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Confining caesium in expanded natural Perlite

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Abstract

We present the potential use of expanded perlite, a metastable amorphous hydrated aluminum silicate, as a permanent medium for the long-term confinement of caesium. A simple loading by mixing an aqueous caesium nitrate solution and expanded perlite at 300 K followed by thermal annealing leads to 96% sintering. The formation of pollucite, CsAlSi2O6, a naturally occurring mineral phase, appears as the crystalline phase embedded in a glassy phase. Leaching tests on the resulting glass-ceramics reveal a very low Cs departure of 0.5 mg m−2 day−1. This simple method seems to be a good way to permanently confine caesium issued from recycled nuclear waste.

Keywords: Sintering; Leaching process; Nuclear applications

1. Introduction

The generation of large quantities of environmentally unfriendly nuclear waste, for example those with long-lived radioactivity, sets to humans a great problem for now and the future. Ways of separating and storing these materials are of constant debate and promote the search for efficient methods of finding containers for their safe storage for long-term periods spanning over thousand years. One of the nucleids under debate in the France is radioactive 137 caesium issued from nuclear fuel recycling process in the La Hage plant. Our interest has been in finding suitable host for the storage of caesium and its decayed products. The present paper reports the use of expanded perlite on a laboratory level for trapping natural caesium.

Borosilicate glass, glass-ceramics, caesium-loaded zeolites and pollucite have been shown to be potential candidates for storage. Pollucite, CsAlSi2O6, can accommodate up to 40 wt.% Cs into its structure thus producing a highly dense waste form. Furthermore, the measured solubility of pollucite is lower while the leakage of Cs from pollucite are three orders of magnitude less than those for silicate glasses.1 Therefore, various synthetic methods have been considered for pollucite using self-propagating high-temperature synthesis (SHS),2 reaction between gaseous Cs and fly ash,3 hydrothermal,4 arc melting technique,5 ion exchange in zeolites6 and crystallization from a glass.7 Moreover, pollucite is a naturally occurring mineral with a three-dimensional structure that contains and irreversibly traps more Cs.8 The resulting samples have been the subject of several characterizations in view of their possible application as waste form such as thermal expansion coefficient measurements,9 leaching tests,7 sinterability, effect of beta radiation effects10 and radiogenic transmutation.11 The possibility to obtain pollucite from zeolites8 stimulates the research of other aluminosilicate minerals. These results are very encouraging and have driven us to search for other related materials and here, we focus on a metastable amorphous hydrated aluminum silicate, perlite.

Perlite with typical average chemical analysis 71–75 wt.% SiO2, 12.5–18 wt.% Al2O3, 4–5 wt.% K2O, 1–4 wt.% Na2O and CaO, traces of metal oxides and 2–5 wt.% water12 is the naturally occurring mineral issued from rapid cooling of undersea igneous rocks. However, perlite is mainly used in its expanded form which is obtained by flame spreading of powered mineral. It results in an extremely light material looking like glass pearls with a large void in the core. Expanded perlite is fire resistant and acts as an excellent thermal and acoustical insulator. The open structure of expanded perlite can also be exploited for the insertion of waste mineral cations (for example: Cs) from aqueous solutions. Due to the strong excess of silicon compared to aluminum, the formation of a glass ceramics is expected with pollucite as the crystalline phase. The localization of Cs in pollucite will potentially decrease its leakage.5 Finally, expanded

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perlite is a ready-to-use material with a very reasonable cost. All these considerations stimulate the study of the use of perlite for the confinement of Cs. Our works were carried out on natural caesium in order to estimate the stability of perlite matrix upon water leaching who can occurs in deep underground deposits.

2. Experimental

2.1. Synthesis

Expanded perlite was produced by Thermal Ceramics de France who produce flame resistant panel. An aqueous caesium nitrate, CsNO₃ (Aldrich, 99%), solution was dropped in expanded perlite in a 5:1 weight proportion in order fully cover the apparent volume of powder and homogeneously disperse caesium. Different amounts of CsNO₃ were dissolved in a constant volume of water in order to achieve various Cs₂O/perlite weight ratios between 5 and 50 wt.%. The resulting samples were then dried at 423 K to remove water. In order to check the influence of the porosity, similar samples have been elaborated starting from thoroughly ground powders of the expanded perlite. Dense pellets for the leaching tests have been prepared using different techniques: (1) pre-calcination at 1373 K for 1 h to reduce the sample volume, grinding, shaping into pellets and sintering at 1293 K for 1–3 h, (2) starting from ground perlite, followed by direct caesium loading and direct sintering at 1293 K in a crucible with a small load, and (3) sintering using the flash technique. Samples were sintered at 1123 K for 5 min in a Spark Plasma Sintering (SPS) furnace, Dr. Sinter 2080 (Sumitomo Coal Mining). Before sintering, expanded perlite powders were directly loaded into cylindrical graphite die with inner diameter of 8 or 20 mm and then put into the sintering chamber that was evacuated to 3 Pa. The control of the heating rate was made by a computer program system and the temperature was recorded by a thermocouple fixed on a small hole at the external surface of the graphite die. A pressure of 25 MPa was applied by manual operation and not released until the beginning of cooling stage. A constant heating rate of 100 K min⁻¹ was applied during each sintering cycle. Contrary to borosilicate glass used to dissolve caesium and other nuclide (R7T7 COGEMA) we also try to obtain a castable glass. But, moulding from the liquid was not possible due to its high viscosity even at temperatures as high as 1823 K.

2.2. Characterizations

Chemical analyses were performed by ICP-MS technique with a Plasma Quad ICP/MS instrument equipped with Meinhard nebulizer. Operating conditions were single ion monitoring at m/z 133; plasma gas: 14 l min⁻¹, auxiliary gas: 1 l min⁻¹, nebulizer gas: 0.75 l min⁻¹. Cs solutions used for calibration were daily prepared on a mass basis by non-consecutive dilutions of a CPI stock certified solution (1 g l⁻¹ in Cs) with 2.5% nitric acid (Suprapur Merck). The leaching solutions were diluted 10 times with 2.5% nitric acid (Suprapur Merck) before analysis. TG-DTA experiments were performed on a TA instrument under air at a 5 °C min⁻¹ heating rate. The leaching tests have been performed either in static or dynamic mode (Soxhlet apparatus) at 353 K with various surface/volume (S/V) ratios. X-ray diffraction patterns were collected on a Siemens D5000 diffractometer using monochromatized Cu Kα₁ radiation (λ = 1.5406 Å). SEM observations have been done on a Jeol 6700F apparatus.

3. Results and discussion

3.1. Chemical analysis

In order to evaluate the reproducibility of the composition of perlite, chemical analyses have been performed on the five samples received from the Sardinia quarry by Thermal Ceramics at 4 months intervals. As shown in Table 1, the chemical composition is quite consistent, the largest variation concerns Al₂O₃. Besides the other elements usually found in perlite, Al, Na, K and Ca, we also note the presence of a significant amount of iron. The whole content differs from 100% and it has been related to the presence of water and other minor elements.

The caesium content of samples heat treated at different temperatures between 873 and 1823 K indicates no Cs loss upon melting for all loading between 5 and 50 Cs₂O wt.%. This means that Cs is perfectly trapped in the matrix.

3.2. Thermal analysis

A typical TG-DTA trace is shown in Fig. 1 for a 27.7 wt.% CsNO₃ (20 wt.% Cs₂O) sample. At low temperature, a first weight loss, around 2–3 wt.%, has been attributed to adsorbed water. The main weight loss takes place between 713 and 913 K and corresponds to the nitrate decomposition. Surprisingly, an increase of the decomposition temperature is observed with increasing initial CsNO₃ content. Moreover, the weight loss deviates from the value expected from the decomposition of CsNO₃ into Cs₂O. The observed value fits with the formation of a mixture of Cs₂O₃ and Cs₂O₄. At low CsNO₃ contents, the weight loss corresponds to the formation of Cs₂O₄ whereas the value expected for the formation of Cs₂O is observed for the 50 wt.% Cs₂O sample. These results are perfectly reproducible. The fact that the nitrate is dispersed inside the pores may be a reason for the unexpected thermal behaviour.

![Fig. 1. TG-DTA traces of a 20 wt.% Cs₂O sample.](image-url)
3.3. X-ray diffraction

X-ray diffraction patterns, recorded on powders annealed at different temperatures and containing different amounts of Cs, reveal the crystallization of pollucite, CsAlSi$_2$O$_6$ (Fig. 2). The pattern is usually characterised by a high background and a broad hump around 2θ = 25°; the first is related to the presence of an amorphous phase and the second from poorly crystallised silicate. The resulting diffraction pattern has been compared with the corresponding JCPDS file No. 88–0055. No significant shift of the lines appears. This means that the other alkali cations in the original perlite have not been inserted (or at very minor levels) in the pollucite network. The presence of pollucite is evidenced by sharp peaks. Moreover, the presence of pollucite also depends on the Cs content and on the annealing time and temperature. To promote crystallization of pollucite, longer annealing times are needed with decreasing Cs content. For example, at 1373 K, 2 and 12 h are needed for 50 and 25 wt.% Cs$_2$O, respectively. Furthermore, the effect of grinding perlite yields a decrease of the crystallization temperature; for example for the 50 wt.% sample it is reduced to 973 K and requires only 2 h.

3.4. Sintering

The sintering temperature of 1293 K has been chosen as it yields a good densification without deformation of the sample. For these experiments the powder was annealed at 1373 K and reground before to be shaped by pressing in 25 mm × 3 mm round pellets. At higher temperatures, the viscous flow promotes the deformation. After annealing, the colour of the samples turn from a dirty white to brown, in relation with the oxidation of the small amount of Fe$^{2+}$ to Fe$^{3+}$. In view of a possible application, and in order to explore ways of shaping and sintering samples loaded with radionuclide solution, we have considered the possibility of a one-step sintering process, particularly avoiding the pre-heating steps. As described in Section 2 the grinded expanded perlite is loaded with caesium solution in an inconel crucible and a very low load is applied over the powder. Drying, decomposition of nitrate and sintering take place successfully in the same heat treatment at 1293 K. However, the for 50 wt.% caesium load the crystallization of pollucite at low temperature (973 K) inhibits full densification. Another way to promote full densification but at low temperature and therefore reduce the size of pollucite crystal was the flash sintering technique. It leads to a full sintered sample at 1073 K. The densification of both samples, as received and grounded powder, exhibit densities better than 96% of the flash values.

3.5. Leaching experiments

Preliminary experiments performed on powdered samples yield a removal rate of Cs around a few micro grams per m$^{-2}$ day$^{-1}$ at room temperature. Therefore, other experiments either on powdered or sintered pellets have been performed. The influence of the S/V ratio has been tested on a powder, 10 wt.% Cs$_2$O annealed at 1293 K for 1 h, with a specific surface area around 1 m$^2$ g$^{-1}$, under static mode at 353 K. Different quantities of powders were inserted in a closed container with a constant volume of 80 ml water in order to achieve S/V ratios of 12.5, 125 and 250 cm$^{-1}$. Ten milliliters of aliquots were then taken at regular intervals and replaced by the same volume of deionized water. As shown in Fig. 3, the removal rate of Cs is always very slow with values of 2 to 7 × 10$^{-9}$ g m$^{-2}$ day$^{-1}$ after 21 days or more. Moreover, the shape evolution with time is comparable to what is expected for a mechanism involving the formation of a surface gel. The values for the highest S/V ratio are significantly lower and are in agreement with an improved efficiency of the surface gel. A sintered pellet with 10 wt.% Cs$_2$O has been cut in pieces and submitted to the same treatment at 353 K. One piece has been taken at regular intervals for SEM. Observations at regular intervals up to 64 days contact with water (Fig. 4) show the presence of white spots at the surface, whose size and shape are independent of time. EDX analysis of the Cs content at the surface of the samples reveals a constant value close to the expected 10%. No apparent degradation of the glass matrix is observed.
Leaching experiments have also been performed on pellets either in static or dynamic mode. In static mode, for a pellet with 12 wt.% Cs$_2$O, $S/V = 0.025$ cm$^{-1}$ (only the geometric surface of the samples has been considered for working out the rate) at 353 K the removal rate of Cs rapidly decreases and reaches a constant value after 40 days. The estimated removal rate is $1.5$ mg m$^{-2}$ day$^{-1}$ on the 7th day and $0.6$ mg m$^{-2}$ day$^{-1}$ after 42 days. In dynamic mode, perlite pellets from both sintering processes (virgin or ground) were used and the experiments were performed with 450 ml ion-exchanged water with the sample surfaces being around 7 cm$^2$. Both samples give very similar results as shown in Fig. 5. Both processes resulted in very slow removal rates for Cs, with a time evolution in agreement with the surface gel model. The pH monitored on a daily basis reveals no changes from a value close to 6 for both samples and for the two processes.

4. Conclusions

The results give great promises for the use of perlite as a potential matrix for confining natural caesium. We, therefore, propose it as a possible candidate for radioactive waste matrix. Its high stability, low removal rate of caesium in water due to two facts: high stability of the glass matrix against water corrosion, crystallisation of pollucite as the most stable crystalline form of caesium compounds. Moreover, its abundance, lost cost, the non-toxicity of the mineral makes it ideal for the environment and in favourable position as confinement matrix. The irreversible transformation to pollutte at high temperature and high densification achieved by sintering technique are additional bonuses. Current work in our group has extended the range of elements (Rb, Sr, Ba, Ni, Cd, Nd and Eu and I) that can be confined in perlite. Nevertheless, the effect irradiation by confining radioactive cation has to be studied.

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