

Hindawi Publishing Corporation
Advances in Materials Science and Engineering
Volume 2010, Article ID 683429, 8 pages
doi:10.1155/2010/683429

Research Article

Parameters Expediting the Thermal Conversion of Ba-Exchanged Zeolite A to Monoclinic Celsian

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Received 27 July 2010; Accepted 28 October 2010

Academic Editor: David Cann

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Four samples of Ba-exchanged zeolite A, bearing small residual amounts of Na (0.27, 0.43, 0.58, and 0.74 meq/g), were thermally treated in the temperature range 200–1500°C for times up to 28 hours. The same samples were pressed at 30 and 60 MPa to form cylindrical pellets which were thermally treated at 1300°C for 5 hours. All materials were characterized by room temperature XRD. The sequence of thermal transformations that Ba-exchanged zeolite A undergoes (zeolite → amorphous phase → hexacelsian → monoclinic celsian) and the strong mineralizing action developed by Na are confirmed. Pressing the Ba-exchanged zeolite A powder-like samples to obtain cylindrical pellets is found to expedite the sluggish final phase transition hexacelsian → monoclinic celsian. The optimum residual Na content of Ba-exchanged zeolite A for transformation into monoclinic celsian is assessed to be between 0.27 and 0.43 meq/g.

1. Introduction

1.1. Uses of Monoclinic Celsian. Monoclinic celsian ($\text{BaAl}_2\text{Si}_2\text{O}_8$) is used as a refractory, high temperature electrical insulator, or as a substrate for integrated circuits because of its good thermal and electrical properties [1]. Innovative studies suggest the use of monoclinic celsian in aeronautic and aerospace field, as matrix material for reinforcement with continuous ceramic fibers of high temperature structural composites to be used in hot sections of turbine engines [1–11] and in the field of radomes [12, 13].

1.2. Problems in Preparation. The synthesis of monoclinic celsian gives rise to considerable problems [14]. On one hand, it requires high temperatures and/or long reaction times, which give rise to high process costs. On the other hand, the first polymorph to nucleate is hexagonal celsian (hexacelsian) although it is thermodynamically stable at temperatures higher than 1590°C [15]. This behaviour was ascribed to the simpler crystal structure of the high symmetry modification presenting a lower kinetic barrier to nucleation [16].

The early crystallisation of hexacelsian appears to be a serious drawback for two reasons. On one hand, hexacelsian cannot be used as a refractory because it, at 300°C, undergoes a reversible transformation into an orthorhombic form accompanied by a detrimental ($\geq 3\%$) volume change [15]. On the other hand, the transformation of hexacelsian to monoclinic celsian occurs only after prolonged heating (more than 20 hours) at temperatures higher than 1500°C in the presence of mineralizers and monoclinic seed particles [17].

A long-term study [14, 18–25] concerning the thermal transformation of Ba-exchanged zeolites into monoclinic celsian was undertaken. This study started from the work of Subramanian and coworkers [26–29] who crystallised alkaline-earth and alkaline framework aluminosilicates, which belong to the class of the feldspars, by thermal treatment of cation exchanged zeolites at temperatures slightly higher than 1000°C for some hours. The technique proposed by Subramanian and coworkers [26–29] appeared to be an interesting way of avoiding the drawbacks arising in the traditional syntheses of monoclinic celsian [14]. Actually very interesting results were obtained in the study concerning

the thermal transformation of Ba-exchanged zeolites into monoclinic celsian [14, 18–25]. In particular, it was found that, upon a thermal treatment at temperatures higher than 1000°C, Ba-exchanged zeolite A, which after dehydration exactly has the stoichiometric composition of celsian, gave rise to the following sequence of transformations: zeolite → amorphous phase → hexacelsian → monoclinic celsian. Moreover, it was found that thermal treatments at 1100°C for 6 hours [14] and 1300°C for 22 hours [18] were sufficient to transform samples of Ba-exchanged zeolite A, containing 0.58 meq/g and 0.20 meq/g Na (residual amount), respectively, into fully monoclinic celsian. These data reveal that Ba-exchanged zeolite A does appear to be a very reactive precursor inasmuch as the full conversion into monoclinic celsian was obtained at far milder conditions than those reported in the literature (heating for more than 20 hours at temperatures higher than 1500°C) [14].

Nevertheless, this study revealed that the temperatures and duration times of thermal treatments necessary to attain the full conversion of samples of Ba-exchanged zeolite A into monoclinic celsian are strongly dependent on the residual Na content of various zeolite A samples.

1.3. Objectives of this Work. This investigation aims at ascertaining to what extent this parameter affects the thermal transformations that Ba-exchanged zeolite A undergoes. To fulfil this goal, four zeolite A samples bearing 0.27, 0.43, 0.58, and 0.74 meq/g Na (residual content) were prepared by cation exchange and their thermal transformations were investigated by room temperature X-ray diffraction and differential thermal analysis. Such Na residual contents were selected on the basis of the following considerations. In a previous paper, [18] a sample of Ba-exchanged zeolite A with 0.20 meq/g Na residual content was prepared. It was found very difficult to further reduce such Na residual content as 0.12 meq/g Na were shown to be related to NaAlO₂ occluded in its β-cages (sodalite cages), similar to feldspathoid behavior [30]. It appeared very likely that Na arising from the occlusion of NaAlO₂ in the β-cages could not be practically removed and, thus exchangeable Na (0.08 meq/g) still present after the last Ba-exchange was between 1 and 2% of the Na originally present in the zeolite framework (the real initial Na content of zeolite A was 5.60 meq/g [18]). Moreover, it was reported that, for this sample, 22-, 15- and 22-hour thermal treatments at 1300, 1400, and 1500°C, respectively, were sufficient to attain the full conversion of hexacelsian into monoclinic celsian. Thus 0.27 meq/g was selected as it is an Na residual amount slightly higher than the value considered as a limit (0.20 meq/g), for which temperature and time of the thermal treatments necessary to attain the full conversion of hexacelsian to monoclinic are well known. The remaining Na residual amounts (0.43, 0.58 and 0.74 meq/g) were selected as they steadily increase starting from 0.27 meq/g so as to cover the whole range of Na residual amount which could be accepted as an impurity in a celsian based ceramic manufactured by this technique.

This work also aims at attaining a second goal. Actually the thermal transformation of compacts obtained by pressing

powder-like samples of Ba-exchanged zeolite A appears very interesting inasmuch as it exhibits potential for practical applications in the manufacture of celsian based ceramics. Thus the role played by the forming pressure of compacts on the thermal transformation of Ba-exchanged zeolite A is investigated by cold pressing the various powder-like samples of zeolite A at 30 and 60 MPa. These compacts were subjected to the same thermal treatments as the powder-like samples and then investigated by room temperature X-ray diffraction in order to compare these results.

2. Experimental

2.1. Ion Exchange with Ba²⁺. Carlo Erba reagent grade synthetic zeolite 4A (Na₁₂Al₁₂Si₁₂O₄₈·27H₂O) was subjected to various cation exchange operations in order to obtain four samples of Ba-exchanged zeolite A bearing different residual amounts of Na. The samples which will be identified from this point onwards as 1, 2, 3, 4 were obtained according to the procedures hereafter described.

Sample 1. Original zeolite 4A in its Na form was contacted overnight with a warm (60–70°C) [Ba²⁺] = 0.2 N solution at a weight solid/liquid ratio (S/L) = 1/20. The solid was separated from the liquid through filtration and again contacted with fresh exchange solution for a total of nine times.

Sample 2. Original zeolite 4A in its Na form was contacted overnight with a warm (60–70°C) [Ba²⁺] = 0.2 N solution at a weight solid/liquid ratio (S/L) = 1/20. The solid was separated from the liquid through filtration and again contacted with fresh exchange solution for a total of five times.

Sample 3. Original zeolite 4A in its Na form was contacted overnight with a warm (60–70°C) [Ba²⁺] = 0.5 N solution at a weight solid/liquid ratio (S/L) = 1/20.

Sample 4. Original zeolite 4A in its Na form was contacted overnight with a warm (60–70°C) [Ba²⁺] = 0.2 N solution at a weight solid/liquid ratio (S/L) = 1/20. The solid was separated from the liquid through filtration and again contacted with fresh exchange solution for a total of two times.

The exchange solutions (initial pH about 6.3) were prepared using doubly distilled water and Carlo Erba reagent-grade Ba(NO₃)₂ (purity 99.5%). The pH of the suspensions zeolite A- Ba²⁺ solutions ranged between 6.9 and 7.0.

The residual Na content of the different zeolite samples intermediate and total was determined by the following procedure. The zeolite was chemically dissolved in a hydrofluoric and perchloric acid solution and its Na⁺ concentration was determined by atomic absorption spectrophotometry (AAS), using a Perkin-Elmer AAnalyst 100 apparatus. The consistency of concentration measurements related to cation

TABLE 1: Thermal treatments done to all four Ba-exchanged zeolite A samples and the figures in which the corresponding XRD patterns are shown.

Temp (°C)	Thermal treatment ^a (h)
200	2 (Figure 1(a))
500	0 (Figure 1(b))
800	28 (Figure 1(c))
1000	5 (Figure 2)
1100	5, 10
1200	5
1300	5 (Figure 3), 10, 15
1400	5
1500	5

^aThe samples were heated at a rate of 10°C/min up to the reported temperature, kept at that temperature for the reported time, and allowed to cool in air.

TABLE 2: Composition of the products.

Sample	Na (meq/g)	x_{Na} ^a	Na ₂ O w.% ^b	Chemical formula ^{b,c}
1	0.27	0.05	0.84	Ba _{0.950} Na _{0.100} Al ₂ Si ₂ O ₈
2	0.43	0.08	1.33	Ba _{0.921} Na _{0.158} Al ₂ Si ₂ O ₈
3	0.58	0.10	1.80	Ba _{0.894} Na _{0.212} Al ₂ Si ₂ O ₈
4	0.74	0.13	2.29	Ba _{0.866} Na _{0.268} Al ₂ Si ₂ O ₈

^a equivalent fraction.

^b after thermal treatment at 950°C for 2 hours.

^c chemical formula of the product celsian.

exchange operations was checked by determining Ba²⁺ concentration by titration with EDTA using erio T as the indicator and an NH₃ and NH₄⁺ solution at pH 10 as buffer [31].

The resulting powders were washed with doubly distilled water, dried overnight at 80°C and stored for at least 3 days in an environment having about 50% relative humidity to allow water saturation of zeolite.

2.2. Thermal Treatments. Ba-exchanged zeolite A samples, obtained according to the procedures previously described, were subjected to various thermal treatments in a Lenton furnace, which ensures stable temperature to within $\pm 2^\circ\text{C}$, using Al₂O₃ crucibles. The thermal treatments, which are summarised in Table 1, were all done in air atmosphere. They were performed as follows. In some cases, the samples were heated at a rate of 10°C/min up to the fixed temperature and subsequently cooled to room temperature. In the other cases, the samples were heated at a rate of 10°C/min up to the fixed temperature, kept at this temperature for the fixed time, and subsequently cooled to room temperature.

These products were characterised by X-ray diffraction at room temperature using a Philips X'PERT diffractometer with Cu K α radiation and $10^\circ < 2\theta < 60^\circ$ with a step width 2θ equal to 0.02° and 1 s data collection per step.

The various Ba-exchanged zeolite A samples were also characterised by simultaneous differential thermal analysis (DTA) and thermogravimetric analysis (TGA), using a Netzsch thermoanalyser model STA 409 with α -Al₂O₃ as a reference and a 10°C/min heating rate.

The cation composition of the final products of the thermal treatments, summarized in Table 2, were determined as follows. The various samples of Ba-exchanged zeolites A were thermally treated at 950°C for 2 hours in order to obtain the thermal collapse of the microporous zeolite structure and their irreversible dehydration. These thermally treated samples were chemically dissolved in the same hydrofluoric and perchloric acid solution and their Na⁺ concentrations were measured according to the procedure previously described.

2.3. Preparation of the Compacts. The powder-like Ba-exchanged zeolite A samples were used to prepare compacts according to the following procedure. Various zeolite powders were added with 6 weight % doubly distilled water and then pressed with an uniaxial laboratory press at 30 or 60 MPa to obtain cylindrical pellets (diameter = 15 mm, height = 5 mm). These pellets were dried at 105°C for 24 hours. Subsequently, the pellets were thermally treated in the previously described Lenton furnace according to the following thermal cycle:

- (1) heating from room temperature to 500°C with a heating rate of 5°C/min,
- (2) heating from 500 to 1300°C with a heating rate of 10°C/min,
- (3) 5 hours at 1300°C,
- (4) cooling from 1300 to 800°C with a cooling rate of 20°C/min,
- (5) cooling from 800°C to room temperature in air.

Subsequently these pellets were ground to fine powders which were characterised by X-ray diffraction at room temperature according to the modalities previously described.

3. Results

3.1. Cation Exchange. Data concerning cation exchange operations and, thus, cation composition of various zeolite A samples are summarized in Table 2. The residual Na content of various zeolite A samples is reported in the second column, whereas the Na equivalent fraction (x_{Na}) was in the third. These values were calculated with respect to the real initial Na content of Carlo Erba reagent-grade zeolite A. This quantity, which was experimentally determined in previous work [18], was 5.60 meq/g.

In the same Table 2 the Na₂O % weight of samples 1, 2, 3, and 4 thermally treated at 950°C for 2 hours are reported. These values were used to calculate the chemical formula of celsian which can be obtained through the thermal treatments of samples 1, 2, 3, and 4. Such chemical formulae are reported in the final column of Table 2.

3.2. X-Ray Diffraction. XRD patterns of sample 1 thermally treated at 200°C for 2 hours, heated at 500°C, and then cooled down in air atmosphere and thermally treated at 800°C for 28 hours are reported in Figure 1 together with the spectral lines of monoclinic celsian (JCPDS card n. 38-1450).

The thermal treatment at 200°C for 2 hours resulted in the thermal collapse of the microporous zeolitic structure with the consequent formation of an amorphous phase. In the XRD pattern of sample 1 heated at 500°C, and then cooled down in air atmosphere the presence of extremely broad diffraction peaks, which can be ascribed to monoclinic celtsian, may be recorded. XRD patterns of samples 1 thermally treated at 800°C for 28 hours closely resembles XRD diffraction pattern of sample 1, heated at 500°C and then cooled down in air atmosphere, which shows that thermal treatment at 800°C for 28 hours does not result in a sensible increase of the extent to which the crystallization of any phase occurs. The XRD patterns of samples 2, 3, and 4 thermally treated at 200 for 2 hours, heated at 500°C and then cooled down in air atmosphere, and thermally treated at 800°C for 28 hours are not reported as they closely resemble the homologous XRD pattern of sample 1.

The XRD pattern of samples 1, 2, 3, and 4 thermally treated at 1000°C for 5 hours are reported in Figure 2. In the XRD pattern of sample 1 sharp peaks of hexacelsian appear together with peaks of monoclinic celtsian whose intensity is slightly higher than the intensity of peaks of monoclinic celtsian recorded after thermally treating sample 1 at 800°C for 28 hours (Figure 1(c)). The intensity of peaks of hexacelsian decreases with increasing Na residual content of Ba-exchanged zeolite A, attains a minimum in sample 3, and then increases again in sample 4. In Figure 2 and in the following ones, the main diffraction peaks of hexacelsian and monoclinic celtsian are denoted with H and M, respectively. An opposite trend may be detected by examining the intensity of peaks of monoclinic celtsian which increases with increasing Na residual content of Ba-exchanged zeolite A, attains a maximum in sample 3, and then decreases again in sample 4. It must be noticed that in the XRD pattern of sample 4 a very low intensity peak (denoted with S), indicating the presence of a small amount of the sodium aluminosilicate known as low carnegieite (NaAlSiO_4) (JCPDS card n. 33-1203), may be detected. The attribution of the peak at 21° 2θ to low carnegieite was performed on the basis of the following considerations. Zeolite A in its original Na form was subjected to a 5 h thermal treatment at 1300°C which resulted in the crystallization of low carnegieite, whose peak of highest intensity is located exactly at 21° 2θ . The remaining peaks of low carnegieite may not be immediately detected in the Figure 2 on account of the following reasons:

- they superimpose with peaks of monoclinic celtsian or hexacelsian;
- their intensity is too weak due to the low amount of crystallized carnegieite.

XRD pattern of various samples thermally treated at 1100°C for 5 and 10 hours (not reported) and thermally treated at 1200°C for 5 hours (not reported) resembles the XRD pattern of homologous samples thermally treated at 1000°C for 5 hours. The only difference that can be noticed by comparing these XRD patterns is that the intensity of peaks of hexacelsian decreases and the intensity of peaks of

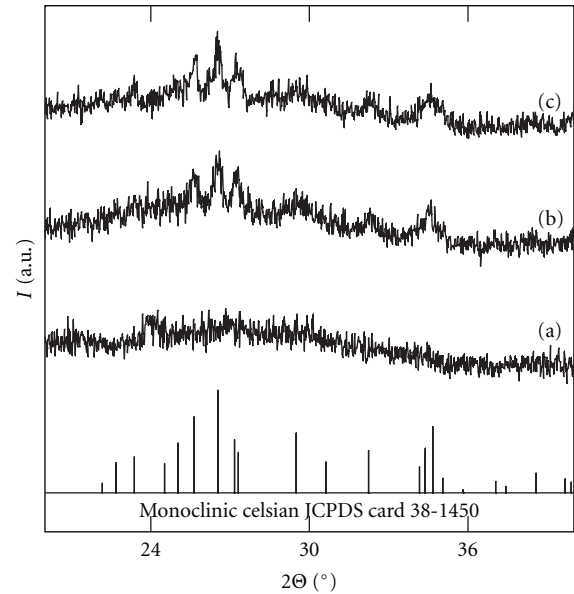


FIGURE 1: XRD patterns of sample 1 thermally treated at 200°C for 2 hours (a), heated at 500°C and then cooled down in air atmosphere (b), and thermally treated at 800°C for 28 hours (c).

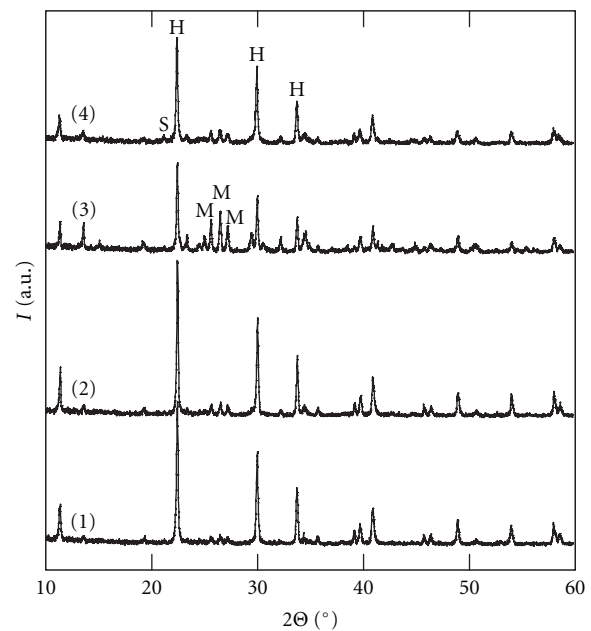


FIGURE 2: XRD patterns of powder-like samples 1, 2, 3, and 4 thermally treated at 1000°C for 5 hours (M = monoclinic celtsian, H = hexacelsian, S = sodium aluminosilicate).

monoclinic celtsian increases with increasing the temperature and duration time of thermal treatment, all other conditions being equal.

XRD patterns of samples 1, 2, 3, and 4 thermally treated at 1300°C for 5 hours are reported in Figure 3. In this same figure, the XRD patterns of the materials obtained by

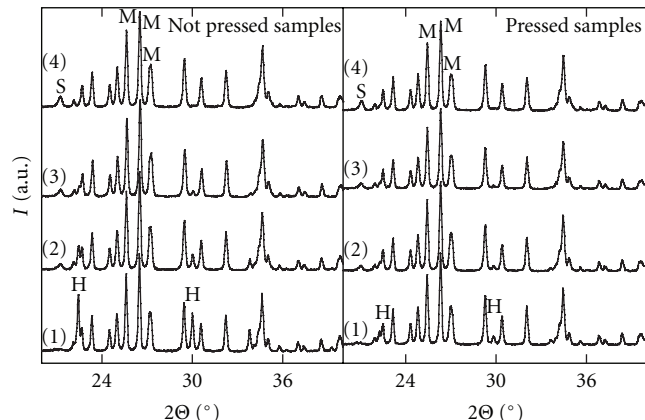


FIGURE 3: XRD patterns of powder-like samples 1, 2, 3, and 4 and of materials obtained by grinding pellets manufactured by pressing samples 1, 2, 3, and 4 at 30 MPa, thermally treated at 1300°C for 5 hours (M = monoclinic celsian, H = hexacelsian, S = sodium aluminosilicate).

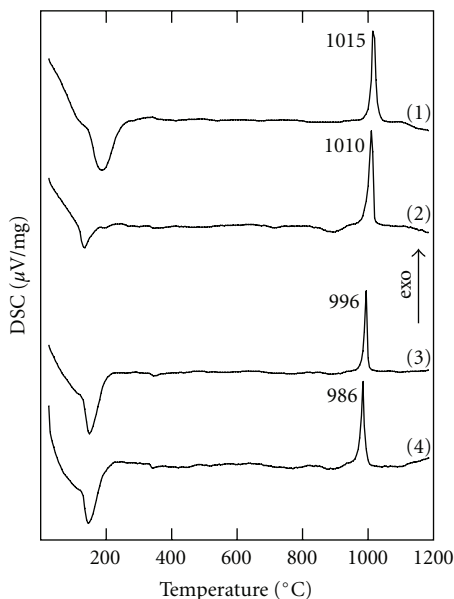


FIGURE 4: DTA curves for the samples 1, 2, 3, and 4.

grinding the cylindrical pellets previously prepared by cold pressing at 30 MPa samples 1, 2, 3, and 4 and then subjected to the thermal cycle described in experimental are reported.

In these XRD patterns, the diffraction peaks of monoclinic celsian appear together with those of hexacelsian. The intensity of peaks of hexacelsian steadily decreases and the intensity of peaks of monoclinic celsian steadily increases with increasing the Na residual content of Ba-exchanged zeolite A samples. In Figure 3 the presence of diffraction peaks of hexacelsian can be hardly detected in XRD pattern of Ba-exchanged zeolite A samples exhibiting the highest Na residual content. It must be noticed that in the X-ray diffraction pattern of samples 2, 3, and 4, a very low intensity peak indicating the presence of an extremely

small amount of sodium aluminosilicate NaAlSiO_4 , may be detected, regardless of the fact that the thermal treatment was performed on a powder or on a previously pressed pellet. Obviously, the intensity of this peak appears to increase with increasing Na content. Finally, it appears noteworthy that the intensity of peaks of monoclinic celsian is higher and the intensity of peak of hexacelsian is lower in the XRD patterns of the powders obtained by grinding the pellets than in the XRD patterns of the powder-like Ba-exchanged zeolite A samples, all other conditions being equal. The XRD patterns of the materials obtained by grinding the cylindrical pellets previously prepared by cold pressing at 60 MPa samples 1, 2, 3, and 4 and then subjected to the thermal cycle described in experimental, are not reported, as they did not sensibly differ from those of the materials obtained by grinding the cylindrical pellets previously prepared by cold pressing at 30 MPa samples 1, 2, 3, and 4.

XRD patterns of powder-like samples 1, 2, 3, and 4 thermally treated at 1300°C for 10 and 15 hours are not reported inasmuch as they resemble the XRD patterns of homologous powder-like samples reported in Figure 3, although they clearly denote that the intensity of peaks of hexacelsian decreases and the intensity of peaks of monoclinic celsian increases with increasing the duration time of thermal treatment, with all other conditions being equal. Even in such XRD patterns, the intensity of diffraction peaks of monoclinic celsian increases and the intensity of peaks of hexacelsian decreases with increasing the Na residual content of Ba-exchanged zeolite A samples. In particular, the presence of diffraction peaks of hexacelsian can be very hardly detected in XRD pattern of samples 1 and 2 thermally treated at 1300°C for 10 hours and of sample 1 thermally treated at 1300°C for 15 hours, whereas completely disappeared in the remaining XRD patterns.

XRD patterns of samples 1, 2, 3, and 4 thermally treated at 1400 and 1500°C for 5 hours are not reported inasmuch as they very closely resemble the XRD pattern of homologous samples thermally treated at 1300°C for 5 hours, thus indicating that the temperature of thermal treatment does not play a crucial role for temperatures higher than 1300°C.

3.3. Thermal Analysis. Figure 4 reports the DTA curves of samples 1, 2, 3, and 4. In all curves a large low-temperature endothermic effect and a sharp high-temperature exothermic effect may be recorded. The large low-temperature endothermic effect, which attains its minimum at about 180°C, is related to the progressive dehydration of zeolite A upon heating followed by the thermal collapse of the microporous zeolite structure with the formation of an amorphous phase. This finding is in agreement with the XRD data which reveals that when thermally treated at 200°C for 2 hours, the various samples result in a completely amorphous phase (Figure 1). The sharp high-temperature exothermic effect, which can be reasonably ascribed to the crystallization of hexacelsian from the amorphous phase, occurs at temperatures, ranging between 986 and 1015°C, which steadily decrease with increasing Na content. Even this finding is in agreement with the XRD data which reveal

that equal thermal treatments of various samples in the temperature range 800–1100°C result in the crystallization of hexacelsian from the amorphous phase and that the extent to which such crystallization occurs steadily increases with increasing Na content. Evidence of the sluggish hexacelsian → monoclinic celsian transition may not be found in the DTA curves of various samples.

4. Discussion

The careful inspection of the data here reported confirms the results of previous studies [14, 18–25]. In particular, the sequence of thermal transformations of Ba-exchanged zeolite A (zeolite → amorphous phase → hexacelsian → monoclinic celsian) and the formation of small crystallites of monoclinic celsian prior to the crystallisation of hexacelsian, which expedite the subsequent conversion of hexacelsian into monoclinic celsian, were confirmed. The strong mineralizing action developed by Na, envisaged in previous work [14, 18–25], is confirmed on the basis of the considerations that will be hereafter reported. Actually small increases of Na residual content result in a steady decrease of the temperature at which the crystallization of hexacelsian from the amorphous phase, arising from the thermal collapse of the microporous zeolite structure, occurs (see Figure 4). Then small increases of Na residual content result in sensible decreases of temperatures and times needed to attain the full conversion of hexacelsian to monoclinic celsian for thermal treatments at temperatures not lower than 1300°C. As far as the thermal treatments in the temperature range 1000–1200°C are concerned, the extent to which the conversion of hexacelsian in monoclinic celsian occurs attains a maximum for sample 3 that exhibits 0.58 meq/g Na residual content. This seeming inconsistency may be explained considering that, in this temperature range, the crystallization of a small amount of low carnegieite NaAlSiO_4 occurs only in sample 4, which exhibits the highest Na residual content (0.74 meq/g), at a temperature as low as 1000°C (see Figure 2). The Na present in this phase does not appear available to promote other phase transformations such as the conversion of hexacelsian to monoclinic celsian. The extent of the conversion of hexacelsian in monoclinic celsian steadily increases with increasing Na residual content when the temperature of thermal treatments is not lower than 1300°C. Actually such thermal treatment promotes the crystallization of a small amount of low carnegieite NaAlSiO_4 even in samples 2 and 3 at 1300°C. Moreover, the strong mineralizing role of Na is confirmed by the fact that for Na residual contents not lower than 0.27 meq/g, the temperature of the thermal treatment, if higher than 1300°C, does not play a crucial role in the thermal transformations of Ba-exchanged zeolite A.

Another point that appears worth to be stressed is that the amount of monoclinic celsian is higher and the amount of hexacelsian is lower in all the materials obtained by grinding the pellets than in the powder-like Ba-exchanged zeolite A samples, all other conditions being equal (see Figure 3). This finding could be interpreted bearing in mind that pressing the powder-like samples of Ba-exchanged zeolite A

to manufacture cylindrical pellets gives rise to an increase in the contact surface among the grains of hexacelsian. This fact could likely make easier and expedite the conversion of hexacelsian into monoclinic celsian in the case this phase transition starts on the surface of grains of hexacelsian rather than in their bulk. An increase in the forming pressure from 30 to 60 MPa does not result in a further increase of the reactivity of the system, presumably because the increase in the contact surface among the grains of hexacelsian, related to such increase in forming pressure, does not occur to a very large extent. These findings are in agreement with the literature, as it was found that forming pressures up to 50 MPa promote a densification process, due to the grains deformation which produces a decrease of porosity [32]. Forming pressures higher than 50 MPa do not further improve the densification process [32].

The increase in the reactivity of the system arising from pressing powder-like samples of Ba-exchanged zeolite A samples to obtain cylindrical pellets has another important consequence. Actually an inspection of the data of this paper reveals that thermally treating at 1300°C for 5 hours samples of Ba-exchanged zeolite A with Na residual content higher than 0.27 meq/g previously pressed at 30 MPa to obtain cylindrical pellets, results in final products almost totally made up by monoclinic celsian with a very low amount of hexacelsian. This finding appears of great interest from the point of view of practical applications inasmuch as the presence of not more than 10% weight of hexacelsian does not result in a sensible reduction of the technological properties of celsian [1]. This fact allows to envisage that thermally treating at 1300°C for 5 hours samples of Ba-exchanged zeolite A with a residual Na content higher than 0.27 meq/g, has potential for practical application in the manufacture of celsian-based ceramics.

5. Conclusions

The results of this work confirm the sequence of thermal transformations that Ba-exchanged zeolite A undergoes (zeolite → amorphous phase → hexacelsian → monoclinic celsian) [14, 18–25] and the strong mineralizing action developed by Na. Moreover, the fact that pressing the Ba-exchanged zeolite A powder-like samples to obtain cylindrical pellets which expedite the sluggish final phase transition hexacelsian → monoclinic celsian is revealed.

Nevertheless, the results of this work infer further interesting considerations concerning the optimum residual Na content of Ba-exchanged zeolite A to be transformed into monoclinic celsian. Actually the results of this work seem to supply strong indication that such optimum Na residual content ranges between 0.27, and 0.43 meq/g. The considerations that suggested this conclusion are the following.

- (1) Na residual contents lower than 0.27 meq/g result in high temperatures and/or long thermal treatment times to transform hexacelsian into monoclinic celsian (as an example the completion of this transformation was attained by thermally treating at 1300, 1400, and 1500°C for 22, 15, and 22 hours,

respectively, a sample of Ba-exchanged zeolite A with a 0.20 meq/g Na residual content) [18].

- (2) Na residual content not lower than 0.43 meq/g may result in the formation of amounts of sodium aluminosilicate NaAlSiO_4 exhibiting low melting temperature, sufficient to negatively affect the good technological properties of celsian-based ceramics.
- (3) On the basis of the results reported in this paper it may be said that a 5-hour thermal treatment at 1300°C of a pellet obtained by pressing at 30 MPa Ba-exchanged zeolite A samples with a residual Na content ranging between 0.27 and 0.43 meq/g results in a product mostly made up by monoclinic celsian which should presumably exhibit the good technological properties typical of celsian-based ceramics. Actually the amount of hexacelsian, (presumably lower than 10% weight), and the very small amount of sodium aluminosilicate NaAlSiO_4 possibly present in the final product together with monoclinic celsian, do not appear able to sensibly affect the good technological properties of celsian-based ceramics.

Acknowledgment

This work was carried out with the financial contribution of the (MIUR) Ministero dell'Istruzione dell'Università e della Ricerca.

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