



# Sustainable alkali-activated binders with municipal solid waste incineration ashes as sand or fly ash replacement

Mohammad R. Irshidat<sup>1</sup> · Nasser Al-Nuaimi<sup>1</sup> · Mohamed Rabie<sup>1</sup>

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## Abstract

The present study investigates the feasibility of using two types of municipality solid wastes incineration ashes, namely, fly ash and bottom ash in the production of sustainable alkali-activated binder. The ashes are collected from the incineration plant and characterized to determine their particle size distribution, specific gravity, chemical composition, and heavy metals content. The ashes are then used as either fly ash or sand replacement with five replacement ratios 0%, 5%, 10%, 15%, and 20% to produce the binder. The produced binder are characterized in terms of strength, workability, density, water absorption, thermal conductivity and stability, chemical composition, and heavy metals content. The results reflect the ability of producing sustainable alkali-activated binder with small dosage of MSWI ashes as either fly ash or sand replacement without negatively affecting its strength, workability, density, and water absorption. The ashes enhance the thermal insulation capability of the binder.

**Keywords** Municipal solid waste · Incineration ashes · Alkali-activated binder · Mechanical properties · Thermal characteristics · Heavy metals

## Introduction

Rapid urbanization and social civilization have accelerated the generation of municipal solid wastes (MSW) all over the world. In 2016, two billion tons of MSW were generated around the world, this number is expected to greatly increase in the future [1]. It has been reported that the annual generation of the MSW in the United States, Europe, China, and India are 238.5, 241, 300, and 238 million tons [2, 3], respectively. The huge generation of the MSW draws the attention of the municipalities and nations toward this troubling issue as it can cause serious problems to the human health and environment. Accordingly, an effective way to recycle these wastes become necessary all over the world. The traditional treatment solutions of the MSW include recycling, landfilling, or incineration [3–5]. Among the

mentioned solutions, landfilling is the conventional technique that globally used to deal with these wastes. However, landfilling MSW owns many drawbacks, such as possible contaminating water and soil, greenhouse gas emission, and limited available spaces for landfill in most countries [4]. These drawbacks and the environmental awareness encourage the nations to look for more practical and sustainable techniques to get rid of these wastes. In this regard, incineration represents a more promising technique to implement. Incineration of MSW not only reduces the volume of the wastes by 90% [6] and kills the viruses and bacteria at such high temperatures [7], but also allows producing energy from waste. The waste-to-energy principle led to change the traditional concept of considering wastes as pollution towards treatment of the wastes as resources [3].

Despite of the above-mentioned benefits of MSW incineration, it produces considerable amount of solid residues, mainly bottom (BA) ash and fly ash (FA). Unfortunately, the heavy metals contained in the wastes migrate into the ashes. Due to the significant presence of these toxic chemicals, many countries consider the municipal solid waste incineration (MSWI) ashes, especially the FA, as hazardous materials [8]. Thus, further treatment of these wastes are highly required. Many treatment techniques of the MSWI

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Mohammad R. Irshidat on leave from Jordan University of Science and Technology (JUST), Jordan.

✉ Mohammad R. Irshidat  
mirshidat@qu.edu.qa

<sup>1</sup> Center for Advanced Materials (CAM), Qatar University,  
P.O. Box 2713, Doha, Qatar

ashes have been utilized including underground storage or landfilling, thermal and chemical treatments. Among the aforementioned techniques, landfilling is the simplest one. However, this technique requires a large space of land [9], and faces the regulations and policies of many countries due to the potential health and environmental concerns [5]. On the other hand, thermal and chemical treatments have been employed to extract the heavy metals from the MSWI ashes. Despite the efficiency of these two techniques, many concerns have been reported in the literature such as the extra energy consumption required by the thermal treatment and the long-term stability issue of the chemical treated ashes [5].

Recently, solidification or immobilization of the heavy metals by incorporating them in the production of alkali-activated (geopolymer) composites were successfully employed as new treatment techniques. During the geopolymerization process, reaction products with 3D network are formulated with the ability of encapsulating the heavy metals inside the matrix [10]. Thus, incorporating MSWI ashes in geopolymer composites production has started to catch research interest. The MSWI ashes were utilized in the preparation of the alkali-activated binders as either partially replacement of the source materials [11] or as the only source material [12]. Diaz-Loya et al. and Ye et al. [13, 14] reported that even though using MSWI FA to replace the source materials decreased the mechanical strengths of the geopolymer, the products met the acceptance criteria for the waste landfilling. Other studies showed that geopolymer with good mechanical properties could be produced even with incorporating the MSWI FA as source materials [15, 16]. Wongsa et al. [17] investigated the effect of partially substitution of source materials with MSWIBA on the strength of fly ash-based geopolymer composites. Their results showed that up to 40% of the fly ash content could be replaced by MSWIBA without negatively affecting the compressive strength of the geopolymer. Ren et al. [18] showed that using silica fume and sodium silicates enhanced the compressive strength of geopolymer composites made by MSWIFA as source materials. In addition to the acceptable mechanical properties of the geopolymer composites made of MSWI ashes as the sole or partial source materials, they also benefits the solidification of heavy metals [5, 17–19].

As the alkali-activated (geopolymer) materials showed excellent potential to serve as heavy metals immobilizing agent, most of the studies focused on investigating the ability of these composites to stabilize these hazardous materials. Little focus has been given to investigate the mechanical properties of the produced geopolymers [3, 4]. Moreover, the MSWI ashes have been mainly used as source materials and the attention was paid to investigate the effect of the synthesis parameters on the heavy metal solidification and stabilization efficiency. The current research focuses

on investigating the ability of using MSWI ashes in geopolymer composites production not only as source materials replacement but also as sand replacement. The novelty of this work comes from two points: (1) little or no studies used the MSWI ashes as sand replacement in geopolymer composites, (2) little studies focused on investigating the fresh and harden properties of alkali activated binders made of MSWI ashes. In this study, MSWI bottom ash and fly ash were used as either sand replacement or fly ash replacement in alkali-activated binder production. The effect of the replacement ratio on the mechanical strengths, workability, density, water absorption, thermal insulation capacity, thermal stability, and chemical composition of the fly ash-based geopolymer composites were explored.

## Experimental program

### Characterization of MSWI ashes

Two MSWI ashes include fly ash and bottom ash were used in this study as received from the incineration plant at the Domestic Solid Waste Management Center (DSWMC) in Qatar. Comprehensive experiments were conducted to characterize these ashes before implemented into the geopolymer mix. The particle size distribution (PSD), particles' shape, specific gravity, chemical composition, and the total concentration of heavy metals of the MSWIFA and MSWIBA were investigated. The PSD is plotted in Fig. 1. It is clear in the figure that the MSWIFA owned finer particles and narrow and less varied PSD than MSWIBA. The size of the MSWIBA particles was close to the size of the silica sand particles used in preparing the alkali-activated binder. The D50 of the silica sand, MSWIBA, and MSWIFA were equal to 550  $\mu\text{m}$ , 500  $\mu\text{m}$ , and 130  $\mu\text{m}$ , respectively. According to the scanning electron microscopic (SEM) imaging

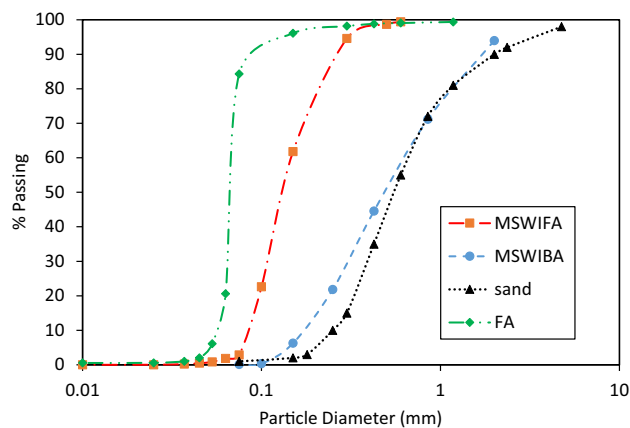
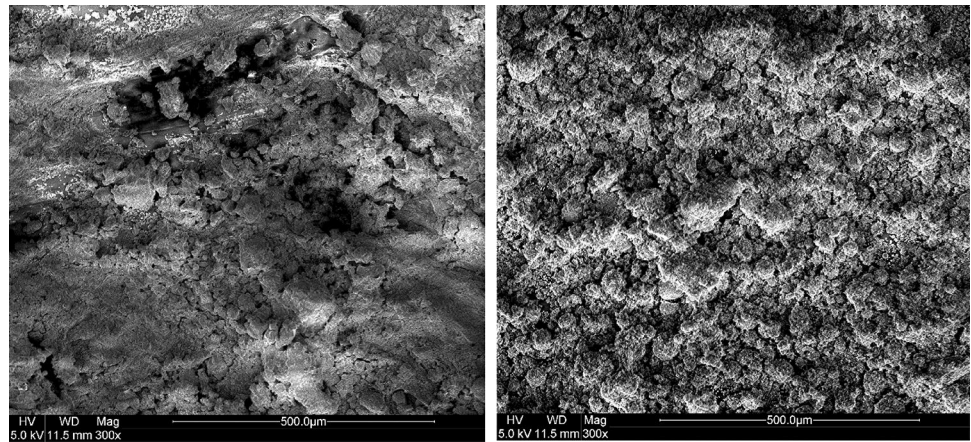


Fig. 1 Particle size distribution of fly ash, sand, and MSWI ashes

**Fig. 2** SEM imaging of **a** MWSIBA and **b** MSWIFA**Table 1** Chemical composition of ashes used in this study

Oxides	MSWIBA (%)	MSWIFA (%)	Fly ash (%)
CaO	46.8	61.8	11.80
Fe <sub>2</sub> O <sub>3</sub>	20	2.4	7.83
SiO <sub>2</sub>	9.6	6.6	49.90
Al <sub>2</sub> O <sub>3</sub>	5.1	1.2	17.10
Na <sub>2</sub> O	4.1	–	0.14
SO <sub>3</sub>	3.7	12.1	0.42
Mgo	2.5	3.4	4.90
K <sub>2</sub> O	0.1	4	0.28
ZnO	2	3.3	–
P <sub>2</sub> O <sub>5</sub>	1.9	1.4	–
CuO	1.1	0.4	–
PbO	0.8	1	–
TiO <sub>2</sub>	0.7	0.1	–
BaO	0.4	0.5	–
SnO <sub>2</sub>	0.2	0.5	–
Sb <sub>2</sub> O <sub>3</sub>	0.2	0.5	–
MnO	0.1	–	–
ZrO <sub>2</sub>	0.1	–	–
CdO	–	0.2	–
Ta <sub>2</sub> O <sub>5</sub>	–	0.1	–
Cr <sub>2</sub> O <sub>3</sub>	–	0.1	–
As <sub>2</sub> O <sub>3</sub>	–	0.1	–

(Fig. 2), both MSWIBA and MSWIFA have irregular non spherical shapes. The SEM analysis was conducted using NOVA NanoSEM 450 device. The specific gravity of the MSWIFA and MSWIBA were found to be 2.25 and 2.07, respectively. X-ray fluorescence (XRF) test procedure was conducted using JSX 3201 M (Jeol) spectroscopy machine to determine the chemical composition of the MSWI ashes. The results are summarized in Table 1. It clear that both ashes have high calcium oxide (CaO) content. Compared to MSWIFA, MSWIBA shows higher contents of alumina, silica, and iron oxide, but a lower content of sulfur trioxide (SO<sub>3</sub>). In addition, some alkali metals, such as sodium and magnesium oxides, and heavy metals such as zinc and lead were detected in both ashes. The presence of heavy metals in the ashes encouraged to conduct ICP-OES analysis to investigate the total concentration of these metals. The traced elements include Cu, Cr, Mn, Ni, Zn, and Pb. The results are presented in Table 2. The amount of heavy metals found in MSWIBA was much higher than that of MSWIFA. Compared to the values reported in the literatures [20, 21], the amount of Zn and Cr founded in MSWIBA were below the minimum values, whereas the amount of Ni and Pb were closer to the minimum reported values than they were to the maximum. In the case of MSWIFA, the amount of Cr, Ni, and Zn were found to be below the minimum values reported in the literature, whereas the amount of Cu and Pb were closer to the minimum reported values than they were to the maximum.

**Table 2** Heavy metals concentrations in the incineration ashes

Sample ID	Concentration (µg/g)					
	Cu	Cr	Mn	Ni	Zn	Pb
MSWIFA	378	30	97	11	2397	813
MSWIBA	5341	115	426	27	1288	1642
(Min–max) <sup>a</sup>	330–5530	120–2026	NA	17–614	3800–16,800	640–5500

<sup>a</sup>Funari et al. [20]

## Materials

The alkali-activated binder was prepared with class F fly ash, silica sand, Sodium Hydroxide (NaOH) and Sodium Silicate ( $\text{Na}_2\text{SiO}_3$ ). In addition, (PC 485, EPSILONE) superplasticizer was used to enhance the workability of the mix. The fly ash was received from SMEET Qatar, and characterized in terms of particle size distribution, chemical composition, moisture content, and density. Figure 1 shows that the fly ash owned finer particles and narrow and less varied PSD than the MSWI ashes. The chemical composition of the fly ash is complying with ASTM C618-12a standards, as shown in Table 1. The density and the moisture content of the fly ash were equal to  $2.23 \text{ g/cm}^3$  and 0.5%, respectively. Locally available silica sand with specific gravity, fineness modulus, and water absorption of 2.56, 2.31, and 1.87%, respectively, was used. The sand conforms to ASTM C778 standard. The Sodium Silicate solution was received from Qatar Detergent Company in Qatar with  $\text{Na}_2\text{SiO}_3$  concentration of 40%. The NaOH solution was prepared using NaOH pellets with purity of 98% that was acquired from a local supplier. The NaOH solution with concentrations of 10 Molar was prepared by mixing 320 g of NaOH pellets in one liter of distilled water. Since the reaction between the NaOH and water is exothermic reaction, the solution was prepared in a big plastic container immersed in a water bath to accommodate the heat of the reaction. The solution was left at room temperature for 30 min to lower its temperature, and was then poured in an airtight glass jar to prevent the reaction with air.

## Mix design and alkali-activated binder preparation

The MSWI ashes were incorporated into the mixes as either fly ash replacement or sand replacement. For each approach, four different replacement ratios were used: 5%, 10%, 15%, and 20% by weight of fly ash or sand. The purpose of the replacement was to investigate the amount of the MSWI ashes that could be incorporated into the mixes without negatively affecting the mechanical and physical properties of the binders. Twenty different mixes were used to prepare the alkali-activated binders, as shown in Table 3. All mixes were prepared with NaOH solution with ten molarity (10 M), fluid to binder ratio of one, and Sodium Silicates to Sodium Hydroxide ratio of 2.0. The specimens were designated by BA or FA, which refer to the type of the incineration ash, followed by letters S or F, which refers to sand or fly ash, and followed by number refers to the replacement ratio.

The alkali-activated binder was prepared by mixing the NaOH and  $\text{Na}_2\text{SiO}_3$  solutions for 2 min in a glass-measuring jar. The MSWI ashes were dry mixed with either sand or fly ash. The activator solutions were placed in the mixing bowl, and fly ash (with or without MSWI ashes) were added and mixed in the mixing bowl for 30 s at low speed. After

that, the sand (with or without MSWI ashes) was added to the mix and mixes for 30 s at low speed. The mixing speed increased to high speed and the mixing continues for 120 s. At this point, the flow table test was conducted, followed by 15 s mixing at high speed. The prepared binder was used to cast specimens with dimensions of  $5 \text{ cm} \times 5 \text{ cm} \times 5 \text{ cm}$  and  $4 \text{ cm} \times 4 \text{ cm} \times 16 \text{ cm}$  to perform the compressive and flexural strength tests, respectively. All geopolymer mortar specimens were cured at a fixed temperature of  $80 \text{ }^\circ\text{C}$  and for a duration of 24 h. After curing, the specimens were tested according to the procedures mentioned in the following section.

## Test procedures

### Mechanical strengths

The mechanical behavior of alkali-activated binder with MSWI ashes was investigated through compressive and flexural strength tests according to the ASTM C109 and ASTM C348 standards, respectively. The specimens were loaded until failure with loading rate of 1.3 kN/s and 0.044 kN/s for compression and flexural tests, respectively. The average strength value of three tested specimens for each mix was reported.

### Workability

Effect of MSWI ashes on the workability of the alkali-activated binder was investigated using flow table test according to the ASTM C1437 standards. The average of four readings for each mix was calculated and reported.

### Density

The effect of MSWI ashes on the density of the produced binder was determined according to the following procedure: three 50 mm cubes for each mix design were placed in an oven for 24 h at  $110 \text{ }^\circ\text{C}$ . The specimens were removed from the oven and cool in an ambient temperature for 2 h, and the dry weight was recorded for all specimens. The oven dry density was calculated by dividing the oven dry weight by the volume of specimen. The average of three density values of each mix were reported.

### Thermal conductivity

Thermal conductivity of alkali-activated binder with various dosages of MSWI ashes as either fly ash replacement or sand replacement was examined using a hot disk thermal constant analyzer machine. The sensor of the machine was sandwiched between two-binder specimens, and then the transient plane source (TPS) method was used to acquire

**Table 3** Alkali-activated binder mix designs

Specimen	Mix ID	NaOH (M)	F/B	Na <sub>2</sub> SiO <sub>3</sub> /NaOH	Fly ash (kg/m <sup>3</sup> )	MSWIFA (kg/m <sup>3</sup> )	MSWIBA (kg/m <sup>3</sup> )	Sand (kg/m <sup>3</sup> )	NaOH solution (kg/m <sup>3</sup> )	Na <sub>2</sub> SiO <sub>3</sub> solution (kg/m <sup>3</sup> )	SP (kg/m <sup>3</sup> )
FA-S-0	M1	10	1	2	711	–	–	1956	171	256	7
FA-S-5	M2				711	98	–	1858			
FA-S-10	M3				711	196	–	1760			
FA-S-15	M4				711	293	–	1662			
FA-S-20	M5				711	391	–	1564			
BA-S-0	M6				711	–	–	1956			
BA-S-5	M7				711	–	98	1858			
BA-S-10	M8				711	–	196	1760			
BA-S-15	M9				711	–	293	1662			
BA-S-20	M10				711	–	391	1564			
FA-F-0	M11				711	–	–	1956			
FA-F-5	M12				676	36	–	1956			
FA-F-10	M13				640	71	–	1956			
FA-F-15	M14				604	107	–	1956			
FA-F-20	M15				569	142	–	1956			
BA-F-0	M16				711	–	–	1956			
BA-F-5	M17				676	–	36	1956			
BA-F-10	M18				640	–	71	1956			
BA-F-15	M19				604	–	107	1956			
BA-F-20	M20				569	–	142	1956			



the thermal conductivity data. The size of specimens was 40 mm × 40 mm × 80 mm, which was sufficient to cover the full area of the sensor. The humidity condition was measured and reported to be 55%RH during the thermal conductivity test.

### Water absorption

Water absorption of alkali-activated binder specimens was measured according to the ASTM C1403. Three 50 mm cubes for each mix design were placed in an oven for 24 h at 110 °C. The specimens were removed from the oven and cool in an ambient temperature for 2 h. The dry weight was recorded for all the specimens, and then the binder specimens were placed in a container. The specimens were partially immersed in water with depth of  $3 \pm 0.5$  mm. The binder specimens' weight were recorded at 0.25 h, 1 h, 4 h and 24 h. After that, the weight was recorded on a daily basis until there were no difference between the two successive weightings.

### Thermal stability

Thermogravimetric analysis (TGA) test was performed to investigate the effect of MSWI ashes on thermal stability of alkali-activated binder. Fragments extracted from selected specimens were first grinded to a size of 45  $\mu$ m then tested using Thermogravimetric Analyzer TGA 4000 PerkinElmer device in the range of 30 °C and 730 °C. The mass loss in the specimens was monitor with respect to the temperature.

### Chemical composition analysis

The effect of MSWI ashes on the chemical composition of the produced binder was examined using X-ray fluorescence (XRF) technique. Small pieces were taken out from different

location of the compression specimens, grinded to powder that pass sieve with 150  $\mu$ m opening, and tested using JSX 3201M (Jeol) spectroscopy device.

### Heavy metals concentrations

The total concentration of the heavy metals in the MSWI ashes and binder specimens was measured by inductively coupled plasma-Optical Emission (ICP-OES). Acid digestion followed by ICP-OES were used for samples elemental analysis. The digestion was optimized using 0.26 g of sample, 6 ml of HCl, 2 ml of HNO<sub>3</sub>, and 2 ml of HF using microwave digestion system (MARS 6). ICP-OES was then conducted for elemental analysis using Optima 7300 DV (PerkinElmer).

## Experimental results and discussion

### Workability

Effect of MSWI ashes on the workability of alkali-activated binder was examined using flow table test. The spread diameter values of different binder mixes are plotted in Fig. 3. The effect of MSWIFA on the workability of the binder was more pronounced in the case of using it as sand replacement than fly ash replacement, as shown in Fig. 3a. For fly ash replacement approach, using 5% replacement ratio increased the workability of the binder by 17%. Any further replacement of the fly ash with MSWIFA up to 20% reduced the workability of the binder to be close to the control mix. In the case of sand replacement approach, using 5% replacement ratio increased the workability of the binder by 22%. Binder with 10% MSWIFA as sand replacement owned similar workability of the control mix. Increase the replacement ratio to be 15% clearly reduced the workability by 18%

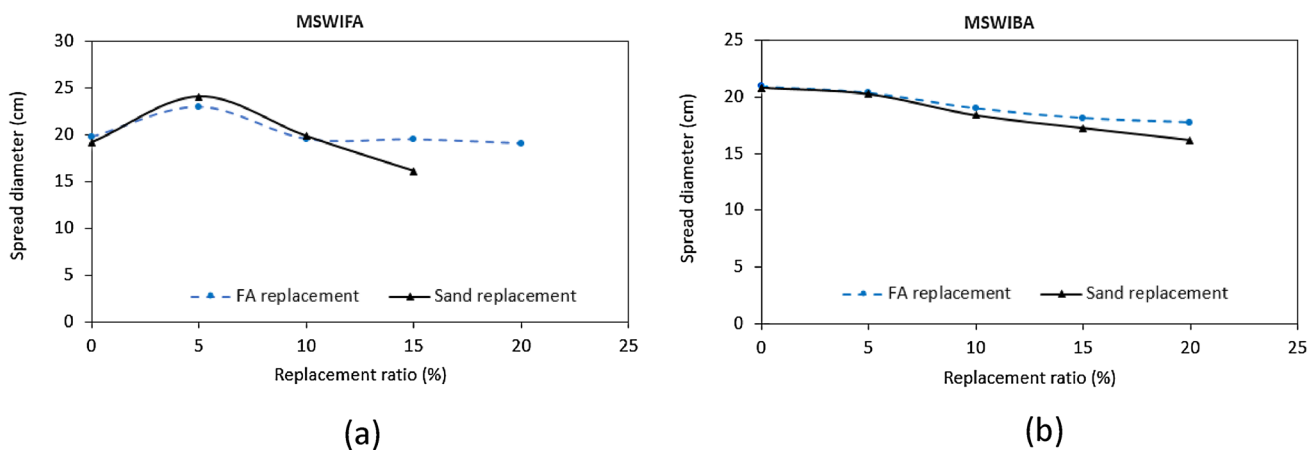


Fig. 3 Flow table results of alkali-activated binder with MSWI ashes

compared to the control specimen. It is good to mention that specimen with 20% sand replacement ratio showed flash setting during mixing process, thus the spread diameter could not be measured.

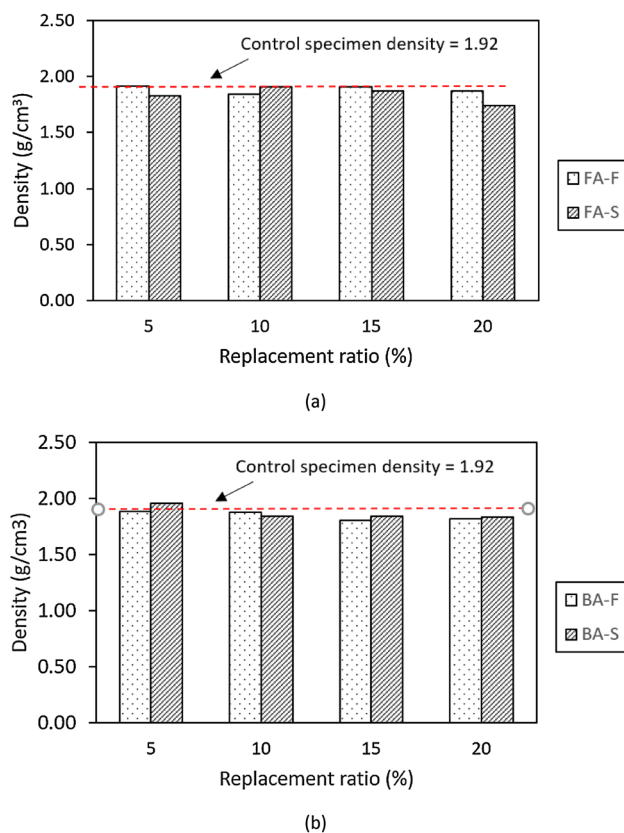
The effect of MSWIBA on the workability of the binder is shown in Fig. 3b. It is clear in the figure that both approaches; sand and fly ash replacement; showed similar effect on the workability of the binder. Replacement 5% of either fly ash or sand content by MSWIBA marginally increased the workability of the control mix by 3%. After that, the workability of the binder decreased by 10% and 18% with increasing the MSWIBA content to be 20% for fly ash replacement and sand replacement approaches, respectively. This reduction might be ascribed to (1) the high water adsorption of the MSWI ashes [22]. The water absorption of the fly ash, MSWIFA, and MSWIBA used in this study was equal to 1.15%, 5%, and 4%, respectively. Or (2) the irregular shape and rough surface of the MSWI ashes [23, 24].

### Density

Figure 4 shows the density of the alkali-activated binder with different dosages of MSWI ashes as either fly ash or sand replacement. It was noted that incorporating the ashes in the binder insignificantly affected its density. The density of the control specimen was equal to  $1.92 \text{ g/cm}^3$ . The density of specimens with different dosages of either MSWIFA or MSWIBA located within 5% of the control specimen. The slight reduction in the density due to the incorporation of the MSWI ashes could be attributed to the lighter specific gravity and porous nature of the ashes [22].

### Mechanical strengths

Effect of incorporating MSWI ashes as either fly ash replacement or sand replacement on the mechanical strengths of alkali-activated binder is discussed in this section. The strength values combined with the reduction ratios and the standard deviation values are summarized in Table 4. The compressive strength of alkali-activated binder without MSWI ashes (control specimen) equals to 40.93 MPa. Replacement up to 10% of the fly ash content with MSWIFA marginally reduced the compressive strength of the binder by 5%. Any more replacement of the fly ash with MSWIFA caused clear reduction in the compressive strength, the reduction reached up to 46% compared to the control specimen in the case of 20% replacement ratio. Similar results were reported in the literature [4, 5]. On the other hand, replacement 5% of the fly ash content with MSWIBA marginally reduced the compressive strength of the binder by 3%. Any more replacement caused clear reduction in the compressive strength. The reduction reached up to 35% compared to the control specimen in the case of



**Fig. 4** Density of alkali-activated binder with different dosages of MSWI **a** FA, **b** BA as either fly ash or sand replacement

20% replacement ratio. The reduction in the mechanical strengths of alkali-activated binder due to fly ash replacement with MSWI ashes could be attributed to the different mineral compositions between the fly ash and the incineration ashes, as shown in Table 1. Compared to fly ash, MSWIFA and MSWIBA contain very less amount of  $\text{SiO}_2$  and  $\text{Al}_2\text{O}_3$ , which contribute to the mechanical strengths of the alkali-activated binder [5, 17, 25]. The low reactivity of the MSWI ashes compared to the fly ash could be another reason beyond the reduction in the compressive strength of the binder specially for high replacement ratio [4]. Figure 5a shows that for fly ash replacement, using MSWIFA is better than MSWIBA. On contrary, for sand replacement, it is better to use the MSWIBA than the MSWIFA especially for replacement ratio more than 10%, as shown in Fig. 5b. However, for small replacement ratio (5%), both ashes showed neglected reduction in the compressive strength of the binder, as shown in Table 4. In addition, the results show that it is better to use the MSWIFA as fly ash replacement rather than sand replacement (Fig. 5c), whereas it is better to use the MSWIBA as sand replacement rather than fly ash replacement, as shown in Fig. 5d. These results could be attributed to the fact that the size of the MSWIBA particles is closer to the size of the sand particles, whereas the size of

**Table 4** Mechanical strengths results

Sample ID	Compressive strength (MPa)	Std	Reduction (%)	Flexural strength (MPa)	Std	Reduction (%)
Control	40.93	0.71		0.59	0.01	
FA-F-5	40.68	0.06	1	0.55	0.01	8
FA-F-10	38.90	0.06	5	0.44	0.01	26
FA-F-15	27.70	1.09	32	0.34	0.02	42
FA-F-20	22.24	0.31	46	0.24	0.00	59
Control	40.93	0.71		0.59	0.01	
BA-F-5	39.54	0.36	3	0.55	0.02	7
BA-F-10	32.79	0.13	20	0.48	0.04	19
BA-F-15	27.13	0.76	34	0.28	0.02	53
BA-F-20	26.57	0.32	35	0.29	0.03	52
Control	40.93	0.71		0.59	0.01	
FA-S-5	41.30	0.49	– 1	0.38	0.01	36
FA-S-10	28.71	0.48	30	0.36	0.01	39
FA-S-15	21.09	0.17	48	0.30	0.03	50
FA-S-20	7.79	0.23	81	0.14	0.03	76
Control	40.93	0.71		0.59	0.01	
BA-S-5	39.81	1.26	3	0.59	0.05	0
BA-S-10	35.03	1.49	14	0.53	0.01	10
BA-S-15	33.34	0.52	19	0.39	0.02	35
BA-S-20	30.59	1.35	25	0.39	0.03	35

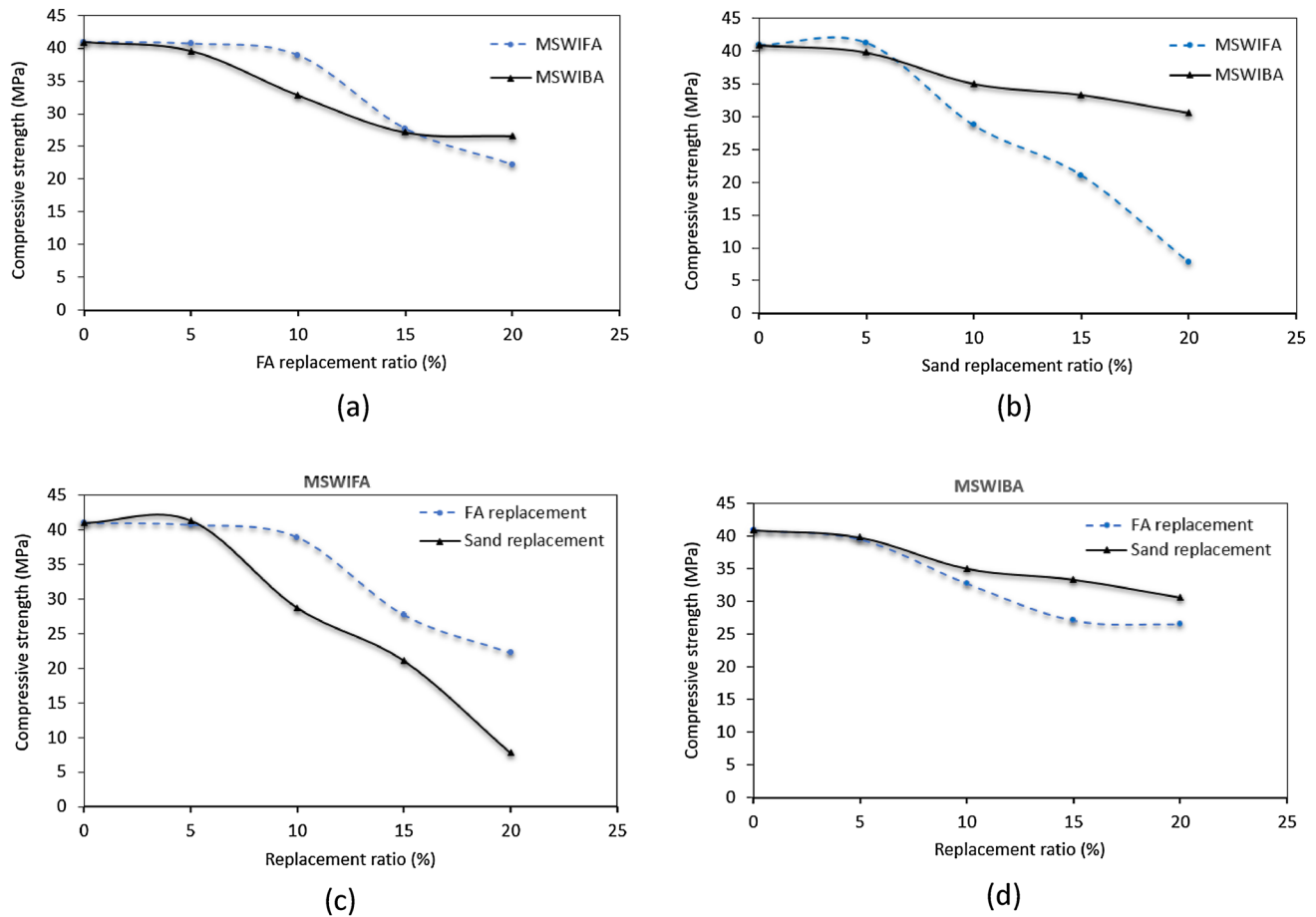
the MSWIFA particles is closer to the fly ash particles [26], as shown in Fig. 1. When the gradation is appropriate, the cemented materials has relatively dense microstructure and superior structural performance [27, 28]. In addition, the size of the particles involved in the mix has a great effect on the strength of the cementitious binders, where the size of the sand particles affects the number and the size of the voids through the matrix as reported in many literatures [23, 29].

The flexural strength of alkali-activated binder without MSWI ashes (control specimen) equals to 0.59 MPa. The results show that using small dosages (5%) of either MSWIFA or MSWIBA as fly ash replacement caused marginal reduction to the flexural strength of the binder by 8% and 7%, respectively, as shown in Fig. 6a. The reduction increased with increasing the dosages of the MSWIFA and MSWIBA to reach up to 52% and 59% of the flexural strength of the control specimens for replacement ratio of 20%, as cleared in Table 4. For sand replacement approach, using 5% of MSWIBA did not affect the flexural strength of the binder, as shown in Fig. 6b. Increasing the dosage to be 10% reduced the strength by 10%. Further increase in the MSWIBA content caused clear reduction in the flexural strength to be 35% in the case of 20% replacement ratio. On contrary, MSWIFA is not good to be used as sand replacement, since using small dosages (5%) caused significant reduction in the flexural strength of the binder by 36%, as shown in Table 4.

### Thermal insulation capacity

Effect of MSWI ashes on the thermal insulation capacity of alkali-activated binder was investigated using thermal conductivity test. The results are presented in Fig. 7. It is clear that sand or fly ash replacement with small dosages of MSWIFA (up to 10%) slightly affected the thermal insulation capacity of the binder. No changes in the thermal conductivity were monitored in the case of fly ash replacement, whereas insignificant increase by 6% and 4%, where noticed in the case of 5% and 10% sand replacement, respectively. For high replacement ratio, significant reduction in the conductivity was noticed especially for sand replacement. Maximum reduction was equal to 21% in the case of 15% sand replacement. On contrary, clear enhancement in the thermal insulation capacity of the alkali-activated binder was noticed in the case of MSWIBA incorporation as either fly ash or sand replacement. Figure 7b shows that presence of MSWIBA reduced the thermal conductivity of the binder for all replacement ratios and for both fly ash and sand replacement. The reduction in the conductivity values in the case of fly ash and sand replacement started at 17% and 14%, respectively, for 5% replacement ratio and increased up to 34% and 29% in the case of 20% replacement ratio. Similar results reported in the literature [30, 31]. It was also noted that specimens with MSWIBA owned lower thermal conductivity than those with MSWIFA. This could





**Fig. 5** Compressive strength of alkali-activated binder with MSWI ashes

be attributed to the high porosity and low density of the BA compared to the FA [32].

### Water absorption

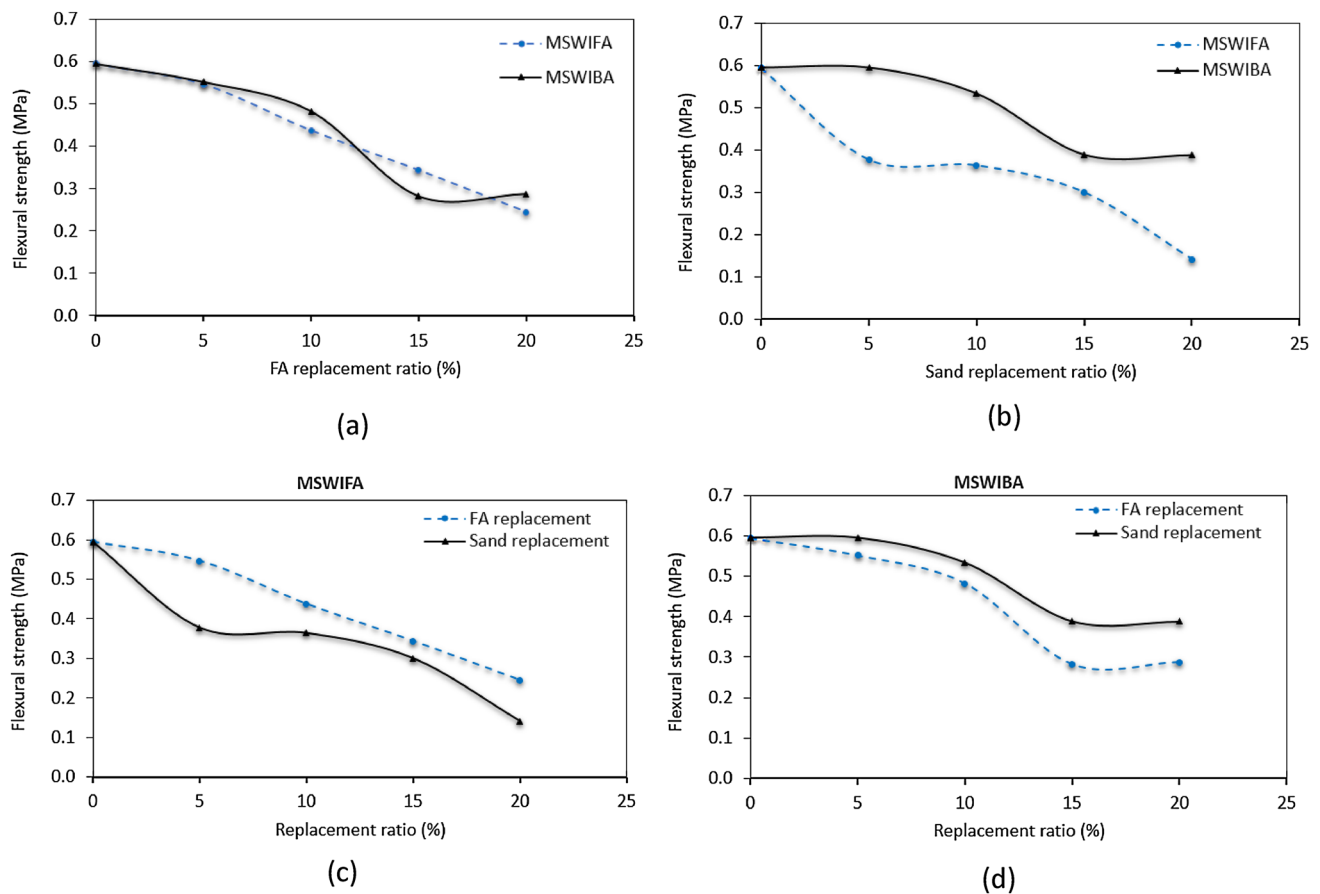
Effect of MSWI ashes on the water absorption capability of the alkali-activated binder was investigated, and the results are presented in Fig. 8. It was noted that the presence of MSWI ashes in the binder increased its water absorption capability. This result attributed to the higher water absorption of the MSWI ashes compared to both fly ash and sand [23]. For both approaches (fly ash and sand replacement), the increase in the water absorption was clear in the specimens with MSWIFA, as shown in Fig. 8a, b, but it was insignificant in the specimens with MSWIBA, as shown in Fig. 8c, d. In addition, the water absorption capability increased with increasing the MSWI ashes dosages. Similar results are reported in the literature [5, 33]. Specimens with MSWIFA showed higher water absorption capability than those with MSWIBA due to the facts that the MSWIFA owned smaller particle size and higher surface area than MSWIBA. Finally, it was noted that all specimens reached the plateau stage

with constant absorption after 24 h, reflected the high rate of absorption of the binder at early ages.

### Thermal stability

Specimens with best mechanical strength results (5% replacement ratio) were selected to perform the TGA test to compare the replacement approach for both MSWI ashes. Thermogravimetric (TGA) profiles and their corresponding derivative function (DTG) curves of the selected alkali-activated binder specimens are presented in Fig. 9. The figure gives an indication about the effect of adding MSWI ashes to the binder as either fly ash or sand replacement on the weight loss of the specimens when exposed to elevated temperature up to 730 °C.

Figure 9 reveals that all TGA curves could be divided into four stages. Stage one (S1) was located between 0 to 120 °C. The weight loss in this stage was associated with the evaporation of the free water presented on the surface and porosity of the specimens [34, 35]. The second stage (S2) was located between 120 to 230 °C. The weight loss in this stage could be attributed to the evaporation of the chemically



**Fig. 6** Flexural strength of alkali-activated binder with MSWI ashes

bound water [34, 36], or was associated with the presence of NASH gel [37]. After the second stage, the rate of the weight loss stabilized between 230 to 630 °C (S3). Another weight loss was captured between 630 and 730 °C (S4). This weight loss could be attributed to the dehydroxylation and recrystallization of geopolymer specimens [35].

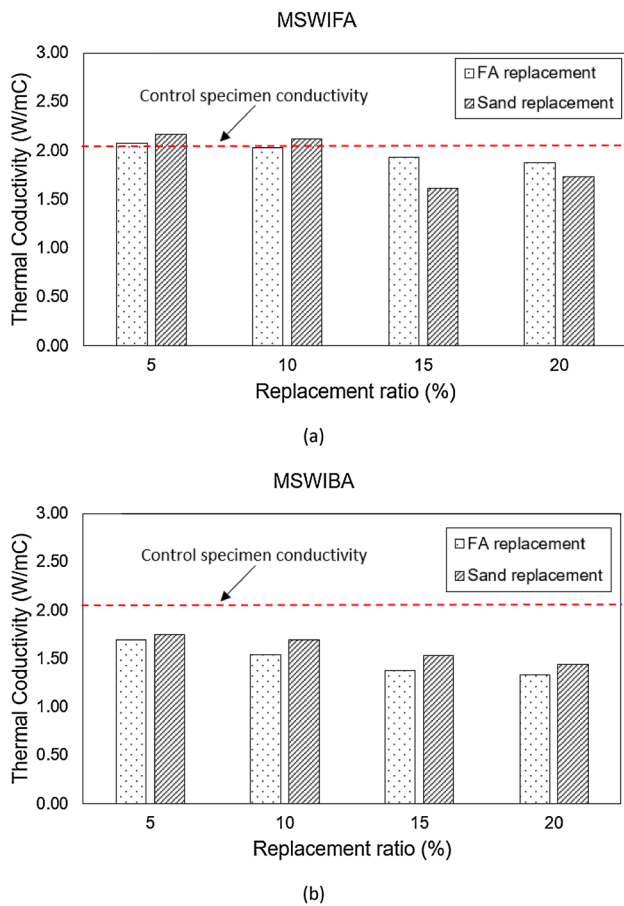
It is clear in Fig. 9a, b that the total weight loss of the fly ash-based geopolymers did not exceed 8%, which agreed with the literatures [38, 39]. It is also clear that the effect of the fly ash replacement on the total weight loss of the binder was more pronounced than the sand replacement for both types of the ashes, as shown in Fig. 9a, b.

In addition, the first peak in the DTG curves of alkali-activated binder with MSWIFA got higher and wider than that of binder with MSWIBA, as shown in Fig. 9c. This result could be attributed to the following two reasons. (1) The fact that specimen with 5% MSWIFA as sand replacement contains more calcium (Ca) atoms than specimen with 5% MSWIBA as shown later in the XRF results; therefore, it can hold pore water more strongly [34]. Knowing that presence of Ca in the geopolymer gel is helpful in holding the pore water through the formation of new phases [40]. (2)

The fact that specimen with 5% MSWIFA as sand replacement absorb more water compared to specimen with 5% MSWIBA as shown in the previous section. Thus, evaporation of more free water from the specimen would take place, which caused more increase in the mass loss.

### Chemical composition and heavy metals concentrations

XRF test was conducted to determine the chemical composition of alkali-activated binder with various dosages of MSWI ashes as either fly ash or sand replacement. The results are summarized in Table 5. It is clear in the table that the main oxides presented in the control specimens are SiO<sub>2</sub>, CaO, Al<sub>2</sub>O<sub>3</sub>, and Fe<sub>2</sub>O<sub>3</sub>. These oxides come from the fly ash that used as source material in the binder mix design, as shown in Table 1. Clear reduction in the silica content was noted for specimens with both MSWIFA and MSWIBA. The reduction was more significant in the case of sand replacement than fly ash replacement. The reason beyond that could be attributed to high silica content in the sand used to prepare the mix (silica sand). On contrary, clear



**Fig. 7** Thermal conductivity of alkali-activated binder with different dosages of MSWI **a** FA, **b** BA as either fly ash or sand replacement

increase in the alumina was noted specially for sand replacement specimens, this could be due to the high content of aluminum oxides in the MSWI ashes.

Since some of the heavy metals were observed in the binder by the XRF analysis, ICP-OES analysis was carried out to investigate the total concentration of these metals in alkali-activated binder with MSWI ashes. The traced elements include Cu, Cr, Mn, Ni, Zn, and Pb. The results are presented in Table 6. The concentrations of Zn, Mn, and Cr were the highest among all other traced metals in the binder specimens. However, their concentrations showed different tendency with the increase of MSWI ash content. With the increase of the MSWI ash replacement ratio, the concentration of Cr tended to decrease, whereas the concentration of Zn tended to increase. The variation in the heavy metals concentration within the binders may indicate to possible migration of some of these metals into the binder during the geopolymerization process. The migration of these metals depends on the replacement ratio. The literatures reported that the heavy metals either transferred from the incineration ashes into the alkali-activated binder, physically

encapsulated by the binder, or chemically encapsulated by the binder through their interaction with the geopolymerization products [5, 41].

On the other hand, leaching of heavy metals is a major concern for any potential application of MSWI ashes in construction industry. Due to the material's potential applications, further research into the mobility of these metals through a leaching procedure is highly recommended. However, many published literatures proved the ability of the alkali-activated composites to stabilize and immobilize the heavy metals through the geopolymerization process [5, 17–19], thus reduced the leaching amount of the toxic metals to be within the limits.

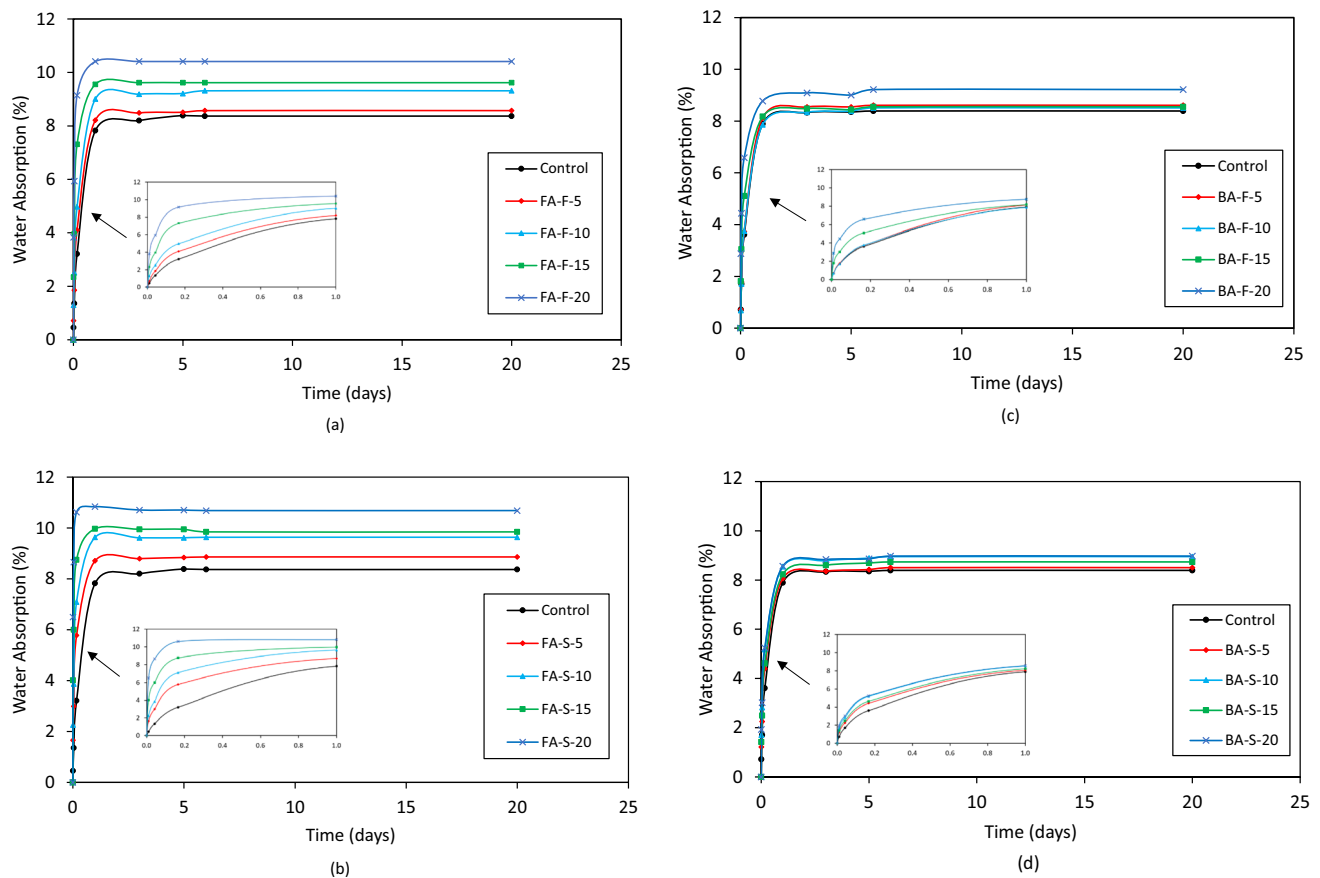
## Recommendation and future work

Investigating the effect of MSWI ashes on the durability of alkali-activated binders is an essential, as it affects the decision to incorporate these wastes in the construction industry. In this regards, it is recommended to investigate the effect of these ashes on dry shrinkage, sulfate acid resistance, and chloride permeability of the binders. In addition, comprehensive study should be conducted to investigate the variation in the chemical composition of the ashes with time and waste source.

## Conclusions

The feasibility of using two types of municipality solid wastes incineration ashes (MSWIA), namely, fly ash and bottom in the production of sustainable alkali-activated binder was investigated in this study. The incineration ashes were incorporated in the binder mix as either fly ash or sand replacement with five replacement ratios 0%, 5%, 10%, 15%, and 20%. The produced binder were characterized in terms of mechanical strengths, workability, density, water absorption, thermal conductivity, thermal stability, chemical composition, and heavy metals content. The main conclusion of this study was that sustainable alkali-activated binder with better thermal insulation capability could be produced with small dosages (5%) of MSWI ashes without negatively affecting the binder strengths, workability, density, and water absorption. The following detailed conclusions could be extracted from the experimental work presented in this research:

1. To produce sustainable alkali-activated binder with MSWI ashes, the incineration fly ash could be used to replace 5% of the fly ash content, whereas the incineration bottom ash could be used to replace 5% of the fly



**Fig. 8** Water absorption of alkali-activated binder with MSWIFA **a** FA replacement, **b** sand replacement and MSWIBA, **c** FA replacement, **d** sand replacement

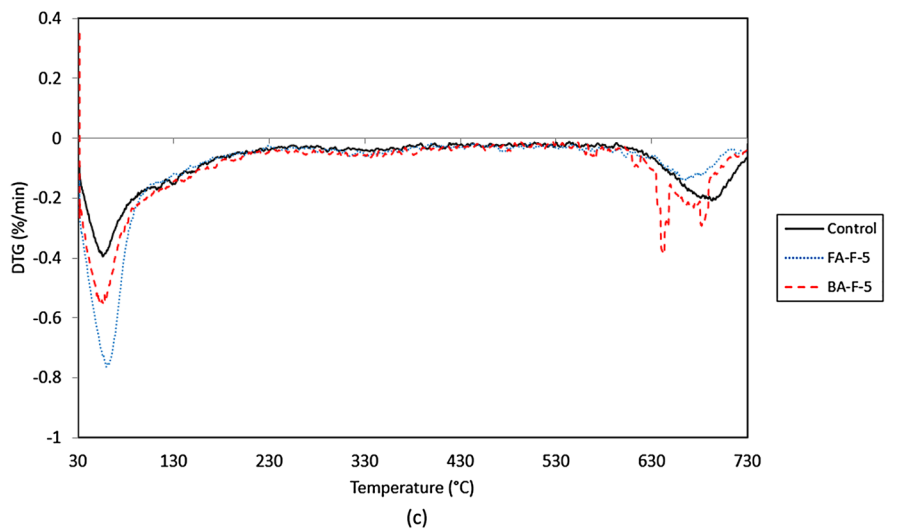
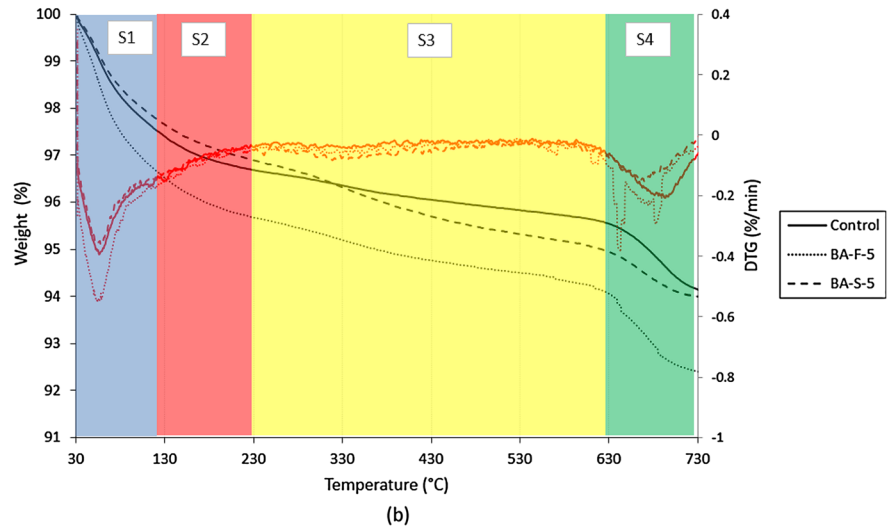
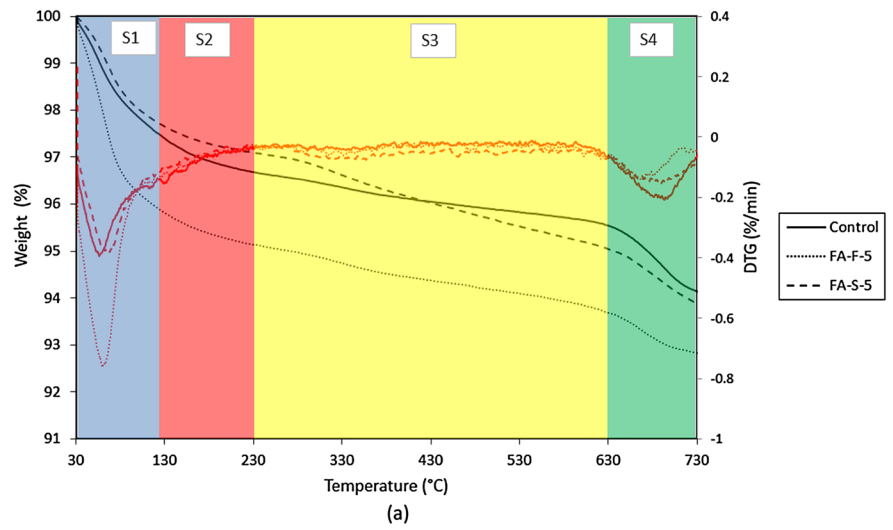
ash content or 10% of the sand content without negative effect on the mechanical strengths of the binder.

- Small dosages of MSWI ashes (up to 10%) slightly affected the workability of the alkali-activated binder, whereas high dosages clearly reduced the workability. The reduction was more pronounced in the case of MSWIBA.
- Incorporating MSWI ashes as either fly ash or sand replacement insignificantly affected the density of the alkali-activated binder.
- MSWI ashes enhanced the thermal insulation properties of the alkali-activated binder. The presence of MSWIBA reduced the thermal conductivity of the binder for all studied replacement ratios and for both fly ash and sand replacement. Small dosages of MSWIFA slightly

affect the conductivity, whereas large dosages clearly enhanced it.

- Presence of MSWI ashes in the alkali-activated binder increased its water absorption capability. The water absorption capability increased with increasing the MSWI ashes dosages. Specimens with MSWIFA showed higher water absorption capability than those with MSWIBA
- The heavy metals presented in the MSWI ashes migrated into the alkali-activated binder. The migration of these metals depends on the replacement ratio. Investigating the mobility of these metals through a leaching procedure is highly recommended before any potential application of the incineration ashes in the construction industry.

**Fig. 9** TGA and DTG curves for alkali-activated binder with 5% MSWI ashes as sand or fly ash replacement





**Table 5** Chemical composition of alkali-activated binder with MSWI ashes (wt%)

Sample	TiO <sub>2</sub> (%)	Al <sub>2</sub> O <sub>3</sub> (%)	SO <sub>3</sub> (%)	CaO (%)	SiO <sub>2</sub> (%)	MgO (%)	MnO (%)	Fe <sub>2</sub> O <sub>3</sub> (%)	K <sub>2</sub> O (%)	Cu (%)	Cl (%)	Zn (%)	Sr (%)	Cr (%)	Sc (%)
FA-F-0	0.6	10.9	1.6	13.3	55.8	4.0	0.2	11.2	1.9	0.0	0.1	0.0	0.1	0.2	0.1
FA-F-5	0.6	10.3	1.9	14.7	55.7	2.9	0.2	10.0	2.3	0.0	1.0	0.0	0.1	0.1	0.1
FA-F-10	0.6	9.7	2.2	15.6	53.7	2.3	0.2	10.6	2.3	0.0	2.3	0.1	0.2	0.1	0.1
FA-F-15	0.7	8.1	3.9	17.7	49.5	0.0	0.2	11.4	2.8	0.0	5.4	0.1	0.2	0.1	0.1
FA-F-20	0.8	8.3	3.5	19.5	45.5	0.0	0.2	12.3	2.9	0.0	6.5	0.1	0.2	0.1	0.1
FA-S-0	0.6	11.2	2.3	13.4	54.4	4.8	0.2	10.6	2.0	0.1	0.1	0.1	0.1	0.1	0.0
FA-S-5	0.8	10.1	3.1	17.8	48.8	0.0	0.2	12.4	2.5	0.0	3.9	0.1	0.2	0.1	0.1
FA-S-10	0.8	8.9	3.0	20.0	44.3	0.0	0.2	12.8	2.8	0.0	6.8	0.1	0.2	0.1	0.1
FA-S-15	0.9	8.0	2.6	22.6	39.9	0.0	0.2	13.7	3.1	0.0	8.2	0.3	0.2	0.1	0.1
FA-S-20	0.9	7.6	2.4	23.7	37.0	0.0	0.2	13.5	3.4	0.0	10.5	0.3	0.2	0.0	0.1
BA-F-0	0.6	10.2	1.6	12.9	57.6	4.8	0.2	9.6	2.1	0.0	0.1	0.0	0.1	0.1	0.1
BA-F-5	0.5	9.9	1.5	11.5	58.6	7.4	0.1	8.1	2.0	0.0	0.1	0.0	0.1	0.1	0.1
BA-F-10	0.8	11.2	2.4	16.7	49.4	2.9	0.2	13.6	2.0	0.0	0.2	0.1	0.2	0.0	0.1
BA-F-15	0.7	11.0	2.0	15.3	48.5	7.1	0.2	12.5	1.9	0.1	0.3	0.1	0.2	0.1	0.1
BA-F-20	0.7	11.3	2.5	16.6	47.5	6.6	0.2	11.9	1.9	0.0	0.4	0.1	0.1	0.1	0.1
BA-S-0	0.6	11.3	1.7	13.2	57.7	3.1	0.1	9.7	2.1	0.0	0.1	0.1	0.1	0.1	0.0
BA-S-5	0.7	11.4	2.2	14.7	51.1	6.0	0.2	11.1	1.9	0.0	0.3	0.1	0.1	0.1	0.1
BA-S-10	0.7	11.2	2.3	16.7	51.7	1.9	0.2	12.2	2.0	0.0	0.5	0.1	0.1	0.1	0.1
BA-S-15	0.9	11.8	2.6	18.4	46.6	1.9	0.2	14.3	1.9	0.1	0.7	0.2	0.1	0.1	0.1
BA-S-20	0.8	12.0	2.2	17.9	47.6	3.9	0.2	12.3	1.7	0.1	0.7	0.2	0.1	0.1	0.1

**Table 6** Heavy metal concentrations in alkali-activated binder with MSWI ashes

Sample	Total concentration (mg/kg)					
	Cu	Cr	Mn	Ni	Zn	Pb
FA-F-0	14.79	234.7	252.6	13.05	54.61	15.73
FA-F-5	16.34	134.2	225.5	9.916	128.7	22.51
FA-F-10	21.32	108.5	236.1	10.38	208.6	32.56
FA-F-15	22.44	110.5	212.3	8.404	269.6	36.82
FA-F-20	26.62	76.31	206.8	8.194	373.5	45.24
FA-S-0	12.45	164.7	195.1	9.526	54.61	4.595
FA-S-5	24.06	91.12	230	9.937	128.7	29.25
FA-S-10	35.39	94.96	225.8	10.26	208.6	51.84
FA-S-15	28.52	51.85	146.8	6.454	269.6	49.13
FA-S-20	53.4	61.00	227.6	11.93	373.5	95.38
BA-F-0	25.2	163.6	244.6	10.75	93.48	17.7
BA-F-5	26.49	176.4	219.5	9.657	98.19	21.04
BA-F-10	126.4	47.1	267.2	12.78	209.4	33.8
BA-F-15	62.82	81.23	244.3	13.27	227.3	36.9
BA-F-20	102.4	59.67	252.9	13.24	340.2	47.2
BA-S-0	11.89	138.9	173.5	8.125	93.48	7.013
BA-S-5	51.43	102.4	190.8	14.94	98.19	21.61
BA-S-10	105.7	125.5	208.9	12.18	209.4	46.28
BA-S-15	101.9	78.79	199.1	10.23	227.3	53.59
BA-S-20	156.3	123.2	242.0	19.82	340.2	92.63

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**Data availability** Data available on request from the authors.

## Declarations

**Conflict of interest** The authors reported no potential conflict of interest.

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