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Microstructure and Mechanical Properties of FA/GGBS-based Geopolymer



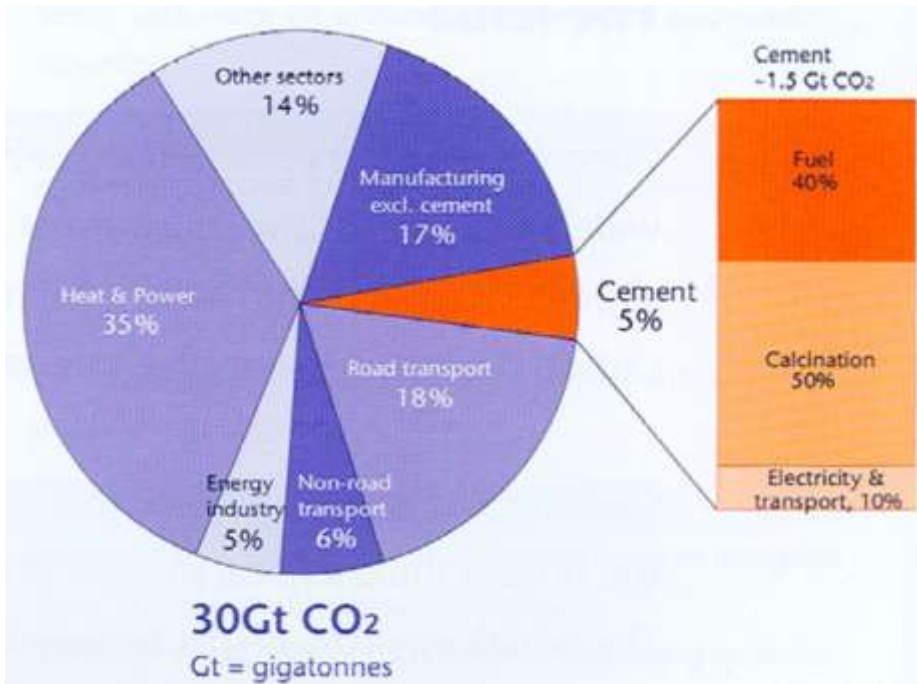
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Cement industry emits high amount of carbon dioxide (CO₂) that accounts for about 5% of global CO₂ emission.



Environmental degradation

can be reduced by



- Partially replace the cement in concrete (E.g. High amount of fly ash used in concrete).
- Develop alternative material (E.g. **Geopolymer concrete**)

Geopolymer cements can be obtained through inexpensive and eco-friendly synthetic procedures.

Fly ash-based geopolymer can cut down about 60% CO₂ emission, compared to OPC (Li *et al.*, 2004)



However...

It has very long setting time and low compressive strength.



Fly ash + ground granulated blast-furnace slag (GGBS)

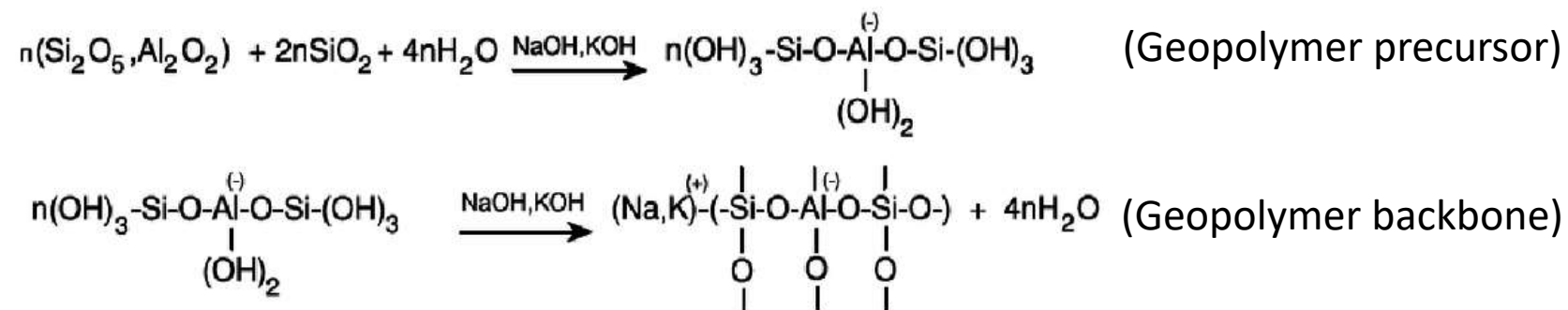
Objective

To investigate effect of GGBS on the microstructure and mechanical properties of fly ash-based geopolymer.

Theory and Application



The term 'geopolymer' was coined in the 1970s by the French scientist and engineer Prof. Joseph Davidovits, and applied to a class of solid materials synthesized by the reaction of an aluminosilicate powder with an alkaline solution.



Brisbane West Wellcamp Airport, Queensland, Australia is the world's largest geopolymer concrete project.

Materials

1. Precursors:

- Fly ash (fineness : 3.550 cm/g^2 , density : 2.24 g/cm^3)
- GGBS (fineness : 4.170 cm/g^2 , density : 2.91 g/cm^3)

2. Alkaline liquids

- Combination of sodium hydroxide (NaOH) 14 M and sodium silicate (Na_2SiO_3).

Table 1. Chemical compositions of raw materials by EDS analysis.

Composition	Fly ash (mass%)	GGBS (mass%)
SiO_2	68.44	34.45
Al_2O_3	20.65	14.06
Fe_2O_3	4.18	0.27
CaO	2.25	43.78
K_2O	1.53	0.23
TiO_2	1.19	0.56
MgO	0.58	5.84
Na_2O	-	0.24
LOI	2.9	0.05

Experimental Method (2)



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Fly Ash



GGBS



Alkaline solution (14 M
 $\text{NaOH} + \text{Na}_2\text{SiO}_3$)

GGBS/FA : 0.15, 0.30, 0.45, 0.60 (By mass)
Solution/Binder : 0.45 (By mass)
 $\text{Na}_2\text{SiO}_3/\text{NaOH}$: 2 (By mass)

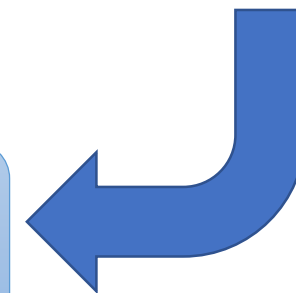
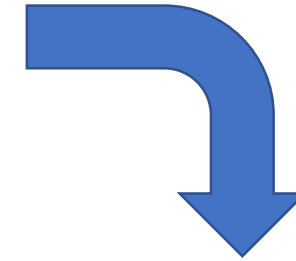


Fresh Geopolymer paste

Characterization and
measurement

Demould and kept in
controlled room
(20°C and 60% RH)
for 7, 14, and 28 days

Casted in cylindrical
plastic mold
(50mmx100mm) and cured
at 70°C for 24 hours





- **Compressive strength test**
- **Scanning Electron Microscopy (SEM)**
- **X-Ray Diffraction (XRD)**
- **Thermo-gravimetric analysis**

Table 2. Compressive strength of geopolymer paste specimens.

Specimen	Compressive strength (MPa)		
	7 days	14 days	28 days
FAS0	24.21 (1.17)	24.7 (1.23)	24.36 (1.43)
FAS15	43.51 (5.74)	44.06 (1.89)	44.38 (3.01)
FAS30	62.92 (5.43)	61.15 (0.83)	56.46 (1.79)
FAS45	77.19 (1.69)	80.39 (6.78)	84.17 (1.79)
FAS60	105.54 (5.55)	100.84 (3.10)	93.36 (7.54)

* () = Standard deviation

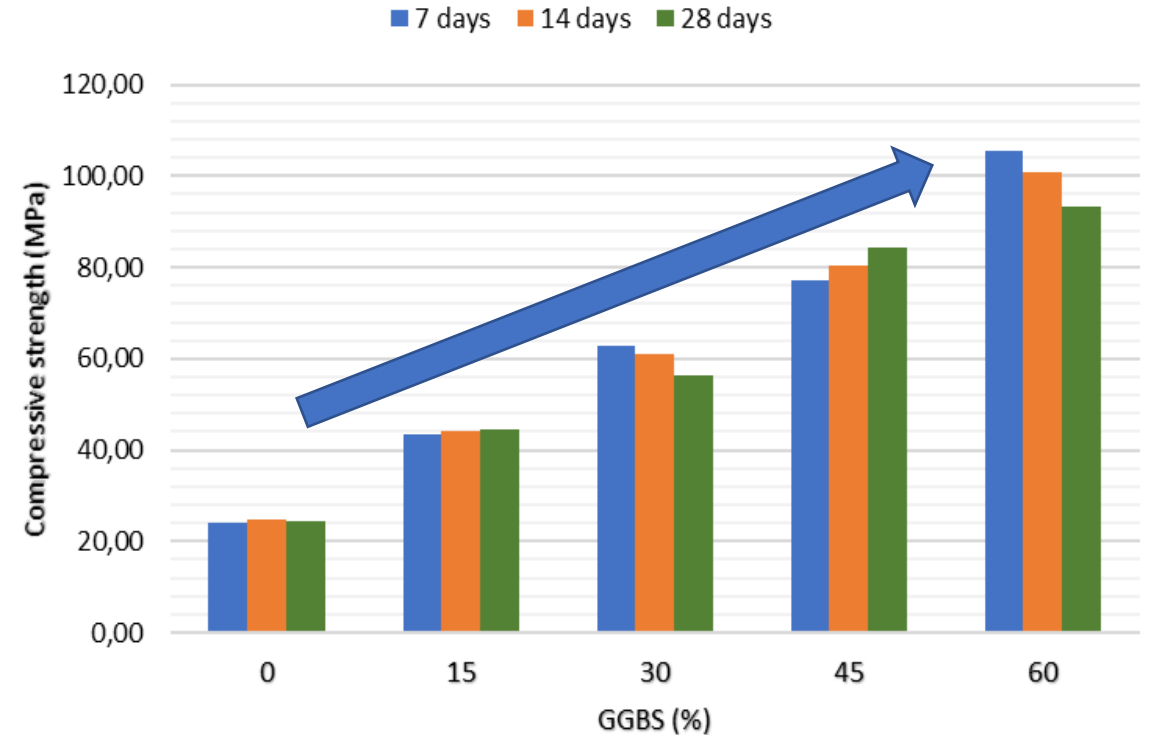
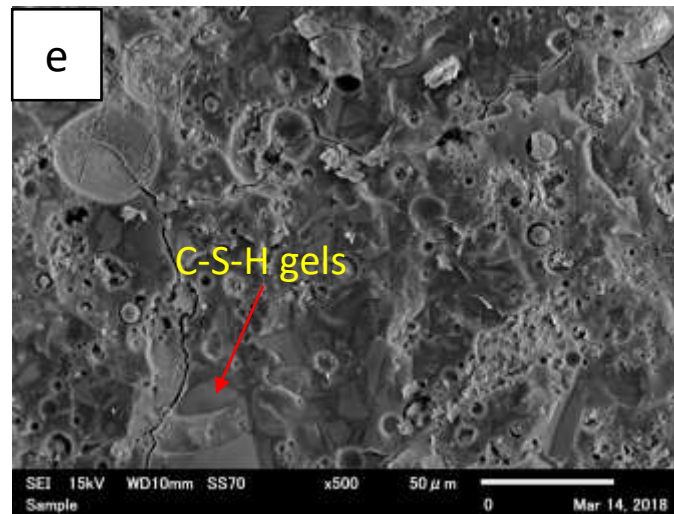
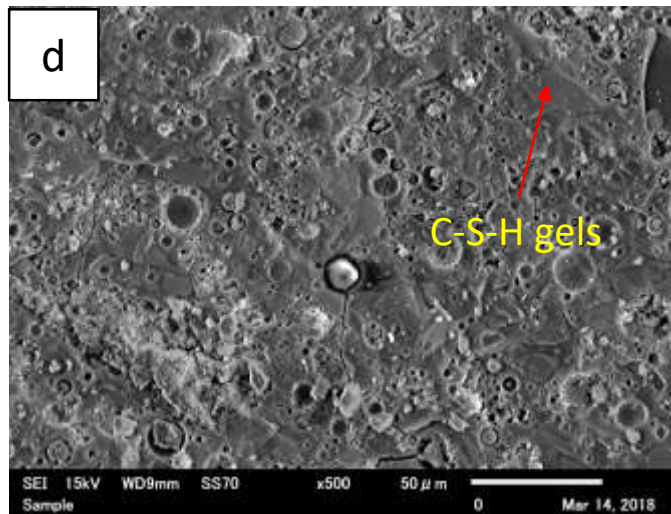
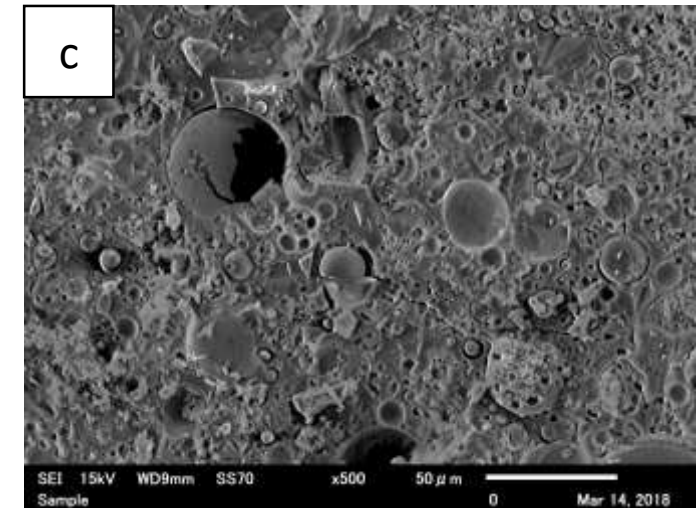
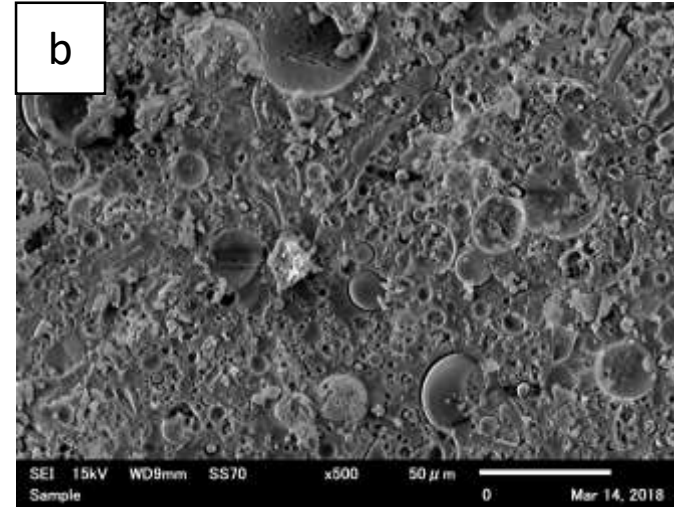
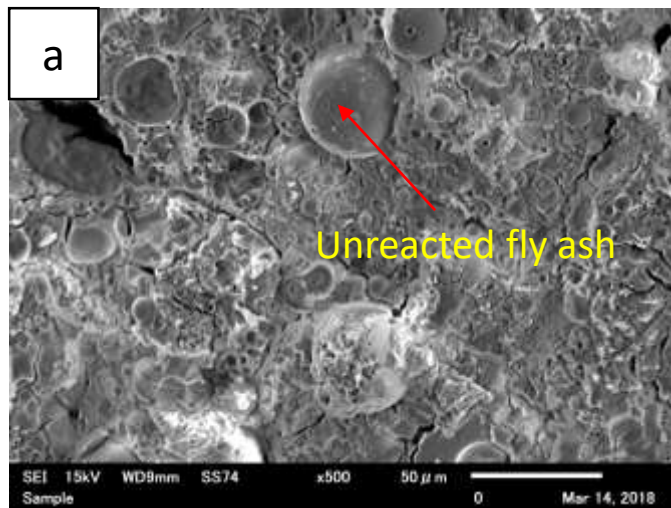


Fig.1. Effect of GGBS on the compressive strength of geopolymer paste specimens.

The compressive strength was found to increase with the increase in amount of GGBS; however, it remained almost constant between 7 and 28 days.



Ca^{2+} and silicon resulting from the dissolution of GGBS react to form a C-S-H gel.

The formation of C-S-H gel helps to form denser structure.

Fig. 2. Micrograph of geopolymer specimens with (a) 0%, (b) 15%, (c) 30%, (d) 45%, (e) 60% of replacement by GGBS.

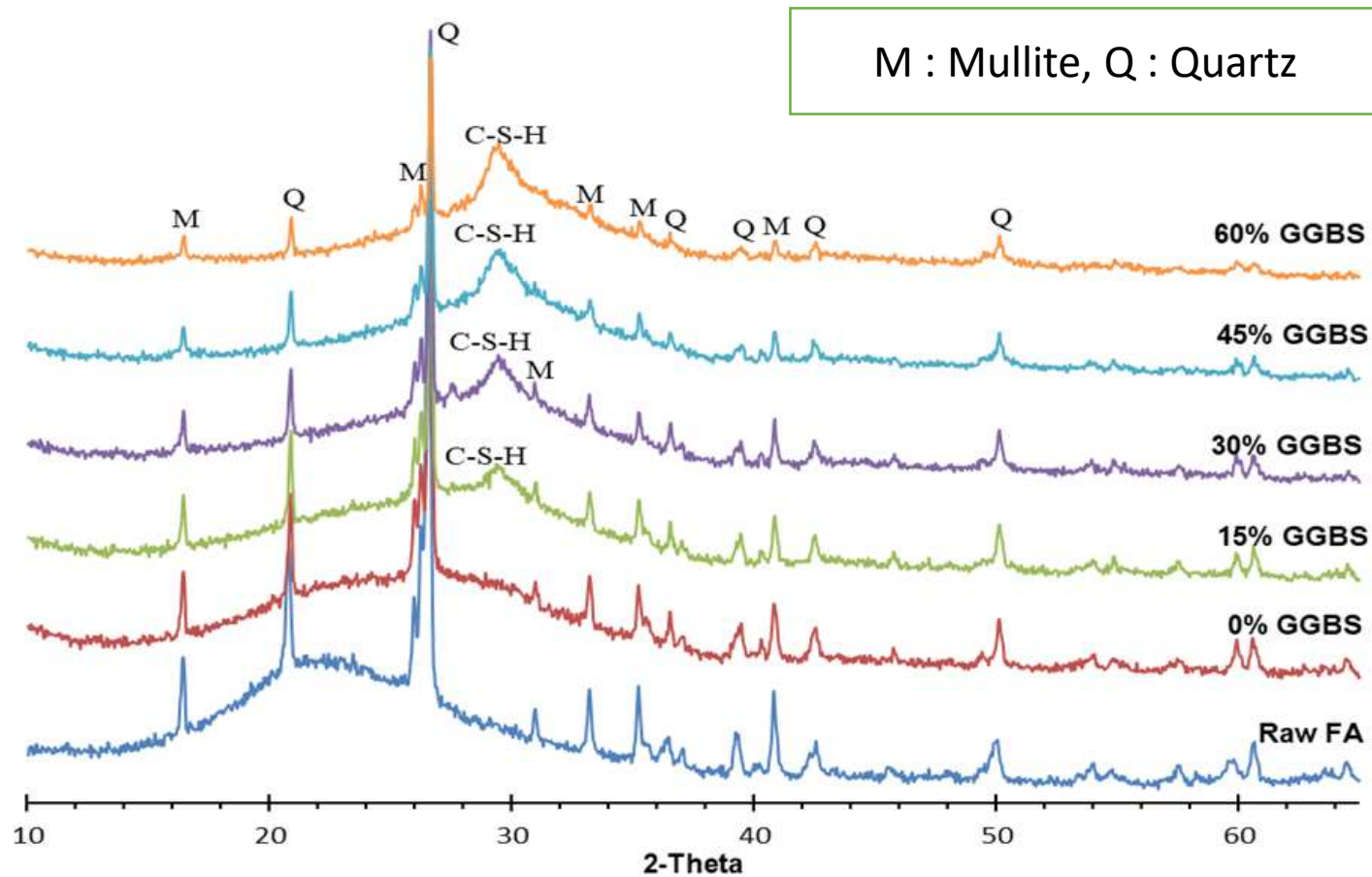
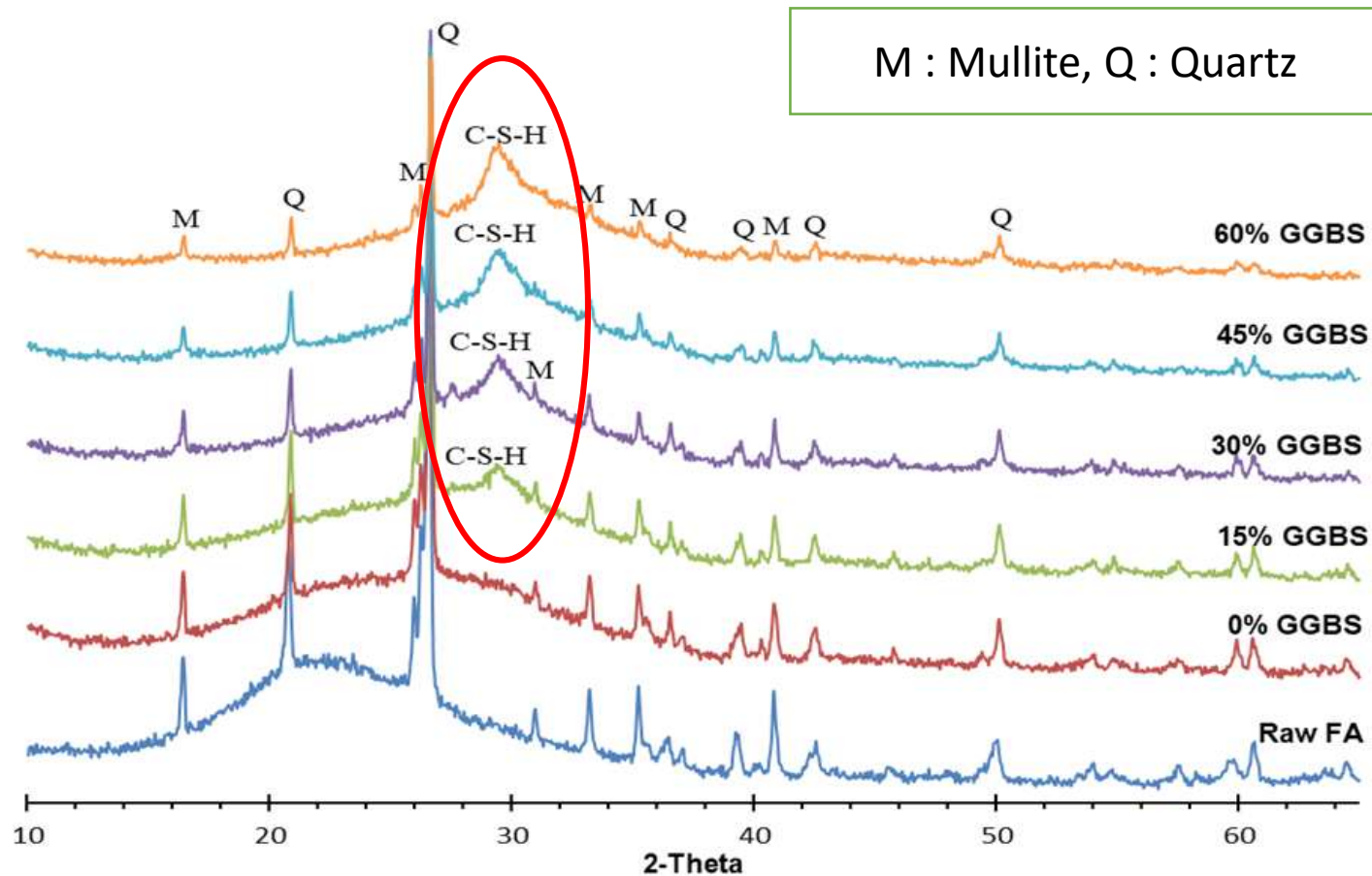


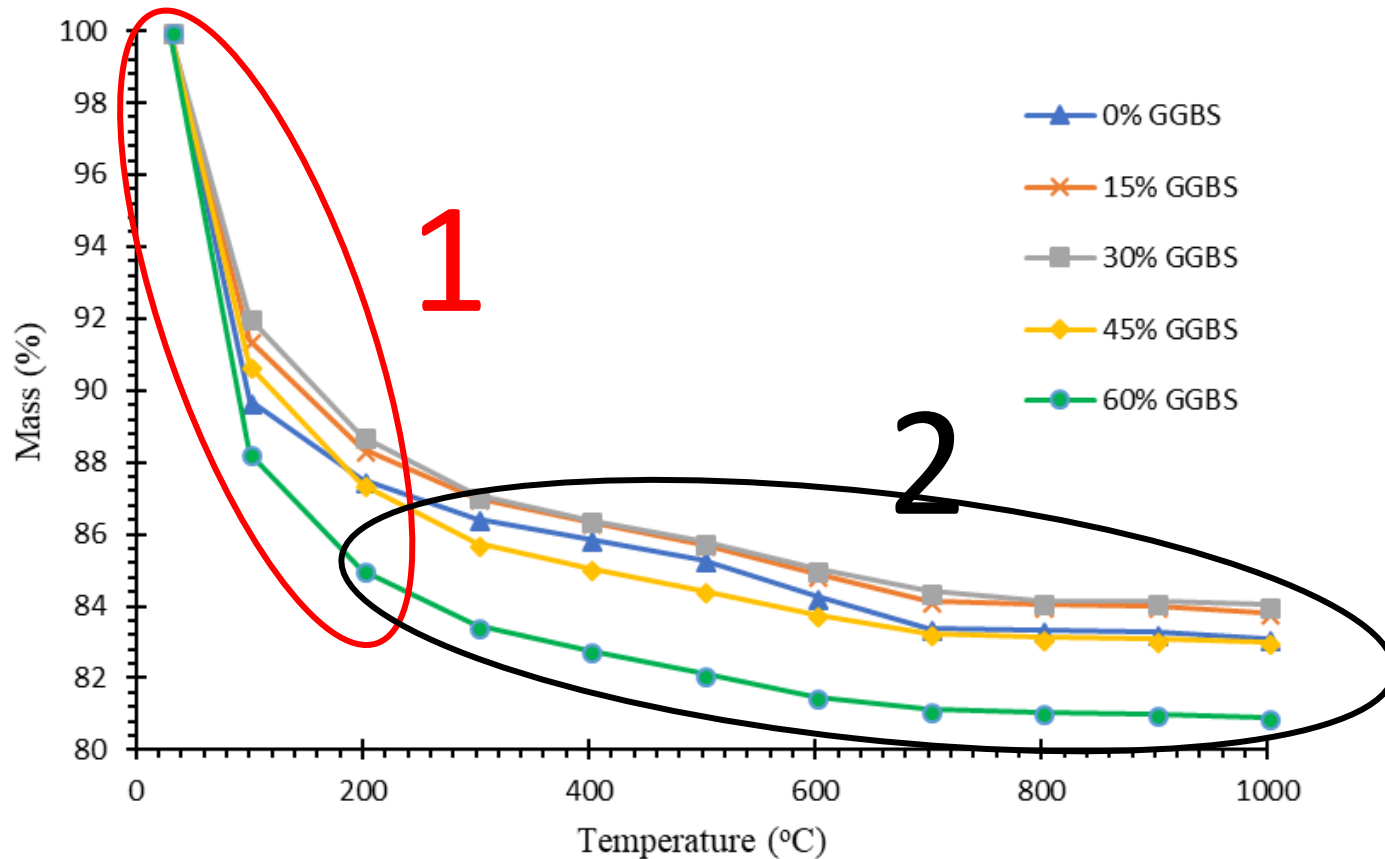
Fig.3. XRD patterns of geopolymer specimens



As **GGBS** was presented in the mixes, the formation of **C-S-H gel** was generated.

The increase amount of GGBS resulting in the increase of C-S-H peak indicates that **the higher the amount of GGBS, the higher the formation of C-S-H.**

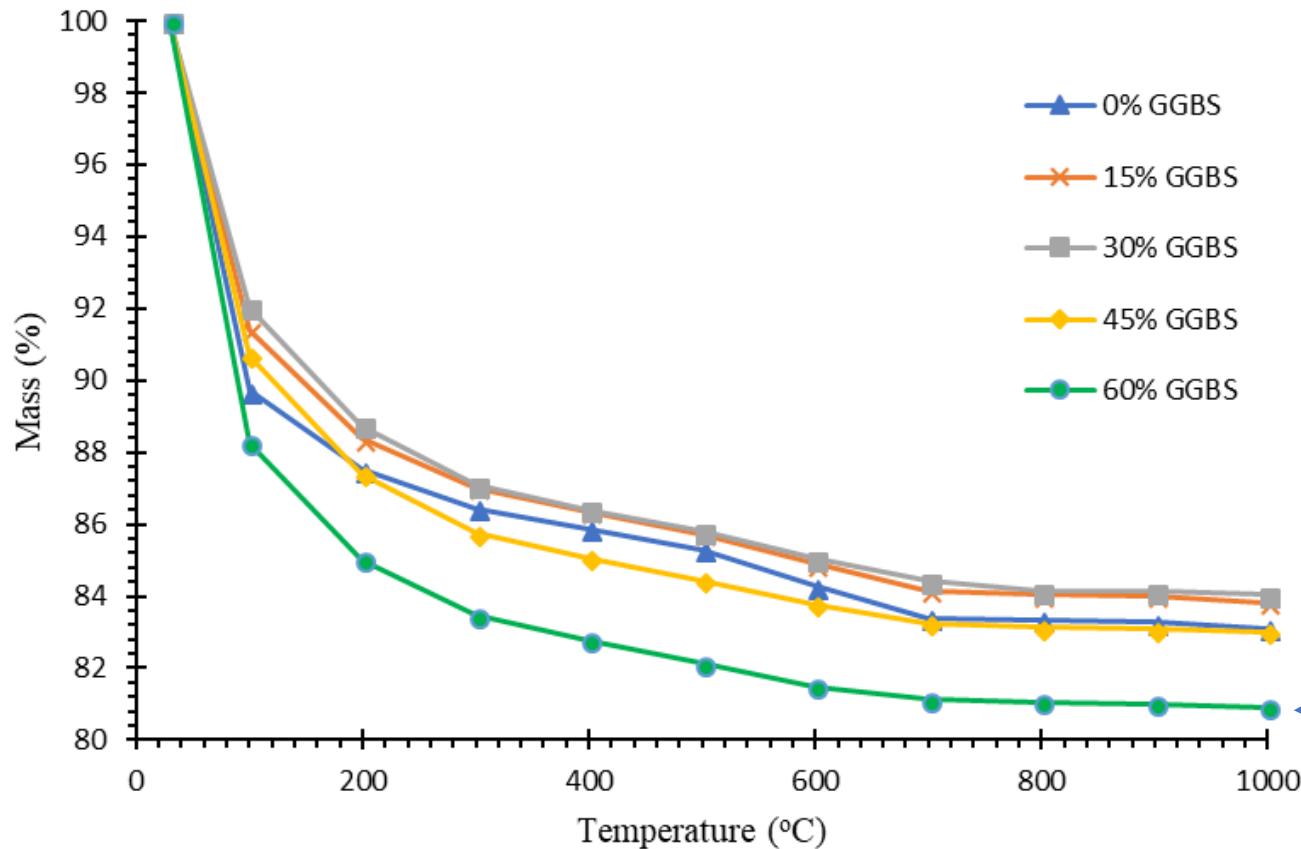
Fig.3. XRD patterns of geopolymer specimens



1. A sharp decrease in mass before 200°C was due to the **evaporation of free water**.
2. Above 200°C, the mass loss is attributed to **dehydroxylation of chemically bound water**.

Fig.4. Thermal stability of geopolymer paste under different dosage of GGBS

Thermo-gravimetric analysis



1. A sharp decrease in mass before 200°C was due to the **evaporation of free water**.
2. Above 200°C, the mass loss is attributed to **dehydroxylation of chemically bound water**.

60% replacement by GGBS may cause **durability problem** when using it in the field.

Fig.4. Thermal stability of geopolymer paste under different dosage of GGBS



1. The addition of **GGBS** in the mixes significantly increased the **compressive strength** of **geopolymer paste** specimens. The highest strength was found at 60% of replacement of **fly ash** by **GGBS**.
2. The **SEM micrographs** show that in the specimens containing GGBS, the **geopolymeric gels** were found to be co-existed with **calcium silicate hydrate (C-S-H) gels**, and thus contributed to the strength development of geopolymer specimens.
3. This study revealed that the **geopolymer cement** made from **fly ash and GGBS** has a potential use as an **alternative binder to replace Portland cement** due to high compressive strength and good thermal stability. However, using too high amount of GGBS (over 45%) may cause durability problem at high temperature due to the C-S-H decomposition.



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Thank you for your attention