

# Potential Use of Sewage Sludge Ash in Lime-Based Materials

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**Abstract:** Recycling of wastes and their by-products is attracting increasing interest worldwide because of the high environmental impact in the cement, concrete and other industries. This work deals with the study of the physico-chemical characteristics of binders based on sewage sludge ashes and lime. In a first step, we used X-ray fluorescence to determine the chemical composition of ash, lime and binders. This allowed us to see that the chemical composition of sewage sludge ash is similar to that of cement. We then used X-ray diffraction to identify the main mineralogical phases in the samples. Compressive strengths of mortars containing 20%, 40%, 60% and 80% of SSA showed that SSA has a long-term positive effect which might be related to a slight pozzolanic activity. The L<sub>4</sub> binder consisting of 80% fine ash and 20% lime has a higher compressive strength than the others. The binder setting start time is greater than that of cement but shorter than that of lime. The study of the thermophysical properties of the L<sub>4</sub> binder shows that it has a higher thermal resistance than cement and clay mortars. Moreover, it heats up less quickly because of its low effusivity compared to the latter two. This analysis highlighted the principal characteristics that must be taken into account to use SSA correctly in lime-based materials.

**Keywords:** Sewage Sludge Ash, Lime, Binder, X-ray Fluorescence, X-ray Diffraction, Thermomechanical Behavior

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## 1. Introduction

Sludge ash is very interesting for many applications because it can be mixed with cement or binders such as lime to give rise to new materials with several attractive physical characteristics such as the increase in resilience, the long-term strength of mortars [1-3]. Their mechanical properties open industrial perspectives for the manufacture of cement and building at lower cost with some ecological contribution. In fact, new environmental challenges and research opportunities arise on the use of these solid wastes as well as their basic properties. Recycling of these wastes and their by-products is attracting increasing interest worldwide because of the high environmental impact in the cement, concrete and other industries. For example, lightweight concrete can be

made from industrial by-products and hazardous solid wastes such as fly ash, sewage sludge ash, bottom ash, slag, etc. Cement can be replaced by 15 to 35% by this waste without significantly affecting the properties of the material obtained [4]. In some studies, replacing cement with 10% or 20% ash from sewage sludge shows an increase in setting time due to the significant presence of phosphorus in sludge ash [5]. Similarly, the compressive strength of the mortars was lower than that of the reference mortar consisting solely of cement [6]. In fact, the addition of these ashes slows down the kinetics of resistance development; however, the long-term strength of the mortar with addition was greater than that of the reference mortar [5]. In the literature, it is reported that these additions can improve sulphate concrete's strength, workability, and also decrease permeability [4]. It has been

observed that the characteristics of ash from sewage sludge are similar to those of ceramics. Their use in ceramics applications for bricks, tiles and ceramic glass considerably improves these materials by densifying them, increasing their strengths [7]. Ceramic panes with ash additions have achieved higher resistances than natural materials such as granite and marble with strong durability properties [8].

The strength activity index (SAI) value of SSA was smaller than that of fly ash; the effectiveness of using SSA in mortar was lesser than that of fly ash [9]. Moreover, the water demand of SSA mortar is higher than that of ordinary cement mortar. S. Pan found that the addition of SSA affected the mortar workability [10]. This phenomenon is primarily due to the porous and irregular morphology of SSA. Consequently, it is difficult to adequately maintain water-to-cement ratio and workability of SSA mortar simultaneously. The application of mechanical grinding to adjust SSA fineness was an effective modification to improve SSA mortar properties [3]. Regarding the problems related to the cost of construction in the building, air pollution such as greenhouse gases and energy expenditure for the production of cement, we propose to study the effect of the integration of the ashes of Sewage treatment plant (STEP) in lime, to produce an environmentally-friendly cement. Indeed, lime can react with silica in the presence of water to produce a hydrated calcium silicate responsible for the resistance of cementitious materials. Since wastewater treatment plants produce large amounts of solid waste such as STEP ashes, they can be recovered and be used as building materials [4-6, 11].

This work deals with the study of the physico-chemical characteristics of binders based on sewage sludge ashes and lime. In a first step, we used X-ray fluorescence to determine the chemical composition of ashes, lime and binders. X-ray fluorescence is also used in ash characterization, analysis of environmental samples, including industrial sites soils to study their contamination by heavy metal [12-15].

Following the X-ray fluorescence of the samples, we studied their mineralogical behavior.

We then used X-ray diffraction to identify the main mineralogical phases present in the samples.

Compressive strength of samples based on sewage sludge ash and lime are also investigated in this study.

The thermal conductivity and effusivity of samples were

determined simultaneously using a transient method.

This analysis highlighted the principal characteristics that must be taken into account to use SSA correctly in lime-based materials.

## 2. Materials and Methods

### 2.1. Origin of Ash

The sewage sludge ash used in this work comes from the Senegalese sewage treatment plant of Camberene located in the Dakar region. The sludge is dewatered, dried and burnt in a fluidized bed incinerator at 850°C. The sewage sludge ash has a gray color, is entrained by the combustion gases and is recovered during the purification (Figure 1).



*Figure 1. Sewage sludge ash.*

It has an apparent density of 0.71 g/cm<sup>3</sup> and an absolute density of 2.4 g/cm<sup>3</sup>.

The lime, as an important material used in the building construction, has an apparent density varying between 0.6 and 0.75 g/cm<sup>3</sup> while its absolute density is in the range of 2.6 and 2.9 g/cm<sup>3</sup>.

### 2.2. Experimental Methods

The experimental methods used for the physical and chemical characterization of ashes are shown in Table 1:

**Table 1.** Experimental methods for the physical and chemical characterization of SSA, and for the study of the activity of SSA in lime-based materials.

Chemical analysis	X-ray fluorescence (XRF) X-ray portable fluorescence Niton XLT900s (P-XRF) for our X-ray analyzes with a measurement time of 350 s. XRF was performed with 100% normalization and full fundamental parameter quantification techniques
Mineralogy	X-ray diffraction (XRD); Cu K $\alpha$ radiation ( $\lambda = 1.54060 \text{ \AA}$ ) 2 $\theta$ step interval of 0.04° (10°-70°) and acquisition time of 1560 s
Effect on binder hydration:	-XRD of mixtures lime-ash-water
Mineralogical study	- Setting time using Vicat apparatus (NF EN 196-3)
Setting time	Compressive strength of 4×4×16 cm prisms (NF EN 196-1)
Effect on mechanical properties of mortars	Hydration times: 3, 7, 28 days; Each value is the average of 6 tests
Effect on thermal properties of mortars	The thermal test samples were also prepared in a mould of dimensions 10 cm × 10 cm × 2 cm. All thermophysical experiments were performed using samples prepared in a mould of dimensions of 10cm×10cm×2cm. The thermal conductivity and effusivity of samples were determined simultaneously using a transient method [16]

For the Thermal test, the hot plate transient method was used in an asymmetrical configuration to determine simultaneously the thermal conductivity and effusivity of samples [16, 17]. The different elements that make up the experimental device are represented in Figure 2.

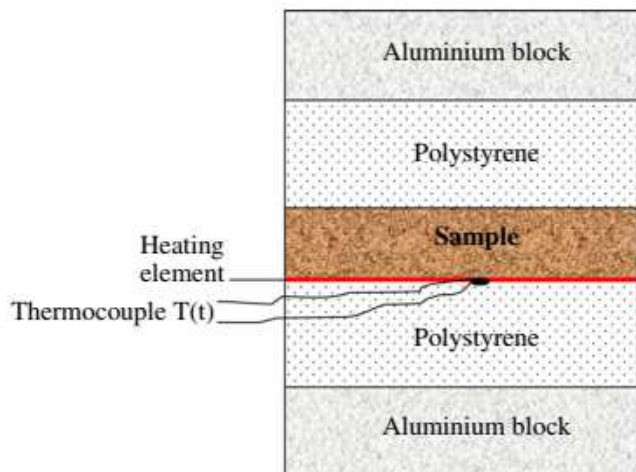


Figure 2. Schema of the experimental hot plate device.

This method was then used in an asymmetrical configuration. It consists in embedding a heating element on which a thermocouple was fixed, between the material to be characterized and a 5 cm thick polystyrene plate.

Since the thermocouple is in contact with polystyrene that is a deformable material, the presence of the thermocouple does not increase the thermal contact resistance between the heating element and the polystyrene. Furthermore, since polystyrene is an insulating material, this thermal contact resistance will be neglected.

The whole set was put between two aluminum blocks with a thickness of 4 cm. A flux step is sent in the heating element and the temperatures  $T_s(t)$  at the center of the heating element is recorded. The processing of the recording of  $T_s(t)$  is realized by supposing that the heat transfer at the center of

the heating element is 1D. The temperature at the level of the aluminum block is constant.

### 3. Results and Discussion

#### 3.1. X ray Fluorescence Analysis

A portable XRF device Niton XLT900s was used to analyze the chemical composition of the sludge ash in terms of major and minor elements (Table 2). Oxides are the major elements and are expressed in%, while the minor elements are given in mg/kg.

Table 2. Results of major (%) and minor (mg/kg) elements using XRF technique of sewage sludge ash contents.

Elements	Sewage sludge ash
SiO <sub>2</sub>	24.1
Al <sub>2</sub> O <sub>3</sub>	6.69
Fe <sub>2</sub> O <sub>3</sub>	12.6
CaO	22.2
MgO	3.90
K <sub>2</sub> O	3.22
TiO <sub>2</sub>	0.66
MnO	0.10
P <sub>2</sub> O <sub>5</sub>	5.92
Pb	357
Cu	1264
Cr	212
Zn	3632
As	16

The chemical composition shows that SSA is mainly of silicon, calcium, iron, aluminum and phosphorus. The amount of CaO and P<sub>2</sub>O<sub>5</sub> is high compared to classical mineral admixtures such as fly ash, silica fume or metakaolin [18-20].

In Table 3 we resume studies from other authors in order to give a comparative asset in term of elemental composition with the samples we have used.

Table 3. Comparison of major (%) and minor (mg/kg) element contents measured with other authors working with the same materials.

Element	Sewage sludge ash PW	(Martin Cyr, 2007)	(Nuria Husillos Rodríguez, 2013)	L. C Morais (2010)	Siew Choo Chin (2016)
SiO <sub>2</sub>	24.1	34.2	3.72	26.39	18.49
Al <sub>2</sub> O <sub>3</sub>	6.69	12.6	2.56	12.16	7.95
Fe <sub>2</sub> O <sub>3</sub>	12.6	4.7	7.58	5.92	14.44
CaO	22.2	20.6	7.62	23.25	4.20
MgO	3.90	1.9	0.60	14.11	0.91
K <sub>2</sub> O	3.22	1.7	0.53	1.32	1.13
TiO <sub>2</sub>	0.66	0.9	0.23	0.81	0.99
MnO	0.10	0.06	0.05	0.11	0.09
P <sub>2</sub> O <sub>5</sub>	5.92	14.8	6.43	8.44	6.56
Pb	357	720	40	-	30
Cu	1264	2483	320	-	130
Cr	212	2636	1270	-	20
Zn	3632	7103	1020	-	810
As	16	23	5.8	-	-

The high variability between values shows that sludge ashes depend on several parameters: sludge composition, treatment conditions such as incineration temperature and additives.

The sum of  $\text{SiO}_2$ ,  $\text{Al}_2\text{O}_3$  and  $\text{Fe}_2\text{O}_3$  represent less than 70% of SSA. It was known that these oxides compose the reactive part of pozzolanic materials but there are significantly lower in SSA than in other classical mineral admixtures [18-19].

The minor elements are much less representative in our ashes when compared with those in Cyr's work [6]. However the concentration of  $\text{SiO}_2$ ,  $\text{Al}_2\text{O}_3$  and  $\text{Fe}_2\text{O}_3$  are similar to both sewage sludge ash used in the present work and sewage sludge ash in ref. [21] of Siew Choo Chin. The concentration of CaO in the former is instead four times higher than that in the latter. The similarity of these ashes with ours is important to highlight as they are both used in the processing of cement and concrete. Indeed, our ashes show some similarities with one type of cement in Senegal having the following chemical composition:  $\text{SiO}_2$  (19.53),  $\text{Al}_2\text{O}_3$  (7.12),  $\text{Fe}_2\text{O}_3$  (2.22), CaO (65.5). This suggests that the sewage sludge ashes presented in this work may be used in civil engineering [22].

In order to assess the hydraulic and pozzolanic activity of SSA, several mixtures: SSA with water and SSA with lime and water were prepared and analysed using XRD and mechanical behavior (Table 4).

**Table 4.** Sewage sludge ash and lime at different proportion.

Mix combinations	Designation
20 wt% Sewage sludge ash+80 wt% lime	L1
40 wt% Sewage sludge ash+60 wt% de lime	L2
60 wt% Sewage sludge ash+ 40 wt% de lime	L3
80 wt% Sewagesludge ash +20 wt% de lime	L4
100 wt% Sewage sludge ash (cboue)	L5

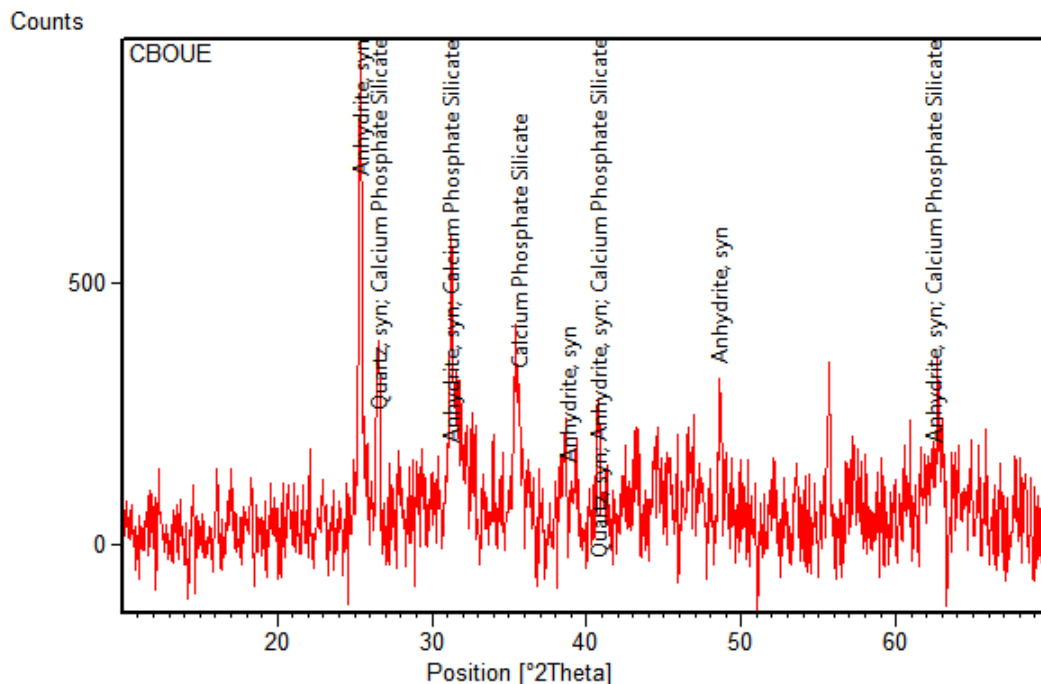
### 3.2. Mineralogical Characterization

The mineralogical characterization was carried out by X-ray powder diffraction (XRD). XRD using a PW1840 diffractometer with  $\text{Cu K}\alpha$  radiation operating at 30 mA and 40 kV.

The analysis was done in the continuous scanning mode with speed of  $0.05^\circ$  per s within the range of  $10^\circ \leq 2\theta \leq 70^\circ$ . The diffractometer is linked to a computer equipped with APD software. For data processing we used X Pert High Score software.

#### 3.2.1. Sewage Sludge Ash

The diffractogram of the sewage sludge ash shows the existence of a complex polyphasic material composed of several crystalline phases and a glass phase [1, 22-23]. Besides glass, the main minerals identified are quartz ( $\text{SiO}_2$ ), anhydrite ( $\text{CaSO}_4$ ), phosphorus calcium silicate ( $\text{Ca}_2\text{SiO}_4\text{-Ca}_3(\text{PO}_4)_2$ ) and minor phases (Figure 3).



**Figure 3.** Diffractogram of the sewagesludgeash Thus the crystallinity of the sewage sludge ash is weak.

This weakness manifests itself in two aspects: The first is the high background and the second is that the intensity of the peaks is low [24].

This high background is due to the amorphous phase [25]. Silicium in the amorphous phase can react with lime and produce new phases.

#### 3.2.2. Lime

Figure 4 shows the diffractogram of lime. The phases identified in lime are portlandite and calcite. Portlandite comes from the slaking of lime. The appearance of calcite could be due to the carbonation of portlandite exposed to air.

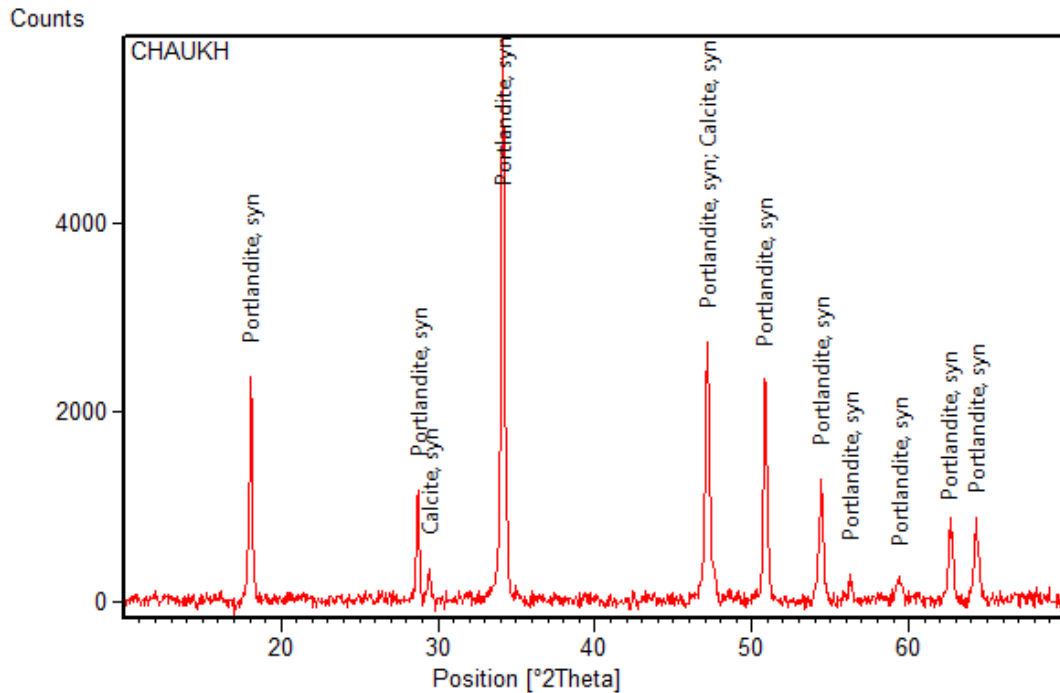


Figure 4. Diffractogram of lime.

### 3.2.3. Mixtures

We have made several formulations to assess the reactivity of ash by mixing it with lime. A fraction of the ash is rapidly soluble (sulfates, aluminum and silica) and can create new-formed products in presence of lime [6].

This is because the lime released during hydration can react with the silicates and aluminates of the binder as in the case of cement. We have built mortars of different formulations to study evolution of compressive strength with

respect to time and then to see how the compressive strength evolves at different times. After crushing these mortars, we analyzed the obtained powders by XRD which revealed the presence of new phases (Ettringite, Gehlenite, Calcium Silicate Hydrate) for certain formulations.

We have represented the diffractograms of  $L_5$  at different timescales: 3 days (L53JCB), 7 days (L57JCB) and 28 days (L528JCB) in Figures 5 to 7.

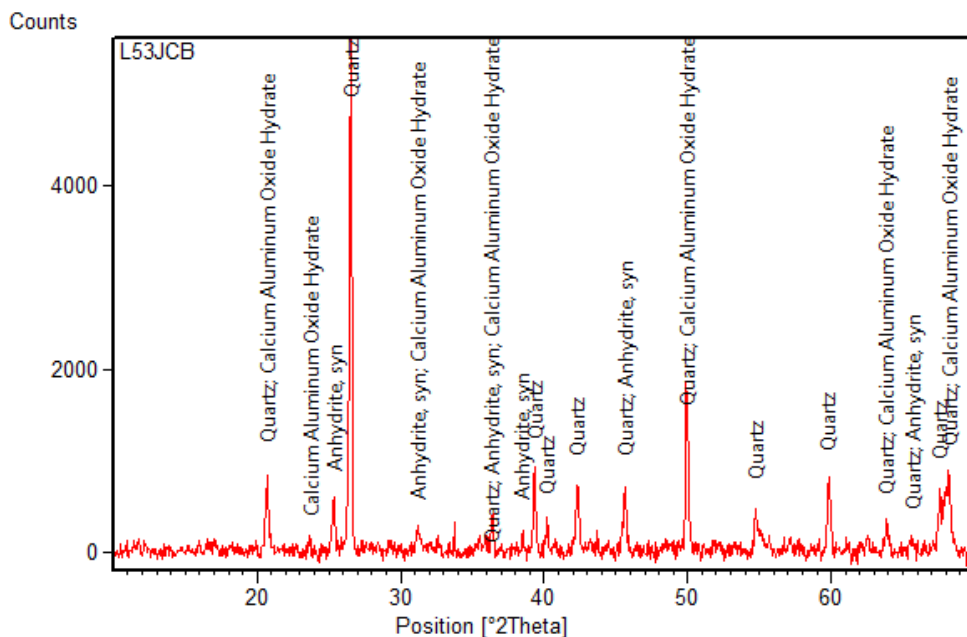


Figure 5. Diffractogram of L53JCB.



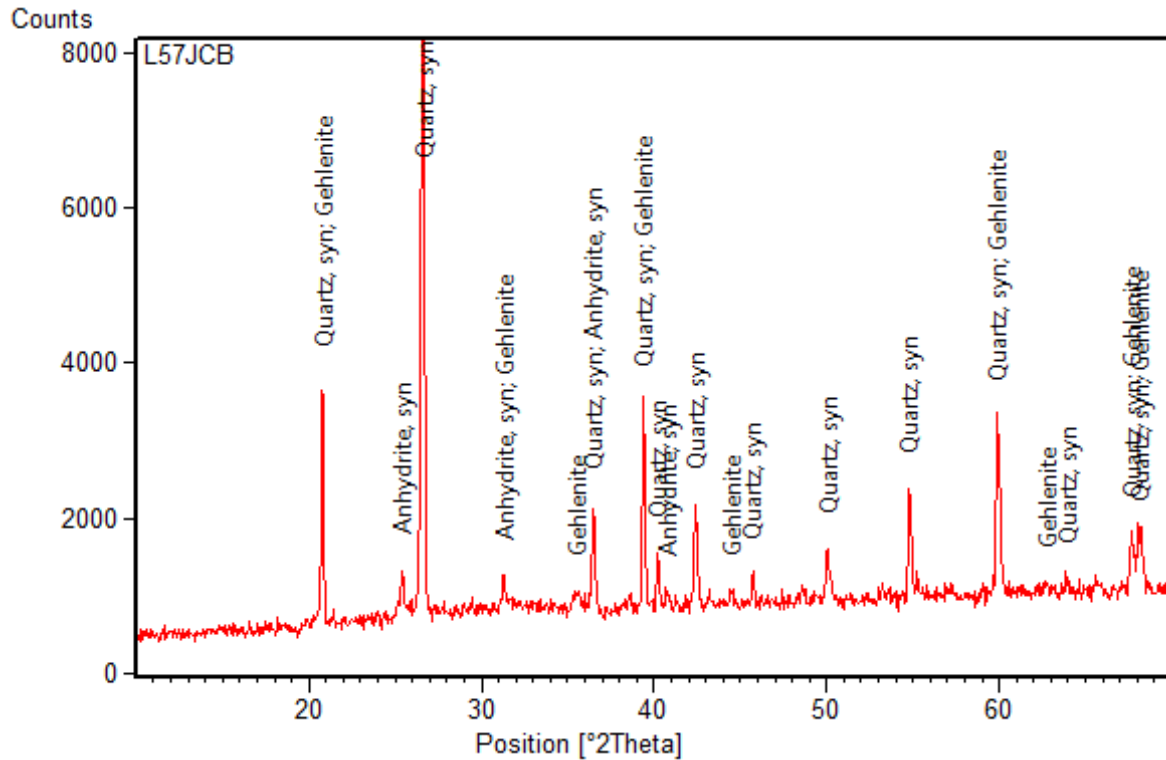


Figure 6. Diffractogram of L57JCB.

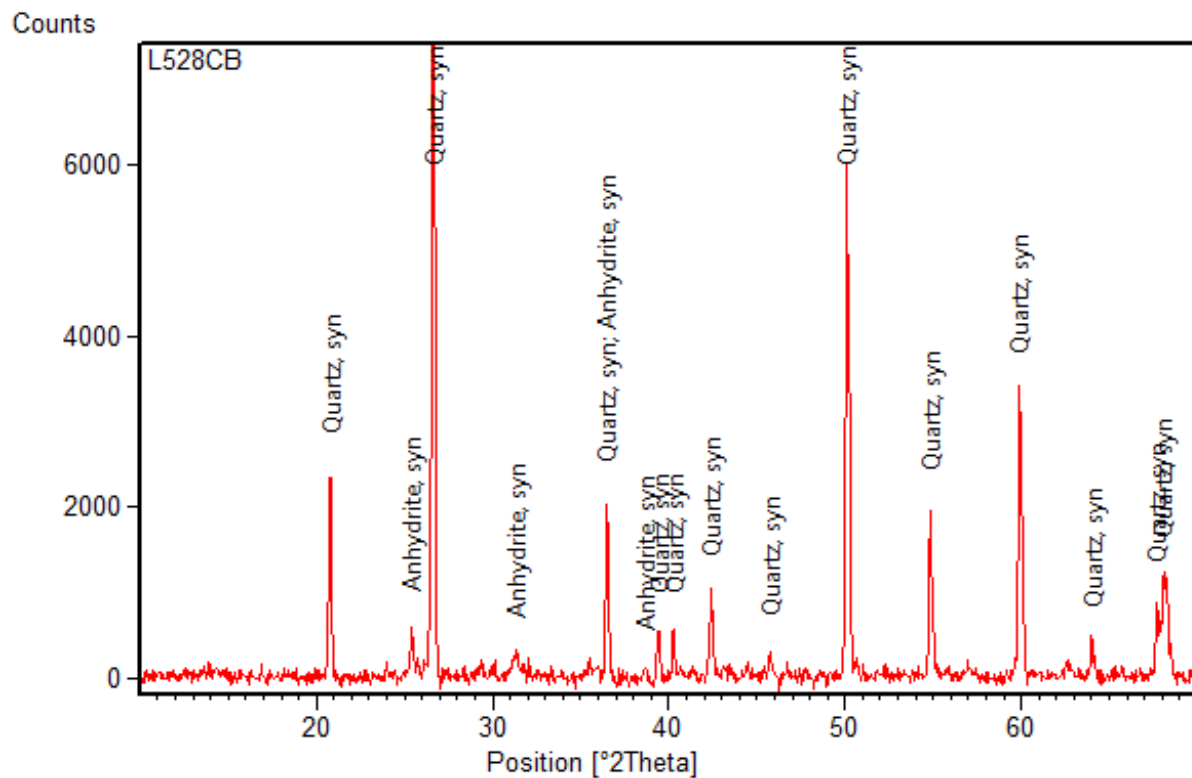
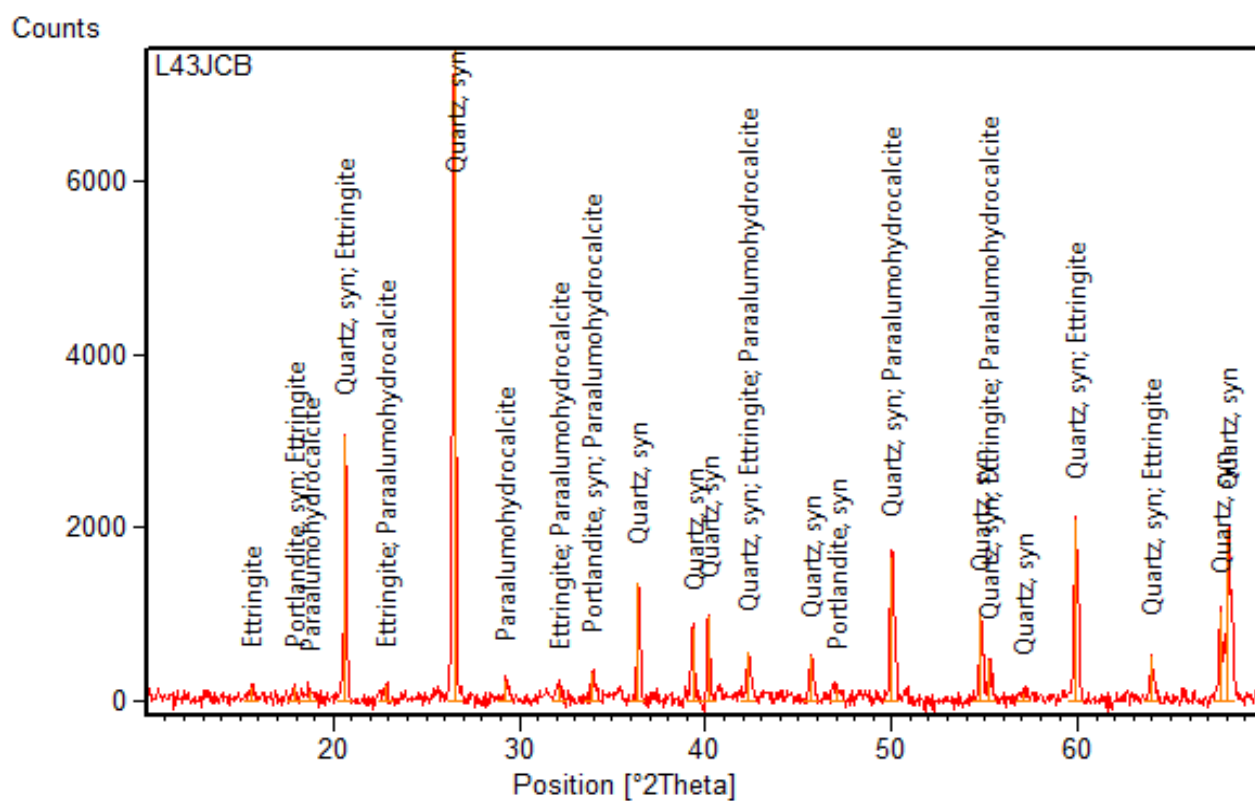
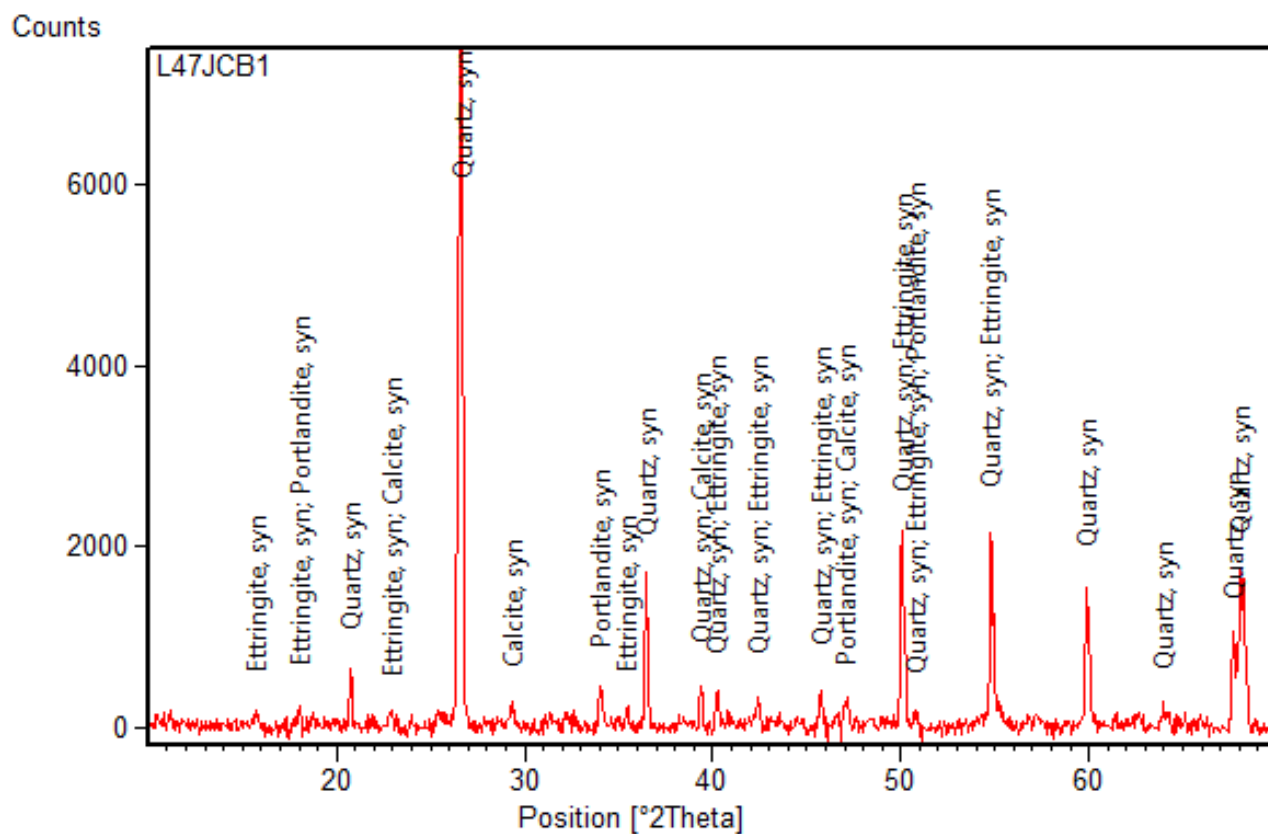


Figure 7. Diffractogram of L528JCB.

We find practically the same phases as those observed for the sewage sludge ash. This shows that ash alone hydrated has no reactivity under water. In addition, no hardening is not observed so we can say that the ash has no hydraulic character [6].

We present the diffractograms of the  $L_4$  binder at different time frames (3 days, 7 days, 28 days) at the level of Figures 8 to 11.



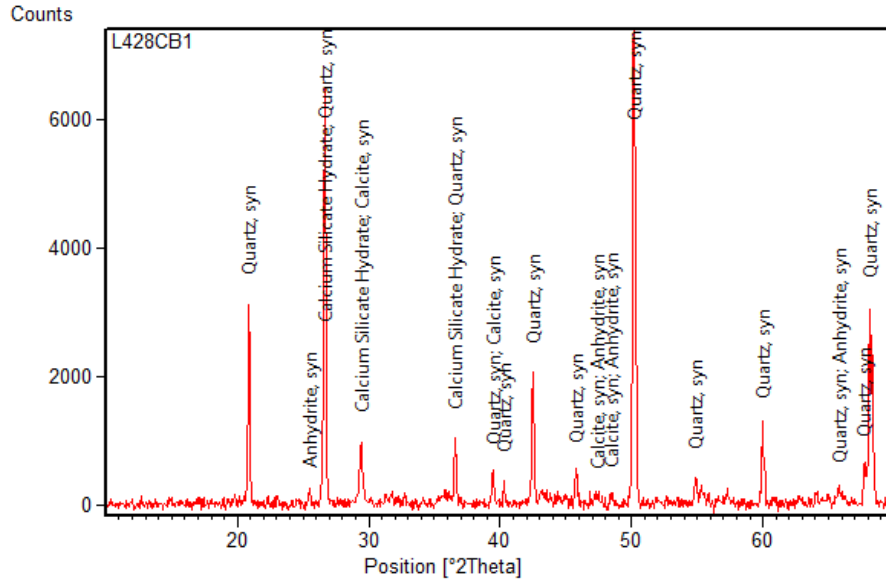
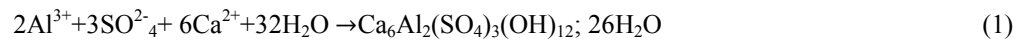


Figure 10. Diffractogram of L428JCB.

The  $\text{CaCO}_3$  that was detected in the  $L_4$  binder after hydration at 3 and 7 days comes from the carbonation ( $\text{CO}_2$  uptake) of calcium hydroxide. This formation of  $\text{CaCO}_3$  does not produce new phases of hydration, but changes the speed of the hydration [26-27]. Part of the  $\text{Ca}(\text{OH})_2$  portlandite formed is due to the hydration reactions of binder.

Gypsum, a hydrated anhydrite is a mineral added in the



The presence of gypsum in the ashes is beneficial because it no longer needs to be added to form a binder based on ash and lime.

We distinguish in the binder  $L_4$  the gradual appearance of hydrated calcium silicate CSH at 28 days responsible for the resistance of cementitious materials.

In addition to the formation of ettringite, we also distinguish the appearance of aluminates and hydrated calcium carboaluminates, as well as hydrated gehlenite, after progressive disappearance of lime, therefore these phases

clinker to regulate the setting of Portland cement. In our binders, the gypsum peak observed in the binder has disappeared in favor of ettringite. In fact, it reacts with calcium aluminates in order to regulate the setting by forming ettringite ( $\text{Ca}_6\text{Al}_2(\text{SO}_4)_3(\text{OH})_{12} \cdot 26\text{H}_2\text{O}$ ) in the following equation:

evolve over time (3, 7 and 28 days).

The other mineral phases that have appeared come from the hydrating reactions of the silicates, aluminates and their combinations.

### 3.3. Mechanical Behavior

We have shown the compressive strengths of mortars made with different binders Figure 11.

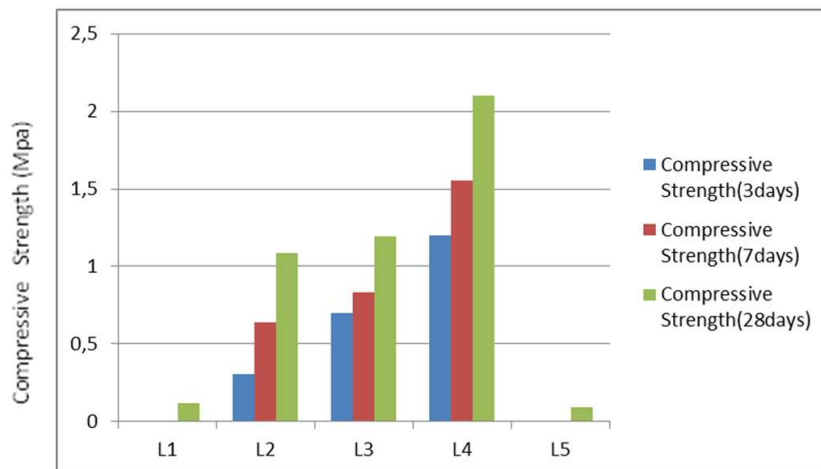


Figure 11. Evolution of the compressive strength as a function of the ash percentages and the age of the mortars.



Overall, the compressive strength increases over time. That at 28 days is more important than that of 7 days itself more important than that of 3 days. The reactions within the binders therefore continue.

The binder  $L_4$  has a compressive strength greater than 1 MPa at 3 d and at 7 d, the values are respectively 1.2 MPa and 1.6 MPa. At 28 days its compressive strength is 2.2 MPa. Likewise, its compressive strength at 3 days is higher than that of  $L_2$  and  $L_3$  at 28 days. This shows that the  $L_4$  binder gives the best compressive strength whatever the age.

From the analysis of the diffraction results of the different formulations, we observed that ettringite can only form on sample  $L_4$ , therefore its formation contributes to the increase in compressive strength of this binder.

This ettringite formation is explained by the equation (1) [1]. In addition to the formation of ettringite, a distinction is also made between the appearance of hydrated calcium aluminates and carboaluminates, as well as hydrated gehlenite, after gradual disappearance of the lime according to the formulation used.

The hydration reaction allowing the passage of the binder paste from the liquid state to the solid state.

It is a complex process in which the main binder compounds react to form new insoluble compounds which cause the material to set and gradually harden. The hydration mechanisms occur simultaneously but to better support our study we will study them separately.

#### Hydration aluminates

With 100% ash ( $L_5$ ) no resistance is observed because the hydration of the ash alone has not given a new phase that participates in compression resistance. Thus the ash alone does not have a hydraulic character.

When the percentage of ash is low, the amount of silica, alumina and gypse is not sufficient to react with calcium hydroxide. This is the origin of the zero resistance observed for sample  $L_1$ .

As the percentage of ash increases, compression resistance increases, which shows that there are new phases formed that are responsible for the strength of the lilibys  $L_2$ ,  $L_3$  and  $L_4$  binders of the material.

The reactions of aluminates can take place in several stages:

#### Step 1:

The aluminates react with the sulphate in the gypsum to form ettringite (AFt phase) (hydrated calcium trisulfoaluminate):  $((CaO)_6(Al_2O_3)(SO_3)_3, 32H_2O)$  [6]. This reaction is strongly exothermic. A layer of hydrates surrounds the aluminates which then form a protective layer around the grain. Thus these results show the activity of the ash in the presence of lime, which mechanically results in the hardening of the paste.

Mineralogical analysis shows a significant consumption of lime, a decrease or disappearance of certain phases of the rawash and reveals new-formed phases.

Step 2: Transformation of ettringite into hydrated calcium monosulfate

When the gypsum is used up, the sulphate concentration in the solution drops. The ettringite then becomes unstable and dissolves to form hydrated calcium monosulfate (called AFm):  $((CaO)_3(Al_2O_3)(CaSO_4), 12H_2O)$ . Chemical reactions intensify.

#### Step 3: Hydration of residual aluminates

After a few weeks, the ettringite usually becomes unstable and dissolves to form hydrated calcium monosulfate (called AFm):  $((CaO)_3(Al_2O_3)(CaSO_4), 12H_2O)$  [28]. After a month, reactions proceed slowly with aluminate ( $C_3A$ ) and aluminoferrite ( $C_4AF$ ) to form hydrated calcium aluminoferrite. Aluminum ferrite Tetracalcium ( $C_4AF$ ) is significantly slower to react and does not appear significantly until the gypsum is completely depleted.

#### Hydration of silicates

In contact with water, tricalcium silicates ( $Ca_3SiO_5$ ) and dicalcium silicates ( $Ca_2SiO_4$ ) dissolve in the form of ions. These interact with each other and form hydrated calcium silicates (C-S-H) and portlandite ( $Ca(OH)_2$ ). These reactions give off a lot of heat, so they are exothermic and can act as a catalyst for the hydration reaction. In the case of  $C_2S$ , the hydration kinetics are slower and so is the amount of Portlandite formed. The strength of a binder is due to the entanglement of the C-S-H gel. In fact, the C-S-H develop on the surface of the grains of the non-hydrated binder and gradually fill in the capillary interstices between the grains. This slows down, after a few hours, the diffusion of ions and water to the anhydrous components of the system. In our binder, a small amount of hydrated calcium silicate was observed on the binder  $L_4$  at 28 days. This weakness could be due to the non-presence of these silicates in our ashes [29].

### 3.4. Setting Time

The setting start time measured with the Vicat apparatus is 5h30min while for cement it is of the order of 2h. There is therefore a delay in setting the binder which can be attributed to the presence of phosphorus and the non-negligible Zinc content because these elements are known as powerful cement retarders [30].

The presence of gypsum in the ashes can also explain this delay in setting. In addition, the lime has a very slow start of setting which is 10 hours.

### 3.5. Thermophysical Properties

The results on thermal tests allowed us to obtain the thermal conductivity and effusivity of binder:

#### Thermal conductivity ( $\lambda$ )

The thermal conductivity reflects the ability of a material to transmit heat by conduction. The test gave an average thermal conductivity ( $\lambda$ ) of 0.33 W / m. ° K for binder  $L_4$ .

For a cement mortar:  $\lambda = 1.4$  W / m. ° K [31].

#### Thermal effusivity ( $E$ )

It indicates the capacity of materials to absorb (or restore) more or less quickly heat up.

The test results gave an average thermal effusivity of

639.0356 J / m<sup>2</sup>.s. ° K for our mortar.

For a cement mortar, the effusivity is:  $E = 1754.99 \text{ J / m}^2 \text{ S.}^\circ \text{K}$

The values of the thermal characteristics of some mortars are given in Table 5.

*Table 5. Comparison of thermophysical properties.*

Mortar	Average thermal conductivity (W/m. °K)	Average thermal effusivity (J/m <sup>2</sup> .S.°K)
Based on lime and ash (Present work)	0,33	680,05
Cement based mortar [31]	1,40	1754, 99
Clay-based mortar [32]	0,35	705

We find that the thermal conductivity of mortar made from ash and lime is lower than that of mortars made from cement on the one hand and clay on the other. This shows that ash and lime mortar is more resistant to conduction heat transfert hancement and clay mortars.

Also, the thermal effusivity value of lime and ash mortar is lower than that of cement and clay mortar. Thus, lime and ash-based mortar absorbs heat less quickly than cement and clay mortars. This mortar when used in the construction industry will develop better thermal characteristics than the latter two.

## 4. Conclusion

This paper aimed to improve our knowledge of the characteristics of SSA and its effect on the properties of lime based materials.

This allowed us to see that the chemical composition of sludge ash is similar to that of cement.

The main phases identified on lime are portlandite and calcite.

In the ash we identified quartz (SiO<sub>2</sub>), anhydrite (CaSO<sub>4</sub>), phosphate calcium silicate (Ca<sub>2</sub>SiO<sub>4</sub>-Ca<sub>3</sub> (PO<sub>4</sub>)<sub>2</sub>) and minor phases.

The ash alone mixed with water has not developed mechanical strength so it has no hydraulic character.

When they are mixed with lime in well-determined proportions to form binders, compressive strengths have been observed.

Binder L<sub>4</sub> formed from 80% ash and 20% lime gave the best compressive strength at 28d.

The binder setting start time is greater than that of cement but shorter than that of lime. This delay in setting is attributed to the presence of the elements phosphorus and zinc on the ashes as well as gypsum

The thermal conductivity of mortar made from ash and lime is lower than that of mortars made from cement and clay. This shows that the ash and lime mortar is more resistant to heat transfer by conduction than the latter two.

Mortar made from ash and lime absorbs less heat than cement and clay mortars.

The binder has good thermal characteristics and could be used as a plaster or filler wall in the construction industry.

This material used in the construction industry will therefore present better thermal characteristics than cement or clay based materials.

As perspective, other basic knowledge about dimensional stability, water demand, leaching, morphology, nature of the bonds should be investigated. Use setting accelerators to

reduce the observed set delay. Add plasticizers and additives to increase the compressive strength of the material.

## Conflicts of Interest

There are no conflicts to declare.

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