

DEVELOPING GEOPOLYMER COMPOSITES USING DRY MIXING TECHNIQUE

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INTRODUCTION

Sustainability Issues:

- Each ton of cement production-one ton of carbon-dioxide, 1 kg of sulphur dioxide (SO₂), 2 kg oxides of nitrogen (NO_x) and 10 kg dust into the atmosphere (Zhang et al., 2018)
- Shortage of Landfill sites

Optimum Solution:

Geopolymer concrete (GPC)- novel form of concrete, synthesized by the alkali activation of source materials (aluminosilicate rich materials) (Davidovits, 1991).

On Site Feasibility Problems:

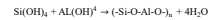
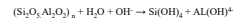
- Highly corrosive alkaline solution-based reagents
- Heat Curing

Feasible Solution: Dry Mixing Technique

- Powder-based reagents: required in less quantity
- Source Materials: aluminosilicate rich materials
- No need of heat curing

OBJECTIVE

- Develop cement free binder for the production/development of sustainable engineered composites.
- Geopolymer Technology



Geopolymerisation (Komnitsas, 2011)

EXPERIMENTAL WORK

Table 1- Mix Proportions for Geopolymer Composites

Binder*	Mix Designation**	Activator	Activator/ Binder	Activator component ratio	Water/Binder	HRWRA***
FA(C)+ GGBS	M1A1	Ca(OH) ₂ + Na ₂ SiO ₃ .5H ₂ O	0.09	Na ₂ SiO ₃ .5H ₂ O/ Ca(OH) ₂ =2.5	0.35	0
	M1A2	Ca(OH) ₂ + Na ₂ SO ₄	0.12	Ca(OH) ₂ / Na ₂ SO ₄ =2.5	0.35	0.01
FA(F)+ GGBS	M2A1	Ca(OH) ₂ + Na ₂ SiO ₃ .5H ₂ O	0.09	Na ₂ SiO ₃ .5H ₂ O/ Ca(OH) ₂ =2.5	0.3	0
	M2A2	Ca(OH) ₂ + Na ₂ SO ₄	0.12	Ca(OH) ₂ / Na ₂ SO ₄ =2.5	0.35	0
GGBS	M3A1	Ca(OH) ₂ + Na ₂ SiO ₃ .5H ₂ O	0.09	Na ₂ SiO ₃ .5H ₂ O/ Ca(OH) ₂ =2.5	0.35	0
	M3A2	Ca(OH) ₂ + Na ₂ SO ₄	0.12	Ca(OH) ₂ / Na ₂ SO ₄ =2.5	0.35	0.02

All numbers are mass ratios of binder

*Binder denotes supplementary cementitious materials (SCMs)

** Mix Designation: M-mix, A-activator

*** HRWRA: Poly-carboxylate based super-plasticizer



Figure 1-Sample Preparation



Figure 2-Ambient Temperature Curing Regime

Figure 3-(a) slump flow spread, (b) setting time test, (c) compressive strength test

RESULTS AND CONCLUSIONS

Table 1-Density and Compressive Strength

Binder*	Mix Designation	Water/Binder	Days	Density (g/cm ³)	Compressive Strength (MPa)
FA(C)+ GGBS	M1A1	0.35	7/14/28/56	2.12/1.97/2.03	37.5/36.6/47.8/55.45
	M1A2	0.35	7/14/28/56	2.14/2.02/2.02	35/44.7/56.3/64.2
FA(F)+ GGBS	M2A1	0.3	7/14/28/56	1.85/1.86/1.86/1.87	20.5/25.6/34.05/34.12
	M2A2	0.35	7/14/28/56	1.75N.A.	8.4N.A.
GGBS	M3A1	0.35	7/14/28/56	2.08/2.02	26.15/22.4/29.3/34.1
	M3A2	0.35	7/14/28/56	2.03/2.00/2.00	26.75/37.8/34.1/31.77

Table 2-Workability

Binder*	Mix Designation	Water/Binder	HRWRA**	Avg. Flow Dia. (mm)	Relative Slump
FA(C)+ GGBS	M1A1	0.35	0	195	2.8
	M1A2	0.35	0.01	165	1.7
FA(F)+ GGBS	M2A1	0.3	0	170	1.89
	M2A2	0.35	0	N.A.	N.A.
GGBS	M3A1	0.35	0	175	2.06
	M3A2	0.35	0.02	200	3

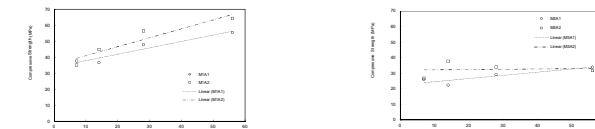


Figure 4- (a) Influence of two types of activators on mix-1, (b) Influence of two types of activators on mix-3

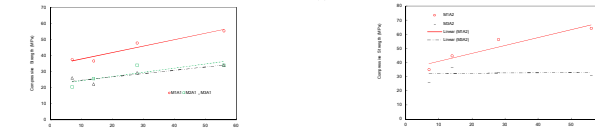


Figure 5- Effect of activator on different source materials (a) A1, (b) A2

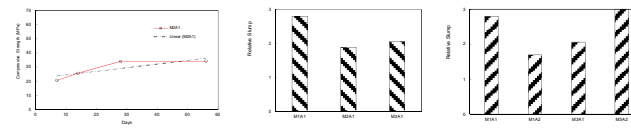


Figure 6- Compressive strength for mix (M2A1)

Figure 7- (a) Relative slump of different mixes for activator A1 (b) influence of activators (A1 and A2) on relative slump of different mixes

- The mix combination (M1A2) achieved the highest compressive strength of 64.2 MPa at 56 days.
- The mix M1A1 exhibited a comparable compressive strength of 55.45 MPa at 56 days and higher slump flow than mix M1A2.
- The initial and final setting time of the mix M1A1 was determined to be as 119 minutes and 259 minutes respectively.
- M1A1 was designated as the best performing mix based on slump flow and compressive strength characteristics.

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