Effect of Elevated Temperature on Alkali Activated Slag and Fly Ash Based Geopolymer Concrete

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Alkali-Activated Slag (AAS) and Fly Ash (FA) based geopolymer concrete a new blended alkali-activated concrete that has been progressively studied over the past years because of its environmental benefits superior engineering properties. Geopolymer has many favorable characteristics in comparison to Ordinary Portland Cement. Many base materials could be utilized to make geopolymer with the convenient concentration of activator solution.

In this study, the experimental program composed of two phases; phase on divided into four groups; Group one deliberated the effect of sodium hydroxide molarity and different curing condition on compressive strength. Group two studied the effect of alkali activated solution (NaOH and Na2SiO3) content on compressive strength and workability.

The effect of sand replacement with slag on compressive strength and workability was explained in group three. Group four studied the effect of slag replacement with several base materials Fly Ash (FA), Ordinary Portland Cement (OPC), pulverized Red Brick (PRB), and Meta Kaolin (MK).

Phase two contains three mixtures from phase one which had the highest compressive strength. For each mixture, the fresh concrete test was air content. In addition the hardened concrete tests were the compressive strength at 3, 7, 28, 90, 180, and 365 days, the flexural strength at 28, 90, and 365 days, and the young’s modulus at 28, 90, and 365 days. Moreover; the three mixtures were exposed to elevated temperature at 100°C, 300°C, and 600°C to study the effect of elevated temperature on compressive and flexural strength.

Keywords: Alkali activated slag (AAS), Geopolymer concrete, Ground granulated blast furnace slag (GGBS), fly ash (FA).

1. INTRODUCTION

Concrete is known as a significant contributor to the emission of greenhouse gases. The cement industry is the second largest producer of the greenhouse gas [1]. Because Portland cement is the main binder in concrete construction, a search for more environmentally friendly materials is undertaken. One possible alternative is the use of alkali-activated binders using industrial by-products [2]. The most common industrial by-products used as binder materials are fly ash (FA) and ground granulated blast furnace slag (GGBFS). GGBFS has been widely used as a cement replacement material due to its latent hydraulic properties, while fly ash has been used as a pozzolanic material to enhance the physical, chemical and mechanical properties of cements and concretes [3].

Recent research has shown that it is possible to use 100% fly ash or slag as the binder in concrete by activating them with highly alkaline solutions just like Sodium or Potassium Hydroxide and Sodium or Potassium Silicate. Mechanical, physical and micro structure of geopolymer concrete is affected by the source material composition, alkali content and liquid/solid ratio [4].

Previous research concluded that liquid sodium silicate gave the highest flexural and compressive strengths of AAS mortars. Another important factor was the concentration of the activator, followed by curing temperature, and the fineness of slag [5]. The choice of the type of activator and dosage is very important in AAS concrete with the higher dosage resulting in higher strength and the higher silicate modulus of the activator resulting in higher strength [6]. The curing temperature should be maintained in between at 40-60°C temperature value to avoid cracks [7]. Flexural strengths are around 10.5% of compressive strength which is slightly higher than that of OPC concrete [8]. Geopolymers are generally believed to perform better than the conventional concretes in fire, due to their ceramic-like properties [9]. However, published research in fly ash-based geopolymer is limited in the area of fire resistance and responses to elevated temperatures. Geopolymer increases in strength after elevated temperature exposure [10]. The fire resistance of geopolymer is likely to be superior to OPC concrete which loses most of its strength after elevated temperature exposure above 800°C. This paper presents an investigation into workability (slump) and compressive strength (at 7 and 28 days) of AAS concrete with different concentration of NaOH (molarity) at 8M, 10M, 12M, and 14M at three different curing conditions. Also, it studied the effect of using different base materials such as OPC and FA on workability and compressive strength of AAS and FA based geopolymer concrete. Moreover, studying the long term properties of AAS concrete and FA based geopolymer concrete such as compressive strength, flexural strength and the effect of elevated temperature on compressive and flexural strength.

II. EXPERIMENTAL PROGRAM

The experimental program was composed of two phases; phase one consisted of fifteen mixtures which were divided into three groups; group one studied the effect of sodium hydroxide molarity and different curing conditions on compressive strength at 7 days and 28 days.
Group two studied the effect of sand replacement by slag on workability (slump) and compressive strength at 7 days and 28 days. Group three studied the effect of slag replacement by FA and OPC on workability (slump) and compressive strength at 7 days and 28 days. Phase two contained three mixes from different group in phase one which had the highest compressive strength. For each mixture, the hardened concrete tests were implemented and included the compressive strength at 3, 7, 28, 90, 180, and 365 days, the flexural strength at 28, 90, and 365 days. Moreover, the three mixes were exposed to elevated temperature at 100°C, 300°C, and 600°C to study the effect of elevated temperature on compressive and flexural strength.

### A. Materials

Ground Granulated Blast Furnace Slag (GGBFS) which is a by-product of the steel industry in Egypt was used instead of cement for producing the alkali activated slag concrete (AAS). Ordinary Portland cement (OPC) CEMI 42.5N was used in some geopolymer mixes with different replacement ratios of (GGBFS). Also, fly ash (FA) was used in some geopolymer mixes with different replacement ratios of (GGBFS). Chemical compositions of cement and geopolymer source materials are illustrated in Table I.

Table I Chemical Composition of Cement and Source Materials of AAS and FA based geopolymer concrete

| Composition by weight (%) | GGBFS | OPC | FA |
|---------------------------|-------|-----|----|
| SiO₂ | 36 | 21.7 | 42.4 |
| Al₂O₃ | 9.48 | 6.3 | 27.4 |
| Fe₂O₃ | 0.8 | 3.4 | 6.54 |
| CaO | 37.7 | 64.5 | 12 |
| MgO | 4.51 | 1.86 | 3.24 |
| SO₂ | 2.29 | 1.77 | 1.71 |
| Na₂O | 0.83 | 0.28 | 0.62 |
| K₂O | 0.47 | 0.54 | 0.68 |

The alkali activator used in preparing AAS and FA based geopolymer concrete mixes consists of two solutions; sodium hydroxide (NaOH) and sodium silicate solution (Na₂SiO₃). The alkali solutions were mixed with (NaOH/Na₂SiO₃) ratio 1:3.

All mixes were produced with potable fresh water, dolomite coarse aggregate of 10mm maximum aggregate size and the fine aggregate was siliceous sand with fineness modulus of 2.9.

### B. Concrete mixes

A mix ratio 1:1:2 (1 slag: 1 fine aggregate: 2 coarse aggregate) had been used to study the workability and the compressive strength at 7 and 28 days. Group one mixtures are shown in Table II, Group two mixtures are shown in Table III and Group three mixtures are shown in Table IV.

### C. Mixing procedures

The concrete mixing procedures used in the study are as follows. Firstly, the base material and fine aggregate were mixed dry until the mixture is thoroughly blended and is uniform in colour. Next, some of activator solution was added to the mix gradually until obtaining a homogenous mixture. After that, the coarse aggregate was added. Then, the remaining activator solution was added to the previous mix until the coarse aggregate was uniformly distributed throughout the batch. Finally, mixes were cast in the molds.

### D. Specimens curing

For steam curing after 24 hours of casting, all specimens were de-moulded and cured in steam for 3 days at 60°C. For oven curing the geopolymer concrete specimens were de-moulded and then heated in oven for 24 hours at a temperature of 60°C and for air curing, the specimens were left in air until the day of testing.

### E. Concrete tests

Slump test, compressive strength, and flexural strength were conducted on specimens as shown in Fig. 1. Slump tests were conducted according to standard specification ASTM C143/C143M-03[11]. Compressive strength of the different mixes was determined experimentally on cubes (100x100x100) mm. The compressive strength tests were conducted according to standard specification BS EN 12390-3:2009[12].

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**Table II:** Mixes studying the effect of sodium hydroxide molarity and different curing conditions.

| Mix Name | Slag (kg/m³) | Fine Agg. (kg/m³) | Coarse Agg. (kg/m³) | NaOH (kg/m³) | Na₂SiO₃ (kg/m³) | Molarity |
|----------|--------------|-----------------|-----------------|-------------|----------------|-----------|
| M1-8     | 556          | 556             | 1112            | 56          | 167            | 8         |
| M2-10    | 556          | 556             | 1112            | 56          | 167            | 10        |
| M3-12    | 556          | 556             | 1112            | 56          | 167            | 12        |
| M4-14    | 556          | 556             | 1112            | 56          | 167            | 14        |

**Table III:** Mixes studying the effect of sand replacement with slag.

| Mix Name | Slag (kg/m³) | Fine Agg. (kg/m³) | Coarse Agg. (kg/m³) | NaOH (kg/m³) | Na₂SiO₃ (kg/m³) |
|----------|--------------|-----------------|-----------------|-------------|----------------|
| S-0%     | 556          | 556             | 1112            | 56          | 167            |
| S-20%    | 667          | 445             | 1112            | 56          | 167            |
| S-50%    | 834          | 276             | 1112            | 56          | 167            |

**Table IV:** Mixes studying the effect of slag replacement with OPC and FA.

| Mix Name | OPC (kg/m³) | Slag (kg/m³) | Fine Agg. (kg/m³) | Coarse Agg. (kg/m³) | Na₂SiO₃ (kg/m³) | NaOH (kg/m³) |
|----------|-------------|--------------|-----------------|-----------------|----------------|-------------|
| OPC-5%   | 28          | 528          | 1112            | 56              | 167            | 56          |
| OPC-10%  | 56          | 500          | 1112            | 56              | 167            | 56          |
| OPC-15%  | 83          | 473          | 1112            | 56              | 167            | 56          |
| OPC-20%  | 111         | 445          | 1112            | 56              | 167            | 56          |
| FA-10%   | ....         | 500          | 1112            | 56              | 167            | 56          |
| FA-20%   | ....         | 445          | 1112            | 56              | 167            | 56          |
| FA-50%   | ....         | 278          | 1112            | 56              | 167            | 56          |
| FA-100%  | ....         | 556          | 1112            | 56              | 167            | 56          |
By using beams (100x100x500)mm flexural strength tests were conducted according to standard specification ASTM C78/C78M-18[13]. For the effect of elevated temperature, three mixtures were casted into cubes (100 × 100 × 100) mm and beams (100 × 100 × 500) mm. The unloaded cubes were placed in oven for 1 hr., and 2 hr., after reaching the required temperature at 100°C, 300°C, 600°C. Half of the samples were left to be gradually cooled, where the other half were suddenly cooled by emerging in water. All samples were tested in compression and flexure strength.

![Figure 1: Concrete Mechanical Tests](image)

| Group No. | Mix name | Slump (mm) | Compressive strength (MPa) 7 days | Compressive strength (MPa) 28 days |
|-----------|----------|------------|-----------------------------------|-----------------------------------|
|           |          | Air        | Oven | Steam | Air | Oven | Steam |
| Group 1   | M1-8     | ---        | 9    | 9.2   | 15  | 10   | 11    | 17    |
|           | M2-10    | ---        | 8    | 9     | 28  | 8.5  | 10    | 30    |
|           | M3-12    | ---        | 10   | 11    | 32  | 11   | 11.8  | 34    |
|           | M4-14    | ---        | 10.5 | 16.5  | 34  | 11.2 | 17    | 35    |
| Group 2   | S-0%     | 50         | ---- | ----  | 32  | ---- | ----  | 34    |
|           | S-20%    | 60         | ---- | ----  | 35  | ---- | ----  | 42    |
|           | S-50%    | 80         | ---- | ----  | 31  | ---- | ----  | 33.8  |
|           | FA-10%   | 70         | ---- | ----  | 34  | ---- | ----  | 37    |
|           | FA-20%   | 80         | ---- | ----  | 38.9| ---- | ----  | 42    |
|           | FA-50%   | 130        | ---- | ----  | 51  | ---- | ----  | 54.4  |
|           | FA-100%  | 210        | ---- | ----  | 26.5| ---- | ----  | 29    |
|           | OPC-5%   | 55         | ---- | ----  | 34  | ---- | ----  | 38    |
|           | OPC-10%  | 60         | ---- | ----  | 37  | ---- | ----  | 42    |
|           | OPC-15%  | 70         | ---- | ----  | 39  | ---- | ----  | 45    |
|           | OPC-20%  | 80         | ---- | ----  | 42  | ---- | ----  | 50    |

- **Effect of sodium hydroxide molarity and different curing conditions on compressive strength.**

The effect of molarity of sodium hydroxide (NaOH) on compressive strength of AAS concrete is shown in Table-V. Fig. 2&3 show that increasing the molarity from 8M to 10M increased the compressive strength by 76.5%. Also increasing the molarity from 10M to 12M increased the compressive strength by 13%. Increasing the molarity from 12M to 14M increased the compressive strength by 3%. It can be noted that increasing the molarity of NaOH from 8 M to 14 M gradually increases the compressive strength of AAS concrete. M3-12 was selected as the basic mixture to study the effect of other parameters for two reasons; firstly increasing the molarity from 12M to 14M increased the compressive strength by 3%, which can be regarded as a small increasing value, and the cost of alkaline liquid is economical when concentration of NaOH solution 12M compared with 14M. Secondly, the required compressive strength was achieved [14, 15]. Fig. 2&3 show that the compressive strength increased by a value ranging from 55% to 200% at steam curing compared with oven curing and the compressive strength increased by a value ranging from 7% to 50% at oven curing compared with air curing.
The results show that steam curing at 60°C for 3 days lead to the highest compressive strength for all samples, then oven curing at 60°C for 24 hours and the least values are related to air curing. The results are consistent with previous research [5]. From the previous results, it can be noted that the air curing values were near to the oven curing value. This may be due to insufficient duration of the 60°C of one day to complete the polymerization reaction. Swanepoel et al. [16] stated that combination of 60°C temperature and 48 hours curing time gives the optimum strength for the binder. The steam curing substantially assists the chemical reaction that occurs in the geopolymer concrete. Both curing time and curing temperature influence the compressive strength of AAS concrete. So the steam curing is selected as the basic curing to study the effect of other parameters.

Fig. 2: Effect of sodium hydroxide molarity and different curing condition on compressive strength of mixes at 7 days.

Fig. 3: Effect of sodium hydroxide molarity and different curing condition on compressive strength of mixes at 28 days.

- **Effect of sand replacement by slag on compressive strength and workability.**

  The mixtures S-0%, S-20% and S-50% have slump 50 mm, 60 mm and 80 mm respectively. It can be noted that 20% sand replacement increased the slump by 20% compared to S-0% and 50% sand replacement increased the slump by 60% compared to S-0% as shown in Fig. 4. Thus, increasing sand replacement by slag increased the slump.

Fig. 4: Effect of sand replacement by slag on slump

Fig. 5: Effect of sand replacement by slag on compressive strength

- **Effect of slag replacement by FA and OPC on compressive strength and workability.**

  Fig. 6 shows that the slump of the mixture by 10% fly ash increased by 40% compared to the mixture with 0% fly ash. The slump of mixture with 20% fly ash increased by 60% compared to the mixture with 0% fly ash. The slump of the mixture with 50% fly ash increased by 160% compared to mixture with 0% fly ash and the slump of the mixture with 100% fly ash increased by 320% compared to the mixture with 0% fly ash. The increase of slump value with increase of fly ash content in the mixture is consistent with previous studies [17, 18] This can be attributed to the spherical shape of fly ash particles compared to the angular shape of slag particles[17].
Figure 7 shows that the compressive strength of mixture with 100% fly ash decreased by 15% compared to the mixture with 0% fly ash. The compressive strength of the mixture with 50% fly ash increased by 60% compared to the mixture with 0% fly ash. The compressive strength of the mixture with 20% fly ash increased by 23.5% compared to the mixture with 0% fly ash. The compressive strength of the mixture with 10% fly ash increased by 9% compared to the mixture with 0% fly ash. The previous research showed that the compressive strength increased with increasing the amount of fly ash up to 50%, then reduced again at 100% fly ash. The increase of the compressive strength may be attributed to the formation of C-A-S-H gels, which would reduce the porosity and condense the microstructure of AAS matrix [19, 20]. The reduction of compressive strength may be attributed to geopolymer containing higher fly ash content, thus highly porous structure resulting in lower compressive strength [21].

Figure 8 shows that the slump of mixture OPC-5% increased by 10% compared to the mixture S-0%, while the slump of mixture OPC-10% increased by 20% compared to the mixture S-0%, slump of mixture OPC-15% increased by 40% compared to the mixture S-0% and slump of mixture OPC-20% increased by 60% compared to the mixture S-0%. Thus, the slump value increased with the increase of OPC content in the mixture.

Figure 9: shows that the replacement of slag by 5%, 10%, 15% and 20% OPC increased the compressive strength by 6%, 16%, 22%, and 31% after 7 days and increase by 12%, 23.5%, 32% and 47% respectively after 28days compared with the control mix without slag replacement. The addition of OPC modifies the products and the pore structure in hardened cementitious materials. Increasing the OPC content results in denser structure by reducing the pore size in concrete, due to GGBS reacts with calcium hydroxide to form extra C-S-H gel in the concrete matrix[22].

Table- VI: Concrete mixtures used in phase one.

| Mixture | Slag kg/m³ | Fly ash kg/m³ | OPC kg/m³ | Fine Agg. kg/m³ | Coarse Agg. kg/m³ | NaOH kg/m³ | Na₂SiO₃ kg/m³ |
|---------|------------|---------------|-----------|----------------|-------------------|------------|--------------|
| S-20%   | 667        | ----          | ----      | 445            | 1112              | 56         | 167          |
| FA-50%  | 278        | 278           | ----      | 556            | 1112              | 56         | 167          |
| OPC-20% | 445        | ----          | 111       | 556            | 1112              | 56         | 167          |

- Development of compressive strength.

Table-VII illustrates the compressive strength development of the tested three mixes.
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Table- VII: Compressive strength development

| Mixture | 3days | 7days | 28days | 90days | 180days | 365days |
|---------|-------|-------|--------|--------|---------|---------|
| S-20%   | 22    | 35    | 42     | 45     | 46      | 46      |
| FA-50%  | 40    | 51    | 54.4   | 61     | 61.5    | 63      |
| OPC-20% | 29    | 42    | 50     | 54     | 55      | 56      |

As shown in Fig. 10 for the mixture S-20%, the compressive strength value at 3 days was 63% of the compressive strength at 7 days. The compressive strength at 7 days was 83% of the compressive strength at 28 days. The compressive strength at 90 days increased by 7% more than the compressive strength at 28 days. The increase in compressive strength after 90 days was small.

For mixture FA-50%, the compressive strength value at 3 days was 78% of compressive strength at 7 days. The compressive strength value at 7 days was 94% of compressive strength at 28 days.

The compressive strength at 90 days increased by 12% more than the compressive strength at 28 days. The increase in compressive strength after 90 days was small.

For mixture OPC-20%, the compressive strength value at 3 days was 69% of compressive strength at 7 days. The compressive strength value at 7 days was 84% of compressive strength at 28 days.

The compressive strength at 90 days increased by 8% more than the compressive strength at 28 days. The increase in compressive strength after 90 days was small.

The previous results showed that the compressive strength of tested concretes increased steadily at 28 days but after that, the increasing rate of compressive strength became much slower. The chemical reaction of the geopolymer is a rapid polymerization process about 70% of its strength is gained within the first 3-4 hours of curing [23, 24].

Table-VIII: Flexural strength development

| Mixture | 28days | 90days | 365days |
|---------|--------|--------|---------|
| S-20%   | 3.9    | 4.1    | 3.7     |
| FA-50%  | 6.9    | 7.5    | 8       |
| OPC-20% | 4.2    | 4.8    | 5       |

For mixture OPC-20%, the flexural strength increased at 90 days by 14% more than the flexural strength at 28 days and increased at 365 days by 4% more than the flexural strength at 90 days. It can be seen that the mixture FA-50% has the highest value of flexural strength compared to S-20% and OPC-20%.

Table-X shows the effect of elevated temperature on compressive strength for S-20%, FA-50%, and OPC-20%. As shown in Fig. 12, the mixture S-20% at air cooled regime heating to 100°C for one and two hours led to compressive strength increase about 15% and 10%, respectively.

On the other hand, heating to 300°C for one and two hours led to compressive strength reduction of about 2% and 9.5%, respectively. Moreover, heating to 600°C for one and two hours led to compressive strength drop of about 7% and 26% respectively. At water cooled regime heating to 100°C for one and two hours led to compressive strength increase of about 7% and 2%, respectively.

On the other hand heating to 300°C for one and two hours led to compressive strength loss of about 7% and 16%, respectively. Moreover heating to 600°C for one and two hours led to compressive strength reduction of about 26% and 28.5%, respectively. The reduction in compressive strength at 300 and 600°C may be due to the thermal discordance between aggregates and geopolymer matrix which led to formation of internal stresses then cracks which deteriorated the concrete. [26, 27].

Fig.10: Compressive strength developments

- Development of Flexural strength

Table-VIII illustrates the flexural strength development of the three tested mixes. As shown in Fig.11 for mixture S-20%, the flexural strength increased at 90days by 5% more than the flexural strength at 28 days and decreased at 365 days by 10% less than the flexural strength at 90 days. The higher amount of slag leads to micro-cracks inside the concrete which may have reduced the flexural strength of concrete at 365days [25].

For mixture FA-50%, the flexural strength increased at 90days by 8.7% more than the flexural strength at 28days and increased at 365days by the same ratio more than the flexural strength at 90 days.
As illustrated in Fig. 14, the mixture OPC-20% at air cooled regime heating to $100^\circ C$ for one and two hours led to compressive strength increase of about 10% and 4%, respectively. On the other hand, heating to $300^\circ C$ for one and two hours led to compressive strength reduction of about 18% and 20%, respectively. Moreover, heating to $600^\circ C$ for one and two hours led to compressive strength loss of about 28% and 36%, respectively. At water cooled regime heating to $100^\circ C$ for one and two hours led to compressive strength increase of about 6% and 2%, respectively. On the other hand heating to $300^\circ C$ for one and two hours led to compressive strength loss of about 22% and 25%, respectively. Moreover, heating to $600^\circ C$ for one and two hours led to compressive strength increase of about 32% and 40%, respectively. The reduction of compressive strength on concrete may be a result of the release of free water and physically bound water [28, 29]. Siddique and Kaur [30] showed that the residual compressive strength decreased at the blended GGBFS and OPC concrete when the increase in temperature was up to $350^\circ C$.

For all three mixes the increase of compressive strength at $100^\circ C$ may be attributed to complete geopolymerization in the matrix. Moreover, the reduction at water cooled regime compared to air cooled regime may be attributed to the sudden cooling by water form micro cracks resulting from the non-uniform thermal field, in addition coarse aggregates would absorb water and expand [31].
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- Effect of elevated temperature on flexural strength

Table XI: effect of elevated temperature on flexural strength

| Mixture  | flexural strength (MPa) | Air cooled | Water cooled |
|----------|-------------------------|------------|-------------|
|          | 100°C 2hr | 100°C 2hr | 300°C 2hr | 300°C 2hr | 600°C 2hr | 600°C 2hr |
|          | 1hr     | 2hr      | 1hr     | 2hr      | 1hr     | 2hr     |
| S-20%    | 3.9     | 4.5      | 4.2     | 2.8      | 2.4     | 2.4     |
| FA-50%   | 6.9     | 8.1      | 6.7     | 6.7      | 5.9     | 5.9     |
| OPC-20%  | 4.2     | 4.7      | 4.3     | 4.1      | 3.42    | 3.42    |

As illustrated in Fig. 15, the mixture S-20% at air cooled regime heating to 100°C for one and two hours led to flexure strength increase of about 17% and 2% respectively. On the other hand, heating to 300°C for one and two hours led to flexure strength reduction of about 3% and 14.5%, respectively. Moreover, heating to 600°C for one hour led to flexure strength drop of about 55% and for two hours led to flexure strength reduction of about 58%, respectively. At water cooled regime heating to 100°C for one and two hour led to flexure strength increase of about 10% and 2%, respectively. On the other hand, heating to 300°C for one and two hours led to flexure strength loss of about 5% and 21%, respectively. Moreover, heating to 600°C for one and two hours led to flexure strength decrease of about 29% and 31%, respectively.

As illustrated in Fig. 16, the mixture FA-50% at air cooled regime heating to 100°C for one hour led to flexure strength increase of about 17% and for two hours led to flexure strength loss of about 3%. On the other hand, heating to 300°C for one and two hours led to flexure strength reduction of about 3% and 14.5%, respectively. Moreover, heating to 600°C for one hour led to flexure strength drop of about 55% and for two hours led to flexure strength reduction of about 58%. At water cooled regime heating to 100°C for one and two hours led to flexure strength reduction of about 4% and 23%, respectively. On the other hand, heating to 300°C for one and two hours led to flexure strength loss of about 38% and 39%, respectively. Moreover, heating to 600°C for one and two hours led to flexure strength decrease of about 60% and 62%, respectively.

As illustrated in Fig. 17, the mixture OPC-20% at air cooled regime heating to 100°C for one and two hours led to flexure strength increase of about 12% and 2% respectively. On the other hand, heating to 300°C for one and two hours led to flexure strength reduction of about 2% and 19%, respectively. Moreover, heating to 600°C for one and two hours led to flexure strength decrease of about 14% and 24%, respectively. At water cooled regime heating to 100°C for one and two hour led to flexure strength increase of about 10% and 2%, respectively. On the other hand, heating to 300°C for one and two hours led to flexure strength loss of about 5% and 21%, respectively. Moreover, heating to 600°C for one and two hours led to flexure strength decrease of about 29% and 31%, respectively.

IV. CONCLUSIONS

From the analysis and discussion of test results obtained from this research, the following conclusions can be drawn:

1. Higher concentration [molar] of sodium hydroxide solution increased the compressive strength of AAS concrete. Mixture with 12 molar was selected as the basic mixture to study the effect of other parameters.
2. Exposure of samples to steam curing at 60°C for 3 days leads to the highest compressive strength for all samples compared with air curing and oven curing.
3. Increasing the sand replacement ratio by slag to 20% by weight of sand increased the compressive strength by 30% more than the mixture without sand replacement.
4. Increasing the sand replacement ratio by slag to 50% by weight of sand decreased the compressive strength by 2%, less than the mixture without sand replacement.
5. Increasing the sand replacement by slag increased the workability.
6. Increasing the slag replacement ratio by fly ash up to 50% of total base material increased the compressive strength up to 54.4MPa, while slag replacement by 100% fly ash decreased the compressive strength by 15% less than the mixture with slag only.
7. Increasing the slag replacement by fly ash increased the workability.
8. Increasing the slag replacement ratio by OPC up to 20% increased the compressive strength up to 50MPa.
9. Increasing the slag replacement by OPC increased the workability of AAS concrete mixtures.
10. The development rate of compressive strength after 28 days was small compared to its value beyond 28 days.
11. Exposure of samples to elevated temperature at 100°C show a certain amount of increase in compressive and flexural strength for all tested mixes regardless of the cooling regime.
12. Exposure of samples to elevated temperature at 300°C decreased the compressive and flexural strength of mix S-20% and OPC-20% while increased the compressive strength of FA-50% only regardless of the cooling regime.
13. Exposure of samples to elevated temperature at 600°C decreased the compressive and flexural strength of all mixes.
14. The performance of the samples under gradual cooling regime (Air cooling) is better than the sudden cooling regime (water cooling).
15. The performance of samples subject to 1 hour of heating in terms of compressive and flexure strength is better than the samples subject to 2 hour heating.
16. Mixture of slag replacement by 50% fly ash (FA-50%) had the best performance in elevated temperature compared with mix S-20% and OPC-20%.

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