Production of agroindustrial ashes with pozzolanic activity via acid leaching, conjugated burning and ultrafine grinding

Guilherme Chagas Cordeiro  
Mônica Nunes Lemos  
Kristian Vinco Xavier  
Charles Prado Ferreira de Lima

Abstract

This paper aims to study the production and the physical and chemical characterization of rice husk, sugarcane bagasse, corn straw, and bamboo leaf ashes to use as pozzolans. The ash production steps basically consisted of raw material collection, chemical treatment by citric acid leaching, conjugated burning (thermal processing) and ultrafine grinding (mechanical processing) of the produced ashes. The results indicated that all ashes could be classified as pozzolanic additions based on specific standards. The high pozzolanicity of the ashes was confirmed by testing the electrical conductivity, mechanical performance index, and hydration kinetics of cement pastes by isothermal calorimetry. Therefore, the production processes used in this work were effective in producing highly reactive pozzolanic materials.

Keywords: Pozzolan. Agroindustrial ash. Ultrafine grinding. Conjugated burning. Acid leaching.

Resumo

Este trabalho objetiva estudar a produção e as características físicas e químicas de cinzas da casca de arroz, do bagaço da cana-de-açúcar, da folha de milho e da folha de bambu com vistas à aplicação como pozolanas. As etapas de produção das cinzas consistiram basicamente de coleta e preparação das matérias-primas, tratamento químico por lixiviação com ácido cítrico, queima conjugada (beneficiamento térmico) e moagem ultrafina das cinzas produzidas (beneficiamento mecânico). Os resultados indicaram que todas as cinzas atenderam aos requisitos de normas específicas e foram classificadas como adições pozolânicas. A elevada pozolanicidade das cinzas foi confirmada em ensaios de condutividade elétrica, de índice de desempenho e pelas mudanças na cinética de hidratação de pastas cimentícias por calorimetria isotérmica. Portanto, os mecanismos de beneficiamento utilizados na produção das diferentes cinzas foram eficazes para geração de materiais pozolânicos altamente reativos.

Palavras-chave: Pozolana. Cinza agroindustrial. Moagem ultrafina. Queima conjugada. Lixiviação ácida.
Introduction

Civil construction is characterized by a set of processes that have significant environmental impacts due to high energy consumption and natural resources, in addition to greenhouse gas emissions and waste generation. One of the main environmental problems of civil construction is associated with the Portland cement production, which is the most used construction material around the world (GARTNER, 2004). Civil construction-related activities were responsible for around 2.6% of greenhouse gas intensifying emissions in Brazil in 2016 (AGÊNCIA FAPESP, 2016), especially CO₂, which is released in large quantities during the production of Portland clinker.

Considering this, using industrial and agroindustrial wastes in partial replacement of Portland cement has been a viable alternative to mitigate the impacts of cement production, enabling the reuse of these waste materials (GANESAN; RAJAGOPAL; THANGAVEL, 2007). Pozzolanic and filler materials are used as mineral additions in commercially available cements worldwide. Pozzolans are defined as reactive aluminosilicates (AₓSᵧ), which can be amorphous (e.g. silica fume) or partially crystalline (e.g. metakaolin). Calcium silicate hydrate (C-S-H) and calcium aluminate hydrate (C-A-H) are the main products of the pozzolan reaction with portlandite (CH) in the presence of water (H), which are shown in Equation 1. The precise composition of the CSH and CAH varies with the reaction conditions and the presence of other components (GARTNER, 2004). Calcium aluminosilicates (C-A-S-H) can also be formed in reactions involving pozzolans with significant amounts of amorphous alumina (RICHARDSON; GROVES, 1993).

\[ A_{x}S_{y} + mCH + nH \rightarrow C-S-H + C-A-H \]  

Studies on pozzolans have shown improvements in cementitious properties regarding rheological behavior (CORDEIRO; TOLEDO FILHO; FAIRBAIRN, 2008), durability (MALHOTRA; MEHTA, 1996; CORDEIRO; TOLEDO FILHO; FAIRBAIRN, 2008; HOPPE FILHO et al., 2014) and mechanical performance (MALHOTRA; MEHTA, 1996; CORDEIRO; TOLEDO FILHO; FAIRBAIRN, 2008; CORDEIRO et al., 2018). This is due to the increased formation of hydrates because of pozzolanic reactions (MALHOTRA; MEHTA, 1996; SOUZA et al., 2014) and the filler effect on cement pastes provided by the fine or ultrafine particles of mineral addition (LAWRENCE; CYR; RINGOT, 2003; CORDEIRO et al., 2008).

Pozzolans from waste generally need different processes to optimize their reactivity. In this respect, it is worth mentioning using thermal (SUGITA, 1994; SABIR; WILD; BAI, 2001; CORDEIRO; SALES, 2016; ROSSIGNOLO et al., 2018), mechanical (CORDEIRO; TAVARES; TOLEDO FILHO, 2016) and chemical (REAL; ALCALÁ; CRIADO, 1996; KRISHNARAO; SUBRAHMANYA; JAGADISH KUMAR, 2001) processes used in the pozzolan production. Concerning chemical processes, Vayghan, Khaloo and Rajabipour (2013) showed that waste treatments with acid leaching can enhance the performance of ash in cementitious systems by increasing the concentration of amorphous silica due to the removal of metallic oxides, such as Al₂O₃, Fe₂O₃, CaO, MgO, Na₂O, and K₂O.

In this context, the present paper aims to study the production and main physical and chemical properties of ashes produced from sugarcane bagasse, rice husk, corn straw, and bamboo leaf to use as highly reactive pozzolans. Choosing these wastes is justified by the wide supply of bagasse, rice husk, and corn and bamboo leaves in Brazil, estimated at around 20 million tons in 2019 (INSTITUTO...; 2020; JUDZIEWICZ et al., 1999), and by the predominant presence of silica in its chemical compositions (ZHANG; MALHOTRA, 1996; MARTIRENA HERNÁNDEZ et al., 1998; CORDEIRO et al., 2008; ADESANYA; RAHEEM, 2009; VILLAR-COCIÑA et al., 2011).

Materials and methods

Rice husk ash (RHA) was used as a control pozzolan since this material has high pozzolanic activity (MALHOTRA; MEHTA, 1996; TASHIMA et al., 2012), especially when it is produced from burning acid-leached husk (REAL; ALCALÁ; CRIADO, 1996; KRISHNARAO; SUBRAHMANYA; JAGADISH KUMAR, 2001). RHA was produced from a rice husk sample collected in the city of Mogi das Cruzes-SP (Brazil). Samples of bagasse and bamboo leaf were collected in the city of Campos dos Goytacazes-RJ (Brazil) and used for producing sugarcane bagasse (SBA) and bamboo leaf (BLA) ashes, respectively. The corn straw sample used to produce ash (denominated CSA) was obtained from the city of Castelo-ES (Brazil).
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Figure 1 shows a flowchart with the sequence of activities used to produce the three ashes. After collection, all raw materials were washed in water to eliminate any impurities from the collection environment, followed by oven-drying at 100 °C for 24 h. To produce RHA, CSA, and BLA, the raw materials were subjected to chemical treatment via leaching with a 10% anhydrous citric acid solution (C₆H₈O₇). The leaching of the materials was carried out for 72 h at room temperature at a proportion of 60 g of raw material to 1 L of acid solution. Afterwards, each sample was rinsed continuously with deionized water and then filtered with qualitative filter paper until pH neutralization. Finally, the leached material was oven-dried at 100 °C for 24 h. These conditions were adapted from Cordeiro, Vieira and Lopes (2017) after changing the type of acid (HCl in the previous study).

After leaching, the rice husk, corn straw, and bamboo leaf were subjected to conjugate burning, which consisted of burning in two stages according to the procedures suggested by Andreão et al. (2019). In the conjugate burning, the leached materials were initially burned in a pilot kiln adapted from Sugita (1994) using the biomass as fuel to reduce the amount of organic matter (volume reduction) and energy savings. This preliminary autogenous burning was not controlled, although the temperature was continuously monitored with thermocouples to ensure that it did not exceed 800 °C, a temperature beyond which silica crystallization may occur (DELLA; KÜHN; HOTZA, 2002). Figure 2 shows the typical temperature evolution over 10 h of burning in the pilot kiln. In this case, it can be observed that the maximum temperature observed was 646 °C and the maximum heating rate was approximately 18 °C/min, which occurred in the first hour of burning. Subsequently, the three ashes generated in the first stage were re-burned in a muffle furnace at 600 °C under oxidizing environment, with a heating rate of 10 °C/min and residence time of 15 min, as shown in Figure 2. Figure 3 shows the CSA appearance after burning in the pilot (Figure 3a) and muffle kilns (Figure 3b).

The production of SBA was carried out using the same processes previously presented, except for the leaching step with citric acid, which was carried out between the two burning stages (Figure 1). This change occurred due to the reduced SBA yield compared to the other studied ashes. In addition, it was interesting to simulate in the laboratory the production of an ash simulating that from the particulate collector in sugar and alcohol plants (CORDEIRO; ANDREÃO; TAVARES, 2019). This behavior was verified by burning the bagasse in pilot kiln that presented an average yield of 2.3% by mass. Yields of RHA, CSA, and BLA in the first burning were 22, 13, and 23%, respectively.

Figure 1 - Flowchart of production processes of ashes

- Sugarcane bagasse
- Rice husk
- Corn straw
- Bamboo leaf

Water washing

Oven drying at 100 °C for 24 h

Acid leaching

Burning - Pilot kiln (Tₘₐₓ ≈ 650 °C)

Acid leaching

Burning - Muffle kiln (600 °C for 3 h)

Ultrafine grinding (D₅₀ ≈ 10 μm)

- SBA
- RHA
- CSA
- BLA
After burning, the four ashes were ground in a high-energy attrition mill (Union Process) in order to obtain products with similar granulometries and $D_{50}$ around 10 μm (CORDEIRO; ANDREÃO; TAVARES, 2019). The grinding conditions were adapted from Andreão et al. (2019), as follows:

(a) operating speed of 500 rpm;
(b) feed mass of 100 g;
(c) 3.5 kg of spherical grinding bodies (2 mm diameter) of zirconia; and
(d) grinding time of 30 min.

The oxide composition was obtained by semi-quantitative analysis using an X-ray fluorescence spectrometer (Shimadzu EDX-720). Loss on ignition was obtained in accordance with NBR NM 18 (ABNT, 2012). The particle size distribution was determined by laser diffraction testing (Malvern Mastersizer 2000), with 15 min of dispersion in deionized water. The pore structure and BET specific surface area of the ashes were...
obtained in a nitrogen adsorption analyser (Micromeritics Tristar II 3020), with a sample treatment at 120 °C under vacuum for 12 h. The mineralogical composition was investigated by X-ray diffraction using a Rigaku Miniflex 600 equipment, operating with Cu-Kα radiation (λ = 0.15406 nm), 40 kV voltage, 15 mA current, 0.02° step, and 5°/min angular speed. Data were collected in a 20 angular interval between 8 and 50°. The quantification of the crystalline phases was performed by Rietveld refinement (LIU; KUO, 1996) using PDXL v. 2 software. The crystallographic information was obtained from Jorgensen (1978). Amorphous content was determined using anatase as the internal standard.

The pozzolanic activity of the ashes was evaluated by electrical conductivity of samples in alkaline solution according to the method proposed by Luxán, Madruga and Saavedra (1989). For this, a suspension was prepared with 98.7 mg of calcium hydroxide, 70 g of deionized water, and 1.75 g of sample. The conductivity variation was monitored for 20 min at 40 ± 1 °C using the Alfakit AT 230 conductivity meter. It should be noted that previous results of electrical conductivity variation obtained for sugarcane straw ashes showed an excellent correlation from the results of the modified Chapelle (CORDEIRO; VIEIRA; LOPES, 2017).

Pozzolanic activity was also investigated by the mechanical performance index based on NBR 5752 (ABNT, 2014). The performance index is obtained from the relationship between the compressive strength at 28 days of a mortar with a 25% cement replacement by the mineral addition and the strength of a reference mortar. The mortar mix-design was established for water-binder (cement and ash) and sand-binder ratios of 0.48 and 3.0, respectively. A class G cement (ABNT, 2006) was used in this test. The chemical composition, density, and Blaine fineness of cement are presented in Table 1. Table 2 displays the contents of superplasticizer (ether carboxylic-based with 28.9% oven-dried residue and 1.12 g/cm³ density) used in each mix to keep the same consistency for all mortars. For each mix, 4 cubic specimens of 50-mm edge were moulded and were kept in the molds for 24 h in a humid environment. After this period, the specimens were demoulded and kept immersed in a lime-saturated solution for 28 days. The axial compression tests were performed using a Shimadzu UHI-500kN testing machine, with a loading rate of 0.5 mm/min.

The hydration kinetics in cementitious pastes was evaluated using isothermal conduction calorimetry tests. In this case, pastes were made with 10 and 20% (in mass) cement replacement by each type of ash, besides a reference paste (without ash). Class G cement (Table 1), deionized water, and ether carboxylic-based superplasticizer were used. The paste mix-design was defined based on a water-binder ratio of 0.4. Table 3 shows the content of superplasticizer used in making the same consistency pastes. The pastes were named P-REF (reference) and P-X-Y, where X was the type of ash and Y the cement replacement used (% by mass).

### Table 1 - Oxide composition, loss on ignition, density, and Blaine fineness of cement

| Oxide     | % in mass |
|-----------|-----------|
| SiO₂      | 16.15     |
| K₂O       | 0.85      |
| Al₂O₃     | 3.23      |
| CaO       | 69.10     |
| SO₃       | 3.60      |
| P₂O₅      | –         |
| Fe₂O₃     | 5.66      |
| Na₂O      | 0.22      |
| MnO       | 0.10      |
| TiO₂      | –         |
| Na₂O₉eq*  | 0.78      |
| Loss on ignition | 1.09 |
| Density   | 3.17 g/cm³ |
| Blaine fineness | 290 m²/kg |

*Note: Na₂Oₑₒ₉ = Na₂O + 0.658 x K₂O.*
Table 2 - Superplasticizer (SP) content of mortars

| Mortar | M-REF* | M-RHA | M-SBA | M-CSA | M-BLA |
|--------|--------|-------|-------|-------|-------|
| SP (%) | 0.065  | 0.844 | 0.504 | 0.546 | 0.546 |

Note: *% solids in relation to binder mass; and 
**M-REF - reference mortar; M-X mortar containing the ash X (RHA, SBA, CSA, and BLA).

Table 3 - Superplasticizer content of pastes

| Paste   | P-REF | P-RHA-10 | P-RHA-20 | P-SBA-10 | P-SBA-20 | P-CSA-10 | P-CSA-20 | P-BLA-10 | P-BLA-20 |
|---------|-------|----------|----------|----------|----------|----------|----------|----------|----------|
| SP (%)  | 0.03  | 0.10     | 0.21     | 0.05     | 0.10     | 0.07     | 0.13     | 0.07     | 0.13     |

Note: % solids in relation to binder mass.

All pastes were mixed following the same procedure, which consisted of homogenization of deionized water and superplasticizer, followed by manual mixing of the liquid part and binder material with a spatula for 30 s. Next, the mixing was concluded in a portable electric mixer for 30 s at 260 rpm, followed by 60 s at 600 rpm. Immediately after the mixing end, samples of each mix were inserted into the Calmetrix I-CAL 2000 isothermal calorimeter. The pastes were monitored for the heat release profile for 72 h at 25 ± 0.1 °C.

Results and discussion

Table 4 shows the oxide composition and loss on ignition of different ashes, which showed a predominant presence of silica (SiO$_2$) in their compositions (around 90%). On the other hand, there were low levels of contaminating oxides in the four ashes, such as K$_2$O, CaO and SO$_3$. These compounds are highly undesirable in cement mixes, mainly because they cause durability problems, such as alkali-aggregate reactions (MEHTA; MONTEIRO, 2014). The SO$_3$ content was lower in the four ashes than those observed for cement (Table 1) and the limit of 4% established for pozzolans in NBR 12653 (ABNT, 2015). The same can be observed in relation to NBR 16697 (ABNT, 2018), which limits the SO$_3$ content for commercial cements in Brazil to 4.5%. The high concentration of silica in the ashes and the low levels of contaminating oxides highlighted the influence of acid leaching in the purification process of the studied materials. Similar results were previously observed (KRISHNARAO; SUBRAHMANYAM; JAGADISH KUMAR, 2001; VAYGHAN; KHALOO; RAJABIPOUR, 2013; CORDEIRO; SALES, 2016; CORDEIRO; VIEIRA; LOPES, 2017). The values of loss on ignition values, also presented in Table 4, showed low levels of carbonaceous material in the ashes after two-stage burning. All values were significantly lower than the maximum content of 5% established by NBR 12653 (ABNT, 2015). The presence of residual carbonaceous material in the ashes reduced the concentration of silica and also promoted a delay in cement hydration reactions (CORDEIRO; BARROSO; TOLEDO FILHO, 2018).

Particle size distribution curves shown in Figure 4 revealed similar granulometries for RHA, SBA, CSA, and BLA. These results showed the efficiency of the grinding process in giving similar sizes to the four ashes, which was the necessary condition for an accurate evaluation of the pozzolanic effect of the ashes since the pozzolanic activity is strongly influenced by the particle size (CORDEIRO et al., 2011; CORDEIRO; KURTIS, 2017). The small differences among the ash granulometry for the same grinding condition indicated different grindability, which is characteristic for agroindustrial silica-based ashes (RIBAS et al., 2014; CORDEIRO; TAVARES; TOLEDO FILHO, 2016). All ashes presented $D_{50}$ values close to 10 µm and lower than the characteristic value of the cement used (14.9 µm). It should be noted that the $D_{50}$ of the four ashes were similar to those studied by Andreão et al. (2019) for sugarcane bagasse samples with strong contamination by quartz produced in an attritor mill.

Gas adsorption isotherms of the ashes are shown in Figure 5. It can be observed that the adsorption and desorption isotherms presented a similar trend for the four ashes, which is characteristic of meso and macroporous materials according to the IUPAC classification (THOMMES et al., 2015). This behavior is typical of a highly porous sample, as are the agroindustrial ashes for example, whose pore network has been significantly affected by high-energy grinding (VIEIRA et al., 2020). The results of the BET specific surface area, shown in Table 5, confirmed the presence of porous particles in the ashes, mainly RHA, CSA, and BLA. SBA was slightly penalized by the difference in its production process, which was necessary to adapt the characteristics of this ash to those observed for ashes originated from particulate collectors in sugar and alcohol plants. It is worth noting that the high values BET specific surface area observed for RHA, CSA, and
BLA were due to acid leaching of raw materials before burning and grinding procedures. The specific surface area values are in accordance with those observed for agroindustrial ashes reported by Della, Kühn and Hotza (2002), Cordeiro, Vieira and Lopes (2017) and Vieira et al. (2020).

Table 4 - Oxide composition and loss on ignition of ashes (% in mass)

| Oxide   | RHA  | SBA  | CSA  | BLA  |
|---------|------|------|------|------|
| SiO₂    | 92.25| 90.80| 89.11| 91.00|
| K₂O     | –    | 2.32 | 2.10 | –    |
| Al₂O₃   | 2.92 | –    | –    | –    |
| CaO     | 0.10 | 0.90 | 1.60 | 3.95 |
| SO₃     | 1.33 | 2.28 | 2.69 | 2.55 |
| P₂O₅    | –    | –    | 0.90 | –    |
| Fe₂O₃   | 0.29 | 1.60 | 0.30 | 0.31 |
| Na₂O    | 0.10 | –    | –    | 0.10 |
| MnO     | –    | 0.10 | –    | –    |
| TiO₂    | –    | –    | 0.10 | 0.10 |
| Loss on ignition | 3.01 | 2.00 | 3.20 | 2.00 |
X-ray diffractograms of the ashes are shown in Figure 6. A typical silica amorphous halo can be seen between 20 angles of 15 and 30°, indicating that the burning and grinding procedures were adequate to maintain the silica originally present in the plants in an amorphous state. This characteristic is fundamental for pozzolanic production and was previously observed for rice husk (REAL; ALCALÁ; CRIADO, 1996; KRISHNARAO; SUBRAHMANYAM; JAGADISH KUMAR, 2001; DELLA; HÜHN; HOTZA, 2002), sugar cane straw (CORDEIRO; VIEIRA; LOPES, 2017), and elephant grass (CORDEIRO; SALES, 2016) ashes. In fact, the four ashes were predominantly amorphous and did not contain crystalline phases with significant content. Based on Rietveld’s refinement, SBA was the only ash to present quantifiable phases with quartz and hematite contents of 1.8% (± 0.5%) and 0.2% (± 0.1%), respectively.

Results of electrical conductivity tests for all ashes in calcium hydroxide solution are shown in Figure 7. Initially, there was a considerable drop in conductivity for the RHA, CSA, and BLA samples. The conductivity variation increased during the tests, which was an indication of the continuity of reactions between ashes and calcium hydroxide, especially the CSA. At 2 min of testing, the electrical conductivity variation values for RHA, CSA, and BLA were 5.78, 3.73, and 5.91 mS/cm, respectively, as shown in Table 5. According to the classification proposed by Luxán, Madruga and Saavedra (1989), the 2-min variations indicated that the three ashes showed good pozzolanic since ΔC values after 2 min of test were significantly greater than 1.2 mS/cm. Similar electrical conductivity results have been reported by and in studies with rice husk (FENG et al., 2004) and cane straw (CORDEIRO; VIEIRA; LOPES, 2017) ashes, which were both generated after treating of raw materials with hydrochloric acid leaching. SBA had ΔC of 0.74 mS/cm and, therefore, was classified as a medium pozzolanic material (0.4 mS/cm < ΔC < 1.2 mS/cm). The mean pozzolanicity of SBA was also observed for other bagasse ash assessed using the same methodology (ANDREÁO et al., 2019).

The mechanical performance index (PI) values are shown in Table 5 and confirmed the pozzolanic character of four ashes produced. In this case, all ashes presented PI values higher than 100%, especially in RHA, CSA, and BLA, which corroborated the electrical conductivity test results. The compressive strength gain in mortars containing ashes could be attributed to characteristics such as high content of amorphous silica, reduced loss on ignition and high BET specific surface area. Moreover, the reduced ash particle size certainly contributed to the increase in mechanical strength by increasing packing density of mixes (CORDEIRO et al., 2008; CORDEIRO; KURTIS, 2017). SBA also achieved a satisfactory PI, however not as expressive as that observed for other ashes. SBA PI was in line with electrical conductivity results and with previous studies that indicated an intermediate pozzolanic activity for sugarcane bagasse ashes in mechanical tests (CORDEIRO et al., 2018). The better performance of SBA produced in this work in relation to different samples in the literature could be attributed to the reduced contamination of SBA by quartz and the citric acid leaching process, which provided the elimination of contaminating oxides, especially K₂O (ANDREÁO et al., 2019).

Figure 8 shows the heat flow curves of reference mix and pastes containing RHA, SBA, CSA, and BLA. The curves were defined based on the binder mass and provided a clear view of the effect of each ash on hydration kinetics. The enlarged graphs of heat flow curves up to 8 h of hydration are also shown in Figure 8. Overall, all pastes exhibited the typical hydration behavior of a cementitious paste, with a first peak due to the initial contact between water and cement (which occurred outside the calorimeter and was not discussed in this work), induction, an acceleration period with maximum flow at the characteristic peak associated with C₃S hydration and a secondary peak corresponding to the C₃A reaction (BULLARD et al., 2011), followed by a progressive deceleration of reactions over time.

| Ash  | D₅₀ (µm) | BET specific surface area (m²/kg) | ΔC* (mS/cm) | PI** (%) |
|------|----------|----------------------------------|-------------|----------|
| RHA  | 10.5     | 63,232                           | 5.78        | 136      |
| SBA  | 9.2      | 13,332                           | 0.74        | 107      |
| CSA  | 6.8      | 91,630                           | 3.73        | 148      |
| BLA  | 7.6      | 199,000                          | 5.91        | 140      |

Note: *ΔC - variation of electrical conductivity (LUXÁN; MADRUGA; SAAVEDRA, 1989); and **PI - pozzolanic performance index (ABNT, 2014).
Figure 6 - X-ray diffraction patterns of RHA, SBA, CSA, and BLA (peaks intensities in arbitrary unit)

Figure 7 - Electrical conductivity curves of ashes during 20-min testing
The results indicated that the addition of RHA promoted a slight reduction in the induction period for the two cement replacement contents evaluated, despite the opposite effect generated by the higher superplasticizer consumption (SAKAI et al., 2006), as indicated in Table 6. It should be noted that the amounts of superplasticizer used in pastes with RHA were significantly higher than that used in the reference (around 7 times higher). The reduction of induction period in pastes with RHA was due to dilution effects and, mainly, heterogeneous nucleation associated with this mineral addition (LAWRENCE; CYR;
RINGOT, 2003; CORDEIRO; KURTIS, 2017). This behavior was also observed in BLA pastes, which proved the high reactivity of this ash. On the other hand, in spite of promoting an increase in heat flow in relation to the reference, pastes with SBA and CSA presented induction periods similar to that observed for the reference. This greater heat flow in the first hours of hydration may be associated with the presence of K₂O in SBA and CSA mixes, which promoted a significant increase in the availability of alkalis in pastes with these ashes, as shown by the alkali equivalent (Na₂Oeq) values of cementitious materials used in different pastes (Table 7). Similar results were previously observed by Andréao et al. (2019) and Cordeiro, Andréao and Tavares (2019) for pastes containing bagasse ashes. In addition, the extremely high BET specific surface area of CSA may have caused a reduction in the initial hydration rate due to the high-water demand by pozzolan (KORPA et al., 2008), even after correcting the consistency of pastes with the superplasticizer. It is interesting to note that there was an increase in the secondary peak (relative to the formation of monosulfate - AFm) in the ash pastes compared to the P-REF curve. This effect may be related to the heterogeneous nucleation of hydrates formed on the ultrafine pozzolanic particles (LAWRENCE; CYR; RINGOT, 2003) and also to a significant reduction in SO₃ content (TAYLOR, 1997) in the binder of different blended pastes in relation to the reference, as shown in Table 7.

Figure 9 shows the accumulated released heat curves over 72 h of hydration. The accelerating effect of hydration reactions promoted by RHA and BLA was even more evident in the analysis of the accumulated heat of pastes, especially during the first day of hydration. On the other hand, the paste with 20% of SBA presented the lowest value of accumulated heat (Table 6). This showed the greater influence of the dilution effect compared to the other physical and chemical effects promoted by ash. The reduced values of the SBA specific surface area and pozzolanic activity compared to the other ashes also contributed to the reduction of accumulated heat for P-SBA-10 and P-SBA-20 in relation to the reference.

Table 6 - Parameters of isothermal calorimetry of different pastes

| Pasta      | Induction offset time (h:min) | Maximum heat flow (mW/g) | Released heat at 72 h (J/g) |
|------------|------------------------------|--------------------------|----------------------------|
| P-REF      | 03:24                        | 3.07                     | 267.3                      |
| P-RHA-10   | 02:24                        | 2.93                     | 265.9                      |
| P-RHA-20   | 02:30                        | 3.00                     | 260.5                      |
| P-SBA-10   | 03:00                        | 3.06                     | 264.2                      |
| P-SBA-20   | 03:30                        | 2.87                     | 245.7                      |
| P-CSA-10   | 03:40                        | 3.03                     | 268.8                      |
| P-CSA-20   | 03:50                        | 2.69                     | 252.9                      |
| P-BLA-10   | 01:58                        | 2.91                     | 267.9                      |
| P-BLA-20   | 03:12                        | 2.53                     | 260.5                      |

Table 7 - SO₃ content and alkali equivalent in terms of Na₂O (Na₂Oeq) of paste binders

| Paste     | P-REF | P-RHA-10 | P-RHA-20 | P-SBA-10 | P-SBA-20 | P-CSA-10 | P-CSA-20 | P-BLA-10 | P-BLA-20 |
|-----------|-------|----------|----------|----------|----------|----------|----------|----------|----------|
| SO₃       | 3.60  | 3.37     | 3.15     | 3.47     | 3.34     | 3.51     | 3.42     | 3.50     | 3.39     |
| Na₂Oeq (%) | 0.78  | 0.71     | 0.64     | 0.85     | 0.93     | 0.84     | 0.90     | 0.71     | 0.64     |

Note: % of Na₂Oeq in relation to mass binder.
Conclusions

Based on this experimental research, the following conclusions could be drawn:

(a) the ashes produced by the confluence of different production methods presented typical properties of pozzolanic materials. From an efficient calcination process, the oxide composition of ashes showed high levels of amorphous silica (SiO$_2$) and reduced values of loss on ignition;

(b) the predominant presence of amorphous silica and high BET specific surface area allowed the classification of RHA, CSA, and BLA as ashes of good pozzolanicity and SBA as a material of medium pozzolanic activity, according to the electrical conductivity testing. The reactivity of the four ultrafine ashes was confirmed in the performance index evaluation, where the minimum value obtained of 107% was significantly higher than the minimum established by standard (90%); and

(c) the partial cement replacement by ashes (10 and 20% by mass) promoted small changes in hydration kinetics of pastes during the first 3 days. The samples with greater reactivity (RHA and BLA) promoted a significant acceleration of reactions, while SBA generated an increase in induction period for the 20% cement replacement level.

Thus, it can be concluded that acid leaching combined with two-stage burning, and high-energy grinding procedures were efficient for producing pozzolans from biomass wastes. Finally, it should be noted that the leaching procedure can be carried out with acid effluents within biomass-based plants (e.g. vinasse, produced in ethanol distilleries). This practice would prevent the generation of new waste and contribute to the production of biomass ash with a more controlled chemical composition. Future research should be carried out with leaching using other types and different concentrations of acids, in addition to biomass samples of biomass from different sources.

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