

ELECTRICAL PROPERTIES OF FLY ASH GEOPOLYMER COMPOSITES WITH GRAPHITE CONDUCTIVE ADMIXTURES

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ABSTRACT. Construction materials with increased electrical conductivity could be possibly used in health monitoring of structures (stress, deformation, damages), their maintenance or traffic monitoring. The aim of this study was the application of functional filler and its influence on the electrical properties of the alkali-activated fly ash matrix. The graphite powder was added to the reference material in the amount of 2–10%. Besides the assessment of the critical amount of filler necessary to achieve a percolation threshold in the structure of the composite, the effect on the electrical properties of the matrix (resistance, capacitance, conductivity) was determined. The optimal amount of filler was also determined with respect to the changes in microstructure of the binder and its mechanical properties.

KEYWORDS: Fly ash, geopolymer, graphite, electrical properties, mechanical properties, microstructure.

1. INTRODUCTION

Recently, one of the field in materials and structures research is the concept of advanced smart (or self-sensing) material. Its composition design is based on the content of conductive fillers that enhance its electrical properties. For example, smart composite with higher electrical conductivity can be used for the assessment of structural health in terms of stress monitoring, localization of cracks and damages, humidity and temperature changes etc. [1]. There is a wide range of different functional fillers to use: steel fibres, carbon black, graphite powder, carbon fibres (CF), carbon nanotubes (CNF), graphene, nickel powder.

So far, this approach has been studied predominantly in traditional cement concretes and composites technology. Alkali-activated materials are binders derived from the reactions of raw materials with a high content of amorphous aluminosilicate phase with alkaline activator solution. Alkaline activator (most often in form of alkali hydroxides and/or alkali silicates) is required for the appropriate conditions for the dissolution rate and the formation of new phases [2]. The first theoretical background of alkaline activation was developed by Glukhovskiy in 1959 [3]. The term “geopolymer” was firstly used by prof. Davidovits and refers to a material formed by almost exclusively highly coordinated units of aluminosilicate structures forming regular polymeric three-dimensional structure [4].

Compared to clay-based geopolymers, fly ash provides more favourable rheology of the fresh binder [5], strength of the hardened binder can be enhanced by thermal treatment. Properties of alkali-activated binders are comparable or superior to Portland cement binders [6]. Fly ash geopolymers are generally more durable because of higher chemical and thermal resistance [7]. Moreover, the geopolymer production using mostly secondary raw materials do not require

an exploitation of natural resources and high energy consumption that significantly contribute to lower emissions of CO₂ compared to Portland cement [8].

According to Hanjitsuvan et al. [9, 10], the electrical conductivity of fly ash geopolymer matrix is affected by the NaOH concentration in activator solution, the frequency spectrum and liquid activator/ash ratio (L/A). Payakaniti [11] studied properties of geopolymer incorporated with CF stating the percolation threshold was reached by the geopolymer with 0.5% CF content, this value was increasing with growing L/A. This CF content was also responsible for the best mechanical performance. Vaidya et al. [12] tested fly ash geopolymer concrete beams and cylinders with 0.4% CF while exposed to bending and compressive stress to assess the changes of electrical resistance. Regarding the application of CNT in geopolymer concrete, Saafi et al. [13] states that the optimal dosage is 0.5% which ensures the improvement of both electrical and mechanical performance, higher percentage leads to less effective dispersion and thus cluster formation within the matrix. In another study, the author used graphene as a conductive filler [14]. Only 0.35% addition of graphene resulted in significant increase of strength and modulus of elasticity accompanied by lower porosity and 209% increase of conductivity.

This paper is focused on the application of graphite powder as a conductive admixture in fly ash geopolymer mortars considering its impact on selected electrical properties, strength and changes in binder microstructure.

2. MATERIALS AND METHODS

The geopolymer binder was produced by mixing of fly ash (FA) with commercial sodium silicate solution with SiO₂/Na₂O = 1.6. The fly ash is a product of black coal combustion and its chemical composition is given in Table 1. Quartz sand with a maximum

	SiO ₂	Al ₂ O ₃	Fe ₂ O ₃	CaO	MgO	S _{total}	Na ₂ O	K ₂ O
(%)	49.82	24.67	7.5	3.91	2.68	0.91	0.7	2.78

TABLE 1. Chemical composition of fly ash.

	REF	G2	G4	G5	G6	G8	G10
Fly ash (g)	350	350	350	350	350	350	350
Sodium silicate (g)	280	280	280	280	280	280	280
Graphite powder (g)	-	7	14	17.5	21	28	35
Sand (g)	1050	1050	1050	1050	1050	1050	1050
2% Triton X-100 (g)	-	7	14	17.5	21	28	35
1% Lukosan (g)	-	3	6	7.5	7.5	7.5	7.5
Water (g)	35	30	30	32	32	32	32

TABLE 2. Mix composition of FA geopolymer mortars with graphite powder.

grain size of 2.5 mm was used as aggregate. Graphite powder PMM 11 was used as a conductive filler in 2, 4, 5, 6, 8 and 10% wt. of the fly ash and its dispersion within the matrix was supported by the addition of dispersing agent (2% Triton X-100). Defoaming agent (1% Lukosan) was introduced to reduce the air content in fresh and hardened binder.

The raw mix composition is presented in Table 2. Each of the mixes was produced following these steps: at first, the fly ash was mixed with water glass. Then, the graphite powder and dispersing agent were added and homogenized with the binder and a small amount of water, if needed. Thereafter, each fraction of aggregate (from fine to coarse) was added and mixed properly. The defoaming agent and remaining amount of water were added at the very end of the mixing procedure.

The mortars were cast into prismatic moulds (40 × 40 × 160 mm) and covered with a plastic sealant to avoid moisture loss. After 2 hours in the ambient conditions, the specimens were heated at 40 °C for 24 h. After demoulding, the hardened specimens were stored in the laboratory conditions (22 ± 2 °C, $\varphi = 45 \pm 5\%$) till the age of testing.

The electrical and mechanical properties (compressive and flexural strength) were tested at the age of 56 days. The prepared prismatic samples were characterized by impedance spectroscopy in the range of 40 Hz to 1 MHz using an Agilent 33220A sinusoidal signal generator and an Agilent 54645A dual-channel oscilloscope. The output voltage of the signal generator was 5.5 V. The input values for the electrical capacity and the resistance of the oscilloscope were 13 pF and 1 M Ω , respectively. In order to perform impedance analysis, the prismatic specimens were placed between parallel brass electrodes (30 × 100 mm) so that a distance between electrodes was 40 mm. Conductivity measurements were performed in the range of 10 to 3510 MHz. The microstructure and porosity of the geopolymers were evaluated using Micromeritics Pore-

sizer 9310 and scanning electron microscope (SEM) Tescan MIRA3 XMU.

3. RESULTS AND DISCUSSION

Electrical resistance of all samples decreased with higher frequency applied, as can be seen from Fig. 1. The lowest resistance was observed in case of 10% graphite content (G10), at all frequencies. The differences in resistance curves are clearly visible up to 10 kHz. Above this frequency, the resistance of all samples fluctuate and the curves overlap. At 100 Hz, the resistance of reference sample was 159.80 M Ω . The resistance of G5 dropped to 43.78 M Ω and 0.85 M Ω only at G10.

Changes of resistance and conductivity at chosen frequency are shown in Fig. 2. While the resistance decreased continuously, we registered an exceptional increase in conductivity of specimen G10 from 0.38 to 6.31 mS·m⁻¹. These results suggest that geopolymer with 10% of graphite may have reached the percolation threshold.

The capacitance of geopolymers with various graphite content as a function of frequency is shown in Fig. 3. Up to 6% of graphite, the changes in capacitance are comparable, but the capacitance of G8 and G10 shows the most notable increase over all frequencies. Capacitance of all samples except G10 decreased with higher frequency. The capacitance of G10 first dropped from 222 to 122 pF (40–180 Hz), within the frequency 180–10 000 Hz it was about 125 pF and at higher frequency applied it was decreasing again.

The changes of mechanical performance depending on the graphite content are displayed in Fig. 4. The flexural strength was not negatively influenced by the graphite addition and fluctuated around 5 MPa at all graphite concentrations. On the contrary, the compressive strength gradually decreased. While the reference sample without graphite reached 23.5 MPa, the samples with 2% graphite content (G2) reached 17.7 MPa and 10% graphite content (G10)

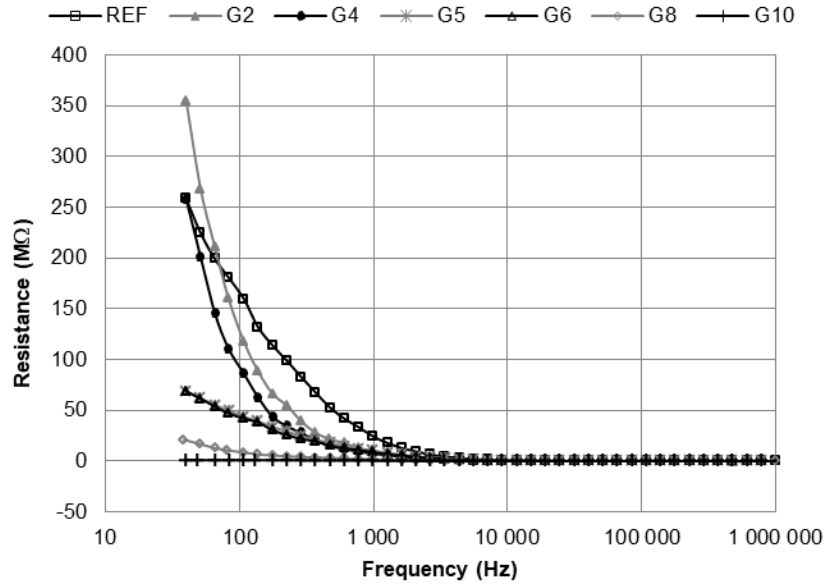


FIGURE 1. Variation in electrical resistance of FA geopolymers with different graphite content.

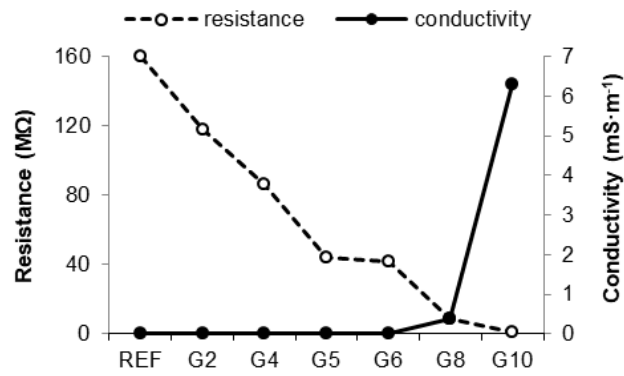


FIGURE 2. Resistance (at 100 Hz) and conductivity (at 100 MHz) of FA geopolymers with different graphite content.

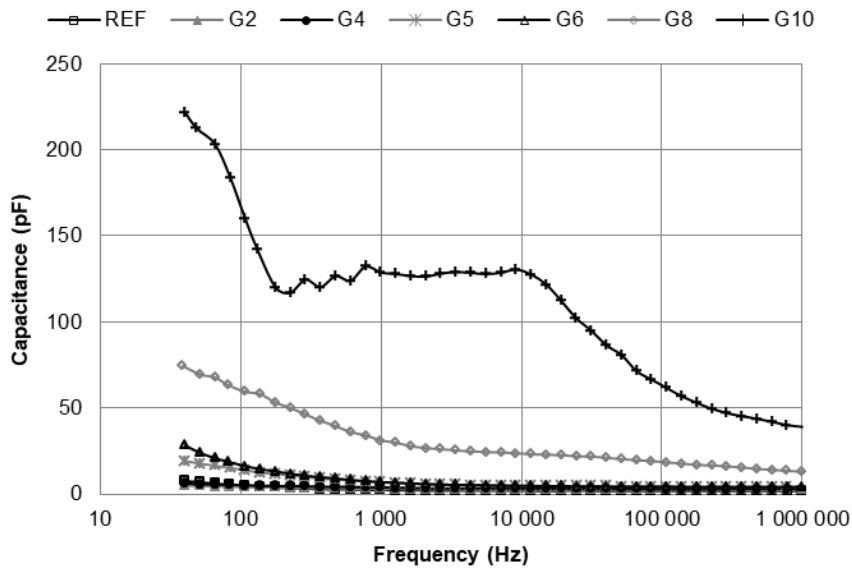


FIGURE 3. Variation in capacitance of FA geopolymers with different graphite content.

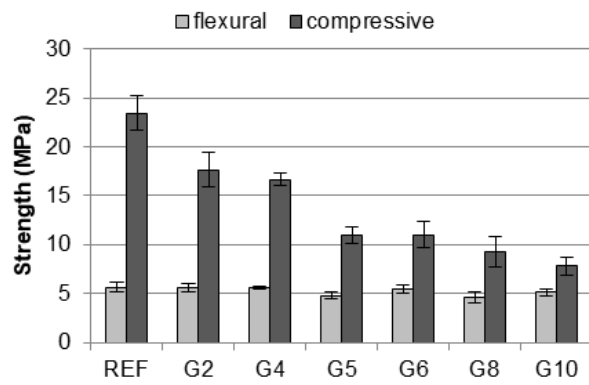


FIGURE 4. Mechanical properties of FA geopolymers with different graphite content.

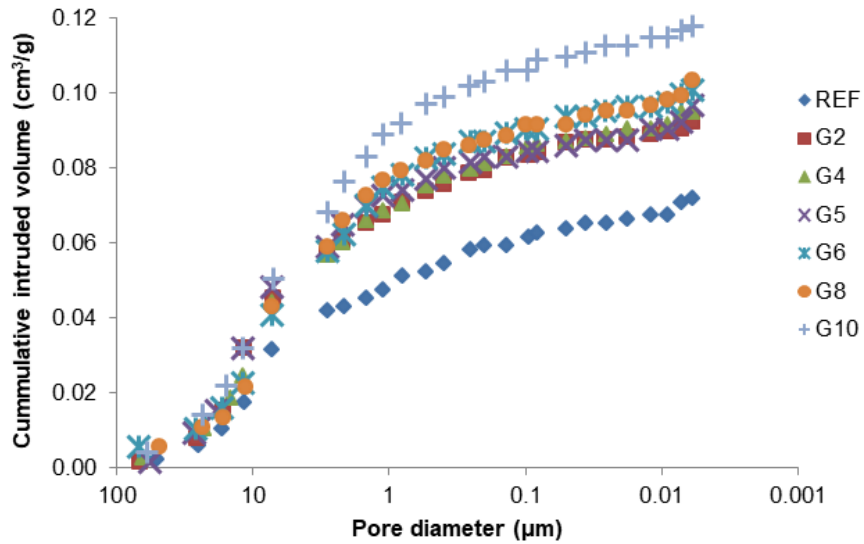
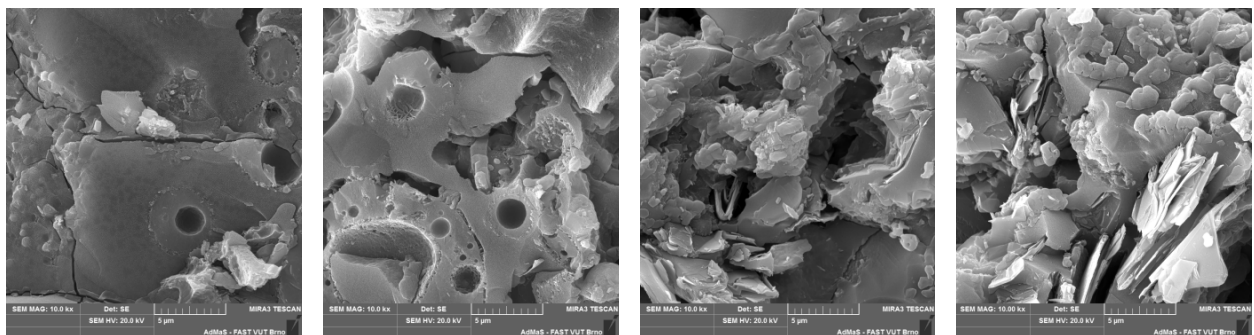


FIGURE 5. Cumulative intruded volume of FA geopolymers with different graphite content.



(A) . REF.

(B) . G2.

(C) . G5.

(D) . G10.

FIGURE 6. SEM images of REF, G2, G5 and G10 geopolymers.

only 7.8 MPa, which is equal to 25% and 65% drop of strength, respectively.

Regarding the microstructure of the geopolymer binder, even 2% of graphite caused an increase in total porosity of the samples (Fig. 5). While the geopolymers with a graphite content of 2–8% did not exhibit considerable differences, we observed a further increase in pore volume of G10 which is more than 60% higher than the reference sample.

In Fig. 6 we can see the nature of chosen binders in SEM images. Reference sample is characterized by homogeneous amorphous binder phase whereas number of pores and inhomogeneities increased with the amount of conductive filler.

4. CONCLUSIONS

The results mentioned above confirm that graphite powder can be applied to fly ash geopolymer matrix to enhance its electrical properties. Higher graphite content resulted in reduced electrical resistance, especially in frequency up to 1 000 Hz. Likewise, the samples with high graphite content exhibited an increase in capacitance. The best performance in terms of electrical properties was observed in case of geopolymer with 10% graphite content providing significant increase in conductivity, which may refer to achieving the percolation threshold. However, the presence of conductive filler in such concentration deteriorated the mechanical performance in compression by more than 65%; flexural strength of all samples remained constant regardless of the amount of graphite powder. Despite the use of the defoaming agent, the microstructure of all geopolymers with graphite was characterized by higher pore volume which is in a good accordance with reduced compressive strength. We can conclude that graphite powder in fly ash geopolymer allows to develop a conductive network within the matrix but also causes generation of additional pores during mixing. The negative impact of graphite powder on the compressive strength is apparently attributed to a weak bond between the matrix and graphite particles because of nonpolar character of graphite surface. Low toughness of graphite itself can further contribute to the strength loss at high concentrations. Possible ways to reduce these impacts include more efficient defoaming or improved adhesion of graphite particles to the matrix by partial modification of its structure with hydrophilic substituents.

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REFERENCES

- [1] B. Han, L. Zhang, J. Ou. *Smart and Multifunctional Concrete Toward Sustainable Infrastructures*, chap. Self-Sensing Concrete, pp. 81–116. Springer, Singapore, 2017. DOI:10.1007/978-981-10-4349-9_6.
- [2] C. Shi, D. Roy, P. Krivenko. *Alkali-Activated Cements and Concretes*. CRC press, 2003. DOI:10.1201/9781482266900.
- [3] V. D. Glukhovskiy. *Soil silicates (Gruntosilikaty)*. Budivelnik Publisher, Kiev, 1959.
- [4] J. Davidovits. Properties of geopolymer cements. In *First international conference on alkaline cements and concretes*, pp. 131–149. 1994.
- [5] L. Provis, J. S. van Deventer. *Alkali Activated Materials*. Springer, 2014. DOI:10.1007/978-94-007-7672-2.
- [6] P. Duxson, A. Fernández-Jiménez, J. L. Provis, et al. Geopolymer technology: the current state of the art. *Journal of materials science* **42**(9):2917–2933, 2007. DOI:10.1007/s10853-006-0637-z.
- [7] A. Fernández-Jiménez, I. García-Lodeiro, A. Palomo. Durability of alkali-activated fly ash cementitious materials. *Journal of Materials Science* **42**(09):3055–3065, 2007. DOI:10.1007/s10853-006-0584-8.
- [8] K. P. Mehta. Reducing the environmental impact of concrete. *Concrete international* **23**(10):61–66, 2001.
- [9] S. Hanjitsuwan, S. Hunpratub, P. Thongbai, et al. Effects of NaOH concentrations on physical and electrical properties of high calcium fly ash geopolymer paste. *Cement and Concrete Composites* **45**:9–14, 2014. DOI:10.1016/j.cemconcomp.2013.09.012.
- [10] S. Hanjitsuwan, P. Chindaprasirt, K. Pimraksa. Electrical conductivity and dielectric property of fly ash geopolymer pastes. *International Journal of Minerals, Metallurgy, and Materials* **18**(1):94–99, 2011. DOI:10.1007/s12613-011-0406-0.
- [11] P. Payakaniti, S. Pinitsoontorn, P. Thongbai, et al. Electrical conductivity and compressive strength of carbon fiber reinforced fly ash geopolymeric composites. *Construction and Building Materials* **135**:164–176, 2017. DOI:10.1016/j.conbuildmat.2016.12.198.
- [12] S. Vaidya, E. N. Allouche. Strain sensing of carbon fiber reinforced geopolymer concrete. *Materials and structures* **44**(8):1467–1475, 2011. DOI:10.1617/s11527-011-9711-3.
- [13] M. Saafi, K. Andrew, P. L. Tang, et al. Multifunctional properties of carbon nanotube/fly ash geopolymeric nanocomposites. *Construction and Building Materials* **49**:46–55, 2013. DOI:10.1016/j.conbuildmat.2013.08.007.
- [14] M. Saafi, L. Tang, J. Fung, et al. Graphene/fly ash geopolymeric composites as self-sensing structural materials. *Smart materials and structures* **23**(6):065006, 2014. DOI:10.1088/0964-1726/23/6/065006.