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Mathilde Pelosi, Monique Tillard, David Zitoun. Ge nanoparticles by direct oxidation of Zintl alloys and their electrochemical behavior as anodes of Li-ion batteries. *Journal of Nanoparticle Research*, 2013, 15 (8), pp.1872. 10.1007/s11051-013-1872-2 . hal-00860971

**HAL Id: hal-00860971**

**<https://hal.science/hal-00860971>**

Submitted on 24 May 2022

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# Ge nanoparticles by direct oxidation of Zintl alloys and their electrochemical behavior as anodes of Li-ion batteries

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## Abstract

Germanium nanoparticles (NPs) are obtained by direct oxidation of a solid-state  $\text{Na}_{12}\text{Ge}_{17}$  Zintl phase used as germanium precursor. Alcohols are chosen as oxidizing agents in a room-temperature process. The materials are characterized by X-ray diffraction, transmission electron microscopy, and Raman spectroscopy. The nature of the alcohol influences the final products and their crystallinity. The germanium NPs, tested as anodic materials for lithium ion batteries, display interesting behaviors with initial capacities up to  $1,530 \text{ mAh g}^{-1}$ .

## Keywords

Germanium Nanocrystals Nanoparticles Li-ion Battery

## Introduction

Germanium nanoparticles (NPs) have attracted attention as potential competitors for optoelectronic devices, elaborated by a bottom-up approach. Nano-crystals of germanium have also been studied for the quantum size effect on their optical properties (Heath et al. 1994; Bostedt et al. 2004). Elemental Ge has important applications in transistors, photodetectors, sensors, and batteries. As anodic material for Li-ion batteries, germanium displays a high theoretical capacity ( $1,600 \text{ mAh g}^{-1}$ ) and allows faster lithium diffusivity than silicon (extrapolated to be 400 times higher at room temperature) (Fuller and Severiens 1954). Nevertheless, the use of group 14 elements like Ge, Si, or Sn in anodes has been hampered by the poor cyclability of bulk materials, a drawback that could be overcome at the nanometric scale (Armand and Tarascon 2008). The success in colloidal bottom-up approach synthesis of Ge is more limited than the textbook examples of II–VI and even IV–VI and III–V semiconductors and synthesis pathways deserve further investigations.

Gas-phase syntheses have been reported for the preparation of well-defined germanium nanocrystals (Medeiros-Ribeiro et al. 1998; Min et al. 1996) and nanowires (Morales and Lieber 1998; Wu and Yang 2000; Wang and Dai 2002, 2006; Wang et al. 2005). The common precursors used in chemical vapor deposition are germanes (like  $\text{Ge}_2\text{H}_6$ ), chlorogermanes, or phenylgermanes, which undergo thermal decomposition or hydrogenolysis; in the presence of a metal catalyst, Vapor–Liquid–Solid mechanism leads to the formation of anisotropic nanostructure. These routes have been adapted to the synthesis of Ge nanocrystals in high boiling point solvents (Gerion et al. 2004; Zaitseva et al. 2007; Chockla et al. 2011) or supercritical fluids (Lu et al. 2004; Collins et al. 2010) and to Solution–Liquid–Solid mechanism toward the formation of Ge nanorods (Hanrath and Korgel 2002; Lu et al. 2005a, b).

Examples of solution routes are sparser and involve the reduction of a germanium halide ( $\text{GeCl}_4$  and  $\text{RGeCl}_3$ ) by a strong reducing agent alkalides (Heath et al. 1994), butyl-lithium, naphthalides (Kornowski et al. 1993; Chiu and Kauzlarich 2006), or hydrides (Wilcoxon et al. 2001; Chou et al. 2009), in the presence of a surfactant to limit the NPs growth and coalescence. Very recently, oleylamine has been successfully used as reducing agent in a microwave-assisted synthesis of Ge NPs (Muthuswamy et al. 2013). Alternative routes consist in the thermal

decomposition of chlorogermane (Henderson et al. 2008), germanium alkoxide (Gerung et al. 2006), or  $\text{Ge}\{\text{N}(\text{SiMe}_3)_2\}$  (Lambert et al. 2007; Gerung et al. 2005). As shown recently, the use of  $\text{GeI}_4$  and/or  $\text{GeI}_2$  as precursors allowed a better control over the particle mean size and the size dispersion of the colloidal product (Lu et al. 2005b; Ruddy et al. 2010; Vaughn et al. 2010). The stable colloidal suspension is suitable for spectroscopic studies (Lee et al. 2009) but the surfactant shell impedes its use for electronic/electro-chemical applications.

Therefore, the use of germanium Zintl phases (Ge anionic clusters with alkaline or alkaline earth cations) has been standing as a unique room-temperature oxidative route toward Ge nanostructures. The Kauzlarich group has first developed this pathway by varying the Zintl phase ( $\text{NaGe}$ ,  $\text{KGe}$ , and  $\text{Mg}_2\text{Ge}$ ), the oxidative agent  $\text{GeCl}_4$  or  $\text{GeCl}_2$ , and the capping agent (Grignard reagent or alkyl lithium) (Ma et al. 2008; Taylor et al. 1998, 1999, 2002; Tanke et al. 2003; Chiu et al. 2005). Such Zintl precursors have been successfully used as building blocks for Ge mesoporous materials (Armatas and Kanatzidis 2006a, b; Sun et al. 2006) and anodic materials for Li-ion batteries (Lee et al. 2005; Park et al. 2010). Soft oxidation of Zintl phases also yields a guest-free clathrate and an allotropic form (metastable allo-Ge) of germanium (Guloy et al. 2006; Gruttner et al. 1982; Kiefer et al. 2011). The appealing ability to perform a room-temperature synthesis of germanium nanostructures starting from Zintl precursors, and the need for better comprehension of the reaction pathways, drive us to study the reactivity of germanium anionic clusters under oxidation with mild reducing agents.

In this paper, we report on the preparation of Ge NPs by direct oxidation of Ge Zintl anions by alcohols. The  $\text{Na}_{12}\text{Ge}_{17}$  Zintl phase has been used as germanium precursor either in solid state or after dissolution in ethylene diamine. Transmission electron microscopy (TEM) shows that the chain length of the alcohol and the reaction pathway have some influence on the mean size, the crystallinity, and the size dispersion of particles. The crystallinity has been carefully studied by Raman spectroscopy. The electrochemical tests of NPs as anodic materials for Li-ion batteries emphasize the relationship between synthesis pathway and properties.

## Experimental section

Reagents are handled in a glove box under controlled argon atmosphere. Before the reaction with Ge powder (150  $\mu\text{m}$ , Sigma Aldrich, 99.999 %), the alkali metal (Na, ingots, Merck, 99 %) is scraped to remove surface impurities. Benzyl alcohol ( $\text{C}_6\text{H}_5\text{CH}_2\text{OH}$ , 99 %, anhydrous, Acros-Organics) is used without further purification and anhydrous n-butanol ( $\text{C}_4\text{H}_9\text{OH}$ , Sigma Aldrich) is dried on a Schlenk line. The Zintl phase  $\text{Na}_{12}\text{Ge}_{17}$  is prepared from stoichiometric proportions of the elements mixed in a weld-sealed Ta-container, enclosed in its turn within an evacuated silica or stainless steel jacket to avoid any oxidation at high temperature. The Na/Ge mixture is heated to 1,230 K for 2 days during which the reactor is shaken several times to improve homogenization of the melt. The air-sensitive products of reactions are handled in the glove box.

NPs are obtained by direct oxidation of the solid precursor by an alcohol. In such a reaction, called route I reaction, the alcohol is acting as oxidizing agent as well as solvent. The  $\text{Na}_{12}\text{Ge}_{17}$  powder (100 mg) is slowly added to 10 mL alcohol under stirring in a glove box or on a vacuum line. The reaction time is optimized for completion of the reaction, and the solution is left to stir for 16 h, a time beyond which no homogeneity improvement in the product is observed. At the end of the reaction, the black powder (Ge NPs) is separated by centrifugation and washed with hexane.

NPs are also obtained from a different pathway in the route II reaction: the solid  $\text{Na}_{12}\text{Ge}_{17}$  (100 mg) is initially dissolved in ethylene diamine (10 mL) before adding the alcohol (10 mL). For this alternate route, the total reaction time should be divided into two parts: the time required for the solubilization which is set to 72 h and the oxidation time which is fixed to 16 h (as in the route I reaction). Temperature change from 273 to 473 K has no noticeable effect on the reaction products; therefore, all the experiments are carried out at room temperature.

A modified pathway named route III is used in order to avoid the high-temperature synthesis of the Zintl phase. The elements, Na and Ge, were taken in 12/17 atomic proportions and directly mixed in ethylene diamine before the oxidation reaction.

Note that similar reactions are successfully carried out starting from  $\text{K}_4\text{Ge}_9$ , a Zintl phase having a weaker reducing character (attributed to the weaker negative mean charge per Ge atom (Goicoechea and Sevov 2005) and consequently a lower reactivity toward the alcohol.

TEM is performed with a JEOL 1200 EX II microscope operating at 100 kV accelerating voltage. The X-ray diffraction (XRD) patterns are recorded on a Philips analytical X'pert diffractometer equipped with a copper tube

and a hybrid monochromator (parabolic multilayer mirror and two-crystal, Cu K $\alpha$  radiation). The Raman spectroscopy is carried out using a LABRam Aramis IR<sup>2</sup> spectrometer equipped with a helium neon laser (633 nm). The intensity of the laser beam is estimated to a maximal value of 6 mW very close to the sample. The use of various filters (named D0.6, D1, and D2) allows a reduction of intensity to 200, 70, and 50  $\mu$ W, respectively.

The electrochemical behavior of nanostructured Ge powders is evaluated in Li/LiPF<sub>6</sub> (EC, PC, DMC)/Ge electrochemical cells assembled under argon in a glove box. Electrodes incorporated 85 wt. % active material intimately mixed with 15 wt. % carbon black. A 1 M solution of LiPF<sub>6</sub> in a 1:1:3-mixture of ethyl, propyl, and dimethyl carbonates (Merck Laboratories) is used as electrolyte. Charge–discharge experiments are monitored in standard conditions using a MacPile multichannel galvanostat–potentiostat. Li insertion/ extraction is limited within a 0–2.5 V (vs. Li<sup>+</sup>/Li) window voltage. The polarization between charge and discharge voltage upon cycling did not exceed 0.5 V. The voltage versus composition curves are plotted for the first cycles of galvanostatic experiments performed at a regime corresponding to the exchange of one Li atom in 10 h.

## Results and discussion

Ge NPs have been prepared from the oxidation of a Zintl phase used as Ge precursor (Fig. 1). Alcohols are chosen as oxidizing agents for their especially mild reducing character. Furthermore, they can be easily removed from the nanostructure surface after synthesis. The alcoholic proton reacts with the Zintl phase to give gaseous H<sub>2</sub> and alkoxide (which protonates upon reaction in air). Indeed, the reaction occurs with gas evolution and no other by product has been noticed from <sup>1</sup>H NMR spectrum of the supernatants.

The route I reaction is a direct oxidation of the solid phase by a liquid oxidant also acting as solvent. Instead, route II reaction takes advantage of the special property of Ge Zintl phases to be soluble in basic polar solvents, as for example ethylene diamine. Solubilization releases free Zintl anions in the solution that are ready to react directly with the oxidant in the liquid phase.

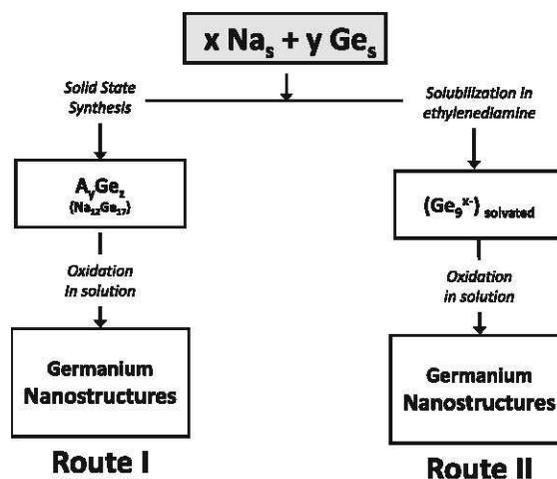


Fig. 1 Schematic representation of the synthetic routes I and II

## Precursor characterization

The solid Zintl phase Na<sub>12</sub>Ge<sub>17</sub> is prepared from the elements by high-temperature synthesis. The profile refinement of its experimental XRD powder pattern (Rietveld procedure in program Jana) leads to good agreement factors R<sub>p</sub>/R<sub>wp</sub> of 1.1/1.8 % (Fig. 2a). Parameters of the monoclinic unit cell (P2<sub>1</sub>/c) converge to a = 22.095(2), b = 12.8090(6), c = 41.580(2) Å, β = 91.341(2), values that agree well with those previously reported (Taylor et al. 1998): a = 22.11, b = 12.80, c = 41.56 Å, β = 91.31°. The structure of Na<sub>12</sub>Ge<sub>17</sub> is characterized by the coexistence of two discrete anionic units, Ge<sub>4</sub><sup>4-</sup> and Ge<sub>9</sub><sup>4-</sup>, packed together with sodium cations (Fig. 2c). Then, the Na<sub>12</sub>Ge<sub>17</sub> Zintl compound can be formally described as [Na<sup>+</sup>]<sub>12</sub>[Ge<sub>4</sub><sup>4-</sup>]<sub>2</sub>[Ge<sub>9</sub><sup>4-</sup>].

Raman spectroscopy is a complementary technique for the full identification of  $\text{Na}_{12}\text{Ge}_{17}$ . According to literature reports (Hoch et al. 2003; Kliche et al. 1992; Somer 2000),  $\text{Ge}_4^{4-}$  unit (symmetry  $T_d$ ) is characterized by three Raman active vibration modes observed at  $274\text{ cm}^{-1}$  ( $m_1$ ),  $207\text{ cm}^{-1}$  ( $m_2$ ), and  $164\text{ cm}^{-1}$  ( $m_3$ ). For  $\text{Ge}_9^{4-}$  anion displaying the  $C_{4v}$  symmetry, only five out of the 20 active modes are observed at  $222\text{ cm}^{-1}$  (intense),  $164$  and  $167\text{ cm}^{-1}$  (medium), and  $241$  and  $188\text{ cm}^{-1}$  (weak).

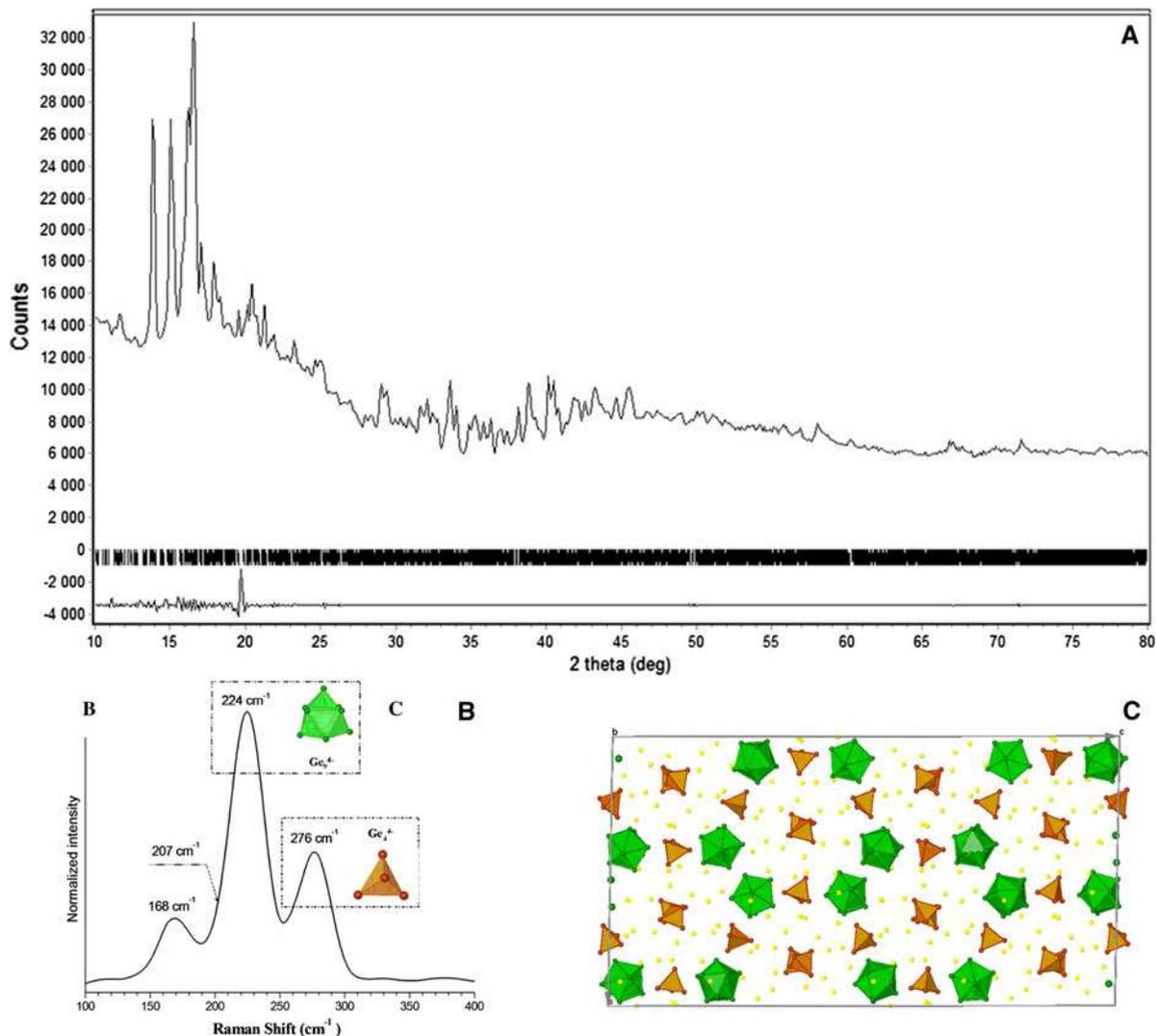


Fig. 2 X-ray diffractogram of  $\text{Na}_{12}\text{Ge}_{17}$  (a), Raman spectrum of  $\text{Na}_{12}\text{Ge}_{17}$  (b) and structure projection along b axis emphasizing the  $\text{Ge}_4^{4-}$  and  $\text{Ge}_9^{4-}$  anionic clusters (c)

The Raman spectrum recorded for  $\text{Na}_{12}\text{Ge}_{17}$  is composed of three bands (Fig. 2b), those at  $276$  and  $224\text{ cm}^{-1}$  are characteristic of  $\text{Ge}_4^{4-}$  and  $\text{Ge}_9^{4-}$  clusters, respectively. The slight asymmetry of the band lying at  $224\text{ cm}^{-1}$  is attributed to overlapping of the  $m_2$  mode of  $\text{Ge}_4^{4-}$  ( $207\text{ cm}^{-1}$ ) with the intense band of  $\text{Ge}_9^{4-}$  ( $222\text{ cm}^{-1}$ ). Finally, the third band at  $168\text{ cm}^{-1}$  is assigned to the  $m_3$  vibration mode of  $\text{Ge}_4^{4-}$  ( $164\text{ cm}^{-1}$ ) which overlaps with the  $\text{Ge}_9^{4-}$  vibration modes occurring in the same domain ( $188$ ,  $164$ , and  $147\text{ cm}^{-1}$ ). Raman spectroscopy brings information that, in addition to XRD results, allows the complete and unambiguous identification of compound  $\text{Na}_{12}\text{Ge}_{17}$ .

## The route I solid–liquid reaction

The route I solid–liquid reaction is carried out using solid  $\text{Na}_{12}\text{Ge}_{17}$  and either benzyl alcohol or butanol as oxidizing agents. Powder XRD analyses of products have shown that  $\text{Na}_{12}\text{Ge}_{17}$  reacts completely during the oxidation reaction. In all cases, products of reaction are crystalline as proved by the electron diffraction patterns indexed within the cubic diamond-type structure of germanium (JCPDS files no. 04-0545,  $a = 5.657 \text{ \AA}$ ).

As clearly illustrated in Fig. 3, particle size depends on the nature of the oxidizing agent: small NPs of  $5.4 \pm 2.3 \text{ nm}$  in diameter are obtained from benzyl alcohol while particles of  $19.1 \pm 8.5 \text{ nm}$  in diameter, with a larger size distribution, are obtained from butanol. In the following, these samples will be referred to as 5 and 20 nm NPs.

Raman spectroscopy has proven its efficiency to characterize these materials, even those being poorly crystallized or amorphous. Actually, Raman lines are characterized by their shape (width, symmetry) and their position, parameters that are strongly correlated with the structure/morphology (crystalline, amorphous) and with the particle size (micrometric, nanometric).

Germanium nanopowders from route I reactions are analyzed by Raman spectroscopy. Their spectra, shown in Fig. 4, present a large and asymmetric band clearly shifted from the position of the Raman line attributed to crystalline germanium (Ge–Ge stretch at  $302 \text{ cm}^{-1}$ ). The Raman band position is dependent on the particle size, shifted toward low wavenumbers when the average particle size decreases, and broadened when the amorphous character increases (Muthuswamy et al. 2013; Schlecht et al. 2004; Gorokhovet al. 2005). After background correction, the experimental Raman spectrum is fitted using two Gaussian functions centered at  $278$  and  $261 \text{ cm}^{-1}$  (sample from benzyl alcohol). The contribution at  $278 \text{ cm}^{-1}$  is attributed to the crystalline core of the particles while that at  $261 \text{ cm}^{-1}$  corresponds to the amorphous part of the sample (surface and/or amorphous NPs). For samples prepared in butanol, Gaussian parameters are found slightly closer to values for bulk Ge, the Raman shifts of  $288$  and  $271 \text{ cm}^{-1}$  are in agreement with an increase in the coherence length of the particles crystalline core. These values of Raman shifts can be compared with  $273 \text{ cm}^{-1}$ , a value reported for nanocrystals (Ma et al. 2008).

## The route II liquid–liquid reaction

Ge-nanosized samples are also prepared by the liquid–liquid route II reaction, which proceeds through the solubilization in ethylene diamine of the  $\text{Na}_{12}\text{Ge}_{17}$  Zintl phase prior to oxidation. In that case, the individualized  $\text{Ge}_4^{4-}$  and  $\text{Ge}_9^{4-}$  Zintl units already present in the solid state are released as free anionic species in the solution. The intense brown–violet color of the solution was indicative of the presence of Ge anionic clusters able to react immediately with the alcohol. The route II reaction takes place in specific conditions (kinetics, mechanism...) that might be significantly different from route I reaction conditions.

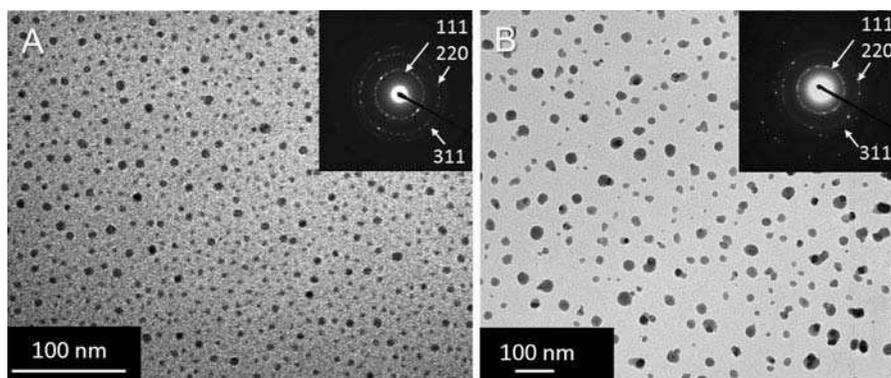


Fig. 3 TEM micrographs of crystalline Ge nanoparticles from route I using benzyl alcohol (a) and butanol (b) as oxidizing agent (insets SAED)

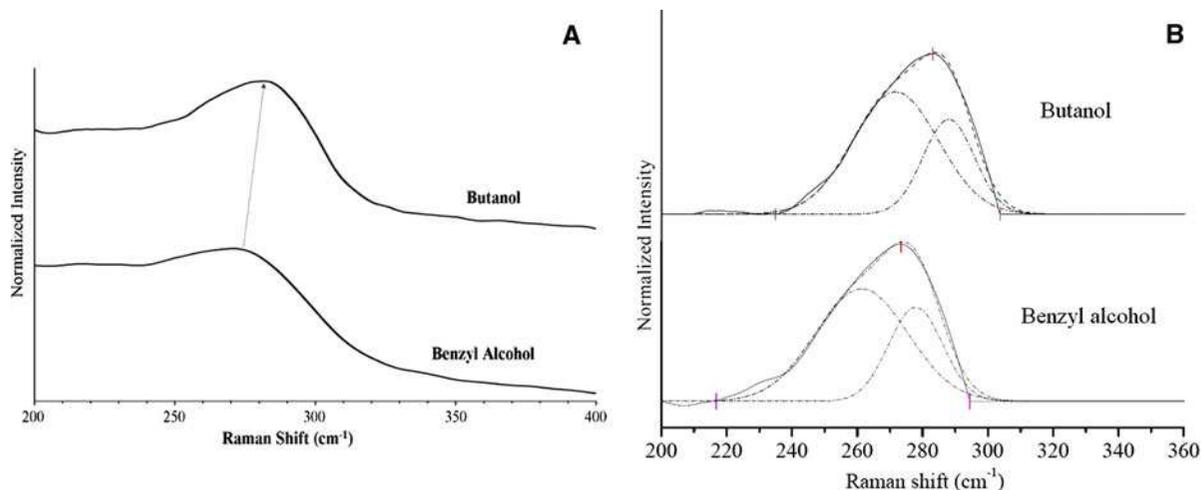


Fig.4 a Raman spectra of Ge nanopowders illustrating the increase in Raman shift (from benzyl alcohol to butanol). b Fitted spectra after background correction

Powders obtained in that way with benzyl alcohol or butanol are studied using Raman spectroscopy. The recorded spectra, typical of Ge nanosized particles, showed large and asymmetric bands with Raman shifts of  $264\text{ cm}^{-1}$  (benzyl alcohol) and  $273\text{ cm}^{-1}$  (butanol). The displacement of the band from  $264$  to  $273\text{ cm}^{-1}$  may be interpreted as the consequence of an increase in the long-distance order within the material, associated with a particle size enlargement or/and a crystallinity improvement. Comparison with the Raman shifts ( $278$  and  $288\text{ cm}^{-1}$ ) observed for route I products indicates that route II products displayed a shorter coherence length.

Sample morphology and particle size distribution are examined by electron microscopy, the TEM micrographs of two samples prepared from benzyl alcohol and butanol are given comparatively in Fig. 5.

Whatever oxidizing agent is used, individualized NPs are found embedded in an amorphous germanium matrix. With benzyl alcohol, particles are below  $7\text{ nm}$  in diameter and amorphous; while with butanol, crystalline particles range between  $10$  and  $15\text{ nm}$ . The difference in morphology between routes I and II products may be attributed to the ability of Zintl anions to react with the alcohol. In the route I reaction, the Zintl anionic units ( $\text{Ge}_4^{4-}$  and  $\text{Ge}_9^{4-}$ ) confined in the solid phase would react rather slowly with the alcohol. Instead, in the route II reaction, these units already present in the solution are immediately available for reaction with alcohol, leading to embedded NPs.

Results acquired from microscopy well agree with findings from Raman spectroscopy and XRD. In route II experiments, as already observed for route I, light alcohols are the most reactive oxidizing agents. Butanol is found to react faster than benzyl alcohol and to produce larger particles with higher crystallinity. The exothermic character of the reaction causes a local elevation of temperature, a factor that could favor the crystallization of Ge particles.

### The route III reaction

In order to avoid the Zintl-phase high-temperature synthesis, elements are directly mixed in ethylene diamine. Though soluble in ethylene diamine, some solid germanium is still present at the bottom of the reactor after stirring the solution during 3 days. The dark-violet coloration of the solution indicated the presence of germanium anions. After filtration, the solution is added to benzyl alcohol or butanol to perform the oxidation reaction. No effect related to the nature of the alcohol can be detected.

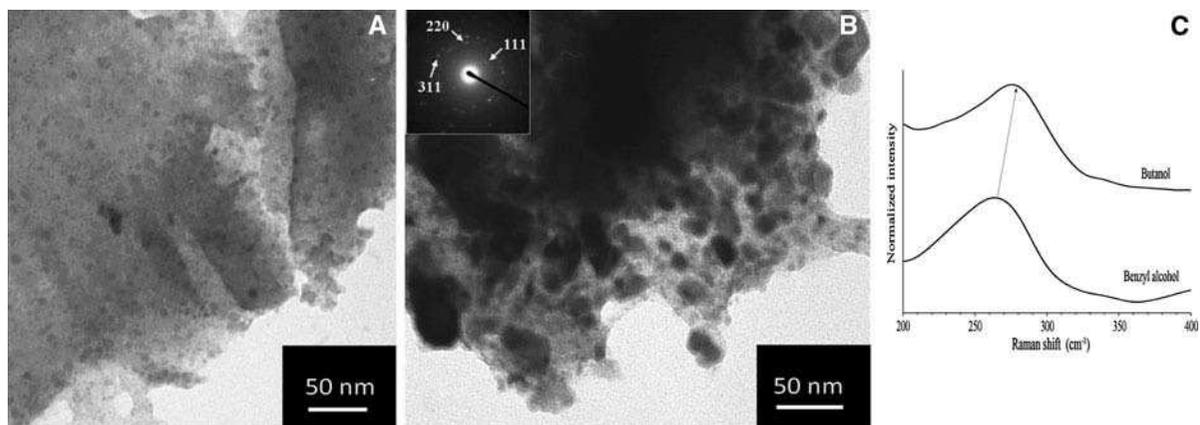


Fig. 5 TEM micrographs of Ge nanoparticles obtained from route II in benzyl alcohol (a) and butanol (b), Raman spectra of the corresponding materials (c)

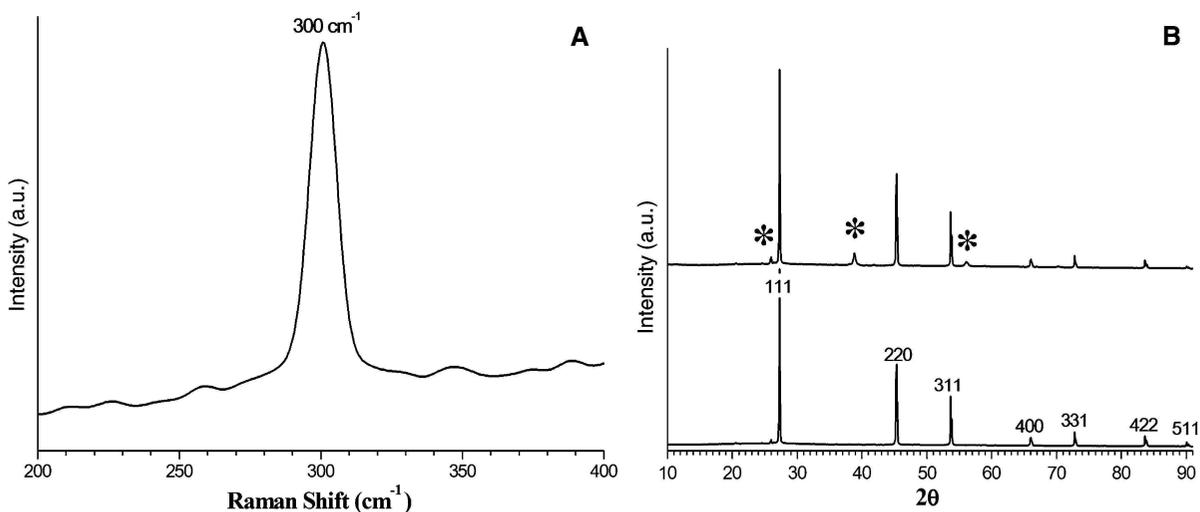


Fig. 6 Raman spectrum for a sample from route III with benzyl alcohol (a) and its XRD powder pattern indexed with cubic Ge (b)

The black powder collected, washed, and dried is analyzed by XRD, its pattern indicated the presence of a small quantity of a sodium hydrogen germanium oxide hydrate, identified as  $\text{Na}_3\text{HGe}_7\text{O}_{16}(\text{H}_2\text{O})_4$  (ICDD no. 01-076-0102), in addition to crystallized Ge. Diffraction lines marked with a star correspond to the germanium hydrate (Fig. 6b).

The presence of germanium hydrate supports the proposed mechanism: Ge anionic clusters can be oxidized either to Ge in anhydrous conditions or to sodium hydrogen germanium oxide hydrate in the presence of residual water (several ppms). The hydrate is easily eliminated by acid wash with dilute nitric acid. Unlike powders from routes I and II, the Ge powder from route III displays very narrow XRD lines suggesting the presence of large crystalline particles. The Raman band is observed at  $300\text{ cm}^{-1}$ , the same position as for Ge commercial powder (150 lm). Nevertheless, with a half-maximum full-width of  $11\text{ cm}^{-1}$  instead of  $4\text{ cm}^{-1}$ , the band is slightly wider. The asymmetry would be assigned to a non-homogeneous size distribution and broadening explained by a reduction in the particle size.

These interpretations are consistent with the large crystalline aggregates of 50–100 nm observed in TEM images for samples prepared either from benzyl alcohol or from butanol (not shown). The observed particle size in product is at least two orders of magnitude below the powder used as reagent, another argument for the reaction mechanism proposed. At first sight, the routes II and III reactions would appear very similar. Nonetheless, they are quite different since their final products do not display close characteristics. It is likely that solubilization in ethylene

diamine of  $\text{Na}_{12}\text{Ge}_{17}$  on one hand, and Na and Ge (in 12/17 atomic proportions) on the other hand, do not produce identical solutions. Raman studies of these solutions could indicate which species were present in the solutions, but they do not give workable data. Finally, no conclusive proof of the formation of Ge Zintl anions in the solution can be provided, anyway consideration of results from route III stresses the need to retain the high-temperature synthesis of the Zintl phase and underlines how complex the reaction media could be.

### Raman spectra and samples stability

For each sample, Raman spectra are first acquired using the lowest intensity for the laser beam (50  $\mu\text{W}$ , D2 filter), and then the intensity can be increased to 75  $\mu\text{W}$  (D1) or 200  $\mu\text{W}$  (D0.6). Depending on experimental conditions, a displacement of the Raman bands is observed that could be correlated with the sample stability under irradiation. To better understand the origin and try to quantify the amplitude of such displacements, Raman experiments are performed for Ge wafers and Ge commercial powder. For these two samples, Raman lines occur at 300  $\text{cm}^{-1}$  in the first spectrum recorded with the lowest intensity (D2 filter).

As shown in Fig. 7, a progressive increase of the beam intensity using successively D1 and D0.6 filters does not induce any change in the band position for the Ge wafer. Instead, the Raman band of the Ge powder is continuously shifted toward weak displacement, as the beam intensity is increased (D1 then D0.6 filter). After a subsequent intensity lowering (D2 filter), the Raman line recovers its initial position. A similar phenomenon has been reported for small silicon crystalline nanowires of about 2 nm in diameter (Gupta et al. 2003). This reversible event results from thermal expansion: crystal cell inflates with the elevation of local temperature due to the beam energy and then contracts upon cooling. In the meantime, the Raman band shifts to low displacement and retrieves its initial position. This effect is only observed for powders and then called “powder effect.” Owing to higher structural constraints, it does not occur for Ge wafers. This phenomenon, sensitive to the sample morphology, would be also associated with a decrease in thermal conductivity caused by the weakening of inter-grain contacts in finely divided powdered samples.

Similar experiments are carried out with the nano-sized Ge powders prepared in this work. As expected, the Ge powder obtained from route III reaction, composed of large particles of \*100 nm, gives rise to the powder effect, like the commercial powder.

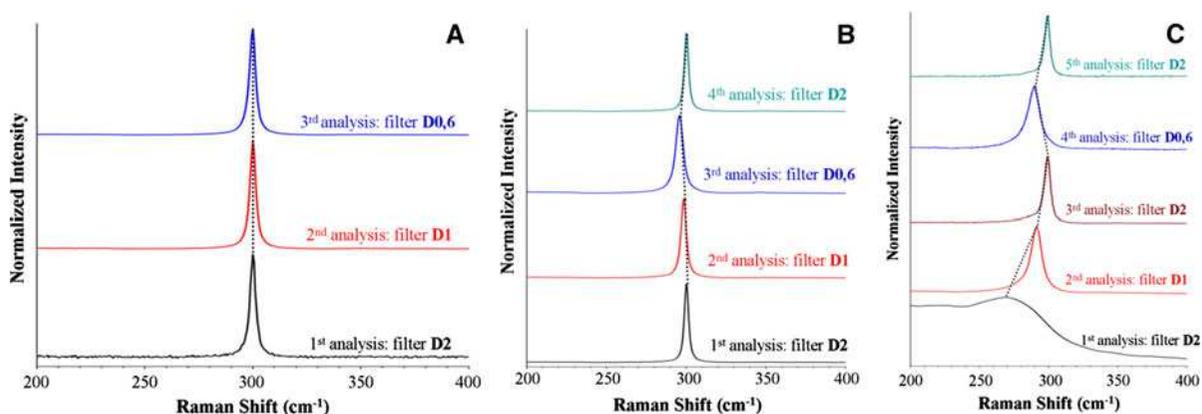


Fig. 7 Raman spectra under different irradiations for Ge wafers (a), Ge commercial powder (b), 5-nm Ge nanocrystals from route I (c)

The Raman band (D2 filter) of particles prepared from route I in benzyl alcohol is very intense (Figs. 4, 7). This intensity is characteristic of a loss in the long-distance order that causes suppression of the selection rules and allows all phonons participation (Richter et al. 1981). The raise in intensity occurs together with a narrowing of the band strongly moved toward high displacement values, sign of some local ordering (coalescence/crystallization). After a subsequent come back to low intensity (D2 filter) the band is observed at 300  $\text{cm}^{-1}$  (Fig. 7). Based on results previously reported for silicon with shifts increasing with the particle size (Iqbal et al. 1980; Morhange et al. 1979), this irreversible phenomenon is assimilated to a thermal annealing.

Particles prepared from route II (16 h reaction) display a Raman band at  $300\text{ cm}^{-1}$ , which shifts reversibly to low displacements demonstrating a powder effect. In case of powders collected after only 2 h of reaction (smaller amorphous particles), displacements of the Raman band initially occurring at  $264\text{ cm}^{-1}$  indicate that the material first undergoes a thermal annealing and then a powder effect.

## Electrochemical experiments

The Ge powders prepared in the present work are tested against a pure lithium electrode in electrochemical cells. Starting with a discharge of the battery, the maximum lithium insertion that can be achieved led to nominal compositions  $\text{Li}_{4.2}\text{Ge}$  and  $\text{Li}_{3.4}\text{Ge}$  for, respectively small particles (5 nm) from benzyl alcohol and large particles (20 nm) from butanol, corresponding to the respective specific capacities of  $1,530$  and  $1,250\text{ mAh g}^{-1}$  (Fig. 8). The limits in stoichiometry reached at the end of the subsequent charges of the cells are  $\text{Li}_{1.7}\text{Ge}$  and  $\text{Li}_{1.2}\text{Ge}$ , leading to irreversible capacities of  $600$  and  $380\text{ mAh g}^{-1}$  that would include solid–electrolyte layer occurring at first discharge. The reversible specific capacity for the complete first cycle is then about  $900\text{ mAh g}^{-1}$  but the capacity rapidly fades upon cycling.

The voltage versus composition curves are represented in Fig. 8 for the first cycles of galvanostatic experiments. The first discharge is characterized by a pseudo plateau representing the continuous insertion of  $\text{Li}^+$  into the electrode (Graetz et al. 2004). After the expected and usual small insertion of lithium in carbon black around  $0.75\text{ V}$ , the Li insertion in Ge particles proceeds through two distinct processes better visualized in the derivative plots  $dx/dV$  as two peaks occurring at  $0.35$  and  $0.18\text{ V}$  (Fig. 8a). During the following charge, the large and intense peak observed at  $0.5\text{ V}$  is associated with lithium extraction. The insertion mechanism of Li in small particles looks more complex, as attested by the additional and not fully separated peaks observed at  $0.5\text{ V}$  in discharge and at  $0.4\text{ V}$  in charge on potential derivative plots (Fig. 8a).

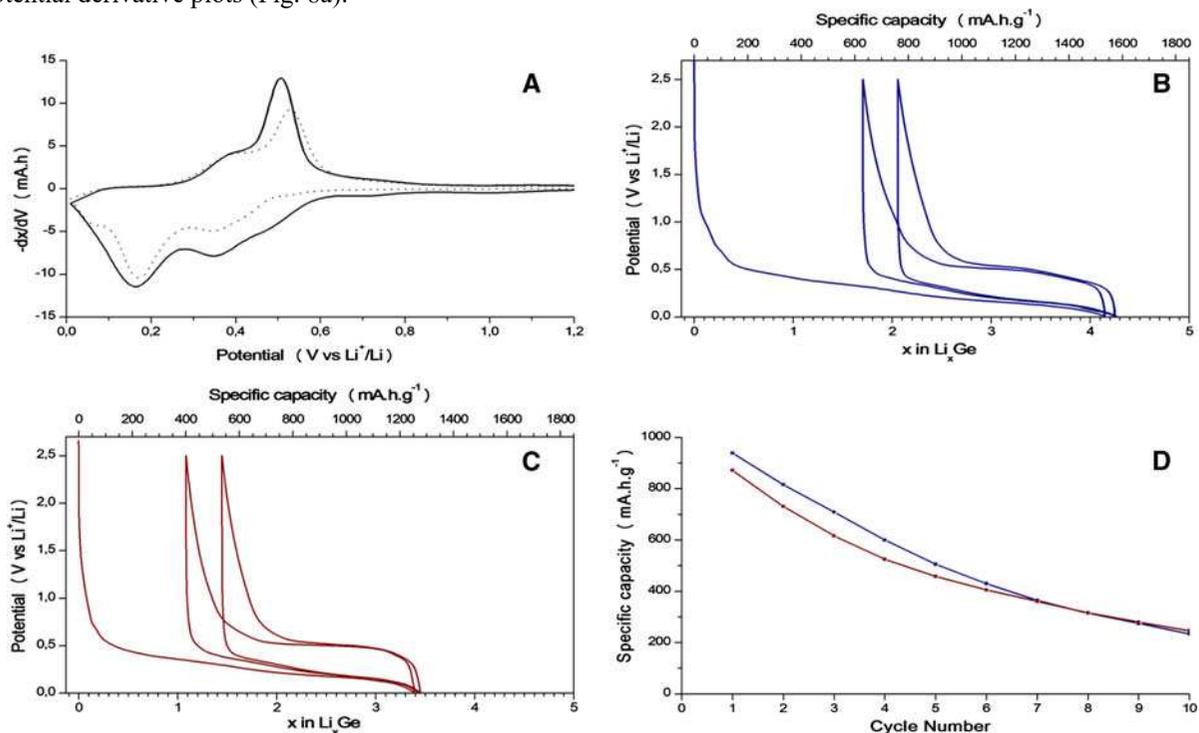


Fig. 8 Derivative plot  $dx/dV$  of small (plain line) and large (dotted line) particles (a), specific capacity for large (b) and small particles (c). d Comparative cycling performances

Despite quite good initial specific capacities, the initial capacity rapidly fades after 10 discharge/charge cycles. Such loss is often encountered for lithium insertion into electrochemically active materials as long as experimental conditions have not been optimized. The factors commonly mentioned to account for cycling degradation are the drop in electric conductivity due to electrode damage upon cycling and the side reactions with electrolyte that are

enhanced by large surface areas of highly divided materials.

Li insertion into larger particles of 50–100 nm prepared from route III also occurs through a pseudo plateau up to a maximum of  $\sim 4\text{Li}^+$  inserted per Ge, leading to a specific capacity of  $1,405\text{ mAh g}^{-1}$ . The subsequent charge does not allow the extraction of the total Li amount, thus the electrode shows a rather large irreversible capacity of  $510\text{ mAh g}^{-1}$ . Nevertheless, the capacity retention is found around 60 % over 10 cycles, which is much better than in case of small particles.

These exploratory experiments clearly evidence that the electrochemical behavior is related to the structure/morphology of the electro-active material. Small-sized particles are found not only to improve the initial capacities but also to increase the irreversible part of the capacity while larger particles are favorable to good capacity retention upon cycling. In these nanosized Ge materials, lithium insertion occurs through a continuous process without steps corresponding to the formation of intermediate intermetallic compounds. Improvement of the electrochemical performances would be reached by finding a compromise and optimizing the particles size. On the other hand, materials conditioning and experimental parameters need to be refined (binder, conducting matrix, exchange rate, potential windows,...) to improve the characteristics and the cycling lifetime of these batteries.

Initial capacities reach without any optimization vary from 1,250 to  $1,530\text{ mAh g}^{-1}$ , not so far from the theoretical capacity of  $1,600\text{ mAh g}^{-1}$  calculated for the nominal composition  $\text{Li}_{22}\text{Ge}_5$ . These capacities are in the same range as initial capacities reported in the literature for other kinds of nanosized germanium (Graetz et al. 2004) like Ge nanowires with an initial capacity of  $1,141\text{ mAh g}^{-1}$  retained over 20 cycles (Chan et al. 2008). Ge amorphous NPs display a capacity of  $1,470\text{ mAh g}^{-1}$  (Lee et al. 2005) while after annealing, the resulting crystallized particles inserted in a carbon matrix display a capacity of  $1,200\text{ mAh g}^{-1}$  of which 88 % is retained over 30 cycles (Lee et al. 2007).

## Conclusion

Nanostructured Ge powders are prepared using three different routes; the two first routes are based on the oxidization by an alcohol of a Zintl phase used as Ge precursor. The third route consists in the in situ Zintl anions formation and oxidation. Results clearly emphasize the crucial and unique role of the Zintl phase as a precursor of small-sized Ge NPs. One-phase reactions (route II) allow preparing crystalline particles embedded in amorphous matrix while two-phase reactions (solid/liquid, route I) yield well-defined isolated NPs. The difference in the product morphology may depend on the ability of the Zintl anions  $\text{Ge}_4^{4-}$  and  $\text{Ge}_9^{4-}$  to react with the alcohol. The free anions in the liquid phase ( $\text{Na}_{12}\text{Ge}_{17}$ /ethylene diamine solution) react more quickly than the anions frozen in the solid  $\text{Na}_{12}\text{Ge}_{17}$  Zintl phase.

The results obtained in the present work show that the size of Ge NPs could be tuned by changing the chain length of the oxidizing agent, butanol and benzyl alcohol leading, respectively, to 5 and 20 nm NPs.

The Ge NPs prepared using these original oxidation reactions are free of organic residues and, when tested in electrochemical cells, exhibited initial capacities up to  $1,530\text{ mAh g}^{-1}$  with a capacity retention reaching 60 %. These results are interesting for Ge anodic materials prepared by solution-phase chemistry in so-far and could be greatly improved by optimization of the electrode preparation.

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