Article

Cement and Clinker Production by Indirect Mechanosynthesis Process

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Abstract: Global cement production has reached 3.9 billion tons. However, the clinkerization process, which is the basis of cement production, is responsible for an approximate annual global CO₂ emission of 2 billion tons. As part of CEMBUREAU’s 5C strategy, the European cement industry aims to achieve carbon neutrality throughout the cement-concrete value chain by 2050. This article is a continuation of the previous article on the indirect mechanosynthesis clinkerization process, which combines mechanical activation (high-energy milling) and thermal treatment at lower temperatures (from 900 °C) than those used for conventional clinkerization to produce clinker. With this process, we manufactured cement and clinker from industrial and laboratory raw mixes, which had to be rectified by adding kaolinite in compliance with the different cement indicators (LSF, SM, AM). The cement and clinker produced by indirect mechanosynthesis (15 min of mechanical activation and heat treatment 900 °C or 1200 °C) were characterized. In order to test the hydraulic properties of the cement produced, cement pastes were made. Mechanical and structural studies were carried out (between 70 and 90% of C₂S). Mechanical tests revealed for 7 curing days, the values of 3.60 and 7.60 MPa at 900 °C and 1200 °C, respectively, in comparison to commercial cements CEM I and CEM III (23.03 and 19.14 MPa).

Keywords: cements; clinkerization; indirect mechanosynthesis; hydraulicity

1. Introduction

According to the European standard NF EN 197-1 [1], “Cement is a hydraulic binder, i.e., a finely ground inorganic material which, when mixed with water, forms a paste which sets and hardens using hydration reactions and processes and which, after hardening, retains its strength and stability even under water”.

The cement is made of Portland clinker, calcium sulfate to control the setting and many minerals additions (slag, fly ashes, pozzolanic materials … etc.) [1]. Cement production starts with extracting and grinding finely raw materials such as limestone and clay to obtain a raw meal. The fine powder is then heated to a sintering temperature of 1450 °C in a cement kiln [2,3]. In 1990, cement production emitted 0.783 kg of CO₂ to produce 1 kg of cement [4]. However, with the effort from academics and the construction sector by introducing different processes to reduce the CO₂ impact. The method used impacts different stages of the cement production process, the most commonly used technique being the substitution of clinker with other materials (fly ash, slag, limestone, etc.) [5–9], the energy efficiency and alternative fuels [4,10–12] and carbon capture and storage [4,10].

Due to all those improvements, in 2017, cement production emitted 0.667 kg of CO₂ to produce 1 kg of cement. However, the effort must continue to achieve carbon neutrality in the construction sector, and this is why the CEMBUREAU launched the “5C approach” to work more actively on the CO₂ emissions and energy consumption of the construction sector including the cement production [13].
Our research project, “clinkerization by indirect mechanosynthesis” aims to reduce the cement production process’s energy consumption and CO₂ emissions.

This article is the continuation of the previous paper [14], which is part of our research project mentioned above. This project aims to produce clinker and cement using the mechanosynthesis process.

The mechanosynthesis is a high-energy ball milling process; the material to be milled is put in a container with balls or bars and will be subjected to repetitive welding/fracturing movement. Due to this movement, metastable or nanocrystalline phases can be produced and also transform crystalline phases into amorphous phases [15–17].

With mechanosynthesis, we can reduce the size and combine solids from the micrometer scale of a powder particle. At the same time, it allows the size of the crystallites (the grains composing a particle) to be reduced to a nanometric size, resulting in nanostructured materials.

Mechanosynthesis can be used in three different ways [18]; the first way is called direct mechanosynthesis, which allows the mechanical activation of materials introduced (one or more simultaneously) into the mill for a short or long period. For cementitious materials, direct mechanosynthesis is carried out for the mechanical activation of by-products for the clinker substitution. This activation improves mechanical performance, especially at a young age, and allows more by-products to be added [19–21]. Different studies used this process to activate by-products for cementitious materials [19,20,22–25] or geopolymeric materials [26–28]. Using planetary ball milling for mechanical activating slag and fly ashes and then substituting 50% CEMI cement, Bouchenfa et al. [20] showed that wet milling with the optimal PCA (process control agent) improved the compressive strength of (50% CEMI/50% milled Fly ashes) for 28 and 90 curing days with +2.88% and +11.03%, respectively, than the reference (100% CEMI). Moreover, the authors [20] found that for (50% CEMI/50% milled slag), the compressive strength increased by 16.1%, 20.9% and 40.7% for 7, 28 days and 90 curing days in comparison to reference. The authors attributed this increase to the pozzolanic reaction activity.

Alex et al. [26] studied the influence of the high-energy grinding atmosphere (air or CO₂) on the geopolymers prepared from milled slag. The authors observed that both air and CO₂ milled slag found good geopolymerization behaviour leading to the high compressive strength of the geopolymer products. According to these studies, using mechanosynthetically activated materials shows improved mechanical performance of cementitious or geopolymeric materials. It has also been shown that activated fly ash improves the performance of bio-based mortars [29].

The second way, known as indirect mechanosynthesis, consists of combining mechanochemical activation (short milling time) with another process. The second process can be a heat treatment (mechanically activated annealing (M2A), mechanically activated sintering (MAS)) [30–32] or another milling (double mechanical alloying (dMA)) [15]. The method used in this study is mechanically activated annealing, which is generally used in the production of refractory materials or ceramics [33]. However, according to the state of the art and the information at our disposal, this method has never been used to produce clinker or cement.

The third way is called mechanochemistry, which refers to any process that uses grinding to trigger chemical reactions. The uses of mechanochemistry include all chemical reactions caused by mechanical activation, such as exchange reactions, reduction/oxidation reactions, decomposition of compounds and phase transformations [34,35]. For example, mechanochemistry is used to prepare a ready-to-use pregeopolymer powder that only needs water to prepare a geopolymer paste [36].

This article is a continuation of a previously published paper [14]. In this part of the project, the indirect mechanosynthesis process was carried out for the laboratory preparation of clinker from a premix (limestone/kaolinite) to obtain clinker production at a temperature of 900 °C.
In the previous study, 80% limestone and 20% clay were used without taking into account the different indicators used by cement manufacturers: lime saturation Factor (LSF), Silica Modulus (SM) and Alumina Modulus (AM). The results obtained showed that the clinker was of a belitic nature with a significant amount of lime. The cement showed a false set with a very high heat release. That is why in this study, a correction is carried out by adding raw kaolinite to the industrial premixes, considering cement indicators (LSF, SM, AM) for producing clinker and cement at temperatures of 900 °C and 1200 °C.

Structural (XRD, PSD) and microstructural studies (SEM) of clinker and cement manufactured by indirect mechanosynthesis were carried out. The hydraulic properties of these cements were also tested.

2. Materials and Methods

2.1. Raw Materials

The limestone was provided for this work by OMYA® (Nièvre, France), containing > 97% of calcite (Calcium carbonate CaCO₃). The kaolinite used was supplied by Proclay® (Beaujard site—Poigny, France); this clay is constituted of quartz and kaolinite. The F1lab. is a premix reconstituted in the laboratory with 80% limestone and 20% kaolinite. Concerning the F2ind. and F3ind., these 2 premixes were provided by 2 different cement manufacturers. Calcium sulfate hemihydrate (CaSO₄, ½ H₂O) was used to regulate the hydration of C₃A. In the study of the hydraulicity of the cements prepared by indirect mechanosynthesis, we compared our cements to industrial cements, CEMI 52.5 N and CEM III C 32.5 N-LH (HOLCIM, France).

Table 1 shows the chemical composition of the premixes used to produce clinker and cement by indirect mechanosynthesis.

Table 1. Chemical composition of the industrial and laboratory premix used.

<table>
<thead>
<tr>
<th></th>
<th>CaO</th>
<th>SiO₂</th>
<th>Al₂O₃</th>
<th>Fe₂O₃</th>
<th>MgO</th>
<th>K₂O</th>
<th>TiO₂</th>
<th>SO₃</th>
<th>MnO</th>
<th>Na₂O</th>
<th>LSF *</th>
<th>SM **</th>
<th>AM ***</th>
</tr>
</thead>
<tbody>
<tr>
<td>Limestone</td>
<td>96.79</td>
<td>1.3</td>
<td>0.63</td>
<td>0.22</td>
<td>0.5</td>
<td>0.16</td>
<td>0</td>
<td>0</td>
<td>0.07</td>
<td>0</td>
<td>0.78</td>
<td>0.47</td>
<td>0.86</td>
</tr>
<tr>
<td>Kaolinite</td>
<td>0.5</td>
<td>69.37</td>
<td>24.08</td>
<td>1.54</td>
<td>0.6</td>
<td>0.38</td>
<td>2.85</td>
<td>0.3</td>
<td>0</td>
<td>0</td>
<td>0.78</td>
<td>0.78</td>
<td>1.00</td>
</tr>
<tr>
<td>F1lab.</td>
<td>77.53</td>
<td>14.91</td>
<td>5.32</td>
<td>0.48</td>
<td>0.52</td>
<td>0.2</td>
<td>0.57</td>
<td>0.06</td>
<td>0</td>
<td>0</td>
<td>1.62</td>
<td>2.57</td>
<td>10.99</td>
</tr>
<tr>
<td>F2ind.</td>
<td>73.01</td>
<td>13.83</td>
<td>4.64</td>
<td>3.49</td>
<td>2</td>
<td>1.55</td>
<td>0.23</td>
<td>0.67</td>
<td>0.13</td>
<td>0</td>
<td>1.58</td>
<td>1.7</td>
<td>1.33</td>
</tr>
<tr>
<td>F3ind.</td>
<td>75.44</td>
<td>12.59</td>
<td>4.58</td>
<td>3.48</td>
<td>1.2</td>
<td>1.45</td>
<td>0</td>
<td>0.42</td>
<td>0.06</td>
<td>0</td>
<td>1.76</td>
<td>1.56</td>
<td>1.32</td>
</tr>
<tr>
<td>Calcium sulfate hemihydrate C₃S</td>
<td>40.21</td>
<td>1.41</td>
<td>0.67</td>
<td>0.08</td>
<td>0.7</td>
<td>0.08</td>
<td>0</td>
<td>51.03</td>
<td>0</td>
<td>5.3</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

*: Lime saturation factor; **: Silica Modulus, ***: Alumina Modulus.

2.2. Methods

2.2.1. Indirect Mechanosynthesis

The indirect mechanosynthesis clinkerization process is made by combining 2 techniques: a high-energy ball milling to mechanically activate the premix and heat treatment to perform the clinkerization, as shown in Figure 1.

A planetary ball mill (PM 400, RETSCH, Haan, Germany) was used to perform the mechanical activation. This device has a planetary disc on which four vials are mounted. The vials rotate in the opposite direction of the disc. In this study, we choose to use a ball/powder ratio of 4 and a milling time of 15 min, and the rotation speed of the disc and vials are 400 and 800 rpm, respectively.

Before the clinkerization, the premix modification is performed by adding kaolinite until reaching the LSF, SM and AM 0.97 ± 0.05, 2.00 ± 0.50 and 2.0 ± 0.50 (Table 2) chosen according to the index used in the cement industries and also found in the literature [2,37]:

- LSF (lime saturation factor) = \( \frac{\text{CaO}}{(2.8 \times \text{SiO}_2) + (1.18 \times \text{Al}_2\text{O}_3) + (0.65 \times \text{Fe}_2\text{O}_3)} \)
- SM (Silica Modulus) = \( \frac{\text{SiO}_2}{\text{Al}_2\text{O}_3 + \text{Fe}_2\text{O}_3} \)
- AM (Alumina modulus) = \( \frac{\text{Al}_2\text{O}_3}{\text{Fe}_2\text{O}_3} \)
Figure 1. Illustration of the clinkerization process by indirect mechanosynthesis.

Table 2. Chemical composition of the mixes used for the clinkerization process.

<table>
<thead>
<tr>
<th></th>
<th>CaO</th>
<th>SiO₂</th>
<th>Al₂O₃</th>
<th>Fe₂O₃</th>
<th>MgO</th>
<th>K₂O</th>
<th>TiO₂</th>
<th>SO₃</th>
<th>MnO</th>
<th>LSF *</th>
<th>SM **</th>
<th>AM ***</th>
</tr>
</thead>
<tbody>
<tr>
<td>F1lab. corrected</td>
<td>66.93</td>
<td>20.71</td>
<td>7.3</td>
<td>2.99</td>
<td>0.52</td>
<td>0.23</td>
<td>0.81</td>
<td>0.1</td>
<td>0.05</td>
<td>0.98</td>
<td>2.01</td>
<td>2.44</td>
</tr>
<tr>
<td>F2ind. corrected</td>
<td>65.03</td>
<td>19.94</td>
<td>6.78</td>
<td>3.27</td>
<td>1.85</td>
<td>1.42</td>
<td>0.52</td>
<td>0.63</td>
<td>0.12</td>
<td>0.99</td>
<td>1.98</td>
<td>2.07</td>
</tr>
<tr>
<td>F3ind. corrected</td>
<td>65.32</td>
<td>20.26</td>
<td>7.21</td>
<td>3.2</td>
<td>1.12</td>
<td>1.31</td>
<td>0.38</td>
<td>0.4</td>
<td>0.05</td>
<td>0.97</td>
<td>1.94</td>
<td>2.24</td>
</tr>
</tbody>
</table>

*: Lime saturation factor; **: Silica Modulus, ***: Alumina Modulus.

The heat treatment was performed by a high-temperature bottom loading furnace (BLF 17/3, Carbolite Gero, Sheffield, UK). Different temperatures of 900°C and 1200°C are applied for heat treatment of 60 min.

This study investigated cement production by indirect mechanosynthesis using a different method for controlling C₃A hydration. This method consisted of introducing the calcium sulfate "$C₅S$" into the premix before using the indirect mechanosynthesis (Figure 2) to show that it is possible to produce cement directly with our process without additional milling.

2.2.2. Characterization of Powders (Clinkers & Cements) and Cements Pastes

The chemical composition of all materials used in these studies was conducted by X-ray fluorescence spectroscopy (S2 Ranger, BRUKER, Karlsruhe, Germany). The structural analysis includes an X-ray diffraction diffractometer (D2 Phaser, BRUKER, Karlsruhe, Germany) with a continuous scanning mode and Cu Kα radiation ($\lambda = 0.1541$ nm). For the XRD analysis, we used a 2θ range of 2–70° with a 0.02° increment for each 0.15 s. For the evaluation of the XRD analysis, the software DIFFRAC.EVA (BRUKER, Karlsruhe, Germany) with ICDD PDF 4 (Newtown Square, PA, USA) was employed. For the quantification phases analysis, the DIFFRAC.TOPAS software was used (BRUKER, Karlsruhe, Germany).

The particle size distributions are measured with an LS 13 320 XR laser particle size analyzer from BECKMAN COULTER (Villepinte, France). The dispersion method of samples in liquids with a size distribution between 0.01 and 2000 μm is used. In this study, the liquid used is ethanol to avoid the hydration of the samples studied.

The sample dispersion method is coupled with PIDS (polarization intensity differential scattering) technology for better accuracy for very fine particles.
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To study the mechanical performances of the cements produced by indirect mechanosynthesis, cement pastes were made with the addition of water. Consistency tests were carried out to determine the optimal quantity of water to be added as indicated in the NF EN 196-3 standard. Thus, water/binder ratios from 0.7 to 0.9 were used to prepare the cement pastes, which were then introduced into cylindrical tubes of Φ: 35 mm, L: 70 mm (Error: Reference source not found). The samples were unmoulded after 24 h and stored in the water tank at room temperature for 7 curing days. In addition, commercial cements CEM I and CEM III pastes were made to compare their mechanical performances with our cement. Compression tests were carried out with 300 kN equipment (Syntech, 3R, Montauban, France). For each product mechanically tested, 4 samples were produced with a loading speed of 2.4 kN/s.

For the morphological observation of the clinkers and cement made by the indirect mechanosynthesis, a scanning electron microscope (SUPRA 55VP, Zeiss, Oberkochen, Germany) coupled with an energy-dispersive spectrometer (EDS) (QUANTAX, BRUKER, Karlsruhe, Germany) from the ISTeP laboratory from Sorbonne university was used.

3. Results and discussions

3.1. Clinker Production

3.1.1. XRD

Figure 3 shows the diffraction patterns of the three premixes used in our study. Three crystalline structures in those materials, rhomboidric calcite (PDF#01-083-1762), triclinic kaolinite (PDF#00-003-0058) and hexagonal quartz (PDF#00-046-1045) are found in those premixes.
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Table 3 shows the crystal phase quantifications of the premixes constituent used in this study. It is found that the calcite (CaCO$_3$) is the predominant part, the quartz (SiO$_2$) is present but at a lower intensity, and kaolinite (Al$_2$Si$_2$O$_5$(OH)$_4$) has the lowest presence.

Table 3. Phases quantification (TOPAS) of the premixes used.

<table>
<thead>
<tr>
<th></th>
<th>F1lab. Corrected</th>
<th>F2ind. Corrected</th>
<th>F3ind. Corrected</th>
</tr>
</thead>
<tbody>
<tr>
<td>Quartz</td>
<td>16.50%</td>
<td>15.26%</td>
<td>15.52%</td>
</tr>
<tr>
<td>Calcite</td>
<td>72.95%</td>
<td>81.46%</td>
<td>79.05%</td>
</tr>
<tr>
<td>Kaolinite</td>
<td>10.75%</td>
<td>3.28%</td>
<td>5.43%</td>
</tr>
</tbody>
</table>

Figure 4 illustrates the F1lab. diffraction pattern corrected by the addition of kaolinite. The diffraction pattern of the milled premix Flab1.corrected shows us the presence of some peaks of kaolinite after 15 min of milling. In the previous study [14], at 15 min of milling, the total disappearance of the kaolinite peaks is recorded. This difference may be due to the amount of kaolinite introduce is higher in this current study.

As shown in the previous article, indirect mechanosynthesis produces the main constituent of the clinker (C$_3$S, C$_2$S, C$_3$A, C$_4$AF), as observed in Figure 4. Some quartz peaks remain for the heat treatment of 900 $^\circ$C, but their intensity has significantly decreased, as seen in Tables 3 and 4. The amount of quartz dropped from 16.50% to 7.34%.

For the heat treatment of 1200 $^\circ$C, all quartz peaks disappeared and were quantified at 0.24% with the TOPAS software. For lime (CaO), all peaks have disappeared from the diffraction pattern. Therefore, the quantification of the remaining lime phase is 1.62%.

The other clinkers produced by indirect mechanosynthesis (Figure 5) show the same results when the corrected F2ind. and F3ind. are used. The clinker’s main constituents (C$_3$S, C$_2$S, C$_3$A, C$_4$AF) are also produced for heat treatment of 900 and 1200 $^\circ$C. Besides, it is observed in the diffraction pattern that quartz and lime (CaO) are still present in the clinkers produced at 900 $^\circ$C and disappear entirely for heat treatment of 1200 $^\circ$C.
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The other clinkers produced by indirect mechanosynthesis (Figure 5) show the same results when the corrected F\textsubscript{2}ind. and F\textsubscript{3}ind. are used. The clinker's main constituents (C\textsubscript{3}S, C\textsubscript{2}S, C\textsubscript{3}A, C\textsubscript{4}AF) are also produced for heat treatment of 900 and 1200 °C. Besides, it is observed in the diffraction pattern that quartz and lime (CaO) are still present in the clinkers produced at 900 °C and disappear entirely for heat treatment of 1200 °C.

Table 4 provides the quantification of the phases of all clinkers made by indirect mechanosynthesis with the 3 premixes used.

Table 4. Phases quantifications (TOPAS) of the clinkers produced.

<table>
<thead>
<tr>
<th></th>
<th>F1lab. 15'M</th>
<th>F2ind. 15'M</th>
<th>F3ind. 15'M</th>
</tr>
</thead>
<tbody>
<tr>
<td>(\alpha)-C\textsubscript{2}S</td>
<td>4.87%</td>
<td>12.12%</td>
<td>6.93%</td>
</tr>
<tr>
<td>(\beta)-C\textsubscript{2}S</td>
<td>49.14%</td>
<td>69.43%</td>
<td>42.41%</td>
</tr>
<tr>
<td>C\textsubscript{3}S</td>
<td>2.44%</td>
<td>2.21%</td>
<td>4.94%</td>
</tr>
<tr>
<td>C\textsubscript{3}A</td>
<td>13.59%</td>
<td>5.38%</td>
<td>13.1%</td>
</tr>
<tr>
<td>C\textsubscript{4}AF</td>
<td>19.3%</td>
<td>9%</td>
<td>20.69%</td>
</tr>
<tr>
<td>Lime</td>
<td>3.31%</td>
<td>1.62%</td>
<td>5.5%</td>
</tr>
<tr>
<td>Quartz</td>
<td>7.34%</td>
<td>0.24%</td>
<td>6.43%</td>
</tr>
</tbody>
</table>

Table 4 provides the quantification of the phases of all clinkers made by indirect mechanosynthesis with the 3 premixes used.

It is shown that regardless of the premix and the heat treatment temperature, a belite clinker is produced with a large proportion of \(\beta\)-C\textsubscript{2}S. In addition, with a higher temperature, the production of belite is increased. For the alite (C\textsubscript{3}S) constituent, the amount produced at 900 °C and 1200 °C is very low. For tricalcium aluminate (C\textsubscript{3}A) the production is higher at 900 °C than at 1200 °C. The amount of tetracalcium aluminate ferrite (C\textsubscript{4}AF) was higher with heat treatment of 900 °C.
Figure 5. Diffraction patterns of the milled premix F2ind. and F3ind. corrected and the clinkers produced by indirect mechanosynthesis.
In the current study, the CaO phase is produced less (<6%), contrary to the previous article [14]. Where a significant amount of CaO reached 11%, this can be explained by the kaolinite added to the F1lab. (used in the previous article), which has increased the amount of SiO₂ in the premix and transformed it into another constituent (C₃S, C₂S).

3.1.2. Particle Size Distribution (PSD)

Through mechanosynthesis, the clinker obtained is in powder form, unlike the clinker obtained with conventional clinkerization, which is in the form of nodules. For this reason, a comparison of the PSD of the Clinker F2ind. 900 °C and that of the premix F2ind. and also that of a commercial Portland cement (CEM I) was made.

Figure 6 and Table 5 shows that the clinker produced (clinker F2ind. 900 °C) has a very similar particle distribution to that of commercial cement. However, it is shown that for F2ind. 900 °C clinker, agglomerations are formed, which may be due to high energy milling [20,33,38]. Moreover, it can be attributed to the sintering (heat treatment). As a result of the indirect mechanosynthesis process, the clinker produced is in powder form. This result will impact the reduction in the energy used for the milling clinker when the gypsum is introduced for cement production.

In the case of the PSD of F2ind. 1200 °C clinker, the agglomeration phenomenon is more critical due to the higher heat treatment temperature used in this case (1200 °C) as observed in Table 5, where the D (90) of the cement produced with a 1200°C is higher than for the 900 °C. Therefore, grinding may be necessary for clinkers produced above 900 °C.

![Figure 6. Particle size distribution of CEMI and clinker produced by indirect mechanosynthesis.](image)

| Table 5. Median particle sizes: D (10), D (50) and D (90) of CEM I, F2ind, Clinker F2ind. 900 °C. |
|-----------------|-----------------|-----------------|
|                  | D (10)          | D (50)          | D (90)          |
| CEMI             | 4.25            | 14.47           | 32.88           |
| F2ind.           | 1.36            | 6.35            | 80.22           |
| Clinker F2ind. 900 °C | 3.56            | 23.22           | 78.84           |
| Clinker F2ind. 1200 °C | 0.67            | 15.15           | 120.8           |
3.2. Cement Production

In this section, the results presented are obtained on cements produced from the F3ind. corrected premix.

3.2.1. XRD

Figure 7 shows the diffraction patterns of cements produced by adding calcium sulfates (C\textsubscript{S}) during the milling process (mechanical activation). The presence of the main crystal phases that should be present in clinker and cement can be observed. Furthermore, it is shown that C\textsubscript{2}S, C\textsubscript{3}S, C\textsubscript{3}A and C\textsubscript{4}AF are present in all the cements produced regardless of the temperature of the heat treatment and the amount of C\textsubscript{S} added before the milling.

Figure 8 shows the phase quantifications of the anhydre cement by indirect mechanosynthesis. It is found that our cements have a belitic nature, depending on the amount of C\textsubscript{S} and there is no big variation in the production of C\textsubscript{2}S with heat treatment of 900 °C. However, it is observed that the amount of belite produced increases with the increase in C\textsubscript{S} added with heat treatment of 1200 °C.

It is observed that the production of C\textsubscript{3}S was low whatever the temperature or the amount of C\textsubscript{S} used in order to produce cement. For the production of C\textsubscript{3}A, it was found that the production was also small.

Concerning the production of C\textsubscript{4}AF, a significant amount was obtained for all the cement produced at 900 °C 15.04%, 17.03%, 16.55%, respectively, for 4%, 6%, 10% of C\textsubscript{S} used. For the cement produced at 1200 °C, only the cement prepared with 4% C\textsubscript{S} produced a significant amount of C\textsubscript{4}AF. As for the cement produced with 6 and 10% of C\textsubscript{S} the amount of C\textsubscript{4}AF found was not higher than 2.5% for the two cements.

Figure 9 gives the phase quantifications of the hydrated cement produced by indirect mechanosynthesis after 7 curing days.

The first results observed are the high amount of the belite residues after 7 days of curing, which may be due to the low reactivity of the C\textsubscript{2}S. In literature, it is shown that belite can take up two days of the induction period before the start of hydration and consequently can slow down the production of CSH [39].

It is also noted that small amounts of portlandite are produced during hydration; this can be attributed to the low amount of C\textsubscript{3}S obtained during the production process and the low reactivity of the C\textsubscript{2}S.

It was also observed that the production of ettringite was significant, and the highest production was observed with the cement with 10% C\textsubscript{S} at 1200 °C (29.89%), ettringite production at 900 °C was also highest compared to the two other cements produced at the same temperature. The ettringite production is due to the hydration of two constituents of the clinker C\textsubscript{3}A and C\textsubscript{4}AF. The equation of the hydration of C\textsubscript{3}A [2] is given in the following equations:

\[
C\textsubscript{3}A + 3C\textsubscript{S}H_2 + 26H_2O \rightarrow C\textsubscript{6}A\textsubscript{3}H_{32} \tag{1}
\]

and concerning the equation for C\textsubscript{4}AF hydration with gypsum might be expressed by one of the following equations [38]:

\[
C\textsubscript{4}AF + 4CaSO\cdot 2H_2O + 35\frac{1}{2}H_2O \rightarrow \frac{4}{3}C\textsubscript{3}(A_{0.75}, F_{0.25}) \cdot 3C\textsubscript{S}H_{31} + \frac{2}{3}FH_3 \tag{2}
\]

\[
C\textsubscript{4}AF + 4CaSO\cdot 2H_2O + 35\frac{1}{3}H_2\frac{4}{3}C\textsubscript{3}(A_{0.5}, F_{0.5}) \cdot 3C\textsubscript{S}H_{31} + \frac{1}{3}AH_3 + \frac{1}{3}FH_3 \tag{3}
\]

\[
C\textsubscript{4}AF + 3CaSO\cdot 2H_2O + 29H_2O \rightarrow C\textsubscript{3}A \cdot 3C\textsubscript{S}H_{31} + CH + FH_3 \tag{4}
\]
Figure 7. Diffraction patterns of cement produced by indirect mechanosynthesis at different heat treatments.
Ettringite Cement with 10% of \( C_S \)

Cement with 6% of \( C_S \)

Cement with 10% of \( C_S \)

![Figure 8. Phases quantifications of the anhydric cement produced by indirect mechanosynthesis.](image)

![Figure 9. Phases quantifications of the hydrated cement produced by indirect mechanosynthesis after 7 days of curing.](image)

### 3.2.2. Particle Size Distribution of Cement

Figure 10 shows the particle size distribution of cement produced by indirect mechanosynthesis. The cement is obtained in a powder form for heat treatment of 900 °C as the clinker is shown in Figure 1.

The PSDs analysis shows that the cement produced with 4% \( C_S \) has a multi-modal distribution, where the first range is found between 0.27 mm and 1.45 mm with a maximum peak at 0.68 mm and the second range is located between 2.1 mm to 5.8 mm with a maximum peak at 2.3 mm.
Figure 10 shows the particle size distribution of cement produced by indirect mechanosynthesis. The cement is obtained in a powder form for heat treatment of 900 °C as the clinker is shown in Figure 1.

The PSDs analysis shows that the cement produced with 4% $\text{CS}$ has a multi-modal distribution, where the first range is found between 0.27 mm and 1.45 mm with a maximum peak at 0.68 mm and the second range is located between 2.1 mm to 5.8 mm with a maximum peak at 2.3 mm. The third range is found between 5.8 mm and 31.5 mm, with a maximum peak at 19.6 mm. The fourth range is found between 31.5 mm and 60.52 mm, with a maximum peak at 41.61 mm. Finally, the last range is located between 60.52 mm and 105.93 mm, with a maximum peak at 80.074 mm.

The cement produced with 6% $\text{CS}$ has a monomodal distribution; the range is found between 0.16 mm and 295.53 mm, with a maximum peak at 18.00 mm.

Concerning the cement produced with 10% $\text{CS}$ with a monomodal distribution, the range is found between 0.16 mm and 185.38 mm, with a maximum peak of 19.76 mm.

Concerning the heat treatment of 1200 °C during the indirect mechanosynthesis process, the particles sintered together and created larger cement particles. In this case, the milling is suitable for cement to be used in the best conditions.

3.2.3. Compressive Strength (CS) of the Cement Paste

To study the hydraulicity of the cement produced by indirect mechanosynthesis, cement pastes were made and mechanically tested and compared to two conventional cements.

Table 6 shows the CS of all the pastes tested after 7 days of curing. For the two conventional cements, the CS of the CEM I and III reached 23.03 and 19.04 MPa after 7 days of curing. Concerning the three cements produced at 900 °C with 10%, 6% and 4% of $\text{CS}$ showed fewer performances than the commercial cement. The best results found with our cements are for an addition of 10% of $\text{CS}$ with 3.60 MPa (900 °C) and 7.60 MPa (1200 °C).

Table 6. Compressive strength at 7 days of curing of the cement pastes produced by indirect mechanosynthesis.

<table>
<thead>
<tr>
<th>Cement Paste</th>
<th>Compressive Strength (CS) (MPa)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>900 °C</td>
</tr>
<tr>
<td>10% $\text{CS}$</td>
<td>3.60</td>
</tr>
<tr>
<td>6% $\text{CS}$</td>
<td>2.93</td>
</tr>
<tr>
<td>4% $\text{CS}$</td>
<td>3.67</td>
</tr>
<tr>
<td>CEM I</td>
<td>23.03</td>
</tr>
<tr>
<td>CEM III</td>
<td>19.14</td>
</tr>
</tbody>
</table>

This leak of performance compared to the performances of the commercial cements can be explained by the small amount of the $\text{C}_3\text{S}$ produced during the cement production.
by our process compared to the commercial cement. The small performances can also be
due to the low reactivity of C\textsubscript{2}S for the production of C-S-H and a different PSD.

According to the results obtained in the production of cementitious pastes, it can
be considered that our cements have performances close to those of masonry cement or
hydraulic road binder.

### 3.2.4. Morphology of the Cement Produced

The morphology of the cement produced at 900 and 1200 °C by indirect mechanosyn-
thesis was studied, as shown in Figure 11.

![Figure 11. SEM micrographs showing the particles of the anhydre cement produced by indirect
mechanosynthesis at (a) 900 °C (b) 1200 °C.](image)

The first picture (on the left) is related to the cement produced at 900 °C. Some spherical
particles that may be related to the belite are observed [39]. For cement produced at 1200 °C,
the shape of the belitic particles seems smoother and more abundant.

Figure 12 shows the morphology of the hydrated cement produced by our process of
indirect mechanosynthesis. It is found that the presence of ettringite in the shape of needle
particles, as confirmed by the X-ray diffraction analysis. The EDS analysis confirmed the
ettringite presence, as shown in Figure 12. Some particles in the shape of some hexagonal
particles form are observed and attributed to the portlandite.

It can also be seen from the photo (a,c) that there is still a significant amount of spherical
particles associated with the belite, which confirms the result of the Rietveld quantification.
4. Conclusions

The previous work published in 2020 shows that clinkerization by mechanosynthesis allows us to produce clinker with heat treatment of 900 °C. In this work, we have continued the manufacture of cement and clinker using an indirect mechanosynthesis process by introducing CS before milling.

It was found that the cements produced are belitic cements. Phase quantifications by TOPAS software showed us that the belite is generally around 70%, with a more significant amount produced using 10% CS (92.29%) at 1200 °C. It was also noted that there was a significant production of C4AF in cements produced at 900 °C. The presence of belite is confirmed by the SEM analysis, with spherical particles characteristic of belite particles.

Regarding the mechanical performances of the cements produced by indirect mechanosynthesis, it is remarked that in general, the performances at 7 curing days of the pastes produced with our cements are weaker with 10% CS 3.60 and 7.60 MPa at 900 °C and 1200 °C, respectively, than the pastes produced with commercial CEM I and CEM III cements (23.03 and 19.14 Mpa, respectively).

The structural analysis of the cementitious pastes produced with our process showed that at 7 days, the quantity of remaining belite is high, resulting in a lack of reactivity. In addition, the production of portlandite is also low, which corroborates the lack of reactivity of C2S and the low presence of C3S in our cements. Finally, the important presence of ettringite in our cements is due to the hydration of C3A and C4AF in the anhydrous cement.
The next step of the study will be to work on the tempering of the products after the thermal treatments because we suspect that during the slow cooling of the product, transformations in the different structures are produced.

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**References**