Prediction models of mechanical properties for pet-mortar composite in sodium sulphate aggressive mediums

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Abstract. In this research, an investigation was carried out on the effect of sodium sulphate attack on the durability of composites produced with waste polyethylene terephthalate (PET). Experiments were accomplished on limestone sand and cement mortars where the blended Portland cement was partially replaced by various volume fractions of waste PET particles (6%, 12% and 17%). The test solutions used to supply the sulphate ions and cations were 5% sodium sulphate solution. Compressive strengths measured on specimens were used to assess the changes in the mechanical properties of PET-mortars exposed to sulphate attack at different ages, mainly the Young modulus of elasticity. Based on experimental compressive tests on PET-Mortar composite specimens and there densities, the evolution of Young modulus of elasticity has been analyzed in accordance with normative models given by (ACI-318) and (BS-8110) codes of practice. In addition, a comparative study has been carried out for corrosion resistance coefficients K of unmodified mortar to those modified with waste PET particles. It can be noticed that, for the composite immersed in a corrosive Na2SO4 solution, the corrosion resistance coefficients decrease with the increase of the immersion period. The corrosion sulphate resistance K based on Young modulus before and after immersion of PET-mortar composites is better than that of the control mortar. Therefore, for safety considerations of PET-mortar composites use, ACI 318 is recommended code for design and investigation works. Also, it can be concluded that adding waste PET by volume fractions (6%, 12% and 17%) to blend Portland cement renders this cement more resistant to the sodium sulphate aggressive medium. Therefore, composites materials based waste PET are often presented as the materials of the future because of their potential for innovation and the advantages they offer. In fact, using waste PET as cement substitutes reduces the energy consumption. These modified mortars address problems related to environmental pollution by CO2 emissions, and are used to repair various reinforced concrete structures in sodium sulphate aggressive mediums.

1 Introduction

In the last couple years, waste polyethylene terephthalate (PET) is produced within large amount by plastic industry in all over the world [1] and since PET waste is not biodegradable, it can remain in nature for hundreds of years and causes too many environmental problems.

Various studies [2-6] have been developed so far in order to find ecologic and green ways to dispose of plastics wastes, one of the main solutions proposed by researchers is to incorporate waste PET in building materials technology in order to substitute volumetric amount of cement [7] and/or aggregates [8-10] in concrete and mortars mix-design.

In this research, an investigation was carried out on the effect of sodium sulphate attacks on the durability of composites produced with waste polyethylene terephthalate (PET). Experiments were accomplished on limestone sand and cement mortars where the blended Portland cement was partially replaced by various volume fractions of waste PET particles (6%, 12% and 17%).

Test solutions used to supply the sulphate ions and cations were 5% Na2SO4 solution. Tap water was used as the reference solution. Compressive strengths measured on specimens were used to assess the changes in the mechanical properties of PET-mortars exposed to sodium sulphate attacks at different ages, mainly the Young modulus of elasticity. After compression testing, X-ray diffraction was conducted on some selected surface fractures to investigate microstructural nature of the sulphate attacks.
2 Raw materials

2.1 Cement

The cement used was a blended Portland cement type CPJ-CEM II/A42.5 supplied by Zahana factory, located in western Algeria, with 1022 kg/m³ bulk density; its compressive strength at 28 days was 42.5 MPa. The absolute density of the cement used was 3.15 g/cm³ and its specific surface area measured with the Blaine method was 3532 cm²/g. Its initial and final setting times were 170 and 245 min, respectively. Mineralogical and chemical compositions of cement are listed in Table 1. The chemical composition was obtained using an X-ray fluorescence spectrometer.

Table 1. Chemical and mineralogical compositions of cement (wt.%)

| Chemical compositions | SiO₂ | Al₂O₃ | Fe₂O₃ | CaO | MgO | SO₃ | K₂O | Na₂O | CaO free | LOI |
|-----------------------|------|-------|-------|-----|-----|-----|-----|------|----------|-----|
| SiO₂                  | 20.91| 5.52  | 3.56  | 63.50| 0.64| 2.79| 1.23| 0.13  | 2.35     | 1.19 |

| Mineralogical compositions | C₃S | C₂S | C₃A | C₄AF |
|----------------------------|-----|-----|-----|------|
| C₃S                       | 49.39| 22.97| 8.61| 10.83 |

2.2 Sand

The crushed natural limestone sand was obtained from the quarry of Kristel, in Oran, West Algeria. The maximum size of sand grains was 5 mm. The absolute density and absorption coefficient of crushed sand were 2.53 g/cm³ and 0.5%, respectively. The grading of crushed sand is presented in Table 2, according to standard NF P18-560 [11].

Table 2. Sieve analysis of waste PET-particles and crushed limestone sand.

| Sieve size (mm) | Cumulative passing (%) |
|----------------|------------------------|
| Waste PET      | Sand                   |
| 5              | 99.92                  | 99.83                  |
| 2.5            | 98.16                  | 98.37                  |
| 1.25           | 96.82                  | 65.37                  |
| 0.63           | 55.78                  | 38.3                   |
| 0.315          | 35.48                  | 19.07                  |
| 0.16           | 18.28                  | 8.20                   |
| 0.125          | 9.56                   | 3.325                  |

2.3 Waste Polyethylene Terephthalate

Waste PET bottle granules (PET) used as particles were supplied by TRAMAPLAST PET Bottle Plant, in Tlemcen, Algeria. These particles were obtained by collecting the waste PET bottles and washing them; they are then crushed by granules into machines. In addition, they have an irregular shape and a rough texture surface, which enables the adherence of the particle-matrix. The bulk density of the waste PET particles used was 401.4 kg/m³.

After preliminary tests, waste PET particles of size lower than 1 mm were used in this study. The sieve analysis of waste PET particles was carried out according to standard NF P18-560 [11] and is presented in Table 2.

| Sieve size (mm) | Cumulative passing (%) |
|----------------|------------------------|
| Waste PET      | Sand                   |
| 5              | 99.92                  | 99.83                  |
| 2.5            | 98.16                  | 98.37                  |
| 1.25           | 96.82                  | 65.37                  |
| 0.63           | 55.78                  | 38.3                   |
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| 0.16           | 18.28                  | 8.20                   |
| 0.125          | 9.56                   | 3.325                  |

3 Composite mixing conditions

The mortar manufactured without waste PET particles was first optimized on the basis of its mechanical criteria and was then used as a reference composite. The composites containing waste PET particles were produced in accordance with the results of the works of Benosman et al. [12]. A massic ratio of 3 between sand (S) and cement (C) was respected. Four different mixtures were prepared (the control mixtures without plastic waste and three PET# mixtures including 6%, 12% and 17% waste PET particles by volume). Mixture name of the different composites were: PET0 (without plastic waste), PET6, PET12 and PET17. The water to binder ratio was kept constant at 0.5. So, after pouring fresh material into the molds (EN 196-1), the samples were stored in a room where hygrometry and temperature were controlled for 24 h (98% relative humidity, and 20 ± 1 °C). After removal from the molds, at 1 day of age, mortar specimens were cured in saturated lime water at 20 ± 1 °C, until the time of testing.

4 Test methods of resistance to sodium sulphate attack

The mortar specimens were cured in water saturated with lime at 20 ± 1°C for 28 days before being exposed to sulphate attack. Three specimens of each mortar and composite mixes (40x40x160 mm³) were immersed in two types of solutions: distilled water (reference medium) and 5% sodium sulphate solution, Figure 1. According to the standard ASTM C1012-04 [13], the pH of the sulphate solution should be between 6 and 8 and the solution must be renewed each week, which requires huge amounts of sodium sulphate.

For this, Mehta’s [14] method and that of Siad et al. [15] were adopted; they all recommend the control of the pH within a range of 6.0–8.0 by adding a suitable amount of sulfuric acid solution (0.1 N H₂SO₄), figure 2.
The correction is performed daily during the first weeks of immersion, and then becomes weekly, for the rest of the test. In addition, the aggressive solutions were totally renewed each 12 weeks.

After immersion in sodium sulphate solution (Na₂SO₄) for the required period of time, ASTM C1012-04 [13], the specimens were tested for residual mechanical properties. The Young modulus loss (YML%) is calculated as follows:

$$YML(\%) = \frac{E_{cr} - E_{cs}}{E_{cr}} \times 100 \quad (1)$$

where $E_{cr}$ is the Young modulus of the specimens before immersion (MPa) and $E_{cs}$ is the average Young modulus of the specimens after immersion in sodium sulphate solutions for the required period of time (180 days).

5 Results and discussion

5.1 Prediction models of mechanical properties

The static modulus of elasticity $E$ (Young Modulus) represents one of the most important mechanical characteristics of construction materials (concrete, reinforced concrete, mortars, composites, etc.). This intrinsic property is considered as the basic parameter for the computing strain-stresses in construction structures.

Various countries have been established their design codes based on this empirical relationship between static modulus of elasticity $E$, and compressive strength of plain concrete at 28 days of curing. The ACI code (ACI-318) [16] defines the relationship between elastic modulus of concrete and compressive strength as:

$$E = w^{1.5} \times 0.043 \times f_c^{0.33} \quad (2)$$

The British Code of practice (BS-8110) [17] recommends the following expression for static modulus of elasticity with cube compressive strength of concrete as:

$$E = w^{2} \times 0.0017 \times f_c^{0.33} \quad (3)$$

where, $E_c$: The static modulus of elasticity (E) at 28 days in MPa $f_c$: Compressive strength at 28 days, in MPa $w$: Air dry density of mortar

Based on experimental compressive tests on PET-mortar composite specimens [18], and there densities, graphs in (figures 3,4) below show the evolution of Young modulus evaluated by empirical relationships in accordance to (ACI-318) and (BS-8110) codes [16,17].

In the other hand, Young modulus loss ratios have been studied for several volumetric waste PET rates and for each model of predicted codes. The results of the Young modulus loss using the specimens immersed in 5% Na₂SO₄ solutions (figure 5) showed that there are variations in time and group. However, a decrease in
Young Modulus values of all specimens was observed when Young Modulus of elasticity is computed via ACI-318 model. However, the rate of increase varied slightly within the group.

![Young modulus loss (YML%) with volumetric waste PET rate and prediction codes (BS-8110 & ACI-318) before and after 5% Na₂SO₄ immersion.](image)

It was expected that the mechanical properties loss (YML%) values of modified mortars with waste PET plastic particles would be lower than those of unmodified mortar PET0, by 14.35%, 15.28% and 14.53% for PET6, PET12 and PET17, respectively for the case where the prediction model is BS-8110 and respectively by 11.72%, 13.01% and 11.51% when Young Modulus of elasticity is computed via ACI-318 model. It can therefore be concluded that modified mortars by waste PET plastic particles are resistant to the sodium sulphate aggressive mediums conditions often encountered in the field.

Similarly to experimental results obtained by some previous studies [19, 20] in similar fields, it can be stated that, mortar blended within waste polymer particles present a better and promising composite material in building repair against sulphate aggressive mediums (Na₂SO₄, MgSO₄).

### 5.2 Corrosion sodium sulphate resistance

Furthermore, to compare, effectively, the corrosion resistance coefficients K of unmodified mortar (Eq. 4) to those modified with PET particles, as it was used by Benosman et al. [12] and Jiang et al. [21]:

$$K = \frac{Eci}{Ecs}$$

Where $E_{ci}$ is the Young modulus of composite mortars immersed in corrosive Na₂SO₄ solutions, $E_{cs}$ the Young modulus of the normally cured composite mortars. The corrosion resistance coefficients of the specimens with and without waste PET plastic are given in Table 3.

It can be seen from table 3 that, for the composite immersed in a corrosive Na₂SO₄ solution, the corrosion resistance coefficients decrease with the increase of the immersion period. The corrosion sulphate resistance K based on Young modulus before and after immersion of PET-mortar composites is better than that of the control mortar (PET0).

### Table 3. Corrosion resistance coefficients of PET-mortar composites in 5% Na₂SO₄

| Sodium sulphate attacks | K (BS.8110) 90 days | K (BS.8110) 180 days | K (ACI.318) 90 days | K (ACI.318) 180 days |
|------------------------|---------------------|---------------------|---------------------|---------------------|
| PET0                   | 0.922               | 0.823               | 0.927               | 0.850               |
| PET6                   | 0.970               | 0.856               | 1.000               | 0.900               |
| PET12                  | 0.945               | 0.847               | 0.946               | 0.870               |
| PET17                  | 0.956               | 0.855               | 0.986               | 0.904               |

For all studied cases, corrosion resistance coefficients K computed via BS.8110 code of practice are lower than the ones computed by ACI 318 model. Therefore, for safety considerations of PET-mortar composites use, ACI 318 is recommended code for design and investigation works. It can be concluded that adding waste PET by volume fractions (6%, 12% and 17%) to blended Portland cement renders this cement more resistant to the sodium sulphate aggressive medium. It is evident that the resistance of cement to sulphate aggression is also related to its content in Ca₃[22].

Therefore, composites materials based waste PET are often presented as the materials of the future because of their potential for innovation and the advantages they offer. In fact, using waste PET as cement substitutes reduces the energy consumption. These PET-modified mortars address problems related to environmental pollution by CO₂ emissions, and are used to repair various reinforced concrete structures in sodium sulphate aggressive mediums.Alqahtani et al. [23] reported that recovering plastic waste would reduce the CO₂ emissions by 3.8 million tons. So, these results are in agreement with those stated by these authors [23].

Finally, we can suggest the use of these current composite materials for surface treatment of building and construction façades exposed to marine environment (For example; Hassan II mosque), as well as harbour infrastructures and manufacture of waste water sewer pipes and their accessories (junction mortars).

### 5.3 XRD analysis

Figure 6 exhibits the XRD analysis of PET0 and PET17, after attack by Na₂SO₄ solution. The stacking of various spectra confirms the appearance of:

- Ettringite (E) $Ca₄(Al₃(SO₄)₃(OH)₆)·26H₂O$: 9.73° 2θ;
- Thaumasite (T), $Ca₆[Si(OH)₆]₂(CO₃)₂(SO₄)₂·24H₂O$: 9.73° 2θ;
- Gypsum (G) $CaSO₄·2H₂O$: 11.59°, 20.72°, 23.40° 2θ;
- Calcite (C), $CaCO₃$: from the crushed natural limestone sand (29.41° 2θ);
- Portlandite $Ca(OH)₂$: was completely decomposed by Na₂SO₄ solution resulting from the chemical sulphate reactions [6].
Fig. 6. X-ray diffraction pattern of the specimens under 5% sodium sulphate attacks, after 90 days. (Blue: PET0; Pink: PET17).

6 Conclusions

With reference to the results presented in this paper, the main conclusions which can be drawn are summarized as follows:

- Composite modified mortars by waste PET plastics present a significant resistance to sodium sulphate attack which reduces the consumption energy of modified mortars resulting from PET-cement substitutes. The corrosion resistance coefficients decrease with exposure time to sodium sulphate aggressive medium.

- The mechanical properties of PET-mortar composites computed by ACI-318 and BS-8110 codes decrease with the increasing of PET volumetric substitution rate and this, for all studied cases (before and after immersion in 5% Na₂SO₄ medium). Prediction model proposed by BS-8110 gives always the lowest values of elastic Young modulus which can be recommended for structural design in terms of safety and mechanical reliability.

- Since PET-mortar composite are resistant to sodium sulphate attack (as concluded above), the mechanical properties loss in terms of static Young modulus of elasticity (YML%) of modified mortars are lower than the ones of unmodified mortars (PET0). The optimal minimal values of (YML%) are given by ACI-318 code of practice, for the case of PET-Modified mortars within volumetric rate of 6%, 12% and particularly, 17% (PET17).

- Finally, the better durability properties of PET-mortar composites observed in this study indicate longer life of the repaired structure by using this type of green PET-modified repair materials.

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