Novel masonry grout incorporating high volumes of industrial by-products: microstructure characteristics and pursuance of durability properties

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Abstract

One solution to the high cost and scarcity of building materials is to use alternative and sustainable materials. The study presented herein developed an eco-friendly masonry grout using high volumes of palm oil clinker powder to replace cement and palm oil clinker to replace coarse aggregate. Several batches of grouts with different amounts of these materials were made to determine the technical viability of the grout. Scanning Electronic Microscope (SEM), X-Rays Diffraction (XRD), Energy Dispersive X-Ray (EDX) and Thermogravimetric Analyzer (TGA) analyses were conducted to investigate the microstructure characteristics of the grout, and water absorption, initial rate of absorption, sulphate attack and electrical resistivity tests were conducted to determine its durability. Compressive strength tests were conducted at different curing ages and the drying shrinkage of the grout was monitored for 180 days. The results indicate that the new grout is as good as a conventional grout but with added sustainable and economic benefits. The new grout can be used in masonry construction and can be used to alleviate the inadequate supply of affordable housing.

Keywords Masonry Grout · Palm Oil Clinker (POC) · Palm Oil Clinker Powder (POCP) · Lightweight Aggregate · Sustainability · Microstructure Behaviour and Durability

Introduction

Masonry is a composite material and its constitutive relationship is complex with distinguishing material properties, which are directly related to the properties of its constituents. Modern masonry has two main components: the masonry unit and mortar. Masonry grout is added to the basic system

to increase bearing load capacity and reinforcement is added when the system needs to resist tensile loads.

A current common practice in masonry construction is to use hollow masonry blocks (HMB) or units due to the lightweight of the units, affordability and increased masonry productivity. Although HMB masonry construction is effective in resisting compressive loads, it requires continuity between the units when the structure needs to resist lateral loads. For such cases, the structure needs to be reinforced and reinforcement requires masonry grout to bond the reinforcement to the masonry units.

Several studies have been conducted on masonry grout that contains environmentally friendly materials such as fly ash, bentonite, ground granulated steel slag, water glass [1–3], hydraulic lime [4–6], clay-based grout [7], waste marble powder [8], perlite by-product [9] and metakaolin [6]. Studies on the utilisation of palm oil industrial by-products or palm oil clinker (POC) in masonry grout are, however, very limited [10] even though studies have demonstrated the highly pozzolanic reactive binding characteristics of POC [11, 12].

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POC has been proposed as a replacement for coarse aggregate to produce lightweight concrete and as supplementary cementitious material for enhancing the strength of concrete [12, 13]. Mohammed et al. [14] replaced the coarse aggregate in concrete with POC and investigated the behaviour of slabs and the shear and flexure strengths of beams constructed with POC-concrete. The POC concrete was approximately 18% lighter than the normal concrete and the structural behaviour of the POC-concrete slab was similar to that of normal concrete slabs [14]. The failure mode of the POC-concrete beams was similar to that of conventional concrete beams without shear reinforcement [15]. Even though the modulus of elasticity of the POC-concrete was smaller than that of normal concrete, the crack width of POC-concrete beams was within acceptable limits. The behaviour of POC-concrete beams is essentially ductile, giving ample warning of the imminence failure [16, 17]. Kana-dasan and Razak [18] studied self-consolidating concrete (SCC) using POC to replace the coarse aggregate at various replacement rate. The authors noticed a reduction in workability and determined a decrease in compressive and flexural strengths as the POC content increased. POC also negatively affect the elastic modulus of concrete [19]. Although there is a decrease in strength as the POC replacement rate increases [20], the strength of a 100% POC concrete is still approximately 70% of that of a conventional concrete [21]. Studies have also been conducted on using POC to evaluate engineering and sustainability performance of POC-concrete and the results indicate a positive outcome on cost reduction, reduced carbon emission, and energy-saving [22]. The use of POC as fine aggregate in concrete revealed no significant effect on the compressive strength of the concrete [23]. POC has been used as 100% replacement to the fine aggregates in a flowable mortar and although the strength is 86% of that of the control group, the resistivity is 64% greater than that of the control group. Moreover, life cycle assessment showed a reduction of ecotoxicity category [24].

Ibrahim et al. [25] studied the effect of POC in pervious concrete and reported that the density and compressive strength decreased as the replacement of rate of POC increased. In contrast, the permeability and porosity increased as the replacement rate increased. Considering the advantages of the pervious concrete properties, the authors suggested that a 25% replacement is optimum. A similar study [26] was conducted and the authors reported that the abrasion resistance of pervious concrete decreased with increase POC replacement rate. Despite some small negative effects; overall, the utilization of POC in concrete has been shown to give reasonably good results and that it is suitable for many applications [27].

Industrial by-products have been incorporated in mortar, normal and lightweight concrete, self-consolidating concrete, and fibre reinforced concrete. However, there has

not been a comprehensive investigation on the effects of incorporating POC in masonry grout and there have been very few studies on evaluation of its durability properties. Such an investigation is important due to its possible contribution to the global socio-economic-environmental issue. For instance, the lower-middle-income group of many communities around the world prefer to build structures using materials that are affordable, locally available, ease to handle and to construct with, durable and requires very low maintenance. Masonry fits this material category very well and POC-masonry grout can make masonry even more attractive as a building material. Nayaka et al. [10] focuses the study on comparative physical and mechanical performances of masonry grout mixes prepared of OPC and POCP such as compressive strength, flexural strength, split tensile strength, moduli of elasticity etc., comparative carbon footprint and cost efficiencies.

This study fills an important gap in knowledge by first comparing the binary pastes of OPC and POCP binders in terms of consistency and setting time, physical properties of aggregates in terms of water absorption, angularity number etc. in detail from material properties perspective. Current study tries to understand the durability performance of the masonry grouts prepared using OPC and POCP from both final durability performance of the casted masonry grout mixes and from the perspective of durability due to properties of materials used in the mixes.

Materials and methodology

Materials

Binders

The ordinary Portland cement (OPC) grade 42.5, which met the specifications of ASTM C150–14 [28], was used for the production of masonry grout. POC was procured from local palm oil industry and processed to powder (POCP) as described by Nayaka et al. [29]. The physical properties of the OPC and POCP are summarized in Table 1 while their gradations are shown in Fig. 1. The OPC and POCP particles were heavier than water and the POCP particles were

Table 1 Physical properties of binders

Properties	OPC	POCP
Specific gravity	3.15	2.55
Blain's specific surface Area (m ² /kg)	346	210
LOI	2.01	5.23
D50 (μm)	28.20	47.85

Fig. 1 Grain size distribution of OPC and POCP

about 20% lighter than that of the OPC because of their amount of micropores. The fineness of binders and mineral admixtures are vital as they can modify the aggregate-paste weakest part, i.e., the interfacial transition zone (ITZ) [30]. The amount of non-evaporable water and presence of unburnt carbon in binders, which could influence hydration, was measured based on LOI. The LOI of POCP was about 5.23%, which was 2.6 times more than that of OPC although it is within the limit of 12% in accordance with ASTM C618 – 14 [28].

The chemical composition of the OPC and POCP is presented in Table 2. The POCP was rich in silica, with a content of approximately 60%. The $\text{SiO}_2 + \text{Al}_2\text{O}_3 + \text{Fe}_2\text{O}_3$ (SAF) content of the POCP was approximately 71%. POCP can be classified as a Class-F pozzolan material, which contains a minimum amount of SAF of 70%, a maximum amount of SO_3 of 5%, and LOI of less than 10% [31]. Additionally, the soluble contents of silica and alumina in POCP were within the requirements of ASTM C618-14 [31]. These oxides are

Table 2 Chemical composition of binders

Oxides (%)	OPC	POCP
SiO_2 (S)	16.86	60.29
Al_2O_3 (A)	3.83	5.83
Fe_2O_3 (F)	3.76	4.71
ΣSAF		70.83
CaO	68.14	3.28
MgO	2.04	4.2
P_2O_5	0.04	3.78
K_2O	0.21	7.24
SO_3	4.84	0.31
TiO_2	0.14	0.1
MnO	0.11	0.12
Na_2O	0.03	0.2

the primary contributors to the strength and strength gain of the binder paste [32].

Normal Consistency and Time of Setting The normal consistency and time of setting were determined for a control paste that had only OPC and for a binary paste that had a combination of 70% OPC and 30% POCP. The normal consistency test determined the amount of water for the paste to become plastic; initial setting time established the time between the addition of water to the binder and loss of plasticity of the paste, and final setting time established the time between the addition of water to the binder and attainment of firmness of the paste to resist pressure. The results of the normal consistency tests and the time of setting for the two pastes are presented in Table 3. The OPC-POCP binary paste requires more water than the OPC paste to attain normal consistency, i.e., becoming plastic, can be due to lower density of POCP produces greater paste volume which in turn increases the plasticity of paste [33]. The binary paste attained normal consistency with approximately 22% more water than the OPC paste and the initial and final setting times were delayed by 40 and 72 min, respectively, from that obtained for the OPC paste. The observation of delaying setting times in binary mix might be due to the combined effect of lower cement content [34]. Nevertheless, the determined initial and final setting times for the binary paste were within the limits set forth by ASTM C191 – 14 [35].

Fine and coarse aggregates

Locally available natural sand (N-Sand) was used in the control masonry grout (GC), and in the grouts of phase I; manufactured sand (M-Sand) was used in the grouts of phase II GC(M). POC, shown in Fig. 2, was used as coarse aggregate in lieu of granite in the grouts of phase II. The gradations and particle size distributions of the fine and coarse aggregates are shown in Fig. 3. The size distribution of the aggregates is within the upper and lower limits specified in ASTM C404—14 [36].

The specific gravity, water absorption, moisture content, bulk density, and fineness modulus of the fine and coarse aggregates were also determined. The fine and coarse aggregates were tested according with the specification on ASTM C128-14 [37] and ASTM C136-14 [38], respectively. The

Table 3 Normal consistency and setting times of binders paste

Binders	Normal Consistency (%)	Initial Setting Time (Minutes)	Final Setting Time (Minutes)
OPC	32	70	335
70% OPC + 30% POCP	38	110	407

Fig. 2 Palm oil clinker

physical properties of aggregates are presented in Table 4. The fineness moduli of the natural and manufactured sands were 2.60 and 2.80, respectively. The fineness moduli of the granite and POC were 5.95 and 5.89, respectively.

Water and admixture

Potable tap water was used for the preparation of the masonry grout. The polycarboxylate-based superplasticizer Sika ViscoCrete–2055, which has high workability with

excellent slump retention, was used as an admixture. The dosage used was 1.6% by weight of the binder.

Methodology

A coarse masonry grout was selected in accordance with ASTM: C476 – 14 [39], which is preferable to minimize cost and shrinkage and suitable for infilling of masonry structural elements such as reinforced bond beams and masonry columns. A mix proportion of 1:03:02 (Cement: fine aggregate: coarse aggregate) was selected and the proportion of ingredients was calculated in accordance with ACI 211.2–98 [40]. The study was divided into three phases. In phase I, POCP was used to replace up to 50% of OPC.

In phase I, there were 6 mixes: 1 control mix with 100% OPC and 5 binary mixes with different combinations of POCP-OPC. In phase II, POC was used to replace the coarse aggregate up to 100% of the grout mix GB3 from phase I. In phase II there were 5 mixes: 1 control mix with 100% of granite and 4 binary mixes with different combinations of granite-POC. In phase III, the binary mix GBP2 from phase II was considered for further investigation and comparison to the control grout of phase I; both mixes were modified using a superplasticizer and consolidated using a mechanical vibrator. Since this was a comparative study, instead of the absorbent masonry mould described in ASTM C1019 – 14 [41], the specimens were cast in steel moulds. The test matrices for phases I, II, and III are presented in Table 5, Table 6, Table 7, respectively.

The compressive strengths of the grouts in Phases I and II were determined at the age of 3, 7, 28, and 56 days. The specimens in these phases were cured in water. The

Fig. 3 Grain size distribution of fine and coarse aggregates

Table 4 Physical properties of aggregates

Properties	Granite	POC	N-Sand	M-Sand
	(5- 10 mm)			
Specific gravity (SSD)	2.65	1.81	2.57	2.65
Water absorption (%)	0.2	1.09	1.98	1.65
Moisture content (%)	0.25	0.55	0.60	0.25
Fineness Modulus	5.95	5.89	2.60	2.80
Bulk density (kg/m ³)	1429	852	1604	1678
Aggregate crushing value (ACV) (%)	24.88	43.71	-	-
Aggregate impact value (%)	16.21	39.05		
Angularity Number	8	11	-	-

determined at 28 days. The sulphate resistance of the grouts was studied by using a 5% concentration of MgSO₄ solution at 28, 90, and 120 days. The electrical resistivity of the grouts was determined at 7, 14, 28, and 56 days. Some specimens were cured in water for 72 h to study drying shrinkage, and shrinkage values were obtained for a period of up to 180 days. The details of the tests and reference standards followed in this study is summarized in Table 8. To provide an extensive insight to durability, tests such as water absorption, sulphate resistance, electrical resistance, drying shrinkage are performed. Mechanical test results such as those of compression test in terms of structural efficiency, UPV test [10] and fresh properties' evaluation tests such as slump cone test are

Table 5 Masonry grout mix proportions for Phase I

Mix ID	POCP Replacement (%)	OPC (kg/m ³)	POCP (kg/m ³)	N-Sand (kg/m ³)	Granite (kg/m ³)	w/b ratio
GC	0	230	0	692	487	0.84
GB1	10	207	23	692	487	0.84
GB2	20	184	46	692	487	0.85
GB3	30	161	69	692	487	0.85
GB4	40	138	92	692	487	0.86
GB5	50	115	115	692	487	0.86

Table 6 Masonry grout mix proportions for Phase II

Mix ID	Replacement (%)		OPC (kg/m ³)	POCP (kg/m ³)	M-Sand (kg/m ³)	Granite (kg/m ³)	POC (kg/m ³)	w/b ratio
	POCP	POC						
GC(M)	-	-	230	00	762	487	-	0.82
GB3	30	000	161	69	692	487	-	0.85
GBP1	30	025	161	69	762	365	072	0.86
GBP2	30	050	161	69	762	243	144	0.86
GBP3	30	075	161	69	762	122	215	0.88
GBP4	30	100	161	69	762	000	287	0.92

Table 7 Masonry grout mixtures proportions for phase III

Mix ID	Replacement (%)		OPC (kg/m ³)	POCP (kg/m ³)	M-Sand (kg/m ³)	Granite (kg/m ³)	POC (kg/m ³)	w/b	SP (%)
	POCP	POC							
GC	0	0	230	0	692	487	0	0.6	1.6
GBP	30	50	161	69	762	243	144	0.6	1.6

compressive strengths of the grouts in phase III were evaluated at 7, 28, 56, and 90 days. The specimens in phase III were cured in water and air.

Durability properties were determined for the control and the GBP2 i.e. GBP mixes only. The water absorption of the grouts by immersion and capillary action were

explained in paper briefly, only to accompany the durability studies performed. Only then the inter-relation between a structure's durability performance and mechanical performance can be explained properly.

Table 8 Details of tests, reference standards and properties determined for masonry grout

S. No	Test	Reference Standard	No. of Samples	Type and Size of Specimens (mm)	Properties
1	Preparation of specimens and curing	ASTM C1019-14 ASTM C511 -14	-	-	Fresh State
2	Slump cone test	ASTM C143—14			
	Hardened density	ASTM C642-14	3	Cube 100*100*100	Mechanical
3	Compressive strength	BS EN 12,390—3: 2009			
4	Ultrasonic pulse velocity	ASTM C597—14			
8	Bulk and apparent density, water absorption by immersion	ASTM C642-14	3	Cube 100*100*100	Durability
9	Water absorption by capillary	ASTM C1585-14	3		
10	Resistance to sulphate solution	ASTM C157—14	3		
11	Electrical resistivity	ASTM 1202—14	3	Prism 70*70*300	
12	Drying shrinkage	ASTM C596—14	5	Prism 70*70*300	Time-Dependent

To provide an extensive insight to durability, tests such as water absorption, sulphate resistance, electrical resistance, drying shrinkage are performed. Fresh properties' evaluation tests such as slump cone test are explained in paper briefly, only to accompany the durability studies performed. Mechanical test results such as those of compression test in terms of structural efficiency, UPV test [10] and then the inter-relation between a structure's durability performance and mechanical performance were also thoroughly explained using microstructural behavior by conducting scanning electronic microscope (SEM), X-Rays Diffraction (XRD), Energy Dispersive Analysis (EDX) and Thermo-Gravimetric Analysis (TGA) on the optimum samples of masonry grout and conventional grout.

Results and discussions

Microstructure characteristics

Scanning Electron Microscope (SEM) study

The SEM images of the control and binary, i.e., 30% POCP + 70% OPC + 50% POC (GBP) grouts cured in water (GCW and GBPW) and cured in magnesium sulphate (GCS and GBPS) for 28 days are shown in Fig. 4.

The higher silica and calcium correspond to the binder and POC aggregate. In the grouts cured in water (images GCW and GBPW), the tricalcium silicate hydrates with water to form a dense acicular calcium silicate hydrate (C-S-H) and hexagonal calcium hydroxide (CH) crystal along with the formation of capillary pores and voids. The sulphate-cured grouts (images GCS and GBPS) had alumina

silicate (AS), calcium silicate (CS) and silica, besides the presence of gypsum and brucite.

The analysis of the grouts cured in the magnesium sulfate solution showed that the binary grout paste was denser than that of the control grout because of the irregular shape and rough surface texture of the POC, which induces stronger bond between the aggregates and paste [42]. However, GBPW grout in Fig. 4, was observed the loss of moisture results in shrinkage cracks. The loss of moisture due to moisture absorption at the interface with underlying materials [43].

The GBP grout had large crystals of CH and mono-sulphoaluminate [44]. However, only a limited amount of acicular C-S-H was observed due to delay in the reaction of the di-calcium silicates. Also, spherical dense and partially dense ettringite were observed in the binary grout cured in the sulphate solution due to the presence of SO_3 , which acts like gypsum and reacts with the CS crystals. There were, however, no calcium hydroxide plates in both sulphate solution cured GBP grouts although the ratio of MgO to Fe_2O_3 of 1.06 is well within the limit to avoid expansion [45].

An X-ray diffraction (XRD) analysis was conducted for the control and binary grouts cured in air (AC), water (WC) and sulphate (SC) for 28 and 120 days. The XRD patterns are shown in Fig. 5 at the end of the 28-day curing.

High portlandite (CH) peaks were observed for the GC grouts cured in water, which indicates the presence of excess moisture that caused delays in the conversion of the silicates to C-S-H and CH for strength development. In contrast, the CH peaks were smaller for the grouts cured in air and sulphate. For the GB grouts, the air-cured specimen had high peaks of portlandite; parallel cristobalite peaks were also observed. Similar pattern has been observed for fly ash and

Fig. 4 SEM images of masonry grouts **Fig. 5** XRD patterns of masonry grout mixes at 28 days curing

rice husk ash [46]. A high cristobalite peak was observed for the GB grout cured in the sulphate solution, which could be due to the presence of silica in the POCP and the reaction with the sulphate solution to form gypsum.

Energy Dispersive X-Ray Analysis (EDX)

In case of Masonry grouts prepared for phase I, replacement of OPC with POCP was performed at incremental stages whereas Masonry grouts for phase II and phase III it was accompanied with replacement of granite with POC.

In case of GBP grouts, cured in the sulphate solution shows high cristobalite peak, as seen in XRD analysis. The reason behind this lies in microstructure characteristics of POCP binder and POC aggregates. Energy Dispersive X-Ray Analysis (EDX) (Figs. 6, 7) shows the presence of high content of Silica at the region of interest in POCP binder and POC aggregate respectively. POC aggregate also shows immense macro and micro voids. When GBP grouts were cured in water this influences the moisture absorption. Both POCP and POC contain high amounts of reactive silica content which is responsible for their respective pozzolanic nature. High peaks of reactive silica translate to formation of the initial C-S-H process being delayed. There will be delayed strength gain due to secondary pozzolanic action but the initial hydration process yields less initial concentration of C-S-H and C-H compounds. The pozzolanic activity is further investigated using thermogravimetric analysis.

Thermogravimetric Analysis (TGA)

Figure 8 depicts the TGA mass loss vs temperature curves of control mix and binary mix with 30% replacement of OPC with POCP (GBP mix) at a curing age of 28 days. This analysis provides a deeper understanding of the pozzolanic nature of the materials. At a temperature range of 100–180° C primary dehydration takes place and the formation of ettringite can be observed (Celik et al., 2014; Fares et al.,

2010). The dehydration of hydrated calcium monocarboaluminate initiates at a temp around 200° C and around 300° C it may get completely removed along with free water and water bound in C-S-H gel (Fares et al., 2010); (Cizer et al., 2010).

Dehydroxylation of CH commences at a temp range of 380–500° C which causes mass loss in control mixes (Carmona-Quiroga & Blanco-Varela, 2015; Uchima et al., 2015). This phenomenon was found to be comparatively lower in binary mixes where 30% of the OPC was replaced with POCP for the same temperature range. The reasoning behind can be attributed to the formation of less amount of CH in binary mixes when compared to control mixes due to weaker primary hydration process in binary mix. This effect is even more dominant for binary mixes where 30% of the OPC was replaced with POCP because of the dilution effect (Fares et al., 2010; Karim, Hashim, & Razak, 2016).

For XRD data of binary mix with 30% POCP replacement, cured for 28 days, there was a significant reduction of CH. This is due to the initiation of delayed pozzolanic reaction process in case of binary mix and formation of more C-S-H gel. The TGA curves also show that at temperature range between 400–520 °C the difference in mass loss due to dehydroxylation of CH for control mix was 2.00%, whereas the same for binary mix was found to be 1.25%. This evidently concludes that prolonged curing enables the pozzolanic process and formation of more C-S-H gel in cases of

Fig. 6 POCP binder and structure with EDX

Fig. 7 POC aggregate and structure with EDX

Fig. 8 TGA mass loss vs temperature curves

binary mix because of the higher silica content and lower CaO content in constituent materials i.e., POCP binder and POC aggregates.

Workability and fresh state density

The total water to total binder (cementitious materials) ratio (w/b), the POCP and POC replacement levels, in phase I and II, the fresh state density obtained to meet the required consistency (204 mm – 280 mm) for the grouts are shown in Fig. 9.

In phase I, the w/b ratio for the control grout (GC) was 0.84, while binary grouts with POCP (GB1 through

GB5) had w/b ratios between 0.84 and 0.86. The w/b ratio increased with increasing POCP replacement level, which could be attributed to the presence of unburnt carbon, moderately coarse texture, irregular particle shape and sharp edges of the POPC [47]. On the other hand, the specific gravity of the grouts decreased with increasing POCP levels. The decrease in specific gravity was expected since POPC has lower specific gravity than OPC; the decrease should have been linear since the replacement rate used herein was linear.

As observed for the binary grouts in phase I, the binary grouts in phase II experienced an increase in water demand and a decrease in density as the amount of POC increased.

Fig. 9 Fresh state density and w/b ratio as a function of POCP and POC levels

To attain the required consistency, the POC-incorporated grout mixes required up to 9% more water than that required in the control mix due to the irregular shape, porosity, and higher water absorption of POC [48].

The grouts with POC had greater slump than that of the control grout due to the lower density [49] and better-rolling capability of the POC [50]. The binary grout with 30% POCP and 50% POC was considered an ideal mix based on the fresh state density and minimal presence of voids according to the ASTM C476-14 [39] requirement of compressive strength.

In phase – III, a superplasticizer (SP) was used to reduce the water demand and increase the workability, due to greater surface and dispersion properties of the SP [51], of the binary grout with 30% POCP and 50% POC and of the control grout. The fresh properties of this SP-modified binary grout and the SP-modified control grout are presented in Table 9. To minimize the variables and eliminating confounding results, the same w/b ratio and the same SP percentage were used in both mixes.

Mechanical properties of the grouts

Structural efficiency

The definition of structural efficiency (SE) was introduced for the comparison of control specimens with GBP samples [49]. The SE value was determined by compressive strength over the hardened density of masonry grout mixes of water and air-cured for 28 days as shown in Fig. 10. In Phase I, SE decreased from 0.0107 to 0.0066 in order to replace cement by POCP up to 50 percent. When the substitution of POCP increased, the SE decreased due to weak bonding between cement and POCP molecules [12]. POCP has been shown to be a pozzolanic material because the dilution effects did not produce good bonding at higher volume substitution and because of the delay in the pozzolanic reaction, or POCP acts as filler material.

The air-cured specimens obtained higher SE values than the water-cured specimen due to the higher water content used to achieve the necessary slump value, which caused early hydration reactions. In phase II, a higher SE was found for the control mix in which M-sand was used to substitute N-sand in order to prevent the loss of natural resources. SE decreased from 0.0106 to 0.006 when POCP was used as a binder to replace 30% of cement and M-sand was used as a

Fig. 10 Structural efficiencies of masonry grouts [10]

sustainable material. In comparison, POC substitution for granite varied from 0 to 100 per cent. The decrease in SE was attributed to the fact that the porous POC aggregates caused the specimens to fail, while in the case of granite the failure happened in the interfacial region between the paste and the aggregates. In phase III, control (GC) and ideal mix grout (GPP) had SE value of 0.0169 and 0.0139 respectively when water cured. The air-cured specimens had SE values of 0.0174 for control and 0.0147 for ideal grout mix. There was a 37 – 46% enhancement of SE from phase II to phase III due to the influence of admixture used to improve workability and reduce water demand. Another reason is the use of mechanical vibration on masonry grout mixes.

Ultrasonic Pulse Velocity (UPV)

UPV presents the pulse rate transfer through solid media, and this rate is indirectly related to voids present in the specimen. The UPV method was used to determine the denseness of the control grout and the binary grout with POC aggregate. The UPV measurements on the control and binary grouts are shown in Fig. 11; the measured UPV in the control and binary specimens cured in air for 28 days were 4.42 km/s and 3.81 km/s, respectively. The UPV values of air-cured specimens were greater than those of water-cured specimens at all durations of curing. As aforementioned,

Table 9 Fresh state properties of control and ideal grout mixes

Mix ID	Replacement (%)		w/b ratio	Fresh State Density (kg/m ³)	Slump (mm)	S.P (%)
	POCP	POC				
GC	0	0	0.6	2470	228	1.6
GBP	30	50	0.6	2382	216	1.6

Fig. 11 UPV values of GC and GBP mixes with different curing regimes [10]

Fig. 12 Compressive strength and UPV relationship for phase III

the air-cured specimens had slightly higher compressive strength than that of the water-cured specimens (because curing stopped at 90 days, a longer curing period would most likely reverse the trend).

The results indicate that the presence of moisture, honeycombed POC particles, and type of aggregates influenced the UPV values. According to Malhotra [52] excellent UPV values are greater than 4.5 km/s while good UPV values are between 3.5 and 4.5 km/s. Most measurements fell in the good category, including that for the control grout. For the binary grout, the increase in the UPV values from 28 to 90 days of curing was significant and was due to the increase

in the POCP pozzolanic activity and filling of the voids in the POC aggregate. Thus, in the long term, replacing cement by POCP and granite by POC will have no negative effects on the quality of the grout.

The relationship between the compressive strength and the UPV obtained herein is shown in Fig. 12. The high R^2 value indicates an excellent fit of the equation to the experimental data. Using equation 1, the compressive strength of grouts can be obtained using the non-destructive UPV method.

Durability performance of masonry grout

The durability performance of the grouts was determined by testing samples of the control grout and binary grout with POC aggregate.

Densities and water absorption

The results of the densities and water absorption tests are shown in Table 10. The densities of the binary grout were lower than that of the control mixes due to the lower specific gravity of POCP and POC while the water absorption of the binary grouts was higher because of the porosity of the POC and POCP, which weakens the interfacial transition zone. As aforementioned, the rough texture and porous microstructure of the POCP and POC particles [53] compromise the compactness of the grout mix. The higher water absorption of the POC was expected since lightweight aggregates generally have higher water absorption [54]. Overall, however, all masonry grout specimens in this study are well within the limit of concrete quality specified by Neville [55].

Initial rate of water absorption

The rates of water absorption of the grout mixes were also determined by capillary testing, and the values obtained were different from that of the immersion testing due to the internal microstructures and interfacial transitional zone.

The capillary test determined the initial absorption and the secondary absorption until a constant mass, i.e., a mass variation of less than 0.5%, was obtained for two consecutive

Table 10 Densities and water absorption properties of masonry grout

Curing Age (Days)	Mix ID	Water absorption (%) (24 h)	Water absorption (%) (48 h)	Density (Dry) (kg/m ³)	Density (SSD) (kg/m ³)	Apparent Density (Mg/m ³)
28	GC	4.62	4.68	2304	2417	2448
	GBP	4.71	4.79	2146	2250	2301
90	GC	4.90	5.08	2287	2431	2485
	GBP	5.93	6.01	2140	2251	2298

periods of weighing. The results of the test are presented in Fig. 13.

The initial rate of water absorption (IRA) by capillary action of the binary grout was slightly higher than that of the control grout because of poor interlocking bonds within the POCP and POC aggregate, which causes high porosity and increases absorption. The secondary rate of absorption of the GBP grout was, however, lower as compared to that of the control. Generally, utilization of POCP tends to improve the filling effect of grout which improves the rate of absorption at a later stage [12]. The replacement of gravel with 50% POC resulted in greater compactness due to rough surface texture, which leads to lower secondary water absorption after initial absorption at the transitional zone can be observed in Fig. 13, which agrees with previous studies [56]. Furthermore, lower paste strength results in higher permeability [55].

Resistance to magnesium sulphate solution

Sulphate attack is a chemical breakdown mechanism in which sulphate ions attack components of the binder paste. In this study, 5% of magnesium sulphate solution was used. Magnesium sulphate is an alkaline solution capable of reacting with the components of the masonry grout. POCP contains lower quantity of CaO when compared to conventional OPC. The addition of POCP, a pozzolanic admixture, reduced the C_3A content of the paste; therefore, there was a possibility that the paste was susceptible to sulphate attack. The changes in compressive strength and weight of the grouts are shown in Fig. 14 and Fig. 15, respectively, when the grouts are exposed to magnesium sulphate solution for a period of 28, 90, and 120 days. In Fig. 14, the change

Fig. 13 Initial and secondary rate of absorption of masonry grouts

Fig. 14 Change in compressive strength due to sulphate attack

in compressive strength is shown relative to compressive strength obtained when the grouts were cured in water. In Fig. 15, the change in weight is shown relative to the weight determined when the grouts were cured in magnesium sulfate.

After 28 days of curing, the compressive strength of the control and binary grouts improved approximately 2 to 5%. The improvement could be because of the chemical reaction (combination) between the sulphate ions and C-S-H and because the sulphate ion crystals occupied the empty spaces and continue to form chemical structures that cause the paste to expand and crack. After 90 days of curing, a reduction of approximately 0.5% and 2% in compressive strength were observed for the control and binary grouts, respectively, which indicates that ettringite compounds were formed. The strength was decreased due to the highly porous nature of POC with immense macro and micro-voids as was observed in EDX analysis

Fig. 15 Change in weight due to sulphate attack

in Fig. 7. Moreover, this increased void ratio in the grout to allow quick crack propagation as well as failing rapidly under applied load [57]. The changes in strength were more significant in unexposed dry-cured samples than in exposed samples. This trend was continued for 120 days exposure, there was a reduction in compressive strength of the control grout more than that of the binary grout, which could be attributed to the delayed pozzolanic reaction of POCP binder and the gypsum in the GBP mix controlling the sulphate ions to further form brucite and ettringite. An increase in C_3S content and reduction of C_3A content generate large amounts of CH that reacts directly with sulphate ions and are seen as a rust coating on the aggregates in the internal structure of the samples. The pores and bonding between the aggregates and the binder paste were visible for the grouts cured in air, water and sulphate solution at 120 days. This leads to the formation of gypsum and brucite, in which the portlandite peaks appear in the sulphate exposed samples.

After 28 days of curing, the weight of the control and GBP grouts increased by approximately 0 to 1.67%. The sulphate ion in the space in control (GC) paste does not dry completely and hence there is no significant reduction in mass, although almost the same mass was observed in the exposed binary (GBP) mix.

Overall, this causes a reduction in the cohesiveness of the hydrated cement. Thus, sulphate attack results in the decrease of the mechanical properties and durability of the grout by changing the chemical nature of binder paste [58]. The sulphate resistance of the binary grout compared to that of the control grout is noteworthy at 120 days of exposure (can be seen in Fig. 18). This can be attributed to

the secondary pozzolanic action which translates into reduction in pore and void size and better filler effect of the finer particles.

Grouts being composite material, the durability properties depend a lot on the micro, macro voids, porosity, microstructure of hydrated matrix, inter-facial transition zone and packing of constituent materials. The ITZ zone, being the weak link usually shows the symptoms of durability loss upon sulphate attack. Immersion in sulphate facilitates ettringite formation near this ITZ zone which in turn facilitates higher porosity, thus higher permeability. Though binary paste contains highly porous POC aggregates and thus allows for strength reduction in early days of curing, delayed pozzolanic reaction of POCP controls the formation of ettringite in binary mix under prolonged exposure. Thus, the ITZ zone in case binary paste undergoes less deterioration under prolonged sulphate exposure when compared to that of the control mix.

The dense matrix formation at microstructure level assures necessary bonding between binder and aggregates which accounts for both compressive strength and the durability performance of the grout. The reasoning behind lies at the microstructure characteristics of the materials, used to compose the matrix. The TGA curve analysis along with XRD and EDX analysis performed on POCP binder and POC aggregate explains the pozzolanic action of the binary mix along with the behavioral differences when cured in air, water or sulphate. The XRD patterns for the control and GBP grouts are shown in Fig. 16 and Fig. 17, respectively, for the grouts cured in air and in a sulphate solution for 120 days. For the control grouts cured in air and water, the peaks of portlandite were converted to calcite and C-S-H

Fig. 16 XRD patterns of Control mix at 120 days curing

Fig. 17 XRD patterns of Binary mix at 120 days curing

Fig. 18 shows the microscopic images of grout mixes and the interfacial transition zone

gel formation, and alite and belite compounds formed at scattering angles of 18° and 21°. The alite and belite peaks observed in the binary paste were lower than that of the control paste. In the binary paste, ettringite and brucite formations were observed and are due to the sulphate solution penetration and reaction. The conversion of portlandite to

C-S-H was slow in the binary grout cured in water but it was fast in the control grout cured in water. The sulphate treated binary grouts had smaller portlandite peak than that of the control grout cured in the sulphate solution. The ettringite peaks were higher in the control paste than that in the binary paste, indicating the delay in pozzolanic reaction.

Evidently, the lower CaO content and high silica content in POC aggregates and POC binder-based grout when compared to normal OPC and granite-based grout, facilitates controlled formation of C3A, more C–S–H formation due to the delayed pozzolanic reaction and thus allows for a denser microstructure matrix. This inhibits the detrimental ingress of sulphate ions and controls the formation of ettringite, thus showing better durability performance when compared to control mix (Fig. 18).

Electrical resistivity

The electrical resistivity of a specimen gives an indication of the ability of the specimen to resist chloride ion penetration. The electrical resistivity of the control and binary grouts cured in water for 7, 28, 56 and 90 days were determined according to AASHTO [59]. The results are presented in Fig. 19 and all values obtained, except for those for the grouts cured for 7 days, are well above the threshold value of 12 k Ω cm [60]. The measurements show that the chloride ion penetrability of the control and binary grouts are below the moderate penetrability category at 28 days exposure and below the low penetrability at 90 days exposure.

The results obtained herein agree with those obtained by Kanadasan and Razak [20], who showed that the binary grout resistivity was 16% higher than that of the control grout. A probable reason for the high electrical resistivity is the porous microstructure of the POC aggregates as well as the higher moisture content [61]. Aggregate sizes, however,

were less than 10 mm and therefore formed less interfacial transition zones (ITZ), as shown in Fig. 19.

Drying shrinkage

Drying shrinkage measurements were made for up to 180 days and the results are presented in Fig. 20. The POC powder and aggregate significantly affected the drying shrinkage of the grout. A possible explanation may be the higher paste to aggregate ratio in the binary grout. In addition, lightweight aggregates normally cause higher drying

Fig. 20 Drying shrinkage of the grouts

Fig. 19 Electrical resistivity of grout mixes

shrinkage [55]. The retained water in the POC aggregate evaporates over time, which leads to increased shrinkage [62, 63].

Although the binary grout experienced higher drying shrinkage than that of the control grout, the measured value was within the appropriate limit of 750 microns suggested in ASTM C596 – 01 [64].

Conclusions

Experiments were conducted to determine the effect of substituting Portland cement and coarse aggregate in masonry grout by palm oil industrial by-products at different replacement rate. The hardened state density, compressive strength, and durability values are indicative of the satisfactory performance of the eco-friendly masonry grout developed herein. The following are the broad conclusions from this experiment program:

- The binary grout with POC aggregates requires more water to meet practical workability requirements; nevertheless, the use of superplasticizers can reduce the water demand and enhance the fresh state properties of the grout.
- Although a grout with 40% POCP met the minimum required compressive strength, a grout with 30% POCP is suggested as the ideal grout when considering other factors.
- The TGA curve analysis along with XRD and EDX analysis performed on POCP binder and POC aggregate explains the pozzolanic action of the binary mix along with the behavioral differences when cured in air, water or sulphate.
- The use of POC as coarse aggregate reduces the compressive strength of the grout. A 50% replacement of coarse aggregate by POC, however, still can be used since a grout with such an amount of POC meets the minimum required compressive strength.
- The UPV value obtained for the binary grout with POC aggregates indicates that the grouts are of “good” quality.
- Although water absorption and initial rate of water absorption properties of the binary grout with POC aggregates are slightly higher than that of a conventional grout mix, the values are within the standard limit.
- The binary grout with POC aggregates has a high resistance to sulphate attack and a moderate level of chloride ion penetrability.
- The drying shrinkage of the binary grout with POC was within the appropriate limit.
- Overall, the durability performance of the binary grout with POC aggregate is similar to that of a conventional

grout with added value in terms of materials cost and carbon footprint.

- Palm oil industrial by-products, more specifically powder and clinker, can be used in masonry grout without detrimental effects.

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Declarations

Conflicts of interest statement On behalf of all authors, the corresponding author states that there is no conflict of interest.

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