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SYNTHESIS OF CERIUM OXIDE NANOPARTICLES USING *VITEX* EXTRACT

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The aim of this work was to study composition and antioxidant/reducing properties of *Vitex cannabifolia* leaves extract and to characterize it as a possible active agent for green synthesis of cerium oxide nanoparticles (CeO₂-NPs). The aim of the study was also to prepare CeO₂-NPs and to investigate the particles sizes, texture and morphology. Antioxidant/reducing properties of *Vitex cannabifolia* leaves extract were studied using Folin-Chiocalteu and 2,2-diphenyl-1-picrylhydrazyl (DPPH) tests, composition of the extract was explored by means of laser desorption/ionization time-of-flight mass spectrometry method. The extract was found to possess very high antioxidant/reducing capability, showing fast reduction of DPPH radicals even at 100-fold dilution. The main components of the extract were phenolic acids, flavonoids and terpenes; all these compounds are known to be active reducing and/or stabilizing agents in green synthesis of various nanoparticles. Using the extract, CeO₂-NPs were prepared by means of the procedure that included the reduction of cerium(IV) ammonium nitrate by extract components followed by annealing the precipitate at 600 °C under in air conditions. The particles synthesized were characterized by means of scanning electron microscopy, X-ray diffraction and nitrogen adsorption methods. According to X-ray diffraction and electron microscopy data, CeO₂-NPs had crystalline structure, spherical form and fairly uniform particles size distribution; surface area of the particles was estimated from nitrogen adsorption isotherms as about 30 m² per 1 g. The average crystallite size determined from X-ray line broadening data was about 17 nm, the average particles diameter derived from surface area data was up to ~30 nm. Thus, one can conclude that the particles synthesized are small enough to be suitable for biomedical applications while *Vitex cannabifolia* extract is the effective reagent for green synthesis of cerium oxide nanoparticles.

Keywords: *Vitex cannabifolia* leaves extract, green synthesis, cerium oxide nanoparticles, antioxidant/reducing properties, LDI MS, SEM, X-ray diffraction

INTRODUCTION

The potential application of cerium oxide nanoparticles (CeO₂-NPs) in biological and medical fields is very promising [1–3]. CeO₂-NPs have shown protective effects on human health by reducing ischemic brain damage, inhibiting the progression of some blind diseases, diabetes, Alzheimer's disease, atherosclerosis. Also cerium nanoparticles exhibit antioxidant, antitumor, antibacterial and antiviral properties [3]. They are used as effective UV filters in sunscreen cosmetics, too [4].

Among the methods for synthesis of CeO₂-NPs for biomedical applications, green synthesis using plant extracts is of especial interest [1, 5]. Plant extracts can perform both the function of reducing and stabilizing agents that allows ones to use no toxic organic solvent and reagent and to carry out the reaction under mild synthesis conditions. Thus, synthesized nanoparticles do not contain toxic products and surfactants and can be more stable than the particles produced by traditional methods [1].

Recently, green synthesis of CeO₂-NPs was reported using different plants, such as *Aloe vera*, *Gloriosa superba*, *Acalypha indica* [5–7]. However, the synthesized nanoparticles were generally so large in size that, according to literature, they were not appropriate for biomedical applications [2, 8]. Our previous studies have shown that the *Vitex* genus plant possesses high antioxidant/reducing properties and has a great potential in synthesis of silver nanoparticles. The active ingredients of *Vitex* genus leaves extracts are phenols, terpenoids and monosaccharides; these compounds are well known as reducing agents and are capable to reduce metal salts and to convert them to nanoparticles.

The aim of the work was to study *Vitex cannabifolia* leaves extract as a possible active agent for CeO₂-NPs preparation, to prepare CeO₂-NPs using the extract and to characterize the extract and the particles synthesized.

EXPERIMENTAL

Vitex cannabifolia plants were used for green synthesis of CeO₂-NPs. The biologically

active substances were extracted from the leaves with 70 % ethanol solution according to the procedure described in [9]. 100 ml of 70 % ethanol was added to 1 g of finely chopped leaves, after that the mixtures were placed into steam bath for 30 min. After cooling, the extract was adjusted to the initial volume and filtered.

The total concentration of phenolic antioxidant/reducing agents in the extracts was evaluated by using the Folin–Ciocalteu method. To measure the total phenol index [10], 11.5 ml of water, 5 ml of 20 % sodium carbonate solution, 1.25 ml of Folin-Ciocalteu reagent (Merck) and 6.25 ml of water were consecutively added to 1 ml of extract. The solution was stirred for 30 min, then the absorbance at 750 nm was measured, and the total phenol index was calculated in accordance with the protocol [10]. By comparison of the obtained values of phenol index of the extracts with the corresponding data for ascorbic acid (total phenol index for the 0.5 mM solution of ascorbic acid was equal to 1 [11]), the equivalent concentration of the antioxidant in solutions was estimated.

To determine the total content of flavonoids, we used the method based on the capability of the compounds to form a coloured complex with aluminium chloride. 1 ml of extract, prepared as described above, was placed in volumetric flask, 5 ml of 2 % solution of AlCl_3 (Sigma-Aldrich) in 95 % ethanol was added into the glass, and then 95 % ethanol was added to reach a volume of 25 ml. The mixture was stirred for 30 min and the optical density of the solution at 410 nm was measured. To prepare a blank solution, 0.1 ml of concentrated acetic acid was added to 1 ml of extract, followed by the dilution of the mixture by 95 % ethanol to the total volume of 25 ml. To prepare reference samples, the rutin (Sigma-Aldrich) solutions of different concentrations were prepared, using 95 % ethanol as a solvent [12]. Total content of flavonoids was expressed as the equivalent weight of rutin per 1 g of dried leaves.

Antioxidant/reducing activity of *Vitex* extract was evaluated using 2,2-diphenyl-1-picrylhydrazyl (DPPH) radical scavenging method [13]. 1 ml of original was placed into glass and 2 ml of 70 % ethanol and 2 ml of 0.15 mM DPPH solution were consecutively added to the glass. The solution was shaken at 25 °C for 5–120 min. The change in concentration of stable radicals in reaction mixture during the reaction was determined from

the change in absorption at the maximum of 520 nm as compared to absorption value for control solution. To prepare control solution, 3 ml of 70 % ethanol was mixed with 2 ml of 0.15 mM DPPH solution.

Qualitative analysis of extracts composition was performed by means of laser desorption/ionization time-of-flight mass spectrometry (LDI MS). Mass spectra were recorded in positive and negative-ion extraction mode on Autoflex II mass spectrometer (Bruker Daltonics Inc., Germany) equipped with a nitrogen laser (337 nm). The samples were ionized in the pulse mode: pulse length 3 ns, frequency 20 Hz; maximum energy 65 mJ. Spectra were recorded in the reflection mode using a delayed extraction of 20 ns and accelerating voltage 20 kV. The resulting mass spectra were the sum of 100 individual spectra.

To produce CeO_2 -NPs, 100 ml of 0.01 M ammonium cerium(IV) nitrate solution was added to 10 ml of plant extract. The reaction mixture was stirred for 3 h at 80 °C, cooled to room temperature and stored for 2 days. The precipitate was powdered and annealed at 600 °C for 2 h.

XRD patterns of the nanoparticles were obtained at a DRON-4-07 diffractometer (Firm “Burevestnik”, Russia) with filtered CuK_α radiation in geometry of Bregg-Brentano in 2θ range of 10–80°. Phase identification was performed using the database of JCPDS. The average size of crystallites was calculated from X-ray data according to Scherrer formula.

Scanning electron microscopic (SEM) images were recorded using a Gemini 500 electron microscope (Zeiss, Germany). The samples powder was grind on a standard holder that was pre-coated with a double-sided conductive graphite tape. The remaining powder was blown out from the tape in a stream of dry nitrogen. An aperture of 20 μm was used with an acceleration voltage of 1–2 kV. Secondary electron detector was mainly used, which allowed omitting fast surface charging.

UV/Vis spectra of plant extracts and reaction mixtures were recorded on a Perkin Elmer Lambda 35 UV/Vis double beam Spectrophotometer at 25 °C in the wavelength range of 200–800 nm. Scanning speed was 480 nm/min, pathlength of cuvette (Perkin-Elmer) was 10 mm.

Textural characteristics of CeO_2 -NPs were derived from 77 K nitrogen adsorption isotherms using several standard methods (BET, t -plot,

DH, DR methods). Isotherms were recorded and processed on an AUTOSORB-6B (Quantachrome, USA).

RESULTS AND DISCUSSIONS

The data characterizing the antioxidant/reducing properties of *Vitex cannabifolia* leaves extract (the values of the total phenolic index and the equivalent concentration of ascorbic acid) as well as the data on the flavonoid content in the plant leaves are

presented in Table 1. As one can see from the data, the extract contains high amount of phenols and flavonoids. The extract also possesses high antiradical/reducing properties, as it is shown in the reaction of strongly diluted extract with DPPH radicals (Fig. 1).

The more detailed study of the extract composition was carried out using LDI MS method. The appropriate data are given in Fig. 2 and in Table 2.

Table 1. Content of phenolic antioxidants in the *Vitex cannabifolia* leaves and extract

Sample	Total Phenolic index	Equivalent concentration of ascorbic acid in extract, mmol/L	Content of flavonoids per 1 g of dried leaves, %
Leaves of <i>Vitex cannabifolia</i>	15.0	7.5	1.2

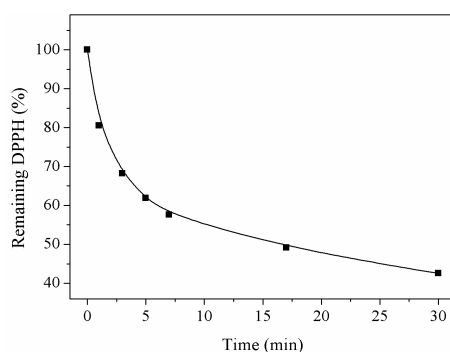


Fig. 1. Inhibition of DPPH radical in reaction with *Vitex cannabifolia* extract. Before measurement the extract was diluted by 100 times

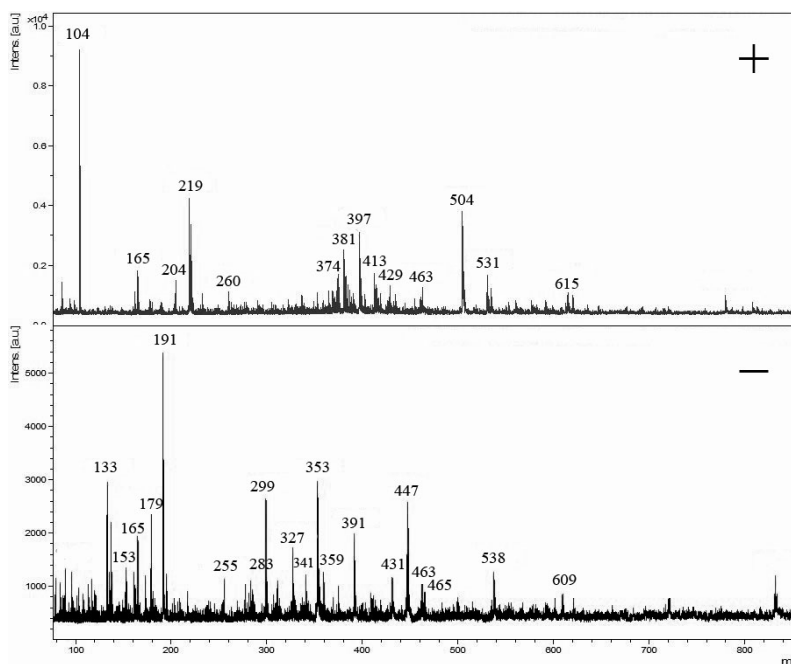


Fig. 2. Fragments of negative (-) and positive (+) ion mode of LDI mass spectra of *Vitex cannabifolia* extract

Table 2. The most abundant ions in the negative and positive modes of mass spectra of *Vitex* extracts

<i>m/z</i>	Compound, ion	<i>m/z</i>	Compound, ion
133 ⁻	Cinnamyl alcohol, [M-H] ⁻	374 ⁺	Casticin, M ⁺
137 ⁻	Hydroxybenzoic acid, [M-H] ⁻	381 ⁺	Lactose / Cellobiose / Maltose / Sucrose, [M+K] ⁺
153 ⁻	Terpinenol, [M-H] ⁻ / Dihydroxybenzoic acid, [M-H] ⁻	397 ⁺	Casticin, [M+Na] ⁺ / Stigmasterol glucoside, [M+1-глюкоза] ⁺
165 ⁺	Eugenol, [M+H] ⁺ / Hydroxycinnamic acid, [M+H] ⁺	413 ⁺	Stigmasterol, M ⁺
179 ⁻	Glucose/Galactose/Fructose/Caffeic acid, [M-H] ⁻	416 ⁺	Methylenedioxy-pentamethoxyflavone, M ⁺
191 ⁻	Citric acid / Scopoletin, [M-H] ⁻	429 ⁺	Vitetrifolin D, [M+Na] ⁺ / Friedelin, [M+H] ⁺
219 ⁺	Caryophyllene oxide, [M-H] ⁺	431 ⁻	Vitexin / Isovitexin, [M-H] ⁻
220 ⁺	Spathulenol, M ⁺	447 ⁻	Orientin / Isoorientin / Cynaroside, [M-H] ⁻
255 ⁻	Hexadecanoic acid, [M-H] ⁻	463 ⁺	Luteolin-glucuronide, [M+H] ⁺
260 ⁺	Cadinol, [M-H+K] ⁺	463 ⁻	Isoquercitrin, [M-H] ⁻
283 ⁻	6-methyl hexadecanoic acid methyl ester / Stearic acid, [M-H] ⁻	465 ⁻	Agnuside, [M-H] ⁻
299 ⁻	Trihydroxydimethoxyflavone, [M-H] ⁻	504 ⁺	Cannabifolin B, [M+H] ⁺
341 ⁻	Myzodendrone / Dicaffeic acid / Lactose / Cellobiose / Maltose / Sucrose, [M-H] ⁻	538 ⁻	Caffeoylmussaenosidic acid, [M-H] ⁻
353 ⁻	Chlorogenic acid, [M-H] ⁻	609 ⁻	Caffeoylisoorientin / Hesperidine / Rutin, [M-H] ⁻
359 ⁻	Chrysosplenol / Centauredin, [M-H] ⁻		

Fig. 2 shows the most abundant ions in the negative and positive modes of mass spectra while Table 2 gives possible interpretation of all the signals identified. For signals attribution, the literature data [14–19] were used.

As one can conclude from the data, the most intensive signals in the mass spectra are due to compounds related to phenolic antioxidants. Indeed, 15 of the 25 most pronounced peaks can be referred to phenolic acids (dihydroxybenzoic acid, hydroxycinnamic acid, caffeic acid, chlorogenic acid) and flavonoids (trihydroxydimethoxyflavone, chrysosplenol/centauredin, casticin, methylenedioxy-pentamethoxyflavone, vitexin/isovitexin, orientin/isoorientin/cynaroside, luteolin-glucuronide, isoquercitrin, caffeoylisoorientin/hesperidine/rutin). Extract also seems to contain citric acid, mono- and disaccharides (glucose/galactose/fructose and lactose/cellobiose/maltose/sucrose), terpenes and terpenoids (terpinenol, cannabifolin B, caryophyllene oxide, spathulenol, cadinol, vitetrifolin D, agnuside, caffeoylmussaenosidic acid). All these compounds are known to be active reducing and/or

stabilizing agents in green synthesis of various nanoparticles.

Fig. 3 shows the SEM image of CeO₂-NPs particles synthesized using *Vitex* extract. The particles seem to have a rather spherical form and fairly uniform particle size distribution. Fig. 4 gives the X-ray diffraction curve; the data confirm crystalline structure of the particles. The average crystallite size determined from X-ray line broadening data is ~17 nm.

Table 3 gives the data on the texture of CeO₂-NPs derived from nitrogen adsorption experiment. The results show that the sample is non-porous powder; the estimated total pore volume and volume of micropores are small enough to be referred to the slits and cavities between the particles.

Average particles sizes may be estimated also from the surface area data. As one can see from the data of Table 3, surface area of the CeO₂-NPs powder is about 26–31 m²/g. Taking into account the bulk density of CeO₂ (7.215 g/cm³), one can estimate the average diameter of the particles as $D = (26 \div 30)$ nm. This value is approximately 1.7 times higher

than the average diameter of crystallites (17 nm), that is the volume of the particle is approximately 5 times higher than the average volume of the crystallite. Perhaps, each particle

consists of 5 or more smaller crystallites, with the slits between the crystallites and the particles of various sizes being the micropores and larger pores.

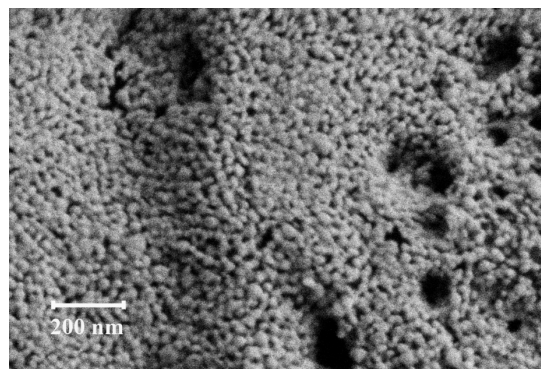


Fig. 3. SEM image of CeO₂-NPs

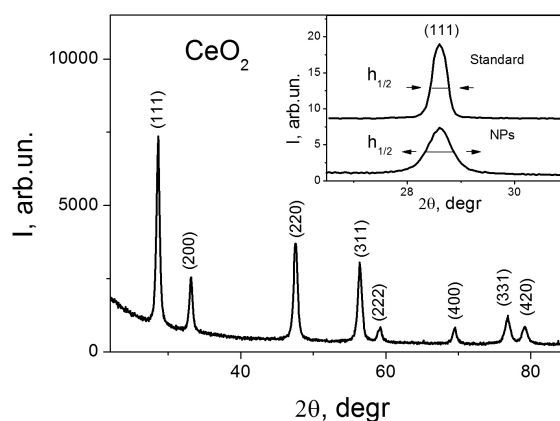


Fig. 4. XRD pattern of CeO₂-NPs

Table 3. Surface area and pore volume for CeO₂-NPs

Sample	BET	<i>t</i> -method	DH method (adsorption/desorption)		DR method	
	<i>S</i> _{BET} , m ² /g	<i>S</i> _{ext} , m ² /g	<i>S</i> _{total} , m ² /g	<i>V</i> _{total} , cm ³ /g	<i>S</i> _{mi} , m ² /g	<i>V</i> _{mi} , cm ³ /g
CeO ₂ -NPs	30	30	26/31	0.012/0.012	13	0.005

*S*_{BET} – surface area by BET method, *S*_{ext} – external surface area by *t*- method,

*S*_{total} and *V*_{total} – cumulative surface area and pore volume by DH method,

*S*_{mi} and *V*_{mi} – micropore surface area and micropore volume by DR method

CONCLUSIONS

The results obtained show that leaves extract of *Vitex cannabifolia* is very effective antioxidant/reducing agent, which can be used for green synthesis of crystalline CeO₂ nanoparticles. The extract contains high amount of phenols and terpenoids; these compounds can act both as reducing agents and as stabilizing

agents preventing particles growth during synthesis. Produced CeO₂ nanoparticles were shown to have crystalline structure, spherical form and fairly uniform particles size distribution. The sizes of the particles do not exceed ~30 nm that make them suitable for biomedical applications.

Синтез наночастинок оксиду церію з використанням *Vitex* екстракту

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Метою дослідження було вивчення складу та антиоксидантних/відновлювальних властивостей екстракту з листя *Vitex cinnabifolia* і характеристика екстракту як можливого активного агента для зеленого синтезу наночастинок оксиду церію ($\text{CeO}_2\text{-НЧ}$). Метою роботи також було одержання $\text{CeO}_2\text{-НЧ}$ і дослідження розмірів, текстури та морфології синтезованих наночастинок. Антиоксидантні/відновлювальні властивості екстракту з листя *Vitex cinnabifolia* було досліджено за допомогою методу Фоліна-Чокальтеу та 2,2-дифеніл-1-пікрилгідразилу (ДФПГ)-тесту, склад екстракту вивчено з використанням методу лазерної десорбційної/іонізаційної часопробної мас-спектрометрії. Знайдено, що екстракт має дуже високі антиоксидантні/відновлювальні властивості, демонструючи швидке відновлення радикалів ДФПГ навіть при 100-кратному розведенні. Основними компонентами екстракту були фенольні кислоти, флавоноїди та терпени; як добре відомо, ці сполуки є активними відновлювальними та/або стабілізуючими агентами в зеленому синтезі різних наночастинок. З використанням екстракту було одержано $\text{CeO}_2\text{-НЧ}$; процедура синтезу складалася з відновлення амонію церію(IV) нітрату компонентами екстракту та наступного відпалу осаду при 600 °С на повітрі. Синтезовані частинки було охарактеризовано методами скануючої електронної мікроскопії, ширококутового рентгенівського розсіювання та адсорбції азоту. Згідно даних електронної мікроскопії та рентгенофазового аналізу, $\text{CeO}_2\text{-НЧ}$ мали кристалічну структуру, сферичну форму і достатньо однорідний розподіл за розмірами; за даними експерименту по адсорбції азоту, питома поверхня наночастинок складала близько 30 м² на 1 г. Середній розмір кристалітів, визначений з даних щодо уширення рентгенівських ліній, становив ~17 нм, середній діаметр частинок, визначений з даних щодо площі їхньої поверхні, не перевищував ~30 нм. Таким чином, можна зробити висновок, що синтезовані частинки мають достатньо маленькі розміри і є придатними для використання в біології та медицині, а екстракт *Vitex cinnabifolia* є ефективним реагентом для зеленого синтезу наночастинок оксиду церію.

Ключові слова: екстракт із листя *Vitex cinnabifolia*, зелений синтез, наночастинок оксиду церію, антиоксидантні / відновлювальні властивості, ЛДІ МС, СЕМ, рентгенофазовий аналіз

Синтез наночастиц оксида церия с использованием *Vitex* экстракта

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Целью исследования было изучение состава и антиоксидантных/восстановительных свойств экстракта из листьев *Vitex cinnabifolia* и характеристика экстракта как возможного активного агента для зеленого синтеза наночастиц оксида церия ($\text{CeO}_2\text{-НЧ}$). Целью работы также было получение $\text{CeO}_2\text{-НЧ}$ и изучение размеров, текстуры и морфологии синтезированных наночастиц. Антиоксидантные/восстановительные свойства экстракта из листьев *Vitex cinnabifolia* были изучены с помощью метода Фоллина-Чокальтеу и 2,2-дифенил-1-пикрилгидразила (ДФПГ)-теста, состав экстракта исследован с помощью метода лазерной десорбционной/ионизационной времяпролетной масс-спектрометрии. Обнаружено, что экстракт обладает очень высокой антиоксидантной / восстанавливающей способностью, демонстрируя быстрое восстановление радикалов ДФПГ даже при 100-кратном разведении. Основными компонентами экстракта были фенольные кислоты, флавоноиды и терпены, которые, как известно, являются активными восстанавливающими и/или стабилизирующими агентами в зеленом синтезе различных наночастиц. С использованием экстракта были приготовлены $\text{CeO}_2\text{-НЧ}$; процедура синтеза состояла из восстановления нитрата аммония церия (IV) компонентами экстракта и последующего отжига осадка при 600 °С на воздухе. Синтезированные частицы были охарактеризованы методами сканирующей электронной микроскопии, широкоугольного рентгеновского рассеяния и адсорбции азота. По данным электронной микроскопии и рентгенофазового анализа, $\text{CeO}_2\text{-НЧ}$ имели

кристаллическую структуру, сферическую форму и достаточно однородное распределение частиц по размерам; по данным эксперимента по адсорбции азота, удельная поверхность частиц составляла около 30 м^2 на 1 г. Средний размер кристаллитов, определенный из данных по уширению рентгеновских линий, составил ~17 нм, средний диаметр частиц, определенный из данных о площади поверхности, не превышал ~30 нм. Таким образом, можно заключить, что синтезированные частицы достаточно малы, чтобы быть пригодными для применения в биологии и медицине, а экстракт является эффективным реагентом для зеленого синтеза наночастиц оксида церия.

Ключевые слова: экстракт из листьев *Vitex cannabifolia*, зеленый синтез, наночастицы оксида церия, антиоксидантные / восстанавливающие свойства, ЛДИМС, СЭМ, рентгенофазовый анализ

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