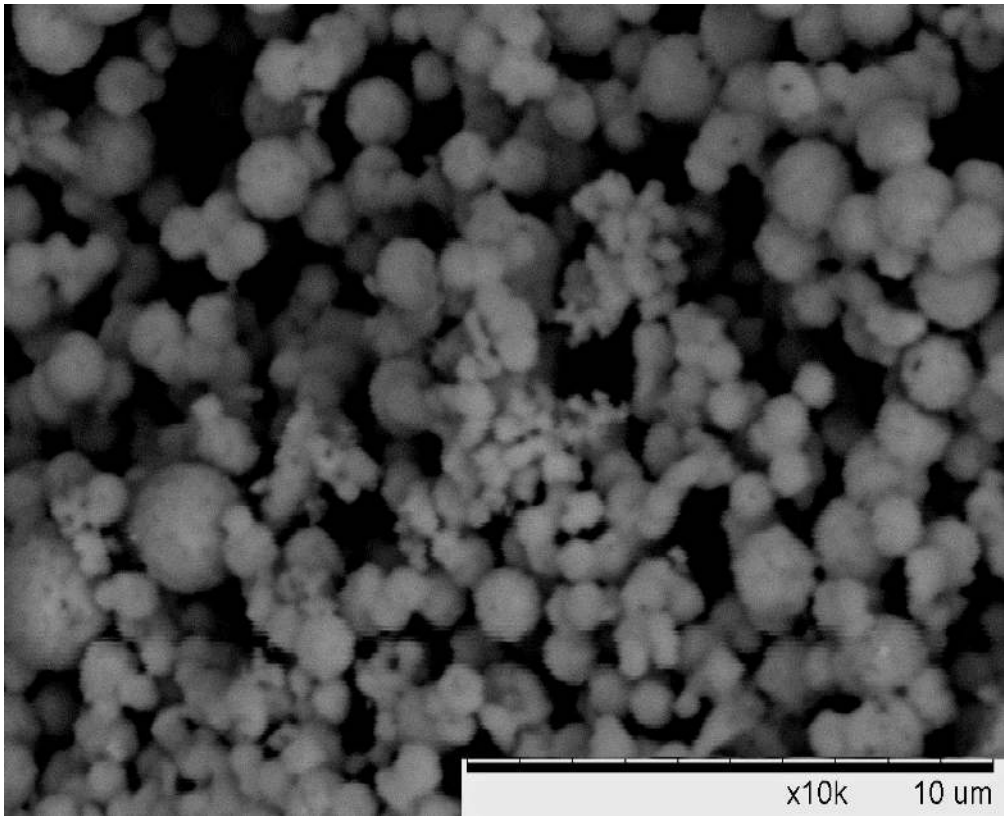
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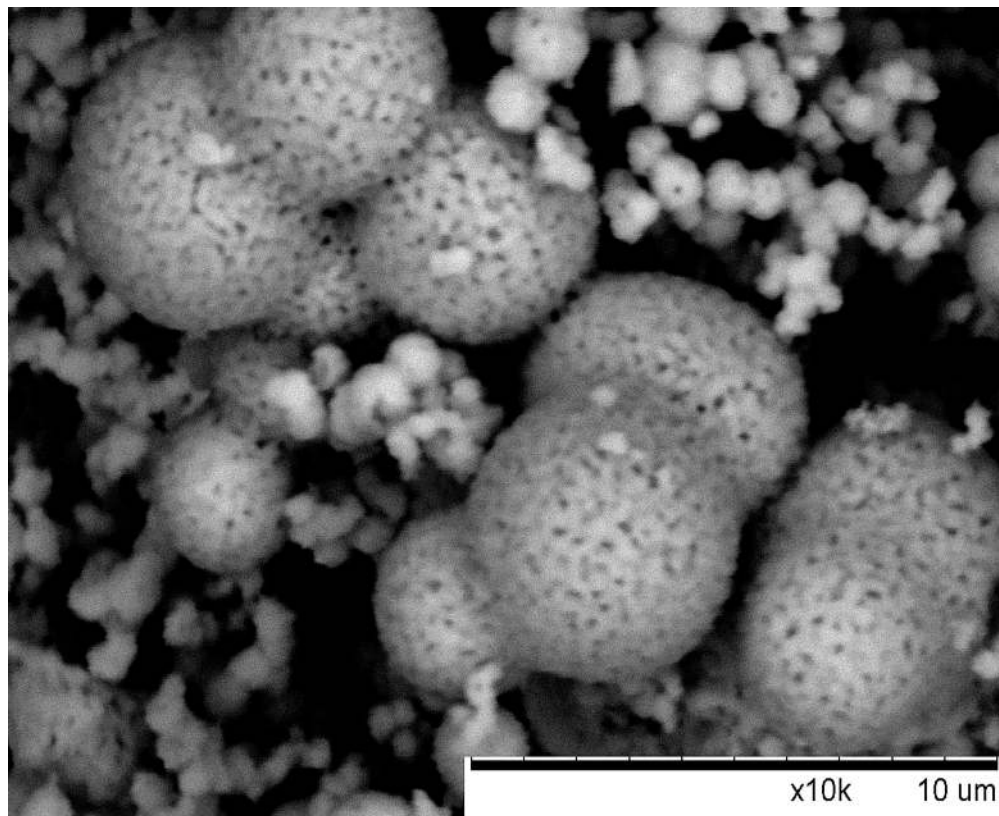
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Obtained from NBI
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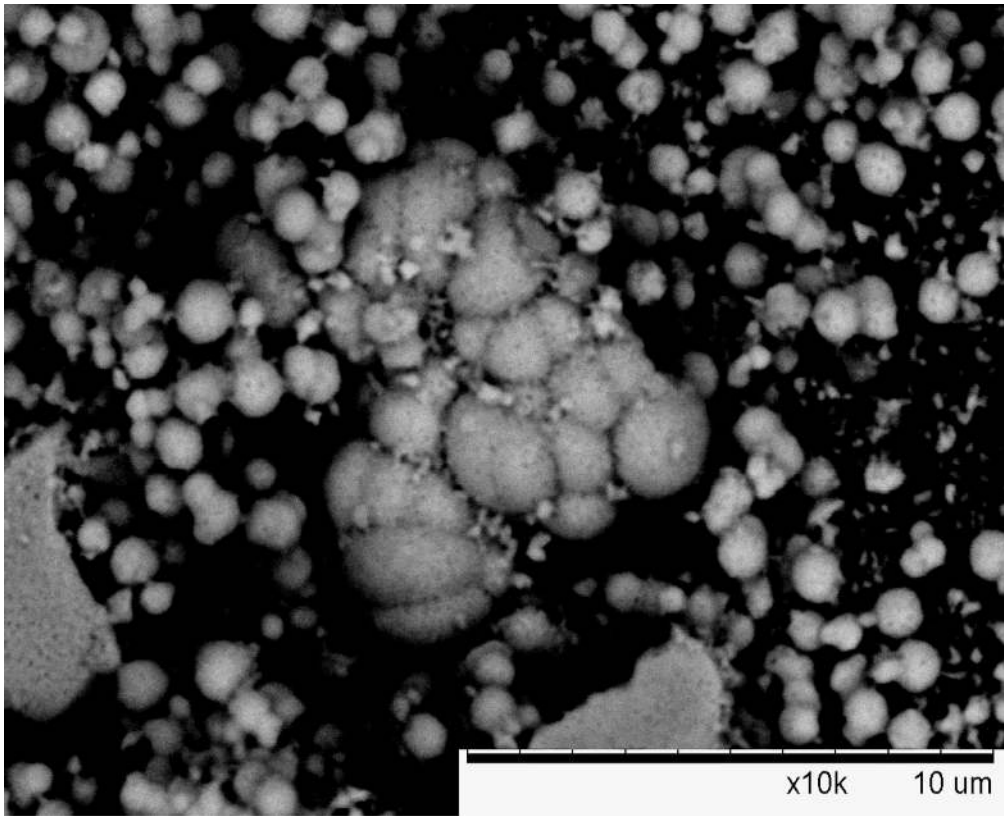
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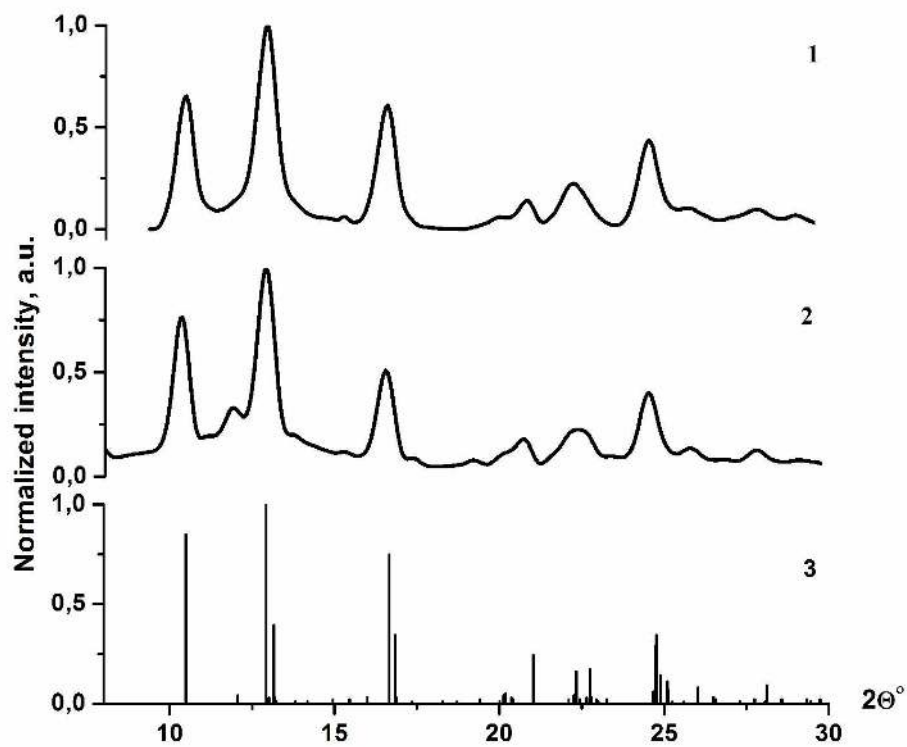
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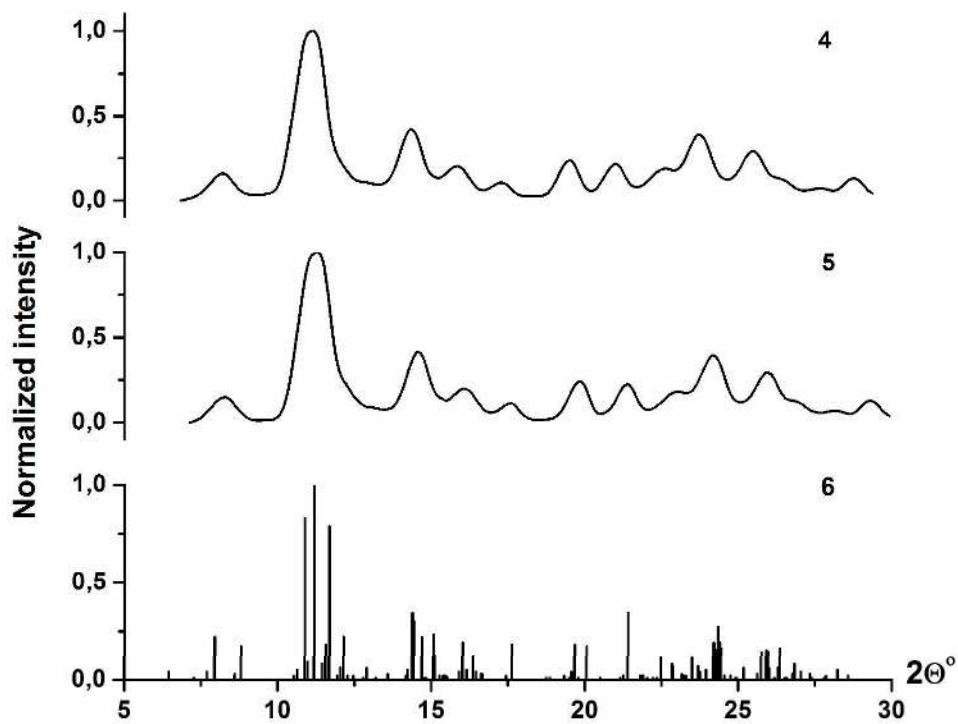


Obtained from NBVI
190x154mm (300 x 300 DPI)



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279x215mm (300 x 300 DPI)

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4 - NbIII, 5 - NbIV, 6 - α -Nb₂O₅ ICDD- JCPDS 37-1468
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