



# The improvement of strength and microstructural properties of fly ash-based geopolymer by adding elemental aluminum powder

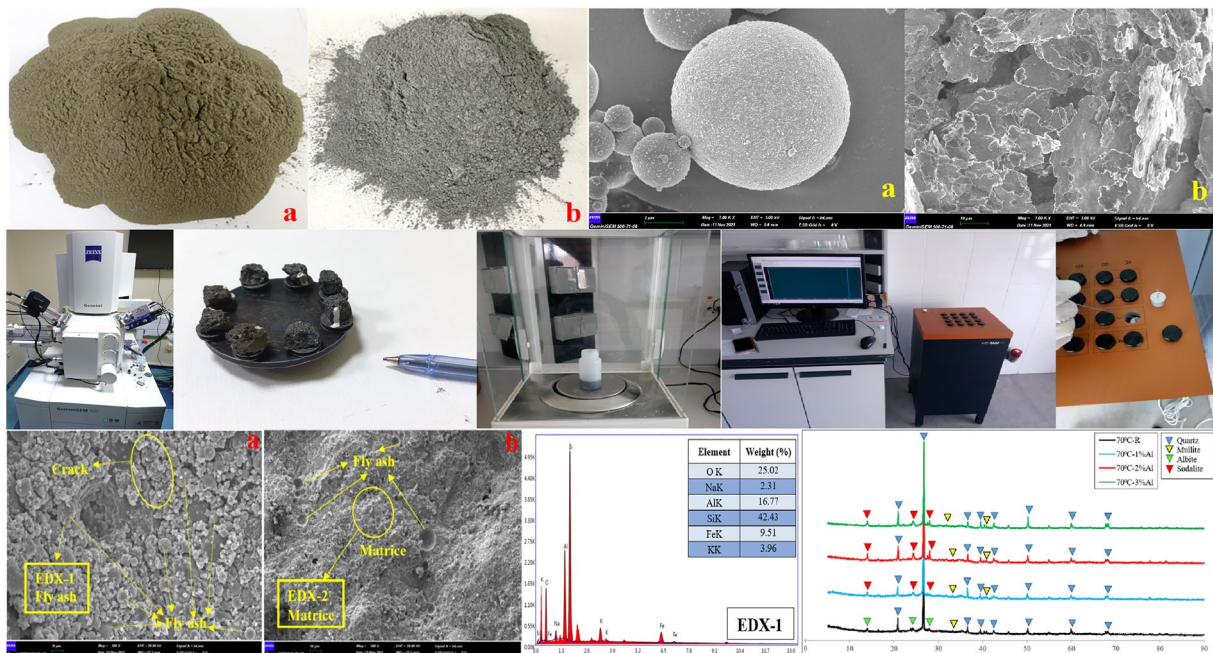
Uğur Durak<sup>1</sup>

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## Abstract

In this study, 1, 2, and 3% micronized aluminum powder were used by replacing with fly ash to improve the properties of F class fly ash-based geopolymer mortar samples. The produced geopolymer mortar samples were heat cured at 60, 70, and 80 °C temperatures for 24 and 48 h. Following heat curing, the unit weights, flexural, and compressive strengths of the hardened geopolymer mortar samples were determined. In addition, XRD, FESEM, EDX, and isothermal calorimetry experiments were carried out to examine the changes in the microstructure with the addition of aluminum in detail. Based on the results obtained, it was observed that the substitution of micronized aluminum powder slightly decreased the workability of the geopolymer mortar samples. Moreover, it increased the flexural and compressive strength significantly (more than double) for 24 h heat curing. The XRD, FESEM, EDX and isothermal calorimetry experiments showed that the substitution of aluminum powder in the fly ash-based geopolymer system increased the geopolymerization reactions and, in comparison to the reference sample, created sodalite minerals in the microstructure. Therefore, it was concluded that the aluminum powder substitution transforms the microstructure into a stronger structure, thus improving the mechanical properties.

## Graphical abstract



**Keywords** Fly ash · Aluminum · Geopolymer · Microstructure · Sodalite

Extended author information available on the last page of the article

## Introduction

Portland cement is one of the most used building materials in the world. Portland cement was used as a binding material in concrete production. It is known that around 4.1 billion tons of Portland cement is produced worldwide in 2020 [1]. In the production of Portland cement, a significant amount of carbon dioxide is released into the atmosphere and high energy consumption is caused [2–6]. According to the research, approximately 0.9 tons of carbon dioxide emissions occur for every one ton of Portland cement produced [7, 8]. With these values, Portland cement production alone is responsible for 5–7% of worldwide carbon dioxide [2, 7, 9–11]. In addition, approximately 5–12% of the energy used in production facilities in the world is spent on Portland cement production [10]. In our world, where the need for shelter is increasing, Portland cement production is increasing day by day, and in parallel to this increase, both carbon dioxide emissions and energy consumption are increasing worldwide. For this reason, researchers are investigating ways to develop new alternative binders in order to eliminate and/or reduce the unfavorable effects caused by the high use of Portland cement.

One of the alternative systems to Portland cement as a binder is geopolymer [12–14]. Geopolymers are a new class of inorganic polymer synthesized by activating of an aluminosilicate source with an alkaline hydroxide or silicate solution, which Davidovits first developed in the late 1970s [15–17]. In the production of geopolymers, waste/inert materials, such as fly ash and blast furnace slag are generally used as binding materials and have low CO<sub>2</sub> emission [6, 14, 18–27]. Therefore, geopolymer systems are known as economical and environmentally friendly products [3, 28, 29]. When the geopolymer systems produced using fly ash are examined in the literature, it is observed that fly ash-based geopolymer systems can gain strength between 20 and 120 MPa with 24–72 h of heat curing at temperatures of 50–120 °C for different alkali activator ratios [3, 23, 30–38].

Moreover, it has been reported that fly ash-based geopolymers are generally more durable and have higher compressive strength than that of Portland cement [39, 40]. However, the energy spent for heat curing conditions is a disadvantage for fly ash-based geopolymer systems. For this reason, researchers have conducted studies in fly ash-based geopolymer systems with nano- and/or micro-particles such as SiO<sub>2</sub> [41–44], Al<sub>2</sub>O<sub>3</sub> [16, 45, 46], TiO<sub>2</sub> [47], CaCO<sub>3</sub> [28] etc. to obtain higher strength at lower temperatures by reducing the heat curing time and/or temperature.

On the other hand, as it is known that the structure of geopolymer systems consists of Si–O–Al bonds [48]. While there is a high amount of Si (50–60%) in class F fly ash used, the Al content is in the range (15–25%). Therefore, the Al amount was thought to be less than that of necessary for complete geopolymeric reaction since each Si atom needs one Al atom. Thus, it was postulated that the number of bonds could be increased by adding some Al to the system to employ extra Si atom. The laboratory study was carried out to prove the above postulation.

Therefore, the influence of the addition of micronized aluminum powder in a fly ash-based geopolymer system was investigated in the current study. By providing extra bond between Si and Al atoms, it was aimed to obtain higher strength and better properties in a shorter time and at lower heat curing temperature by substituting the fly ash with aluminum powder. Within the scope of the study, 1, 2, and 3% aluminum powder was replaced with fly ash. Geopolymer mortar samples activated with sodium hydroxide were heat cured at 60, 70, and 80 °C temperatures for 24 and 48 h. The mini flow workability, unit weight, flexural, and compressive strength of the geopolymer mortar samples after heat curing was measured. Furthermore, field emission scanning electron microscopy (FESEM) examinations, X-ray diffraction (XRD), and Energy Dispersive X-Ray (EDX) analyses and reaction kinetics of geopolymerization were carried out to reveal the effects of aluminum powder on the microstructure. As a result of the study, the intended goals were achieved and it was seen that the physical, mechanical, and microstructure properties of fly ash-based geopolymer mortars could be improved with Al substitution.

## Experiment parameters and test details

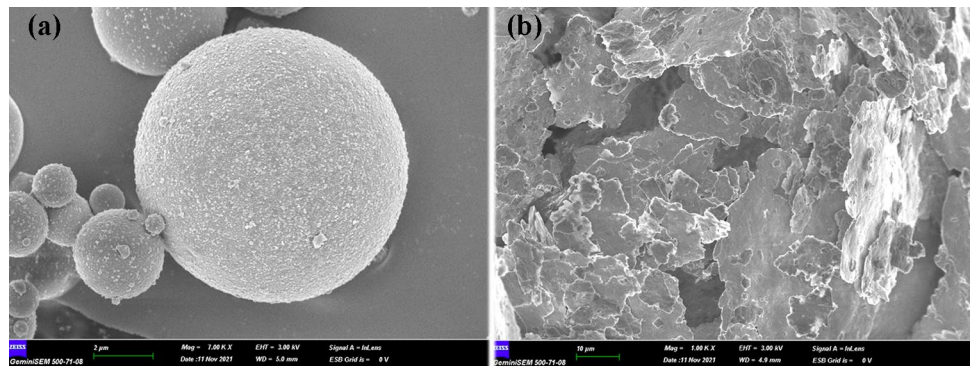
### Materials

Class F fly ash with a specific gravity of 2.34 was used as a binder in the study. Chemical composition, SEM image, and XRD analysis of fly ash are presented in Table 1, Fig. 1, and Fig. 2, respectively. Micronized aluminum powder with a specific gravity of 2.69 was used by replacing with fly ash by weight. The chemical composition, SEM image, and XRD analysis, of the aluminum particles are presented in Table 2, Fig. 1, and Fig. 2, respectively. Solid sodium hydroxide particles were mixed with water and an alkali solution was obtained. Then, the prepared alkali solution was used for the production of geopolymer mortar. Standard Rilem sand

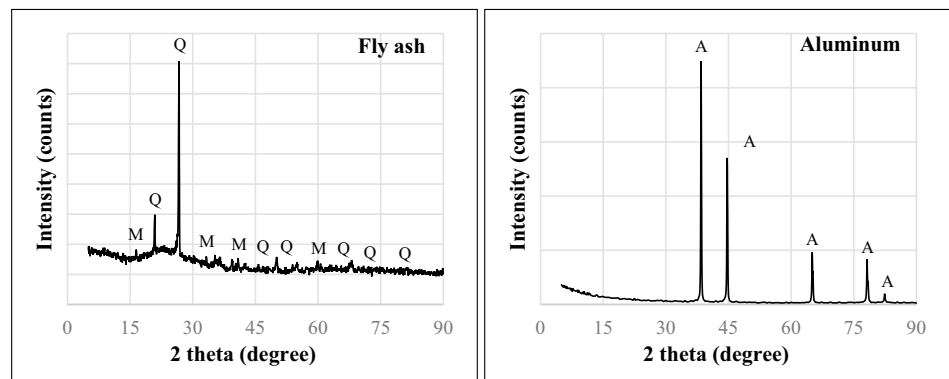
**Table 1** Chemical compositions of fly ash (%)

Oxide	SiO <sub>2</sub>	Al <sub>2</sub> O <sub>3</sub>	Fe <sub>2</sub> O <sub>3</sub>	CaO	SO <sub>3</sub>	Na <sub>2</sub> O	K <sub>2</sub> O	MgO	LOI	Others
Fly Ash	59.11	21.8	8.15	2.91	0.61	1.6	1.8	1.98	0.96	1.08

**Fig. 1** SEM images of fly ash (7000x) and aluminum (1000x) (a fly ash, b aluminum)



**Fig. 2** XRD of fly ash and aluminum (Q Quartz, M Mullite, A Aluminum)



**Table 2** Chemical compositions aluminum powder (%)

Element	Al	Fe	Si	S	Mg	LOI	Others
Aluminum	97.46	0.59	0.40	0.13	0.26	0.95	0.21

**Table 3** Chemical ingredients of NaOH (%)

NaOH	Na <sub>2</sub> CO <sub>3</sub>	Cl	Al	Fe	SO <sub>4</sub>
98.56	1.39	0.02	0.01	0.01	0.01

according to TS EN 196–1 [49] standard was used in the production of the geopolymer mortar. The chemical composition of sodium hydroxide is presented in Table 3.

### Mixture design and detail

In the study, aluminum particles were used by replacing fly ash at the dosages of 0, 1, 2, and 3% by weight. The water/binder ratio was taken as 0.31, and the sand/binder ratio was taken as 3 for the geopolymer mortars. Alkali activator ratios as Na amount were 6, 8, and 10% of fly ash amount in mass basis. The mix proportion of geopolymer mortars are presented in Table 4.

**Table 4** Mix proportion

Specimens	Fly ash	Aluminum	Aggregate	Na <sup>+</sup> ratio	Water	Heat curing condition	Heat curing time
	g	g	g	%	g	°C	Hour
Reference	450.0	0	1350	6-8-10	140	60-70-80	24-48
1% Al	445.5	4.5	1350	6-8-10	140	60-70-80	24-48
2% Al	441.0	9.0	1350	6-8-10	140	60-70-80	24-48
3% Al	436.5	13.5	1350	6-8-10	140	60-70-80	24-48

Fresh geopolymer mortars were prepared by mixing alkali solution with fly ash, aluminum powder, and standard Rilem sand. After the production of geopolymer mortars, the workabilities of the mortars were measured. Then, the fresh geopolymer mortars were placed in molds with which were three cell prisms having the dimensions of  $160 \times 40 \times 40$  (length, depth, and height) mm using vibration table. Geopolymer mortar specimens were heat cured at 60, 70, and 80 °C, for 24 and 48 h in a laboratory oven. The specimens were not sealed during the heating. After heat curing, visual observation showed that there was no significant drying up damage on the samples. Following heat curing, the geopolymer mortar samples were taken out from mold and rested in the laboratory before testing until they reached a temperature of  $23 \pm 2$  °C.

## Testing

### Workability and unit weight

A flow test was carried out according to TS EN 1015–3 [50] specification. According to the relevant standard, the flow diameter value is defined as the average of the flow measurement value in two perpendicular directions of fresh mortar. Following the heat curing process, samples weights were measured and divided by their geometric dimensions and unit weight of geopolymer samples was determined.

### Flexural and compressive strength

A three-point loading test was utilized to measure the flexural strength on three prismatic specimens, according to the TS EN 1015–11 [51] standard for each mixture. After the flexural test, the compressive strength test was carried out using six broken pieces according to the TS EN 1015–11 [51] standard. Flexural and compressive strength results were calculated as an average of the three and six specimens, respectively, for each mixture.

### Microstructural characterization analysis (FESEM and EDX)

After heat curing at 60, 70, and 80 °C, the geopolymer prism specimens were broken and the middle parts were used for the FESEM analyses. The examination was carried out by using Zeiss GeminiSEM electron microscopy. The samples were placed in the device and dried using infrared light for 5 min. The examinations were performed on the fractured surfaces of specimens in a vacuumed environment. Furthermore, EDX analyses were performed during FESEM analysis from designated areas.

### X-ray diffraction (XRD) analysis

After heat curing, the specimens were powdered and sieved under 63 micron. Then, XRD analyzes were carried out on the powder samples obtained using a Bruker AXS D8 Advance device.

### Isothermal conduction calorimetry

The reaction kinetics of the geopolymer mortars with and without Al were analyzed using isothermal calorimetry (TAM Air, TA Instruments Inc.). In the experiment, 0, 1, 2, and 3% Al substituted geopolymer paste samples were produced with containing 10% Na activator. In order to examine the geopolymerization reaction, the paste samples were kept in an isothermal calorimeter device at 70 °C for 48 h.

## Result and discussion

### Workability and unit weight

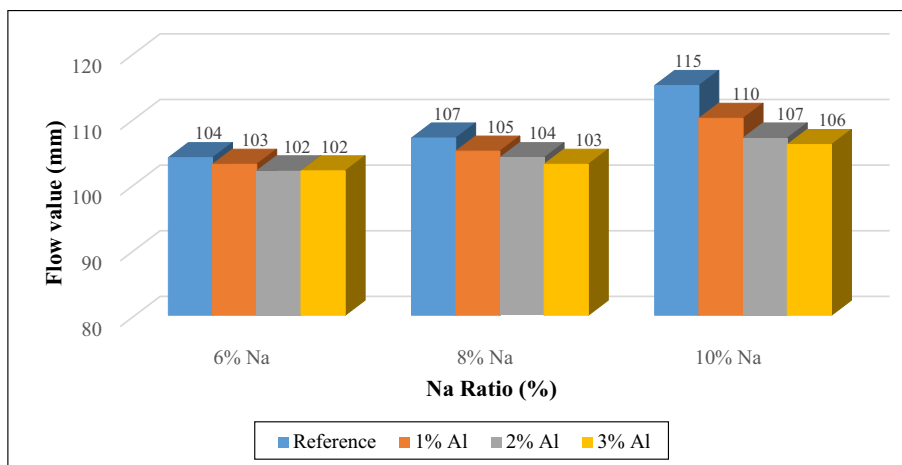
The workability results of the fresh geopolymer mortar samples are presented in Fig. 3. It is a closer observation Fig. 3 showed that the workabilities of fresh reference geopolymer mortars containing 6, 8, and 10% Na<sup>+</sup> were 104, 107, and 115 mm, respectively. Furthermore, the workabilities of fresh geopolymer mortars containing 1% aluminum powder and activated with 6, 8, and 10% Na<sup>+</sup> were 103, 105, and 110 mm, respectively. Moreover, the workabilities of fresh geopolymer mortars containing 2% aluminum powder and activated with 6, 8, and 10% Na<sup>+</sup> were 102, 104 and 107 mm, respectively. Similarly, the workabilities of fresh geopolymer mortars containing 3% aluminum powder and activated with 6, 8, and 10% Na were 102, 103, and 106 mm, respectively.

Comparisons of workabilities made between the reference and aluminum powder containing geopolymer mortar showed that the workability of the geopolymer mortar samples increased with an increased amount of alkali activator. Fly ash particles are surrounded by Na<sup>+</sup> ion. Neighboring fly ash particles surrounded by Na<sup>+</sup> repel each other because the same charged molecule surrounds them. Therefore, agglomeration of fly ash particles is prevented and a workable mixture is obtained. This effect increases with increasing Na<sup>+</sup> content. Similar results have been found from other published articles [28, 52, 53]

However, the addition of aluminum decreased the workability of the fresh geopolymer mortar samples in comparison to the workability of the reference mortar. With the increasing amount of use of aluminum powder, a decrease in the workability values were observed at all sodium activator ratios.



**Fig. 3** Workability of fresh geopolymer mortar

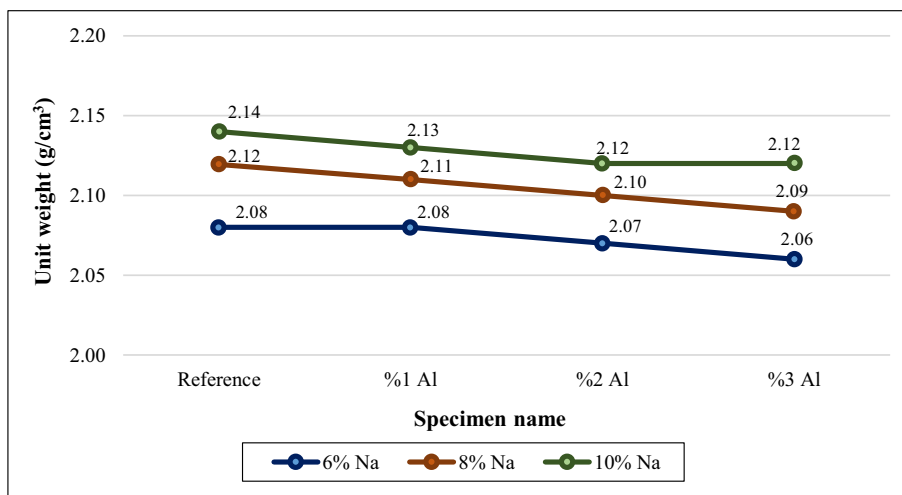


Unit weight values of hardened geopolymer mortar samples are presented in Fig. 4. It is observed that the unit weight values of geopolymer mortar samples increased with the increasing use of activator substitution. For constant amount of aluminum powder, the increase in sodium amount result with higher unit weight. This is explained that more sodium concentration caused better workability thus resulting better compaction of fresh geopolymer mortar. For a constant sodium concentration, the increase in aluminum powder amount in geopolymer mortar gave slightly lower unit weight comparison to control mortar. This is explained by the fact that when fly ash, sodium hydroxide, and aluminum are mixed together, a rapid aluminum reaction starts and hydrogen gas creates a small void inside the mixture. However, since the presence of sand in the mixture increases the volume, this rapid reaction slows down and the amount of void that will be formed becomes insignificant. As a result, the difference between the unit weight of reference geopolymer mortar and geopolymer mortar containing aluminum can be considered negligible.

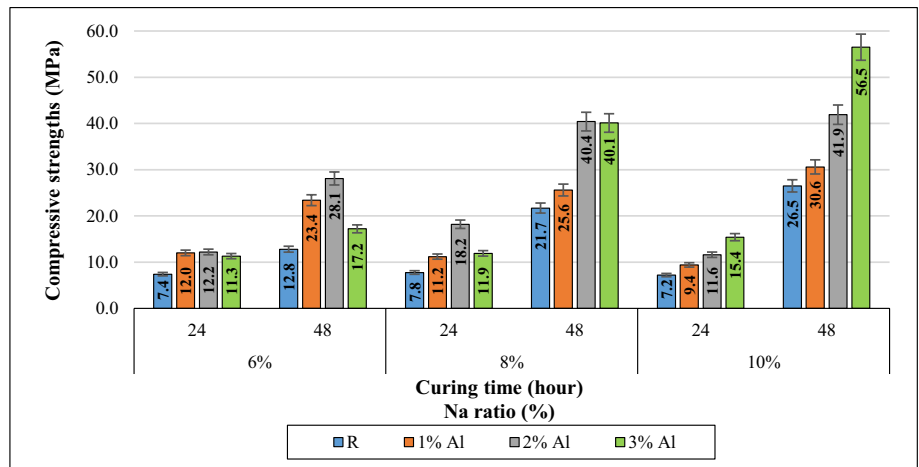
**Compressive and flexural strengths**

The compressive and flexural strengths of 0, 1, 2, and 3% aluminum substituted geopolymer mortar samples, which were heat cured at 60, 70 and 80 °C for 24 and 48 h, are given in Fig. 5, 6, 7 and Fig. 8, respectively. Based on the results, an increase in the compressive and flexural strengths of almost all samples is observed with the increase of heat curing temperature, heat curing time, and amount of activator, separately. In addition, all chemical processes were accelerated with increasing temperature. Increasing the heat curing time and temperature provides a more suitable environment for the formation of geopolymer gels [3, 28]. It is known that all chemical processes take place in time; therefore, increasing curing time increases the amount of geopolymeric gel formation. Moreover, an increase of activator concentration can dissolve more raw materials and increase the amount of geopolymeric gel formed. As a result, compressive and flexural strengths of geopolymer mortars increase [16, 31, 36, 38, 54–57].

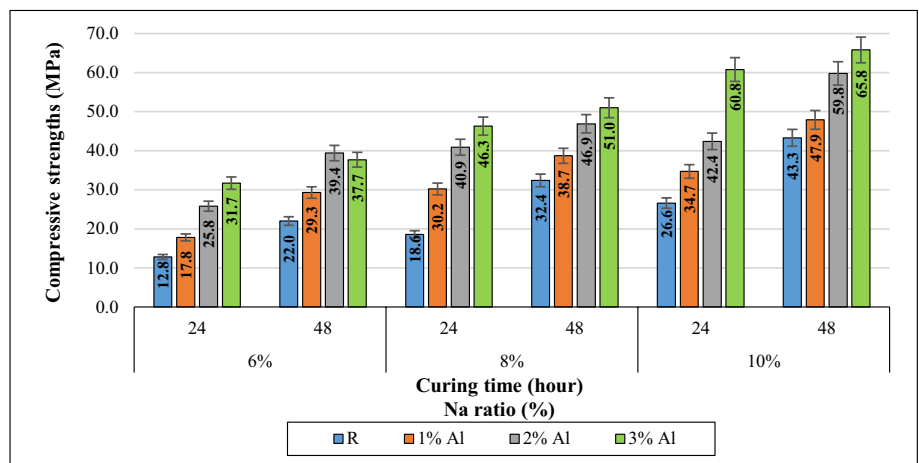
**Fig. 4** Unit weight of mortars



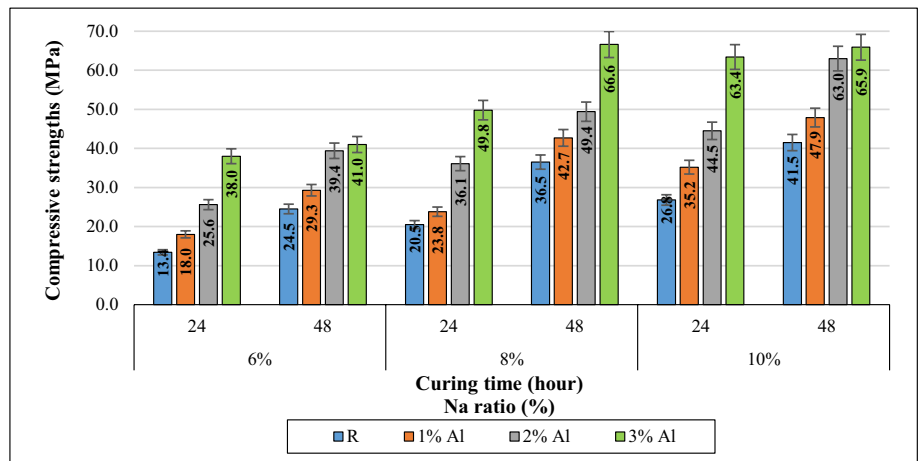
**Fig. 5** Compressive strength of mortars after 60 °C heat curing temperature



**Fig. 6** Compressive strength of mortars after 70 °C heat curing temperature

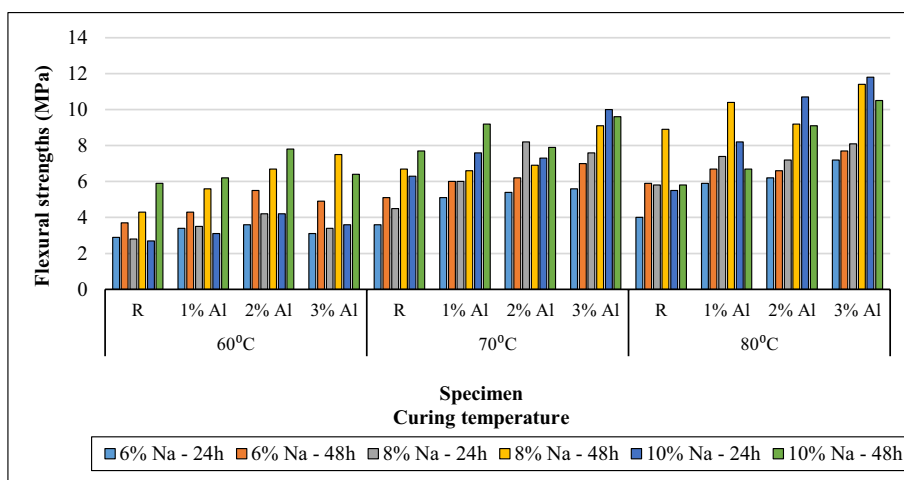


**Fig. 7** Compressive strength of mortars after 80 °C heat curing temperature



However, it was determined that the aluminum particles used by replacing with fly ash in the ratios of 1, 2, and 3% significantly increased the compressive and flexural strengths of the geopolymer mortar samples.

After 60 °C heat curing, the compressive and flexural strengths of the geopolymer mortar samples containing 1, 2, and 3% aluminum were increased by 83, 235, 213, and 30, 56, 74%, respectively, compared to the reference sample.

**Fig. 8** Flexural strength of mortars

Similarly, after 70 °C heat curing, the compressive and flexural strengths of the geopolymer mortar samples containing 1, 2, 3% aluminum were increased by 62, 220, 249, and 42, 82, 69%, respectively, compared to the reference sample. Moreover, in the samples cured at 80 °C, with 1, 2, and 3% aluminum substitution, an increase in compressive and flexural strengths of 34, 91, 284, and 49, 95, 215%, respectively, was observed compared to the reference sample.

According to the strength test results, it was found that an increase of compressive strengths of more than two times can be achieved by substituting aluminum in samples of geopolymer mortar based on fly ash. For example, the compressive strength of 8% Na containing reference geopolymer mixture cured at 70 °C for 24 and 48 h are 18.6 MPa and 32.4 MPa. Extra 24 h heat curing results with 74% increase in compressive strength. On the other hand, the compressive strength of 8% Na containing geopolymer mixture made with 3% aluminum addition cured at 70 °C for 24 h is 46.3 MPa. Addition of 3% aluminum in the geopolymer mixture resulted with 134% increase in compressive strength in comparison to reference geopolymer mixture cured for 24 h. Moreover, the compressive strength of 3% aluminum added geopolymer mortar for 24 h of heat curing was 43% higher than the compressive strength of reference geopolymer mortar heat cured for 48 h. Aluminum addition seems to be more effective in improving strength than that of extra heat curing. With aluminum powder substitution, at the end of the 24 h heat curing period, it was seen that higher strengths could be obtained than the strengths of the reference samples without aluminum addition could reach with 48 h heat curing. Usage of aluminum powder in geopolymer matrix has resulted with significant energy and time-saving.

Although Duxson [58] and Songpiriyakij [59] reported that increasing the Si/Al ratio increases the compressive strength of geopolymer samples, the current study showed that decreasing the Si/Al ratio by addition of extra aluminum powder in the geopolymer mortar mixture remarkably

increased the compressive and flexural strengths of geopolymer mortar more than two times in comparison to the reference mixture.

In addition, an attempt was made to relate compressive and flexural strength of geopolymer mortars, the relationship between flexural and compressive strengths obtained after 24 and 48 h of heat curing is presented in Fig. 9. It was determined that there is a strong relationship between flexural and compressive strengths with  $R^2=0.89$  at the end of the 24 h heat curing period. However,  $R^2=0.69$  for the relationship between flexural and compressive strengths was determined after 48 h of heat curing. This shows that the increase in compressive strength is higher than the flexural strength as the heat curing time increases. In other words, the increase in heat curing time results in a geopolymer mortar with higher compressive strength. However, the structure of geopolymer becomes more brittle with increasing heat curing time and the flexural strengths could not increase as much as compressive strength.

## XRD analysis

XRD analysis results of reference 1, 2 and 3% Al substituted geopolymer powder samples heat cured at 70 °C for 24 h, are presented in Fig. 10. It is observed from the XRD analysis results that quartz ( $\text{SiO}_2$ ), mullite ( $\text{Al}_6\text{Si}_2\text{O}_{13}$ ), and albite ( $\text{NaAlSi}_3\text{O}_8$ ) minerals were detected in the reference geopolymer samples [17, 27]. In addition, sodalite ( $\text{Na}_4\text{Al}_3\text{Si}_3\text{O}_{12}\text{OH}$ ) minerals were detected in geopolymer samples with aluminum substitution. The dominant peak in the structure was the quartz peaks detected around 2 theta 26–27°. In geopolymer samples containing aluminum, quartz and mullite peaks were observed in 2 theta at 21, 26, 27, 36, 39, 43, 50, 60, 68, and 33, 41, respectively, as in the reference sample. In addition, there is a scattered wide peak between 20 and 41 that designates the evolution of N-A-S-H gel with amorphous character [38, 60]. Moreover, it

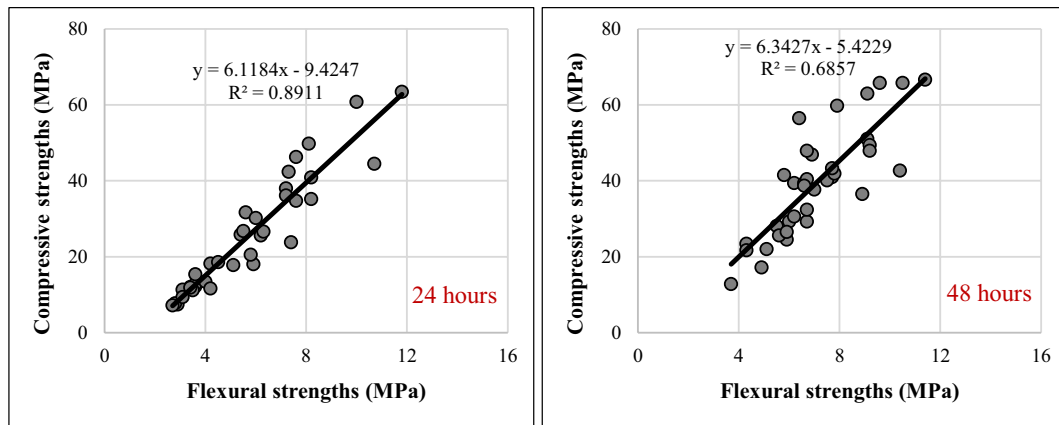
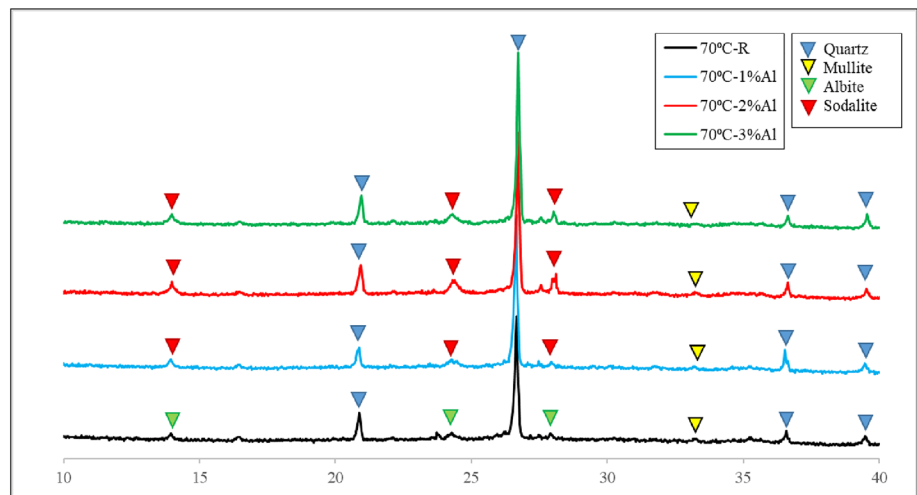


Fig. 9 Relationship between flexural and compressive strengths

Fig. 10 XRD analysis of geopolymer samples (2 Theta (°))



was observed that the Albite peaks detected in 2theta 13.94, 23.73, and 26.65 in the reference sample were transformed into sodalite peaks for geopolymer made with Al substitution. For sodalite peaks, 2theta values were detected 14.02, 24.32, and 26.73 ICDD number of Albite and Sodalite were 00–009–0466 and 01–076–1639, respectively. When the chemical composition of albite ( $\text{NaAlSi}_3\text{O}_8$ ) and sodalite ( $\text{Na}_4\text{Al}_3\text{Si}_3\text{O}_{12}\text{OH}$ ) minerals are examined, it is observed that more aluminum is needed for sodalite formation.

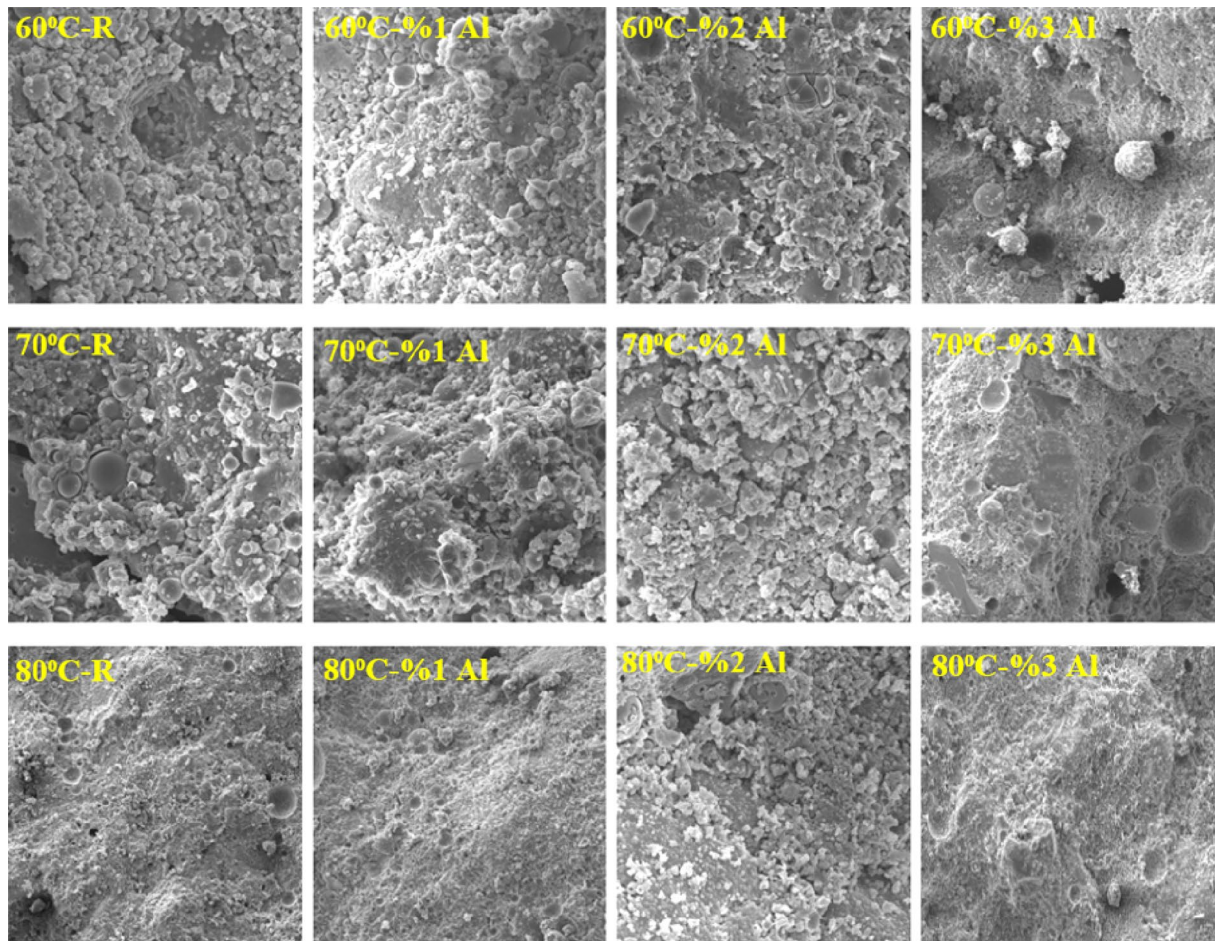
The formation of sodalite minerals instead of albite minerals with Al substitution can be considered as an indication that aluminum powders participate in the geopolymeric reactions. Moreover, it is thought that sodalite formation increases the strength of the geopolymer mortar. Chen et al. [38], studied on ‘The effect on the compressive strength of fly ash-based geopolymer concrete with the generation of hydroxy sodalite’. They stated that at the appropriate amount of sodium hydroxide and aluminum content in the mixture the hydroxy sodalite was formed during heat curing. Thus,

the formation of hydroxide sodalite and N-A-S-H generation leads to high strength. They concluded that the highest compressive strength can be attributed to the generation of N-A-S-H gel and hydroxy sodalites. Moreover, Sathishraj Mani and Bulu Pradhan [61] made a parallel conclusion by stating that the higher compressive strength is attributed to a higher amount of sodium aluminosilicate hydrate (NA-S-H) gel formation through albite phase and predominant zeolite phases such as gismondine and sodalite.

### FESEM-EDX analysis

FESEM images of geopolymer samples are presented in Fig. 11. It is observed from Fig. 11 that the geopolymer samples turn into a compact structure with low porosity due to an increase in the heat curing temperature and aluminum substitution. FESEM pictures of the 60C-R and 80C-R samples are presented in Fig. 12a,b respectively, to examine the influence of heat curing temperature in detail. Many





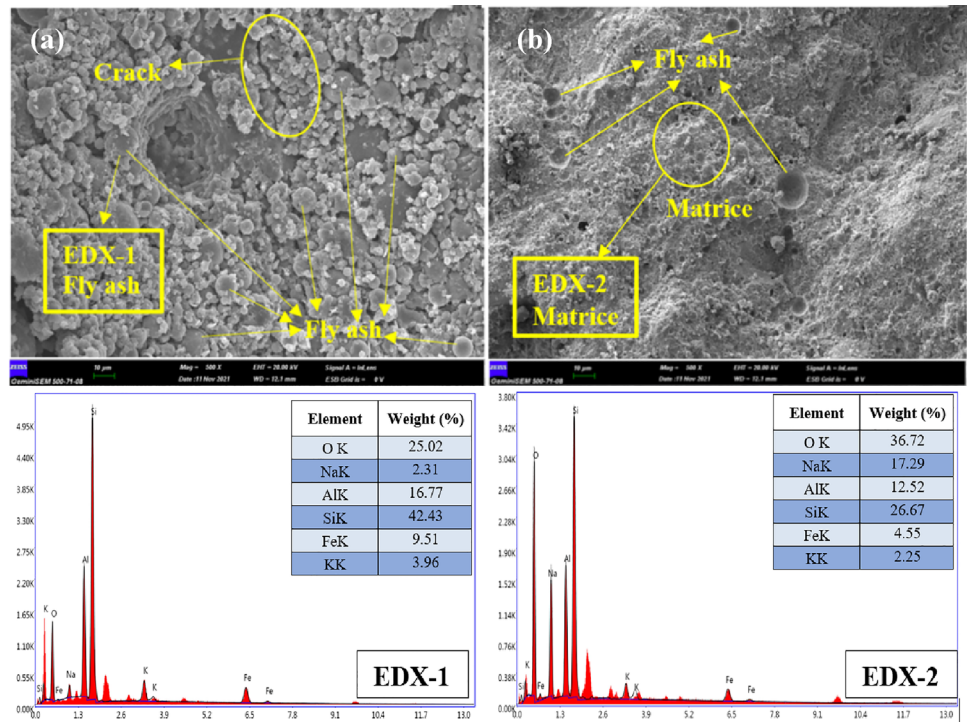
**Fig. 11** FESEM images of geopolymer mortars (500x)

spherical unreacted or partially reacted particles, which are thought to be fly ash particles, are observed in Fig. 12a. The EDX analysis results on these spherical particles are presented in Fig. 12 as EDX-1. According to the results obtained, it was concluded that the chemical composition of these structures is very similar to the chemical composition of fly ash. Therefore, these particles were defined as fly ash. Furthermore, when Fig. 12b (80 °C-R) is examined, it is observed that the amount of unreacted fly ash grains decreases, and the microstructure turns into a more compact structure with an increase in heat curing temperature. The microstructure becomes denser with the increase of heat curing temperature and the higher amount of the reacted fly ash particles are seen as the influencing parameter that contributes to the increase of the strength values due to the effect of heat curing. In the EDX-2 analysis performed in Fig. 12b, the presence of silica, aluminum, and sodium was determined in accordance with the N-A-S-H structure seen in the fly ash-based geopolymer samples [62, 63]. While the amount of silica detected in the matrix structure of the reference sample was (26.67%), the amount of aluminum

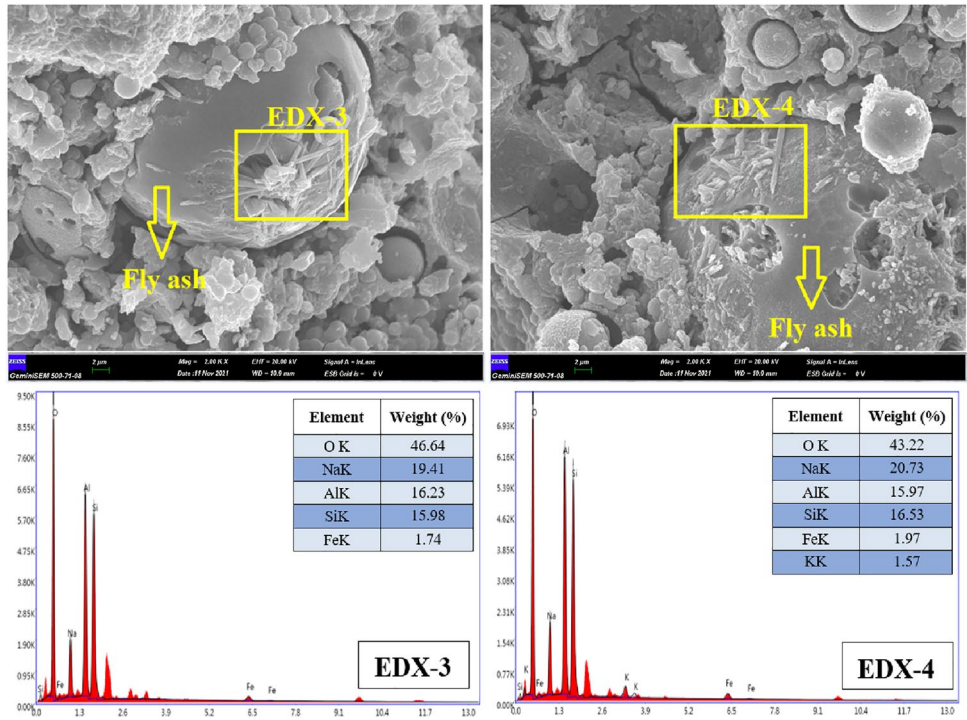
was determined as (12.52%). The rate of Si/Al is found to be compatible with the amount of 59.11% silica and 21.8% aluminum in the chemical composition of fly ash.

On the other hand, to observe the changes in the microstructure with aluminum substitution, the microstructures of the samples with 3%Al substitution were examined under 20,000×magnification. As a result of the examinations, the appearance of the matrix structures in two different regions on the fly ash surface in the microstructure and EDX analyzes are presented in Fig. 13. The needle or rod-like structures observed in Fig. 13 are thought to be sodalite structures (similar sodalite appearances are encountered in the literature [38, 64]), whose presence was also detected in the XRD analysis. The EDX-3 and EDX-4 analyses were carried out to determine the chemical composition of the detected rod-like structures. The obtained results showed that the amounts of aluminum (20.58%) and silica (21.64%) in the chemical composition of the rod-like structure matrices were almost equal. These results are also compatible with the chemical formula of sodalite ( $\text{Na}_4\text{Al}_3\text{Si}_3\text{O}_{12}\text{OH}$ ). The amount of silica

**Fig. 12** EDX and SEM image (500x) of fly ash and geopolymer matrix



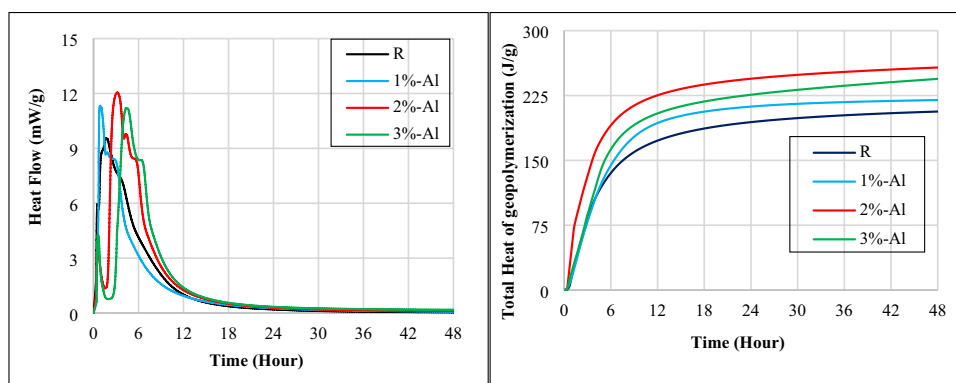
**Fig. 13** EDX and SEM images (2000x) of fly ash and aluminum reaction matrix



detected in the reference geopolymer matrices is more than the amount of aluminum. However, the amount of aluminum in the matrix structure of the samples with aluminum substitutes is almost equal to that of silica is considered as an indication that the aluminum included in

the system participates in the reactions and forms a new structure. Moreover, it is thought that sodalite with rod-like structure, which is thought to be formed in the matrix with aluminum substitution, contributes to the increase in compressive and flexural strengths [38]. Furthermore, it

**Fig. 14** Rate of and cumulative heat of geopolymerization at 70 °C



might be concluded that the strengths increase as a result of the interlocking of the needle-shaped sodalite structures seen in Fig. 13.

### Geopolymerization kinetic

Heat curves of the geopolymerization reaction obtained using reaction kinetic analysis of the control and aluminum powder containing pastes (at 1, 2, and 3% substitution ratio) at 70 °C are given in Fig. 14.

When the heat flow accelerations are examined, it is observed that the heights of rate peaks increased with aluminum substitutions. The peak position also changed in time scale compared to the reference sample (1.8 h), the aluminum containing samples took different times to reach acceleration peaks (1.0 h for 1%-Al, 4.4 h for 2%Al and 3.2 h for 3%Al) In addition, increasing aluminum substitution was elevated the rate of peaks. The maximum peak was obtained from 3% aluminum substitution. It is known that the higher the rate of specimen peak results with a high amount of reaction product, which is associated with sample hardening and strength development [28, 60].

The total heat of geopolymerization curves are presented in Fig. 14. It is observed that the total heat of geopolymerization is 207 J/g for the reference sample. Furthermore, the total heat of geopolymerization for 1%, 2% and 3% aluminum substitution rates are 220 J/g, 243 J/g and 257 J/g. It can be seen that increasing substitution of aluminum, increased the total heat of geopolymerization. It is attributed that the total heat released during the geopolymerization reaction of the samples can give an explanation for both the formation of the reaction product and the strength development [60]. The contributions of aluminum substitution to the rate and the heat of reactions Querydiscussed above are in agreement with compressive and flexural strength results. The significant contribution of aluminum to the strengths of geopolymer mortars was observed for the first 24 h of heat curing. This is found to be in line with the evident effects of aluminum on the rate of heat flows and total heat of geopolymerization curves.

### Conclusions

1. The workability of fresh geopolymer mortars increased with the increase of sodium content. Negligible decreases in workability were observed with aluminum substitution.
2. Increasing the sodium content increased the unit weights of the hardened geopolymer mortar samples. However, increasing the aluminum slightly decreased the unit weights of the hardened geopolymer mortar samples.
3. The increase in sodium content, aluminum substitution, heat curing temperature, and heat curing time increased the compressive, and flexural strength values of the hardened geopolymer mortar samples.
4. More than one hundred percent increases were obtained in the compressive and flexural strength of class F fly ash-based geopolymer mortar samples with aluminum replacement. These increases are attributed to sodalite formation in the microstructure.
5. As a result of XRD analysis, it was observed that albite minerals detected in the reference sample were transformed into sodalite minerals with aluminum substitution.
6. According to FESEM images, it was observed that a rod-like sodalite structure was formed in the microstructure with aluminum substitution. It has been observed that these structures improve the mechanical properties by resulting in better microstructures.
7. It was concluded that the total heat of geopolymerization increased with the substitution of aluminum. This shows that aluminum took part in the geopolymerization reactions, thus resulting in higher strength in comparison to the reference samples.

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## Authors and Affiliations

Uğur Durak<sup>1</sup> 

✉ Uğur Durak  
ugurdurak@erciyes.edu.tr

<sup>1</sup> Civil Engineering Department, Engineering Faculty, Erciyes University, 38039 Kayseri, Turkey