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Abstract: Belite-rich cements hold promise for reduced energy consumption and CO₂ emissions, but their use is hindered by the slow hydration rates of ordinary belites. This drawback may be overcome by activation of belite by doping. Here, the doping mechanism of B and Na/B in belites is reported. For B-doping, three solid solutions have been tested: Ca_{2-x/2}B_{x/2}(SiO₄)_{1-x}(BO₃)_x, Ca₂(SiO₄)_{1-x}(BO₃)_xO_{x/2} and Ca_{2-x}B_x(SiO₄)_{1-x}(BO₄)_x. The experimental results support the substitution of silicate groups by tetrahedral borate groups with the concomitant substitution of calcium by boron for charge compensation, Ca_{2-x}B_x(SiO₄)_{1-x}(BO₄)_x. Otherwise, the coupled Na/B-doping of belite has also been investigated and Ca_{2-x}Na_x(SiO₄)_{1-x}(BO₃)_x series is confirmed to exist for a large range of x values. Along this series, β 'H-C₂S is the main phase (for x $\geq$ 0.10) and is single phase for x=0.25. Finally, a new structural description for borax doping in belite has been developed for β 'H-Ca_{1.85}Na_{0.15}(SiO₄)_{0.85}(BO₃)_{0.15}, which fits better borax activated belite cements in Rietveld mineralogical analysis.

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Reactive belite stabilization mechanisms by boron-bearing dopants

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ABSTRACT

Belite-rich cements hold promise for reduced energy consumption and CO₂ emissions, but their use is hindered by the slow hydration rates of ordinary belites. This drawback may be overcome by activation of belite by doping. Here, the doping mechanism of B and Na/B in belites is reported. For B-doping, three solid solutions have been tested: $\text{Ca}_{2-x/2}\text{B}_{x/2}(\text{SiO}_4)_{1-x}(\text{BO}_3)_x$, $\text{Ca}_2(\text{SiO}_4)_{1-x}(\text{BO}_3)_x\text{O}_{x/2}$ and $\text{Ca}_{2-x}\text{B}_x(\text{SiO}_4)_{1-x}(\text{BO}_4)_x$. The experimental results support the substitution of silicate groups by tetrahedral borate groups with the concomitant substitution of calcium by boron for charge compensation, $\text{Ca}_{2-x}\text{B}_x(\text{SiO}_4)_{1-x}(\text{BO}_4)_x$. Otherwise, the coupled Na/B-doping of belite has also been investigated and $\text{Ca}_{2-x}\text{Na}_x(\text{SiO}_4)_{1-x}(\text{BO}_3)_x$ series is confirmed to exist for a large range of x values. Along this series, α' -C₂S is the main phase (for $x \geq 0.10$) and is single phase for $x = 0.25$. Finally, a new structural description for borax doping in belite has been developed for α' -Ca_{1.85}Na_{0.15}(SiO₄)_{0.85}(BO₃)_{0.15}, which fits better borax activated belite cements in Rietveld mineralogical analysis.

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1. INTRODUCTION

The manufacture of Portland cement on average emits about 0.83 tons of CO₂ per tonne of cement, which is not particularly high for a chemical product. However, because of the enormous volumes of cement used worldwide to make concrete, such emissions amount to about 6% of all anthropogenic CO₂ emissions [1]. Cements based on belite have been proposed as environmentally friendly materials, as the reduction of CO₂-emissions may be as much as 10% for belite Portland cements [2-4] or up to 35% in belite calcium sulfoaluminate cements [5,6]. Belite Portland cements are being already used in “low heat hydration concretes” for construction of large dams and lining of oil wells. However, the massive application of these materials requires the overcoming of some drawbacks such as the low hydration rate of the belite phase and high resistance to be milled [7,8]. Belite calcium sulfoaluminate cements are also being partially used in China [6,9].

Belite (Ca₂SiO₄, C₂S, with some substituting elements) is the major constituent of Belite Portland cements and some calcium sulfoaluminate cements. Currently, borate-activated belite calcium sulfoaluminate cements are undergoing industrial trials under an EU project (<http://www.aether-cement.eu/>) to determine the CO₂ savings. Furthermore, this phase is also the second most abundant constituent of OPC [10]. Stoichiometric C₂S has five polymorphs [11,12] (γ , β , α'_L , α'_H and α) and its temperature evolution is shown in Figure 1. The γ -phase, which is essentially nonreactive with water, is stable at ambient temperature and it crystallizes in an orthorhombic olivine-type structure [13-15]. The β -form is a metastable monoclinic phase at room temperature [16,17]. α'_L and α'_H orthorhombic phases are stable at higher temperatures [18]. The α'_L -polymorph is generally considered a superstructure of the α'_H and it has been reported two possibilities for indexing: doubling the a and c parameters [19] or tripling the b parameter [20-22]. Finally, the highest-temperature polymorph is the α -form whose structure is still under discussion [23-25]. It is noteworthy, that $\beta \rightarrow \gamma$ polymorphic transformation on cooling is disruptive with a change of 12 vol%. This effect, called “dusting”, causes the material to disintegrate spontaneously to a powder [26] and can be avoided by chemical, thermal and mechanical treatments [10,11].

The framework of silicon tetrahedra is very similar among these polymorphs, while the arrangement of calcium cations is slightly different. The crystal structure of γ -C₂S has two calciums in regular six-coordinated oxygen environments. Meanwhile, the crystal structure of β -C₂S has two calciums surrounded by eight oxygens in distorted environments. Furthermore, the crystal structures of α'_H - and α - polymorphs have calcium cations in both eight and nine irregular coordinations [11-16]. It is worth noting that there are experimental evidences that an increase of the calcium coordination number seems to enhance the water reactivity [27]. However, the actual reactivity of each form depends on the kind and amount of dopants employed [28-30]. Therefore, theoretical calculations [31] are needed to properly address this issue.

It is known that the presence of defects or strains in crystalline structures can modify physical and chemical properties, or even causes the stabilization of a high temperature form at room temperature [32,33]. The different type of defects (overall known as microstrains) can be introduced during the material preparation by the formation of solid solutions or by specific thermal treatments [34]. The effects on chemical-stabilizing ions on the stability of β -C₂S have been extensively investigated [11,28,35], and several theoretical predictions have been made about which ions can stabilize β -C₂S (B₂O₃, Na₂O, K₂O, BaO, MnO₂, Cr₂O₃ or their combinations) [29,36-39]. It has been published that foreign oxides, such as MgO, P₂O₅, K₂O, BaO and SO₃, promote the formation of α' -C₂S and increase its hydraulic properties [40,41]. B₂O₃ has also been studied in order to stabilize β -C₂S and α' -C₂S at room temperature [28,42]. Some authors [27,42,43] have demonstrated that the effectiveness of addition of B₂O₃, with no other co-dopants, for stabilizing for α' -forms, is poor. On the other hand, the addition of a combination of dopants, for instance, B₂O₃ and Na₂O, successes to stabilize α' -C₂S [42] although these authors did not state which α' -forms, α'_L or α'_H , was stabilized. Moreover, previous studies reported that borax stabilizes α' -form in a cement matrix [5]. The α -C₂S is much more difficult to stabilize at room temperature, however this phase was stabilized by the addition of alkaline oxides in belite Portland cements [44,45].

The objective of this study is to investigate the mechanism of α' -belite stabilization at room temperature by B₂O₃ and Na₂O additions. An improved structural description of α'_H -C₂S polymorph is reported. The driving force for this research is the need to manufacture, and to understand the production, of highly reactive belite calcium sulfoaluminate cements. The final goal is the need to reduce CO₂ emissions in the cement industry.

2. EXPERIMENTAL SECTION

2.1 Sample preparation. Two types of C₂S series have been synthesized. The first type deals with the C₂S stabilization just by boron doping. Three solid solutions with nominal stoichiometries Ca_{2-x/2}□_{x/2}(SiO₄)_{1-x}(BO₃)_x, Ca₂(SiO₄)_{1-x}(BO₃)_xO_{x/2} and Ca_{2-x}B_x(SiO₄)_{1-x}(BO₄)_x have been tested. In this case, the boron source was boric acid. The second type of series deals with the C₂S stabilization by boron and sodium doping. Two solid solutions with nominal stoichiometries Ca_{2-x}Na_x(SiO₄)_{1-x}(BO₃)_x and Ca_{2-3x}B_{2x}Na_x(SiO₄)_{1-x}(BO₄)_x have been tested. For these solid solutions, borax was used as main dopant source but sodium carbonate and boric acid were also added as correctors to achieve the right stoichiometries for the first and second series, respectively.

All series were prepared using CaCO₃ (99.95%-100.05% from Alfa Aesar) and SiO₂ (99.7% from ABCR). Dopants were added as H₃BO₃ (100% from VWR), Na₂B₄O₇·10H₂O (100% from Aldrich) and Na₂CO₃ (99.999% from Aldrich). Table 1 shows the nominal elemental composition expressed as oxides for each member of each solid solution, excluding CO₂ and H₂O. Raw mixtures were ground for 10 min in an agate mortar with acetone, and preheated at 1000°C for 4 hours (heating rate of 10 °C/min)

in Pt/Rh crucibles. Mixtures were placed on Pt/Rh crucibles and preheated at 1000°C for 4 hours (heating rate of 10 °C/min). After cooling, the mixtures were milled with 2-butanol in a Fritsch ball mill (model Pulverisette 7, 45 cm³ agate vessel containing 7 agate balls with a diameter of 15 mm) during 1 h at 100 rpm with reverse rotation each 20 min, and dried at 100°C in a stove. The resulting powders were pelletized (20 mm diameter) and a second thermal treatment was carried out at 1300°C for 30 min (heating rate of 20 °C/min) and quenched from high temperature with an air flow. The pellets were broken and grounded in WC mortar. The resulting powders were characterized by the techniques described below.

Finally, a laboratory-prepared iron-rich belite calcium sulfoaluminate clinker has also been prepared. This clinker was activated by adding borax (Na₂B₄O₇·10H₂O) (2.0 wt% expressed as B₂O₃) to the raw material to promote room temperature stabilization of reactive belite polymorph (α' _H-form). The preparation and full characterization, including Rietveld quantitative phase analysis, has been already reported [5]. This clinker contains 56.7 wt% of α' _H-C₂S, 31.1 wt% of C₄A₃S̄, 10.1 wt% of C₄AF and 2.1 wt% of CT and it is hereafter named BC SAF-B2.

2.2. X-ray powder diffraction. All members of the C₂S solid solutions were characterized by laboratory X-Ray powder diffraction (LXRPD). Diffraction data were recorded on an X'Pert MDP PRO diffractometer (PANalytical) equipped with a Ge (111) primary monochromator, using monochromatic CuK α ₁ radiation (λ =1.54059 Å) and an X'Celerator detector. The overall measurement time was ~ 4h per pattern to have good statistics over the 5.0° to 140.0° (2 θ) angular range, with 0.017° step size.

2.3. Rietveld analysis. All Rietveld quantitative phase analyses (RQPA) were carried out using the GSAS suite of programs [46]. Final global optimized parameters were: background coefficients, zero-shift error, cell parameters and peak shape parameters using a pseudo-Voigt function [47] corrected for axial divergence [48]. Ca_{1.85}Na_{0.15}(SiO₄)_{0.85}(BO₃)_{0.15} sample patterns were used to obtain an improved structural description for doped α' _H-C₂S polymorph. In this case, the atomic positional coordinates and isotropic atomic displacement parameters (ADP) were optimized.

2.4. Infrared spectroscopy. ATR-FTIR (Attenuated Total Reflectance – Fourier Transform Infrared) spectra were obtained with an ATR accessory (MIRacle ATR, PIKE Technologies, USA) coupled to FTIR spectrometer (FT/IR-4100, JASCO, Spain). All spectra were recorded in the 4000 to 600 cm⁻¹ range at 4 cm⁻¹ resolution and 25 scans were accumulated. The powder samples were placed on a holder of approximately 3 mm of diameter.

2.5. ¹¹B Magic Angle Spinning Nuclear Magnetic Resonance (MAS NMR). ¹¹B (I=3/2) MAS-NMR spectra for three selected samples were recorded in an Avance-400 Bruker spectrometer. The resonance frequency was 128.38 MHz (B₀ = 9.4 T). Spectra were recorded using $\pi/2$ (5 μ s) pulses and 5s intervals between accumulations with the spinning rate of samples being 10 kHz. In all cases, spectral filters used were 125 kHz and the appropriate number of scans was chosen in order to obtain

S/N ratios higher than 30. Measurements were done at room temperature with boric acid as external reference. The error in chemical shift values is estimated to be lower than 0.5 ppm.

2.6. Scanning Electron Microscopy. Small pellets with cylindrical shape for selected compositions were studied in a JEOL JSM-6490LV scanning electron microscope using secondary electrons. EDX measurements were carried out with the OXFORD INCA Energy 350 attachment. This unit has a Si(Li) detector with a super atmospheric thin window (SATW). Samples were coated with graphite.

2.7. Inductively coupled plasma mass spectroscopy. The elemental compositions of samples, after the thermal treatments, were determined by ICP-MS on Perkin Elmer spectrophotometer (Nexion 300D). Previously, samples were digested in an Anton Paar device (Multiwave 3000) by using HNO₃, HCl and HF.

3. RESULTS AND DISCUSSION

3.1. Solid solution mechanisms. Previous studies have shown that boron is an effective dopant in belite to stabilize the β -polymorph [27,43] and even higher temperature polymorphs [28,42]. Furthermore, there is a report mentioning that boron is incorporated in belite as BO_4^{5-} [11]. However, there is not reported mechanism for the boron incorporation within belites. The first issue to be addressed is the boron anion nature within belites, i.e. if the borate group is tetrahedral BO_4^{5-} or planar-triangular BO_3^{3-} . Furthermore, as the B-doping in the silicate framework, SiO_4^{4-} , is aliovalent, it is also a key point to understand the charge compensating mechanism. In order to address these open issues, three nominal solid solutions have been tested: i) $\text{Ca}_{2-x/2}\square_{x/2}(\text{SiO}_4)_{1-x}(\text{BO}_3)_x$; ii) $\text{Ca}_2(\text{SiO}_4)_{1-x}(\text{BO}_3)_x\text{O}_{x/2}$; and iii) $\text{Ca}_{2-x}\text{B}_x(\text{SiO}_4)_{1-x}(\text{BO}_4)_x$ (see Table 1). The first two series are designed for stabilizing with planar-triangular BO_3^{3-} anions. The first one is charge compensated with calcium vacancies and the second with interstitial oxide anions, as in other silicate-based systems [49]. Table 2 shows RQPA results for these two series and chiefly, all members contain large amounts of free calcium oxide, which indicate that the target compositions were not achieved. Furthermore, ATR-FTIR infrared spectra of these two series, Figures S1 and S2, show the presence of BO_3^{3-} units but also BO_4^{5-} -type groups, characteristic vibrations close to 1250 cm^{-1} and 1000 cm^{-1} , respectively [50,51]. The BO_4^{5-} vibration at 1000 cm^{-1} is overlapped with that arising from SiO_4^{4-} . The third mechanism is based on substitution of tetrahedral SiO_4^{4-} groups by tetrahedral BO_4^{5-} units with concomitant boron substitution of calcium for charge compensation, $\text{Ca}_{2-x}\text{B}_x(\text{SiO}_4)_{1-x}(\text{BO}_4)_x$ ($0 \leq x \leq 0.20$). This mechanism is similar to that already reported for aluminum doping in alite, $\text{Ca}_{3-x}\text{Al}_x(\text{SiO}_4)_{1-x}(\text{AlO}_4)_x\text{O}$ [52]. Samples with $x=0.15$ and $x=0.20$ of this series were melted at high temperature, thus these compositions were not further studied. Figure 2 shows raw LXRPD patterns for the $\text{Ca}_{2-x}\text{B}_x(\text{SiO}_4)_{1-x}(\text{BO}_4)_x$ series, up to $x=0.10$, with diagnostic peaks labelled in order to identify a given polymorph. Table 2 gives RQPA for all members up to $x=0.10$ of this series and it shows that the amount of CaO is, for all these x values, lower than 1.0 wt%. Moreover, Table 2 shows that for $x=0.00$ β -polymorph has been partially stabilized, being mainly

due to the milling processes performed to raw materials and to the quenching. The polymorph β -C₂S is present as main phase, i.e. more than 60.0 wt% up to $x=0.05$, meanwhile α' _H-C₂S is the main phase for $x=0.10$. Moreover, ATR-FTIR infrared spectra were also performed and Figure 3a shows the ATR-FTIR infrared spectra of all the members of this series (up to $x=0.10$). These measurements show bands close to 1000 cm⁻¹ assigned to BO₄⁵⁻ stretching vibrations. Furthermore, tiny signals from 1150 to 1350 cm⁻¹, corresponding to BO₃³⁻ (or similar) units, are also observed (see Figure 3a) [50,51]. However, these results do not rule out the formation of this series due to the fact that BO₃³⁻ units (or quite similar groups) can be locally formed where boron cations substitute calcium atoms. Boron is much smaller than calcium and a large displacement within the site is expected which may lead to the formation of a BO₃³⁻-type groups. Variation of the unit cell volume in this solid solution is given in Figure 4 (open symbols) and final refined unit cell values are included in Table S1, as supplementary material. It is relevant that γ -form unit cell volume is constant in its whole chemical field of existence, i.e. dopants are not incorporated within γ -C₂S. On the other hand, β - and α' _H-form unit cell volumes are relatively constant up to $x=0.0125$ and decrease for higher values of x . Taking all these results into account, it can be stated that boron stabilizes high temperature polymorphs by substituting jointly silicon units as BO₄⁵⁻ and calcium cations by B³⁺, as for aluminum doping in alite [52].

On the other hand, borax is a relatively inexpensive raw material for boron (and sodium) doping in belite cements and it is already being used in industrial trials. Furthermore, it is known to stabilize high-temperature forms of belite [12,53-55]. Unfortunately, there are not reports dealing with the charge compensating mechanism for borax, i.e. B and Na, stabilization of belite. As it was discussed above, two possibilities for borate incorporation arises, BO₄⁵⁻ or BO₃³⁻. Therefore, two solid solutions were tested using boron and sodium as dopants: i) Ca_{2-x}Na_x(SiO₄)_{1-x}(BO₃)_x; and ii) Ca_{2-3x}B_{2x}Na_x(SiO₄)_{1-x}(BO₄)_x. In addition to borax, sodium carbonate and boric acid were also added as correctors for achieving the right doping contents, for the first and second solid solutions, respectively. Figure 5 shows raw LXRPD patterns for the Ca_{2-x}Na_x(SiO₄)_{1-x}(BO₃)_x series with diagnostic peaks labelled in order to identify a given polymorph and Table 2 gives RQPA of this series. The first result shown in Table 2 is the absence of free calcium oxide along this series. A second very relevant result is that high temperature polymorphs of C₂S, i.e. β - and α' _H-C₂S, are stabilized as x increases, as expected. Elemental analysis by ICP-MS was performed for all the members of this series to determine the volatilization rate of sodium, see Table S2. The sodium losses, expressed as Na₂O wt%, represented 34% of the total at the lowest Na₂O addition rate but only 15% at the highest addition rate. Therefore, used nominal compositions do not fully represent the stoichiometry of the bulk phases. Furthermore, B₂O₃ contents were also analysed by ICP-MS for the Ca_{2-x}B_x(SiO₄)_{1-x}(BO₄)_x and Ca_{2-x}Na_x(SiO₄)_{1-x}(BO₃)_x series. The measured volatilization rates were negligible with retention rates larger than 95%.

For Ca_{2-x}Na_x(SiO₄)_{1-x}(BO₃)_x series, β -C₂S is present between $x=0.00$ and $x=0.20$. On the other hand, α' _H-C₂S appears from $x=0.025$ up to $x=0.25$, the last composition being single phase. All powder patterns were analyzed by the Rietveld method. Final refined unit cell parameters of each polymorph

and RQPA of all solid solution members are given as supplementary material in Table S3. Figure 4 plots the variation of the refined unit cell volumes along this series (closed symbols). The same conclusions as with the previous discussed series can be drawn for unit cell volume variations within this one: i) γ -form unit cell volume is constant in its whole chemical field of existence, ii) β - and α' -form unit cell volumes are relatively constant up to $x=0.10$ and decrease for higher values of x . However, the contraction of unit cell volume in this series occurs at a smaller pace than in the previous studied one, mainly due to the smaller ionic radii mismatch of the Na/Ca when compared to B/Ca. This unit cell contraction has been previously observed in high belite clinkers doped with borax [5]. ATR-FTIR infrared spectra of this series, Figure 3b, confirms the presence of BO_3^{3-} units and the intensities of their vibration bands, close to 1250 cm^{-1} , increase with x , and the absence of BO_4^{5-} units, as there are not signals around 1000 cm^{-1} .

Figure 6 displays the ^{11}B ($I = 3/2$) MAS-NMR spectra for three selected samples: BCSAF-B2 clinker, $\text{Ca}_{1.85}\text{Na}_{0.15}(\text{SiO}_4)_{0.85}(\text{BO}_3)_{0.15}$ and $\text{Ca}_{1.9}\text{B}_{0.1}(\text{SiO}_4)_{0.9}(\text{BO}_4)_{0.1}$. This figure also shows, as an inset, the ^{11}B MAS-NMR spectra (experimental and calculated) for crystalline $\text{B}(\text{OH})_3$ standard taken from reference [56]. This spectrum shows three maxima arising from a unique trigonal-planar BO_3^{3-} group due to the quadrupolar splitting [57]. The ^{11}B MAS-NMR spectra for BCSAF-B2 and $\text{Ca}_{1.85}\text{Na}_{0.15}(\text{SiO}_4)_{0.85}(\text{BO}_3)_{0.15}$ sample are very similar to that of $\text{B}(\text{OH})_3$ and fully consistent with the presence of trigonal BO_3^{3-} groups. The central signal is formed by two maxima at 19 and 8 ppm and a small shoulder at -2 ppm. A second very small additional signal is evident in the spectrum of $\text{Ca}_{1.85}\text{Na}_{0.15}(\text{SiO}_4)_{0.85}(\text{BO}_3)_{0.15}$ sample which may be related to the incorporation of planar BO_3 units in the side-phase, $\beta\text{-C}_2\text{S}$, see Table 2.

The ^{11}B MAS-NMR spectrum for $\text{Ca}_{1.90}\text{B}_{0.10}(\text{SiO}_4)_{0.90}(\text{BO}_4)_{0.10}$ (see Figure 6 bottom) is more complex in agreement with the infrared data and the charge-compensation mechanism. In addition to at least one (likely two) BO_3 groups, the fitting of the experimental profile requires also the presence of a tetrahedral BO_4^{5-} unit at ~ 0 ppm [58,59]. In any case, it is clear the coexistence of trigonal BO_3^{3-} and tetrahedral BO_4^{5-} units for this composition and that the borax-activated BCSAF clinker contains just trigonal BO_3 groups.

The series $\text{Ca}_{2-3x}\text{B}_{2x}\text{Na}_x(\text{SiO}_4)_{1-x}(\text{BO}_4)_x$ has been also prepared for $0.0 \leq x \leq 0.10$. Sample with $x=0.10$ was melted at high temperature, thus it was not further studied. Table 2 shows RQPA up to $x=0.05$. The proposed stoichiometries do not yield to stabilize high temperature polymorphs of dicalcium silicate as high amounts of rankinite [60] ($\text{Ca}_3\text{Si}_2\text{O}_7$) are formed. Thus, this series was ruled out.

Summarising up this section, we must highlight that several solid solutions have been tested but pure belite phases were only formed for the $\text{Ca}_{2-x}\text{B}_x(\text{SiO}_4)_{1-x}(\text{BO}_4)_x$ and $\text{Ca}_{2-x}\text{Na}_x(\text{SiO}_4)_{1-x}(\text{BO}_3)_x$ solid solutions. However, most of these samples showed co-existence of belite polymorphs.

3.2. Microstructural characterization of $\text{Ca}_{2-x}\text{Na}_x(\text{SiO}_4)_{1-x}(\text{BO}_3)_x$. Figure 5 shows a clear diffraction peak broadening with x , behaviour which was not observed in the remaining series. In order

to quantify this evolution, Figure 7 shows the average full width at the half maximum (FWHM) of the diffraction peaks for the two belite phases (β - and α'_H -forms) as function of the diffraction angle. It can be seen that samples with larger x values present higher FWHM values. This behaviour may be due to (i) small (variable) particle sizes and/or (ii) microstrain evolution. Some members of this series were studied by SEM ($x=0.10, 0.15$ and 0.20) and the average particle size is almost identical, see Figure 8. Consequently, peak shape broadening, mainly for large values of x , is chiefly due to microstrains. These microstrains are likely consequences of the ionic radii mismatch of the B/Si and Na/Ca substitutions. It is interesting to note that there are ‘critical’ doping values where the broadening increases notably, $x=0.10$ for β -C₂S and $x=0.20$ for α'_H -C₂S, see Figure 7.

On the other hand, it is known that surfaces of dicalcium silicate particles present stripes which seem to be lamella-like waves related to $\alpha' \rightarrow \beta$ conversion [26,61]. However, these effects were not observed in any of the samples containing B₂O₃ and Na₂O, independently of the phase assemblage. This result is in agreement with a previous report from independent authors [43]. In this series, room temperature unit cell volume of β - and α'_H -forms are very similar, and as x increases $V_{\alpha'_H}$ is even smaller than V_β , see Figure 4. However, in other studies [25] room temperature unit cell volume of α'_H -form is slightly larger (347.5 Å³) than that of β -C₂S (345.8 Å³). The very small differences between unit cell volumes of β - and α'_H -forms, caused by the addition of borax, is likely a key factor in avoiding the formation of the lamella-like microstructure.

3.3. Crystal structure characterization of α'_H -C₂S. Ca_{1.85}Na_{0.15}(SiO₄)_{0.85}(BO₃)_{0.15} composition was selected to further study the crystal structure of room-temperature stabilized α'_H -C₂S. This choice is based on the trade off between sharp diffraction peaks and the highest content of α'_H -phase. We have used the crystal structure description reported by Mumme et al. [25] as a starting model for the Rietveld refinement. After overall parameters optimization, but keeping fixed the structural parameters, the obtained disagreement factors were $R_{WP}=10.8\%$ and $R_F(\alpha'_H\text{-C}_2\text{S})=4.1\%$. The typical Rietveld plot is shown in Figure 9a.

The structural description of α'_H -C₂S polymorph, space group Pnma, has seven crystallographically independent sites in the asymmetric part of the unit cell: 2 Ca's in general position; 1 Si in special position ($x1/4z$); and 4 O in general positions. Moreover, the disorder in the structure is modeled by the occupation factors which are 0.5 for Ca's and O's in stoichiometric α'_H -Ca₂SiO₄.

Firstly, and keeping the occupation factors of the stoichiometric phase, the atomic positional parameters were optimized. The refinement converged to chemically realistic interatomic distances, and consequently, soft constraints were not included in the refinement. Secondly, ADPs or Debye-Waller factors were optimized. As some oxygen ADP values became negative, all oxygen ADPs were grouped together. This refinement converged to $R_{WP}=6.77\%$ and $R_F(\alpha'_H\text{-C}_2\text{S})=3.02\%$, showing a clear improvement respect to the initial structural description. Finally, nominal cation stoichiometry was included in the refinement by substitution of Ca by Na and Si by B. The occupation factors were kept

fixed to the nominal stoichiometry and positional parameters and ADP factors were optimized. The fit was good enough to refine the oxygen ADPs independently, without any negative Debye-Waller factor. The final key R-factors were to $R_{WP}=6.52\%$ and $R_F(\alpha'_H-C_2S)=2.37\%$. Figure 9b gives the fit for this last refinement. Figure 9 also includes an enlarged view of two selected regions (low and high angle ranges) to highlight the improvement in the fit with the final structural description.

Refined atomic positional parameters and isotropic atomic displacement factors are given in Table 3, and also deposited as a CIF file for RQPA. The final refined unit cell parameters for $\alpha'_H-Ca_{1.85}Na_{0.15}(SiO_4)_{0.85}(BO_3)_{0.15}$ were $a=6.8432(2)$ Å, $b=5.4555(1)$ Å, $c=9.2346(2)$ Å and $V=344.76(2)$ Å³. It is worth mentioning that the phase composition was fixed to the nominal starting cation stoichiometry. However, the oxygen disorder induced by the borate incorporation has not been modelled and neutron powder diffraction is needed to address this feature. The disorder in the SiO_4^{4-} units is large and similar to that already reported [25]. Finally, it is also worth noting that this sample contains 90.1(1) wt% of α'_H-C_2S and 9.9(2) wt% of $\beta-C_2S$.

This structural description was used to perform RQPA of the remaining members of all the series containing α'_H-C_2S , Table 2. Furthermore, this structural description has been used to quantify this phase in belite calcium sulfoaluminate (BCSA) clinkers doped with 2.0 wt% of B_2O_3 , added as borax [5]. This clinker contains three main phases, α'_H-C_2S , cubic [62] $Ca_4Al_6O_{12}SO_4$ (also called Klein salt or ye'elimite) and orthorhombic [63] Ca_2AlFeO_5 . RQPA was performed for this clinker by using the refined crystal structure for α'_H-C_2S obtained in this report (Figure S3b) and the Mumme et al. structure [25] (Figure S3a), using exactly the same refinement strategies. Table 4 includes RQPA of doped BCSA clinker being α'_H-C_2S fitted by the refined crystal description of this work (upper row) and with crystal structure published by Mumme et al. [25] It is observed how agreement indices (R_{WP} and R_F for α'_H-C_2S) are lower, i.e. means better fits, when using the refined crystal structure for α'_H -form and consequently, the weight percentage of this phase has increased. We have evaluated the refined unit cell parameters of this α'_H-C_2S included in a BCSA clinker: $a=6.8263(3)$ Å, $b=5.4684(3)$ Å, $c=9.2658(4)$ Å and $volume=345.88(4)$ Å³. Taking into account this refined volume, it can be assessed that the α'_H-C_2S stabilized in a BCSA clinker by adding 2 wt% of B_2O_3 , added as borax, may have a stoichiometry close to $x=0.10$, see Figure 4, using the $Ca_{2-x}Na_x(SiO_4)_{1-x}(BO_3)_x$ mechanism. Finally, the influence of polymorphism and microstructure in the kinetic of the C_2S hydration are being studied and the results will be reported elsewhere.

4. CONCLUSIONS

The boron and sodium/boron doping of dicalcium silicate has been investigated. In order to understand the substitution of silicon by boron, three solid solutions have been tested: $Ca_{2-x/2}\square_{x/2}(SiO_4)_{1-x}(BO_3)_x$, $Ca_2(SiO_4)_{1-x}(BO_3)_xO_{x/2}$ and $Ca_{2-x}B_x(SiO_4)_{1-x}(BO_4)_x$. The experimental results, in the reported synthetic conditions, support the formation of the $Ca_{2-x}B_x(SiO_4)_{1-x}(BO_4)_x$ series.

This doping mechanism similar to that already reported for aluminum doping calcium silicates. On the other hand, the coupled sodium/boron doping of belite has also been investigated by the preparation of two series: $\text{Ca}_{2-x}\text{Na}_x(\text{SiO}_4)_{1-x}(\text{BO}_3)_x$ and $\text{Ca}_{2-3x}\text{B}_{2x}\text{Na}_x(\text{SiO}_4)_{1-x}(\text{BO}_4)_x$. $\text{Ca}_{2-x}\text{Na}_x(\text{SiO}_4)_{1-x}(\text{BO}_3)_x$ series is formed and $\alpha'_\text{H}\text{-C}_2\text{S}$ is the main phase for x values larger than 0.10 and it is single phase for $x=0.25$. Therefore, boron substitutes silicon as tetrahedral borate anion, BO_4^{5-} , for single-boron doping; but as triangular-planar anion, BO_3^{3-} , for coupled boron and sodium doping. Finally, a new structural description for Na/B doping in belite has been developed for $\alpha'_\text{H}\text{-Ca}_{1.85}\text{Na}_{0.15}(\text{SiO}_4)_{0.85}(\text{BO}_3)_{0.15}$. This structure fits better cement samples including reactive belite prepared by borax addition. This behaviour has also been confirmed by ^{11}B MAS NMR as the borax-activated iron-rich belite calcium sulfoaluminate clinker displays a spectrum typical of trigonal-planar BO_3 groups.

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330 **Figure captions.**

331 **Figure 1.** Polymorphic transformations of stoichiometric Ca_2SiO_4 with temperature.

332 **Figure 2.** 3D-view of a selected range of the LXRPD patterns for $\text{Ca}_{2-x}\text{B}_x(\text{SiO}_4)_{1-x}(\text{BO}_4)_x$ solid solution.
333 Symbols highlight diagnostic peaks for $\gamma\text{-C}_2\text{S}$ (rhombus); $\beta\text{-C}_2\text{S}$ (circle) and $\alpha'_\text{H}\text{-C}_2\text{S}$ (square).

334 **Figure 3.** ATR-FTIR infrared spectra for (a) $\text{Ca}_{2-x}\text{B}_x(\text{SiO}_4)_{1-x}(\text{BO}_4)_x$ and (b)
335 $\text{Ca}_{2-x}\text{Na}_x(\text{SiO}_4)_{1-x}(\text{BO}_3)_x$ series with nominal compositions labelled.

336 **Figure 4.** Variation of the unit cell volumes for $\gamma\text{-C}_2\text{S}$ (rhombus), $\beta\text{-C}_2\text{S}$ (circle) and $\alpha'_\text{H}\text{-C}_2\text{S}$ (square)
337 with nominal composition for $\text{Ca}_{2-x}\text{B}_x(\text{SiO}_4)_{1-x}(\text{BO}_4)_x$ (open symbols) and $\text{Ca}_{2-x}\text{Na}_x(\text{SiO}_4)_{1-x}(\text{BO}_3)_x$
338 (closed symbols) series from Rietveld refinements

339 **Figure 5.** 3D-view of a selected range of the LXRPD raw patterns for $\text{Ca}_{2-x}\text{Na}_x(\text{SiO}_4)_{1-x}(\text{BO}_3)_x$ solid
340 solution with nominal compositions labelled. Symbols highlight diagnostic peaks for $\gamma\text{-C}_2\text{S}$ (rhombus);
341 $\beta\text{-C}_2\text{S}$ (circle) and $\alpha'_\text{H}\text{-C}_2\text{S}$ (square).

342 **Figure 6.** ^{11}B MAS NMR spectra for BCSAF-B2 clinker (top), $\text{Ca}_{1.85}\text{Na}_{0.15}(\text{SiO}_4)_{0.85}(\text{BO}_3)_{0.15}$
343 (intermediate) and $\text{Ca}_{1.90}\text{B}_{0.10}(\text{SiO}_4)_{0.90}(\text{BO}_4)_{0.10}$ (bottom). For the sake of comparison, the top-right
344 inset shows the ^{11}B MAS-NMR spectrum (experimental and calculated) for crystalline $\text{B}(\text{OH})_3$ from
345 reference [56] which displays a unique trigonal BO_3 group.

346 **Figure 7.** Variation of FWHM values with diffraction angle (2θ) for $\beta\text{-C}_2\text{S}$ (left) and $\alpha'_\text{H}\text{-C}_2\text{S}$ (right) in
347 $\text{Ca}_{2-x}\text{Na}_x(\text{SiO}_4)_{1-x}(\text{BO}_3)_x$ series (nominal compositions) obtained from Rietveld refinements using a
348 pseudo-Voigt function.

349 **Figure 8.** SEM micrographs of sintered pellets of $\text{Ca}_{2-x}\text{Na}_x(\text{SiO}_4)_{1-x}(\text{BO}_3)_x$ [nominal composition:
350 $x=0.10$ (a) and 0.20 (b)] series.

351 **Figure 9.** LXRPD Rietveld plot ($\lambda=1.5406 \text{ \AA}$) for the sample with nominal composition
352 $\text{Ca}_{1.85}\text{Na}_{0.15}(\text{SiO}_4)_{0.85}(\text{BO}_3)_{0.15}$ (a) using the crystal structure for $\alpha'_\text{H}\text{-C}_2\text{S}$ reported in reference 25 and (b)
353 using the crystal structure reported here. The insets show detailed views of selected low and high angles
354 regions to highlight the quality of the new fit.
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357 **Table 1.** Nominal elemental composition expressed in weight percentages of oxides for each member
 358 of the proposed Ca_2SiO_4 solid solutions.

Nominal stoichiometry	x	CaO	SiO ₂	B ₂ O ₃	Na ₂ O
Ca₂SiO₄	0.00	65.12	34.88	-	-
Ca_{2-x/2}□_{x/2}(SiO₄)_{1-x}(BO₃)_x	0.025	65.22	34.28	0.51	-
	0.10	65.52	32.40	2.09	-
Ca₂(SiO₄)_{1-x}(BO₃)_xO_{x/2}	0.025	65.36	34.14	0.51	-
	0.10	66.09	31.86	2.05	-
Ca_{2-x}B_x(SiO₄)_{1-x}(BO₄)_x	0.0125	64.93	34.56	0.51	-
	0.01875	64.83	34.40	0.76	-
	0.025	64.74	34.24	1.02	-
	0.05	64.36	33.59	2.05	-
	0.10	63.58	32.27	4.15	-
	0.15	62.78	30.90	6.32	-
	0.20	61.95	29.50	8.55	-
Ca_{2-x}Na_x(SiO₄)_{1-x}(BO₃)_x	0.025	64.78	34.26	0.51	0.45
	0.05	64.43	33.63	1.03	0.91
	0.10	63.72	32.34	2.08	1.85
	0.15	63.00	31.01	3.17	2.82
	0.20	62.24	29.64	4.29	3.82
	0.25	61.47	28.23	5.45	4.85
Ca_{2-3x}B_{2x}Na_x(SiO₄)_{1-x}(BO₄)_x	0.025	63.53	34.48	1.54	0.46
	0.05	61.90	34.06	3.11	0.92
	0.10	58.51	33.18	6.41	1.90

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362 **Table 2.** Rietveld quantitative phase analyses for each member of the proposed Ca_2SiO_4 solid
363 solutions.

Stoichiometry	Nominal value x	$\gamma\text{-C}_2\text{S}$	$\beta\text{-C}_2\text{S}$	$\alpha'_\text{H}\text{-C}_2\text{S}$	CaO
Ca_2SiO_4	0.00	58.9(1)	41.1(1)	-	-
$\text{Ca}_{2-x/2}\square_{x/2}(\text{SiO}_4)_{1-x}(\text{BO}_3)_x$	0.025	3.6(3)	89.2(1)	3.4(3)	3.7(2)
	0.10	3.9(6)	58.2(6)	33.1(8)	4.8(3)
$\text{Ca}_2(\text{SiO}_4)_{1-x}(\text{BO}_3)_x\text{O}_{x/2}$	0.025	3.0(3)	89.8(1)	3.7(3)	3.5(2)
	0.10	3.1(6)	59.9(6)	32.9(7)	4.1(3)
$\text{Ca}_{2-x}\text{B}_x(\text{SiO}_4)_{1-x}(\text{BO}_4)_x$	0.0125	1.9(1)	95.5(1)	2.0(1)	0.7(1)
	0.0187	0.9(1)	83.4(1)	15.6(4)	-
	0.025	0.8(1)	77.1(1)	22.2(4)	-
	0.05	0.7(1)	65.5(2)	33.8(3)	-
	0.10	0.6(1)	15.9(3)	83.5(1)	-
$\text{Ca}_{2-x}\text{Na}_x(\text{SiO}_4)_{1-x}(\text{BO}_3)_x$	0.025	36.4(1)	56.6(2)	6.9(2)	-
	0.05	17.0(2)	56.3(2)	26.7(3)	-
	0.10	-	41.8(2)	58.2(1)	-
	0.15*	-	9.9(2)	90.1(1)	-
	0.20	-	8.4(3)	91.6(1)	-
	0.25	-	-	100.0(-)	-
$\text{Ca}_{2-3x}\text{B}_{2x}\text{Na}_x(\text{SiO}_4)_{1-x}(\text{BO}_4)_x$	0.025 [#]	0.3(1)	69.2(1)	23.8(3)	-
	0.05 ^{\$}	0.5(1)	6.9(4)	67.2(1)	-

364 * Sample used for the structural study of $\alpha'_\text{H}\text{-C}_2\text{S}$. [#] It also contains 6.7(2) wt% of $\text{Ca}_3\text{Si}_2\text{O}_7$. ^{\$} It also
365 contains 25.5(2) wt% of $\text{Ca}_3\text{Si}_2\text{O}_7$.

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368 **Table 3.** Refined atom positions and atomic displacement parameters (ADPs) for α'_H -C₂S in the sample
369 Ca_{2-x}Na_x(SiO₄)_{1-x}(BO₃)_x with x=0.15. Space group: Pnma. Refined unit cell: a=6.8432(2) Å,
370 b=5.4555(1) Å and c=9.2346(2) Å.

Atom	x	y	z	S.O.F.*	ADPs / Å ²
Ca1, 8d	0.3270(2)	0.2850(8)	0.5702(2)	0.4625	0.0260(7)
Na1, 8d	0.3270(2)	0.2850(8)	0.5702(2)	0.0375	0.0260(7)
Ca2, 8d	0.9912(2)	0.2709(9)	0.2938(1)	0.4625	0.0107(6)
Na2, 8d	0.9912(2)	0.2709(9)	0.2938(1)	0.0375	0.0107(6)
O1, 8d	1.0027(7)	0.3247(11)	0.5427(5)	0.5000	0.0374(22)
O2, 8d	0.7555(8)	0.0258(10)	0.7086(7)	0.5000	0.0055(20)
O3, 8d	0.6517(7)	0.1838(13)	0.4439(5)	0.5000	0.0279(19)
O4, 8d	0.6973(9)	0.4868(13)	0.6600(7)	0.5000	0.0182(23)
Si1, 4c	0.7745(3)	0.2500	0.5901(3)	0.8500	0.0175(7)
B, 4c	0.7745(3)	0.2500	0.5901(3)	0.1500	0.0175(7)

371 *Space occupation factor

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375 **Table 4.** Rietveld quantitative phase analyses, expressed in weight percentages, for a doped BCSA
 376 clinker, using the refined structure description for $\alpha'_\text{H}\text{-C}_2\text{S}$ (upper row) and the Mumme et al. structure
 377 [25] (italics). R_WP and R_F for $\alpha'_\text{H}\text{-C}_2\text{S}$ agreement factors are also included.

$\alpha'_\text{H}\text{-C}_2\text{S}$	$R_\text{F}(\alpha'_\text{H})$ (%)	$\text{Ca}_4\text{Al}_6\text{O}_{12}\text{SO}_4$	$\text{Ca}_2\text{AlFeO}_5$	R_WP (%)
58.3(2)	4.36	29.4(1)	12.3(2)	4.77
<i>53.7(2)</i>	<i>5.41</i>	<i>32.1(2)</i>	<i>14.2(2)</i>	<i>5.21</i>

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Figure 1

Figure 2

Figure 3

Figure 4

Figure 5

Figure 6

Figure 7

Figure 8

Figure 9