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Study of physico-chemical and mechanical characteristics of consolidated and unconsolidated cemented paste backfills

1. Introduction

Cemented paste backfill CPB has seen increasing use by mines worldwide, mainly due to its significant cost advantages and the potential for placing the full plant sulphidic tailings in underground stopes (Landriault 2001; Yilmaz 2011). CPB is usually prepared by mixing the tailings, binder, and mixing water to form a composite construction material. After its hardening, the CPB material acts as secondary ground support. The physico-chemical and mineralogical properties of CPB ingredients greatly affect both strength and stability performance (Archibald et al. 1999; Belem et al. 2000; Benzaazoua et al. 2002; 2004; Kesimal et al. 2004; Klein, Simon 2006; Ouellet 2007; Fall, Samb 2008; Yilmaz 2010; Fall et al. 2010; Yilmaz et al. 2011; Cihangir et al. 2012). Uniaxial compressive strength (UCS) is the key index parameter most often used for CPB stability design (Belem, Benzaazoua 2008). The UCS requirements are closely related to the assortment of roles undertaken by CPB. Overall, the target UCS values for CPB vary from 0.25 to 4.35 MPa for a wide range of applications varying from tailings disposal to roof support underground (Hassani, Bois 1992).

CPB design is often based on the evaluated properties of laboratory-prepared backfill material (plastic mould samples). However, experience indicates that it is not easy to ensure that all the physical and mechanical conditions prevailing underground are observed when the CPB sample is prepared, cured, and tested in the laboratory (Servant 2001). Further,

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the mechanical and curing properties of *in situ* CPB material vary noticeably, depending on the methods of preparation, placement, and conditions of the mine environment (le Roux et al. 2005). Previous studies indicate that the UCS magnitudes of *in situ* CPB material are often 2 to 4 times higher than those obtained from laboratory-prepared samples of equivalent batch mix and curing time (Belem et al. 2002; Cayouette 2003). This discrepancy may be attributed to the fact that the hardening of CPB material cured under an effective field stress increases the rate of strength development and the ultimate strength, as reported lately by a number of authors (Belem et al. 2006; Fourie et al. 2006; Grabinsky, Simms 2006; Helinski et al. 2007; Bussière 2007; Yilmaz et al. 2009, 2012; Ercikdi et al. 2010). Additionally, Revell (2004) has shown that scale effects have an impact on CPB strength gain. To the knowledge of the authors of this text, there is no suitable laboratory apparatus or standardized test procedure for CPB that mimics the field mixing, placement, and curing conditions on a laboratory scale. Moreover, only a few studies showed that consolidation both under self-weight and time-dependent surcharge loading could have a notable effect on the overall performance and quality of CPB materials (le Roux et al. 2005; Yilmaz et al. 2008). In existing documentation, there was a lack of information about the characteristics of CPB cured under applied pressure. Accordingly, an understanding of the effect of curing under stress on CPB performance is required for a more reliable and better quality of backfill design.

The main objective of this study was to evaluate the effects of binder content and curing time on CPB hydromechanical performance, as well as to observe the resulting geochemical properties. To achieve this objective, a new laboratory tool named CUAPS (curing under applied pressure system) was used which allowed for the simulation of the *in situ* placement and curing conditions of CPB materials (Benzaazoua et al. 2006; Yilmaz et al. 2010). A comparative analysis of the above-mentioned properties of the CPB cured in both plastic moulds and under effective stress was performed using three different binder contents (3, 4.5, and 7 wt%) and curing times (7, 14, and 28 days).

2. Material and Method

2.1. Characteristics of paste backfill ingredients

Mill Tailings: the tailings sample used in this study was taken from a Canadian gold mine located in the western part of the province of Quebec. The sample's particle size distribution (PSD) was determined using a Malvern Mastersizer laser diffraction-type particle size analyser (Fig. 1), and fit within a typical range of PSD curves of 11 mine tailings sampled from the underground hard rock mines located in the provinces of Quebec and Ontario, Canada (Ouellet 2007).

Table 1 also shows the main physico-chemical characteristics of the tailings sample. It can be observed that the fines content (less than 20 μm) is 44 wt% which corresponds

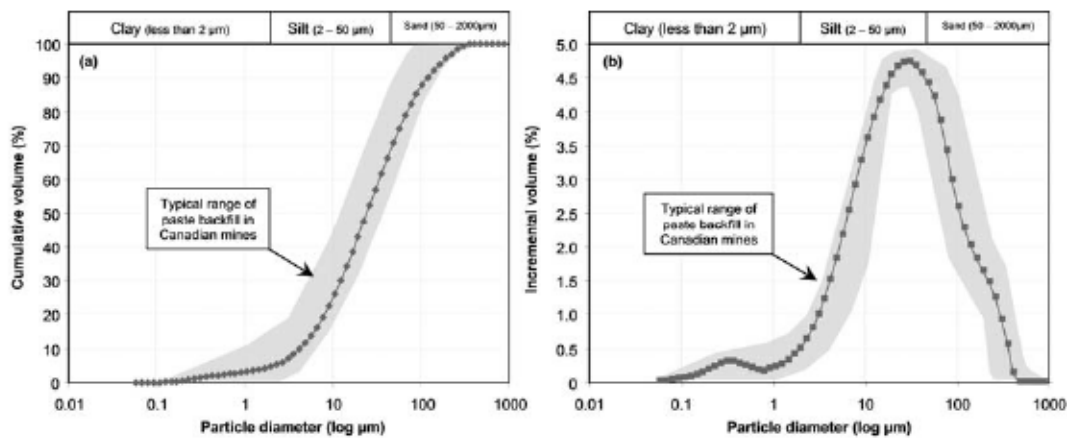


Fig. 1. Particle size distribution curves of the tailings sample used
a) cumulative, b) incremental

Rys. 1. Krzywe składu ziarnowego próbek użytych odpadów
a) skumulowana, b) przyrostowa

TABLE 1

Main physico-chemical characteristics of the tailings sample used

TABELA 1

Charakterystyka podstawowych właściwości fizyko-chemicznych próbek użytych odpadów

Parameter	S_s (m^2/kg)	G_s	D_{10} (μm)	Fines ($<20 \mu m$)	C_u (wt%)	C_c (wt%)	Al (wt%)	Ca (wt%)	Fe (wt%)	S_{total} (wt%)
Tailings	2170	3.7	4.26	44 wt%	8	1.1	2.8	0.57	27.4	20.6

S_s – specific surface,

D_{10} – effective particle diameter.

to medium size tailings, according to Landriault (2001). Tailings are well-graded with a coefficient of uniformity C_u ($= D_{60}/D_{10}$) of 8 and with a coefficient of curvature C_c ($= D_{302}/D_{60} \cdot D_{10}$) of 1.1. Based on the Unified Soil Classification System (USCS), the studied mine tailings are classified as low plasticity silt (ML). The specific gravity G_s of the tailings sample that was measured with a Micromeritics® Accupyc 1330 helium gas pycnometer is 3.7, reflecting a relatively high content of sulphides.

The chemical composition analyses of metals and total sulphur (S_{total}) were conducted on the tailings sample using a Perkin-Elmer® Model Optima 3100 RL ICP-AES (inductively coupled plasma-atomic emission spectroscope) after complete acid digestion by HNO_3 , Br_2 , HF , and HCl . Dilute HCl was used to extract sulfates, and the solution obtained was analyzed by ICP-AES. From the chemical composition analysis carried out on the tailings sample,

it can be observed that the contents of iron Fe, sulphur S, aluminium Al, and calcium Ca were 27.4 wt%, 20.6 wt%, 2.8 wt%, and 0.57 wt%, respectively.

Binding agent: the binder used for the preparation of CPB samples is a blend of ordinary Portland cement or Type I (PCI) and ground, granulated blast furnace slag (Slag). The blending ratio of PCI to Slag is 20/80. Three different binder proportions of 3, 4.5, and 7 wt% were chosen to evaluate the effect of curing conditions on the UCS performance of CPB samples. Proportions were calculated on the basis of weight relative to the total dry tailings. The chemical composition analyses were also conducted on the studied binder samples using ICP-AES method, as explained in the previous section. Table 2 shows the main chemical composition and physical properties of binders.

Mixing water: two types of water namely as-received tailings, pore water, and tap mixing water were chemically analyzed using ICP-AES method. Table 3 summarizes the results of the chemical and geochemical analyses. It can be observed that the pore water is highly aggressive in terms of sulphate content (SO_4^{2-} content around 4880 ppm) but also contains calcium (Ca content about 560 ppm). While Ca comes from the lime added during ore treatment, sulphates can come from the cyanide elimination process (tailings are treated before backfilling) and those produced from sulphide reactivity (Benzaazoua et al. 2004). Additionally, the values in terms of pH, Eh (redox potential), and EC (electrical conductivity) of pore and tap waters were 9.41, 0.147 volt, and 7.42 mS/cm; and 7.82, 0.431 volt, and

TABLE 2

Chemical composition and some physical properties of the binders used

TABELA 2

Skład chemiczny i właściwości fizyczne niektórych używanych spoiw

Parameter	S_s (m ² /kg)	G_s	Al ₂ O ₃ (wt%)	CaO (wt%)	Fe ₂ O ₃ (wt%)	K ₂ O (wt%)	MgO (wt%)	Na ₂ O (wt%)	SO ₃ (wt%)	SiO ₂ (wt%)
PCI	1 580	3.1	4.86	65.76	2.44	0.83	2.21	2.11	3.67	19.51
Slag	3 540	2.8	10.24	31.41	0.55	0.51	11.3	2.01	3.27	36.22
PCI-Slag	2 840	2.9	4.26	42.82	0.64	0.55	6.19	2.03	3.35	30.91

TABLE 3

Chemical and geochemical analysis results of pore and tap waters used as mixing water

TABELA 3

Chemiczne i geochemiczne wyniki analizy porów i odcieków wody wykorzystywanych jako woda zasobowa

Parameter/ element	EC (mS/cm)	pH	Eh (v)	SO ₄ ²⁻ (ppm)	Si (ppm)	Mg (ppm)	Al (wt%)	Ca (ppm)	Cu (ppm)	Fe (wt%)
Pore water	7.42	9.4	0.147	4 883	0.891	1.83	0.212	559	0.286	0.011
Tap water	0.274	7.8	0.431	138	0.901	2.27	0.01	40.9	0.835	0.066

0.274 mS/cm, respectively. The relatively high value of EC within the pore water of as-received tailings indicates the presence of many dissolved conducting ions.

2.2. Experimental program

Experimental apparatus: The CUAPS (Curing Under Applied Pressure System) was developed for simulating field placement and curing conditions of laboratory-prepared CPB materials. The CUAPS apparatus has been described previously by Benzaazoua et al. (2006) and Yilmaz et al. (2010). The CUAPS allows the operator to estimate the more realistic UCS values of the CPB materials cured under constant or variable pressure up to 400 kPa. The operating principle of CUAPS is that CPB is one-dimensionally consolidated by effective stress over time. A Polycarbonate Perspex tube in the CUAPS apparatus forms a component highly resistant to volume change after a series of stresses are applied. Following an increase in effective stress, excess pore water within CPB is expelled through a drainage port located at the bottom of the CUAPS. During curing, drainage water is collected in order to calculate the drainage rate and to analyze the chemical composition for the different recipes.

Backfill sample preparation: the CPB ingredients (i.e. tailings, cement, and water) were thoroughly mixed using a double spiral mixer for about 7 minutes. Right after mixing, the measured slump of all mixtures was set to approximately 7 inches by adding more or less mixing water. The slump test was conducted according to the ASTM C143 standard. The initial water content w_g for all CPB samples was kept constant at 28.2 wt%.

Table 4 lists the initial bulk properties of each CPB batch mix. These values represent volumetric binder content B_v , solids mass concentration C_w , volumetric solids concentration C_v , water-to-cement ratio w/c , specific gravity G_s , void ratio e , wet density ρ , dry density ρ_d , and volumetric water content θ . It can be observed from Table 4 that the w/c ratios decrease from 9.68 to 4.31 when the amount of binder added to the CPB batch is increased from 3 to 7 wt%. Also, ρ_d value was 1810, 1800, and 1790 kg/m³ for 3, 4.5, and 7wt% binder contents, respectively.

TABLE 4

Initial bulk properties of different paste backfill materials prepared for a given batch mix

TABELA 4

Właściwości początkowe (pierwotne) różnych materiałów sypkich do zawiesiny przygotowane dla danej partii mieszanki (zawiesiny)

Parameter	ρ (kg/m ³)	B_v (wt%)	C_w (wt%)	C_v (wt%)	G_s	w/c	e	ρ_d (kg/m ³)	S_r (wt%)	θ
3.0 wt%	2 320	3.8	78	49.1	3.68	9.68	1.04	1810	100	0.5
4.5 wt%	2 310	5.7	78	49.2	3.66	6.55	1.03	1800	100	0.5
7.0 wt%	2 300	8.8	78	49.3	3.64	4.31	1.03	1790	100	0.5

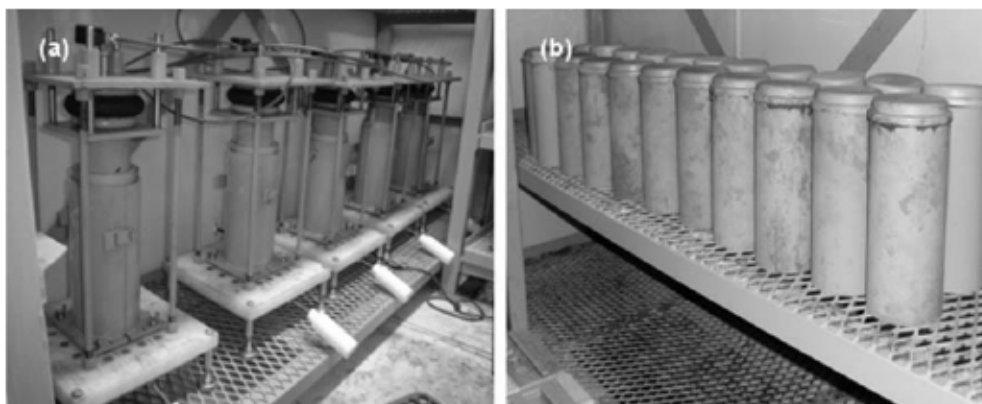


Fig. 2. Photos of (a) CUAPS samples and (b) plastic mould samples curing in a humidity chamber

Rys. 2. Zdjęcia: a) próbki CUAPS, b) próbki z formy ze sztucznego tworzywa w komorze wilgotnościowej

Immediately after mixing, the prepared paste materials were poured into both plastic moulds and transparent polycarbonate round tubes. Tubes and moulds have a shape factor of 2 that corresponds to the height-to-diameter ratio. The CPB samples were then sealed and stored in a damp room maintained at ~80% relative humidity and 24°C ($\pm 2^\circ\text{C}$) in order to mimic curing conditions similar to those observed in the underground stopes of the mine under study. Fig. 2 shows CUAPS-consolidated CPB and plastic mould-unconsolidated CPB samples being cured in a controlled-humidity chamber.

One-dimensional consolidation tests: as part of the present study, one-dimensional consolidation tests inspired by ASTM D2435 and D4186 standard operating procedures were carried out. CPB samples were consolidated using the CUAPS which allows a pressure range of 0–500 kPa. Before running the tests, a seating stress of 15 kPa was applied to provide contact between the sample and loading platen. Then, an example stress sequence of 0.5, 25, 50, 100, 200, and 400 kPa based on a load increment ratio LIR of 1 was applied to the CPB, and the axial deformations during each stress increment were measured with linear displacement variable transducers through a HOBO U12-4 data logger at a time interval of 0, 2, 4, 6, 8, and 10 h. All stresses were incrementally applied during the first day of curing and unloaded down to 0.5 kPa after a desirable curing time.

Unconfined compression tests: A total of 36 CPB samples, 9 CUAPS-consolidated samples, and 27 mould-unconsolidated samples were cured for uniaxial compressive strength (UCS) tests after periods of 7, 14, and 28 days. After curing, all CPB samples were subjected to the UCS tests according to ASTM C39 using a computer controlled MTS 10/GL press which has a nominal loading capacity of 50 kN and a displacement speed of 1 mm/min. The UCS magnitude of each sample was recorded (peak or ultimate stress) along with the elasticity modulus of deformation. For a given binder proportion and curing time, only one-test was done for the CUAPS-consolidated samples, whereas an average of three tests were performed for mould-unconsolidated samples.

Bulk properties determination: after UCS testing, the calculated and/or measured CPB geotechnical index (bulk properties) parameters were determined as follows: gravimetric water content w (%), solids specific gravity G_s , degree of saturation S_r (%), void ratio e (or porosity n), volumetric water content θ , dry density ρ (g/cm^3), solid concentration C_w (%), and specific surface area S_s (m^2/kg). The S_s parameter, based on the BET (Brunauer, Emmett and Teller) method, was evaluated by nitrogen N_2 isotherm adsorption using a Micromeritics® surface analyzer model Gemini III.

Chemical and geochemical analysis: during the one-dimensional consolidation test, a quantity of water was collected from each CUAPS-consolidated CPB sample at a specific time interval. After the acidifying process, analysis of the chemical composition of the collected water was carried out using a Perkin-Elmer ICP-AES (Optima 3100 RL). In addition, the pH, redox potential (Eh), and electrical conductivity (EC) parameters were also measured using a Benchtop pH/ISE meter Orion Model 920A coupled with a Thermo Orion Triode combination electrode (Pt-Ag-AgCl).

3. Results

3.1. Drainage water of consolidated CPB samples

Fig. 3 shows the evolution of cumulative drainage water W_d versus elapsed time for CUAPS-consolidated CPB samples prepared with a binder content of 3, 4.5, and 7 wt% as a function of elapsed time (7, 14 and 28 days of curing). It can be observed from Fig. 3 that the W_d parameter is greatly affected by the hardening process, depending on the amount of binder added to the CPB. The drainage rate decreases with increases in each binder content. For a 8-day curing time, the volume of drainage water for CPB with 3, 4.5, and 7 wt% binder was 19.6 wt% (253.5 mL), 18.8 wt% (242.7 mL), and 17.1 wt% (217.8 mL), respectively. Also, the pressure varying from 0.5 to 400 kPa played a key role in the volume of water collected from the samples. Overall, a pressure of 100, 200, and 400 kPa corresponding to an elapsed time varying between 120 and 600 minutes induced a significant gap in the magnitude of W_d . However, a lower pressure of 0.5, 25, and 50 kPa did not have a big influence on the water collection even at early curing times (up to 7 days).

3.2. Vertical strain of consolidated CPB samples

Fig. 4 shows the evolution of the vertical strain ε_v ($= \Delta H \cdot 100/H_0$) of CUAPS-consolidated CPB samples cured under variable pressures as a function of binder content (3, 4.5, and 7 wt%). For a given applied pressure, Fig. 4 shows that the total ε_v observed from CPB samples containing 3 wt% binder is slightly greater than those observed from CPB samples containing the 4.5 and 7 wt% binder content. For pressure greater than 200 kPa, the settlement behaviour is relatively similar. For a binder content of 3, 4.5, and 7 wt%,

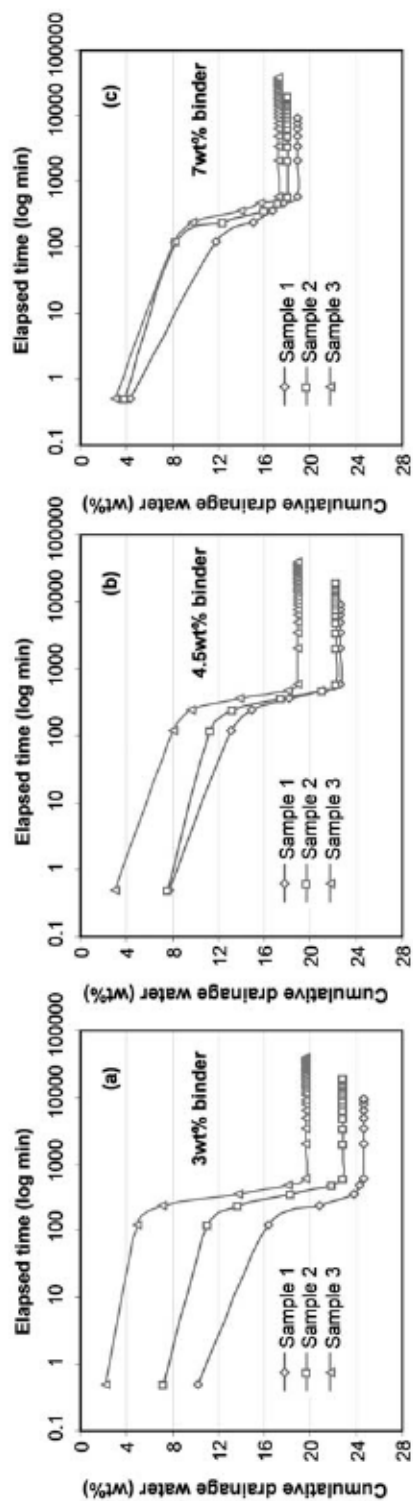


Fig. 3. Evolution of CPB drainage water
a) 3 wt%, b) 4.5 wt%, and c) 7 wt% binder content

Rys. 3. Zmiany drenażu wody z próbek CPB
a) 3% wag., b) 4,5% wag., c) 7% wagowych lepiszcza (spoiwa)

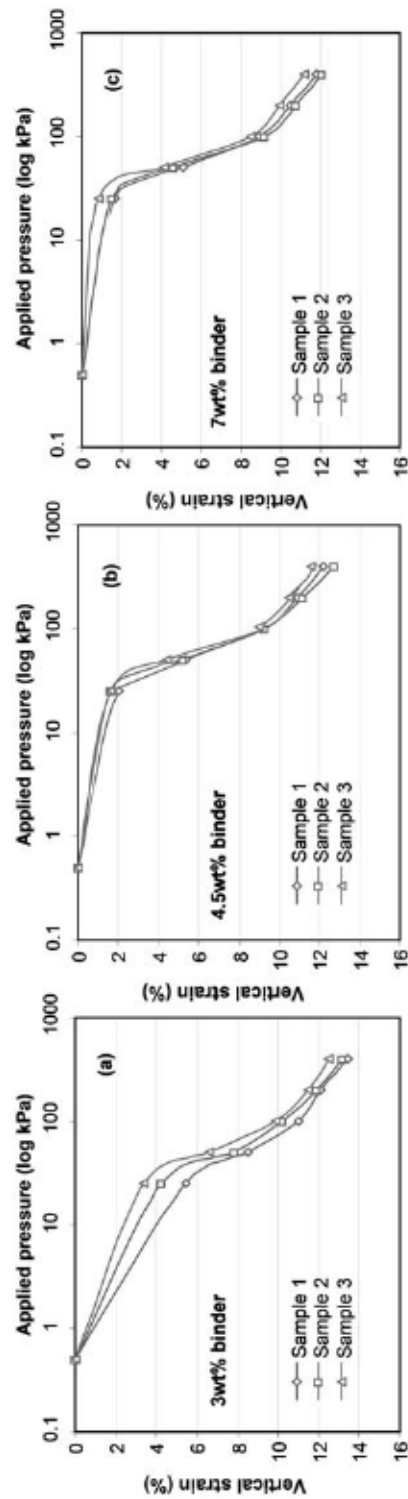


Fig. 4. Variation of CPB vertical strain with applied pressure
 a) 3%, b) 4.5%, and c) 7 wt% binder

Rys. 4. Zmiana pionowej składowej CPB z przyłożonym ciśnieniem
 a) 3%, b) 4,5%, c) 7% wagowych spoiwa

the observed ϵ_v value varies from 12.5 to 13.4%, from 12.2 to 13.3%, and from 11.2 to 12.1 for samples 1, 2, and 3, respectively. The shape of the curves is different particularly for pressure less than 200 kPa. As the amount of binder added to CPB material increases, the ϵ_v value decreases mainly because of binder hydration during the first day.

3.3. Characterization of CPB mechanical strength

Fig. 5 shows the difference in terms of strength development between CUAPS-consolidated CPB samples and plastic mould-unconsolidated CPB samples as a function of binder content and curing time. Strength proportionally increases with binder content as expected. Generally, the CUAPS-consolidated CPB samples systematically produce higher mechanical strength (higher UCS value) than mould-unconsolidated samples, regardless of the curing time. This difference is 57.9%, 64.3%, and 58.2% for 3 wt% binder content; 52.3%, 54.3%, and 51.8% for 4.5 wt% binder content; and 50.5%, 27.1%, and 19.8% for 7 wt% binder content (Fig. 5a, b, c). The highest UCS gap was obtained for CPB with 3 wt% binder

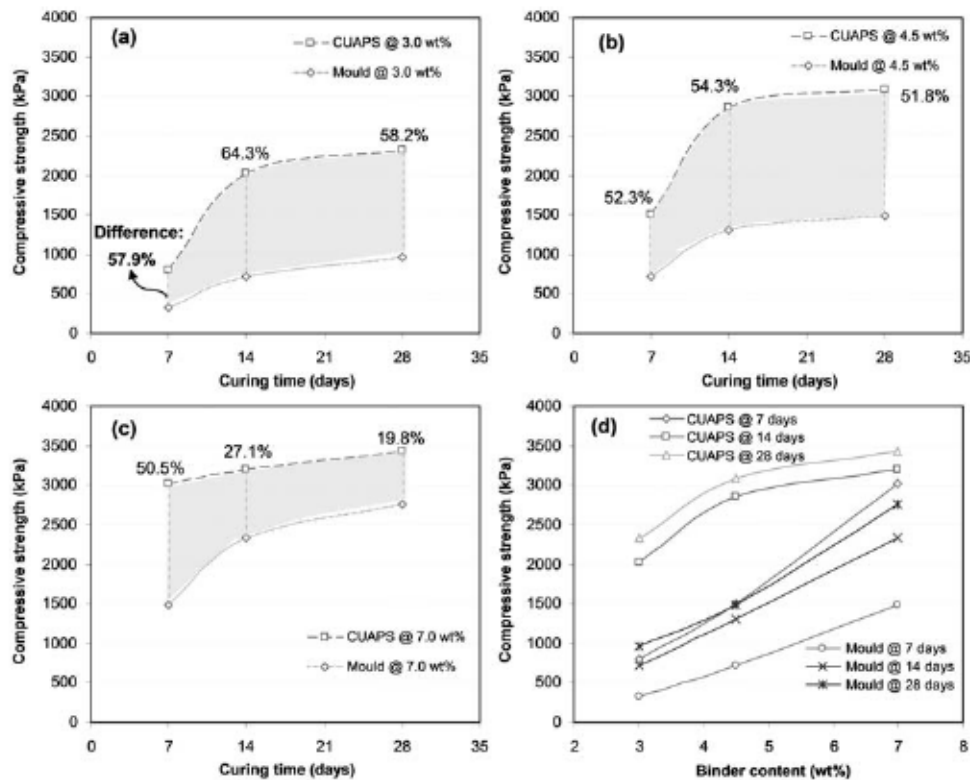


Fig. 5. Comparison of CPB strength
 a) 3 wt%, b) 4.5 wt%, c) 7 wt%, and d) mixed results

Rys. 5. Porównanie siły CPB
 a) 3%, b) 4,5%, c) 7% wag., d) różne wyniki

content. One of the probable reasons for such a high UCS variation could be the high drainage of excess water from these mixes during consolidation which reduces the resultant porosity and permeability. During the drainage, the stiffness of consolidated samples increases as a function of increasing density.

Another reason is that the hardening of CPB materials being cured under different effective stresses greatly contributes to the strength development of backfill. If a series of effective stresses is applied just before the commencement of curing which contributes to CPB hardening, as is being practiced in the present investigation, the rate of strength increase (short-term property), ultimate strength (mid-term property), and durability (long-term property) will be greater compared to the application of pressure after the completion of hydration. It has been experimentally shown from Belem et al. (2002) and Yilmaz et al. (2008, 2009) that the primary bonds in cement in CPB may be broken due to hydrates damage caused by the applied pressure (overloading) in early stages of strength development, which thus reduces the strength and stiffness.

3.4. Characterization of CPB geotechnical index parameters

Fig. 6 shows the variation of the geotechnical index parameters of CUAPS-consolidated and plastic mould-unconsolidated CPB samples containing a binder content of 3, 4.5, and 7 wt% as a function of curing time. The measured geotechnical index parameters are as follows: the gravimetric water content w_g (%), dry density ρ_d (g/cm^3), specific surface S_s (m^2/kg), degree of saturation S_r (%), and void ratio e . There is a notable variation in water content, dry density, and void ratio. The CUAPS samples have a w_g value varying between 18.8–11.2 wt%, a ρ_d value varying between 2.13–2.08 g/cm^3 , and a void ratio e value varying between 0.7–0.57; while mould samples have a w_g value varying between 25.2–17%, a ρ_d value varying between 1.85–1.87 g/cm^3 , and a void ratio e value varying between 0.95–0.71. In addition, the final water content w_g and degree of saturation S_r are highly influenced by the drainage ability of CPB. The highest binder content (7 wt%) exhibits the lowest w_g and S_r . This could basically be explained by the different initial w/c ratios (Table 4) and the amounts of water required for binder hydration.

It was shown by several authors (Belem et al. 2006; Benzaazoua et al. 2004; Yilmaz et al. 2008) that drainage of the excess water existing within the CPB sample contributes to a better hardening process (water trapped during the cement hydration or hydrated phase precipitation as gypsum) for CPB material and, consequently, the reduction of the total void ratio of backfilling. The rate of water drainage is also significantly affected by the settling (i.e. improved consolidation and/or density increase) of the CPB sample being cured under a series of pressure levels during the first step of curing, thus demonstrating higher strengths. Specific surface area S_s is also closely related to the hardening phase formation during curing, and the ultimate mechanical strength. The higher the binder content used, the greater the overall S_s value of the CPB due to hydrate growth. The UCS values increase with the increasing S_s values for a given binder content. Overall, one can say from these tests that the

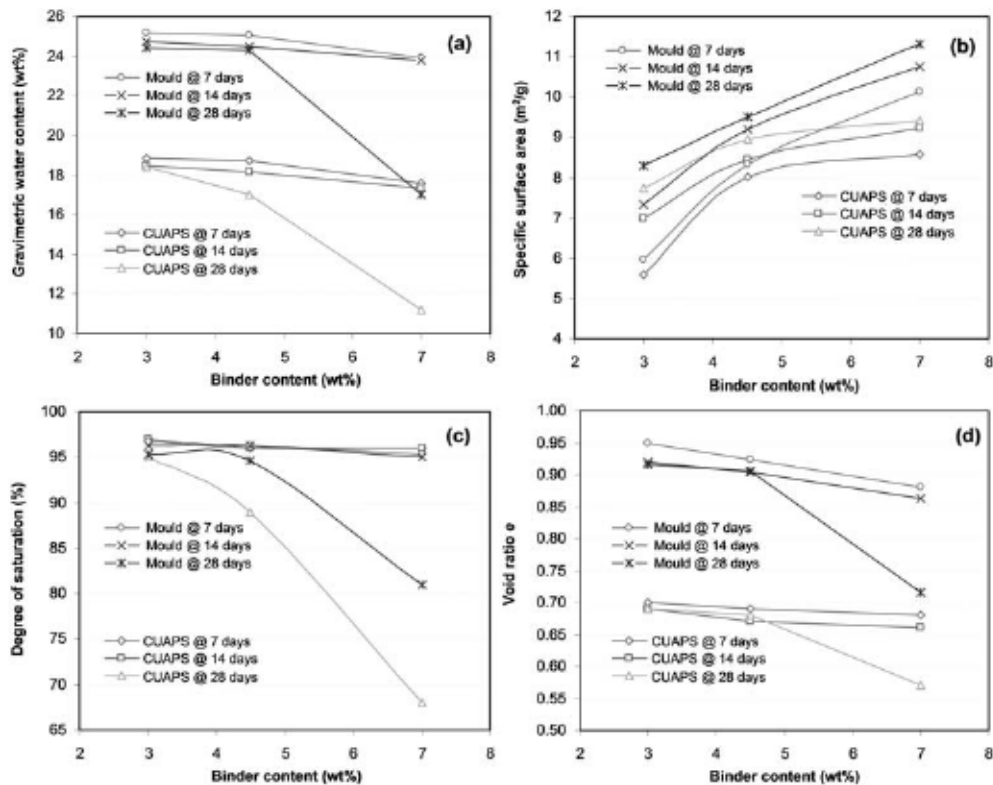


Fig. 6. Evolution of CPB geotechnical index parameters as a function of binder content and curing time

Rys. 6. Zmiana CPB parametrów indeksu geotechnicznego w zależności od zawartości spoiwa i czasu utwardzania

CUAPS-consolidated samples give lower w_g and S_r , and higher UCS than those obtained using plastic moulds due to the variation in the initial void ratio e_0 and the influence of time-dependent consolidation settlement.

4. Discussion

4.1. Correlation between the mechanical strengths

From the compression test results shown in Fig. 7, it seems that a strong correlation exists between the UCS obtained from CUAPS-consolidated samples (UCS_{CUAPS}) and the UCS obtained from mould samples (UCS_{mould}). After normalizing the UCS values for binder content and curing time, it was possible to observe the correlation between the two sets of UCS data. This normalization consisted of multiplying each UCS variable by the binder content/curing time ratio ($UCS \cdot B_w^{0.9}/t$).

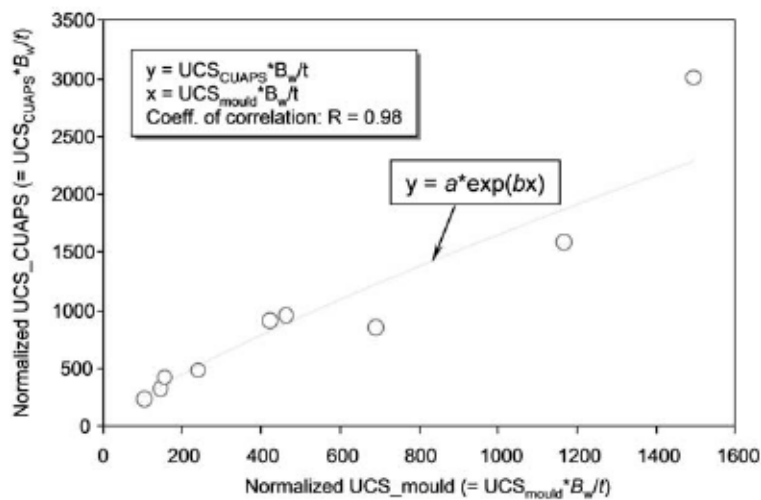


Fig. 7. Correlation between UCS_{CUAPS} and UCS_{mould} based on binder content and curing time

Rys. 7. Korelacja pomiędzy UCS_{CUAPS} a UCS_{mould} uwzględniająca skład spoiwa i czas utwardzania

Fig. 7 shows the relationship between UCS_{CUAPS} and UCS_{mould} . It should be noted that at first approximation, it appears this relationship could be well-described by an exponential function such as $y = a \cdot \exp(b \cdot x)$. Indeed, a coefficient of correlation $R = 0.98$ was obtained, thus indicating the strong relationship existing between the two types of UCS variables. The exponential relationship describing the correlation between these two types of variables (UCS_{CUAPS} , UCS_{mould}) is as follows:

$$UCS_{CUAPS} \cdot \left(\frac{B_{w\%}}{t} \right) = 360.952 \cdot \exp \left(0.001399 \cdot \left[UCS_{mould} \cdot \frac{B_{w\%}}{t} \right] \right) \quad (1)$$

where:

$B_{w\%}$ – binder content (wt%),

t – curing time (day),

UCS values are in kPa.

Finally, the predictive CUAPS-consolidated backfill samples equation from known mould-unconsolidated backfill samples is given as follows:

$$UCS_{CUAPS} \cdot \left(\frac{360.952 \cdot t}{B_{w\%}} \right) \cdot \exp \left(UCS_{mould} \cdot \frac{0.001399 \cdot B_{w\%}}{t} \right) \quad (2)$$

The above equations are specific to the CPB tested and to the loading sequence applied in this study (see the section on one-dimensional consolidation tests). However, ongoing

research on the influence of different pressure loading rates on CPB consolidation characteristics, and consequently on its strength development, is underway. After these tests, it would be possible to develop a more general equation that could be used to improve prediction of CPB strength.

4.2. Chemical analysis of drainage water

Table 5 shows the variation in the chemical analysis of drainage water obtained from CUAPS-consolidated CPB samples during 1-D consolidation tests. Overall, the calcium Ca, silicon Si, and sulphates SO_4^{2-} concentrations of CPB samples containing 3, 4.5, and 7 wt% binder content decreased with increasing curing time. The variation in terms of chemical composition can be explained by the different chemical compositions of both binder and tailings pore water and tap water (see Table 3) used for the CPB mixture, relating to the hydration and precipitation process at early stages. However, the variation is also due to the fact that the denser the cementitious matrix, the more this later acts as a filter of colloidal species from the pore waters. It can be interpreted from Table 5 that for a given binder content, calcium Ca concentration is higher at early ages (7 days) and begins to reduce appreciably when the curing time is increased to 28 days.

The reason for this phenomenon can be explained by the dissolution of Ca-bearing mineral within the cement during early hydration stages. At this stage, Ca remains trapped in porous media. The dissolved Ca is expelled from the sample along with drainage water as a function of pressure. In a similar manner, Si concentration is reduced with increased

TABLE 5
Chemical composition of water collected from CUAPS-consolidated CPB samples

TABELA 5
Skład chemiczny wody pobranej z CUAPS konsolidowanych próbkami CPB

Binder content	Curing (day)	Al (ppm)	Ca (ppm)	Fe (ppm)	Mg (ppm)	Si (ppm)	SO_4^{2-} (ppm)
3.0 wt%	7	0.03	436	0.07	0.08	13.3	6692
	14	0.02	429	0.13	0.04	12.5	6411
	28	0.03	284	0.15	0.05	11.7	4945
4.5 wt%	7	0.01	420	0.06	0.02	15.0	6390
	14	0.02	302	0.12	0.05	14.0	6100
	28	0.03	247	0.17	0.01	13.5	3056
7.0 wt%	7	0.01	412	0.08	0.02	28.0	6279
	14	0.03	251	0.13	0.78	27.5	5572
	28	0.04	194	0.22	0.18	26.7	2885

curing time, but at lower rates than Ca due to the fact that Si occurs as gel phases in solid form. One can also say that sulphate concentration is relatively higher (6692, 6390, and 6279 ppm for 3, 4.5, and 7 wt% binder respectively at 7 days of curing) than the concentration observed from the tailings pore water (4883 ppm, also Table 3). This means that sulphate concentration increases during mixing at early curing ages due to aeration contributing to sulphide oxidation by oxygen intake. Also, the sulphates can come from the sulphated mineral dissolution from the binder.

4.3. Geochemical analysis of drainage water

Fig. 8 shows a variation in geochemical analyses of water collected from the CUAPS-consolidated CPB samples containing a binder content of 3, 4.5, and 7 wt%. The measured parameters are pH, redox potential, (Eh) and electrical conductivity (EC).

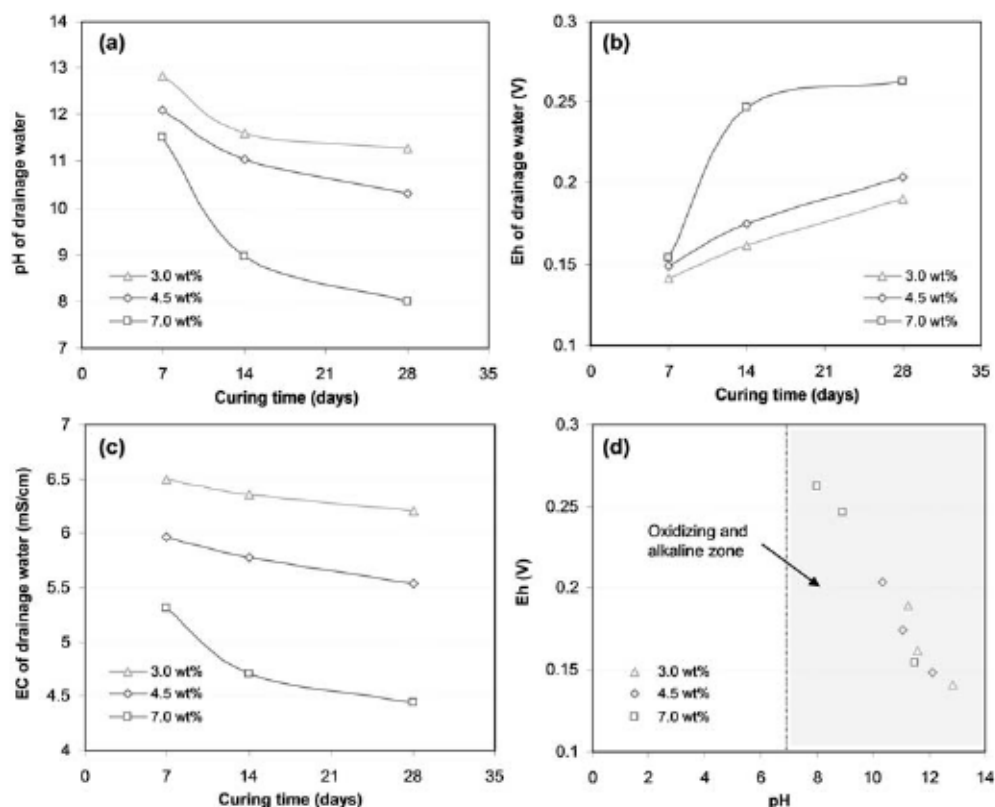


Fig. 8. Geochemical analysis of drainage water
a) pH, b) Eh, c) EC, and d) pH vs. Eh

Rys. 8. Analizy geochemiczne wód z odwodnienia
a) pH, b) Eh, c) EC, d) pH vs. Eh

For all CPB mixtures, as the amount of binder used in the tailings material sample increases, the pH and EC parameter decreases as a function of curing time. However, the Eh value increases with increasing binder content and curing time. In terms of magnitude, the highest pH change (from 11.5 at 7-day curing to 7.9 at 28-day curing) is observed for the sample containing 7 wt% binder content (Fig. 8a). This shows that pH decreases with increasing binder content and curing time. Besides, the Eh values as shown in Fig. 8b indicate that the CUAPS-consolidated sample containing 7 wt% binder acts differently, while a binder content of both 3 wt% and 4.5 wt% act in the same manner. After 7 days of curing, all Eh values are almost the same (0.141, 0.148, 0.154, volts for 3, 4.5, and 7 wt% binder contents, respectively). In addition, pH decreases while Eh increases with increased curing time due to the enhanced reactivity of sulphide during the sample preparation and mixing. Oxidation of pyrite accelerates acidity (i.e. the decrease of pH) and gives rise to a loss of strength over time. The degree of saturation S_r is closely related to the Eh parameter. Eh increases with the decrease in S_r , which contributes to the increase in oxidation by molecular oxygen diffusing through CPB samples. At the highest binder content (7 wt%), the S_r value is much lower compared to other binder contents (3 and 4.5 wt%) and the Eh value becomes higher, as can be seen in Fig. 8b. However, as curing time increases, the gap observed between 7 and 4.5 wt% binder contents increases from 0.17 to 0.25 after 14 days, and from 0.2 to 0.26 after 28 days. The electrical conductivity EC value decreases from 6.5 to 6.2 mS/cm, from 5.9 to 5.5 mS/cm, and from 5.3 to 4.4 mS/cm when curing time is increased from 7 to 28 days (Fig. 8c). In addition, the Eh versus pH curve, as indicated in Fig. 8d, shows that the drainage water exhibits an oxidizing and alkaline medium.

Conclusion

This study compared the results of laboratory testing conducted in order to better understand the hydromechanical and geochemical properties of consolidated (using a new laboratory apparatus called CUAPS) and unconsolidated (using plastic moulds) CPB samples. From the tests performed, one can conclude that the one-dimensional consolidation of CPB samples being cured under a series of pressure increments is essential to a better understanding of events linked to placement and curing conditions occurring in underground stopes during the backfilling process. The CUAPS-consolidated CPBs always give higher mechanical strengths than those obtained from plastic moulds (undrained) for a given binder content and curing time. Geotechnical index parameters show that void ratio is dramatically reduced when CPB samples are prepared using CUAPS cells that allow drainage during a curing time of 7, 14, and 28 days. The reduction in the void ratio gives rise to a stiffer, and therefore more resistant, CPB material. From the chemical composition analysis, it can be noted that the concentrations of sulphate SO_4^{2-} continuously decrease over time, depending on the binder content used in the CPB mixture. Geochemical analyses have also shown that drainage water exhibits an alkaline to neutral oxidizing medium, based on the relationship

between pH and Eh. These results showed that the CUAPS apparatus gives higher CPB mechanical strength than UCS values from laboratory-prepared CPB samples. Consequently, more realistic mechanical strength results can be obtained using this new apparatus. More work is underway to validate the values obtained by the CUAPS apparatus and *in situ* results.

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BADANIA FIZYKOCHEMICZNE I MECHANICZNE WŁAŚCIWOŚCI
SKONSOLIDOWANYCH I NIEKONSOLIDOWANYCH ZAWIESIN NASYCONYCH CEMENTEM

Słowa kluczowe

Odpady do podsadzki, woda w zrobach, indeks geologiczny, efektywne ciśnienie, siła

Streszczenie

W ostatnich latach obserwuje się, że wydajność i jakość próbek zawiesin nasyconych cementem (CPB – *Cement Paste Backfill*) *in situ* są stale niższe niż tych samych próbek przygotowanych w plastikowych formach w laboratorium. Może to wynikać z braku w laboratorium skutecznego narzędzia mogącego naśladować zawiesiny *in situ*, warunki ich utwardzania jak również rozmiary i geometrię próbek. W celu wypełnienia tej luki, w laboratorium opracowano nowe narzędzie o nazwie CUAPS (*Curing Under Applied Pressure System*), wytworzone i wykorzystane do zbadania wpływu ciśnienia na podstawie skutecznego nacisku na właściwości fizykochemiczne i mechaniczne CPB, jak również próbek otrzymanych z plastikowych form. Badania porównawcze przeprowadzono zarówno dla próbek CUAPS jak i próbek otrzymanych w laboratorium, zawierających lepiszcza (cementu) 3,45 i 7% wag. po 7, 14 i 28 dniach utwardzania. Wyniki wskazują, że wydajność próbek konsolidowanych CUAPS są zawsze bardziej prawdziwe (realistyczne) niż próbek otrzymanych w laboratorium, głównie z powodu odprowadzania wody w wyniku konsolidacji. Ostatecznie metoda CUAPS powoduje korzystny wpływ na utwardzenie CPB dzięki zawartości wody (separacja wody od świeżej zawiesiny odpadów z cementu) i połączeniu części wody zasobowej z zawiesiną w zrobach.

STUDY OF PHYSICO-CHEMICAL AND MECHANICAL CHARACTERISTICS OF CONSOLIDATED
AND UNCONSOLIDATED CEMENTED PASTE BACKFILLS

Key words

Tailings backfill, drainage water, geological index parameter, effective stress, strength

Abstract

In recent years, it has been observed that the performance and quality of *in situ* cemented paste backfill (CPB) samples are constantly lower than samples obtained from the same CPB mix poured into laboratory-prepared plastic moulds. This could be well explained by the absence of an efficient laboratory tool capable of mimicking CPB's *in situ* placement, hardening, and curing conditions relating to stope size and geometry. To meet this need, a new laboratory tool named CUAPS (curing under applied pressure system) was manufactured and used to examine the effect of curing under effective stress on physico-chemical and mechanical properties of CPB, along with plastic mould samples. A comparative study was conducted for both CUAPS and mould samples containing a binder content of 3, 4.5, and 7 wt% after curing times of 7, 14, and 28 days. Results indicate that the performance of CUAPS-consolidated samples are always more realistic than those of plastic mould-unconsolidated samples mainly due to water drainage induced by consolidation. CUAPS has witnessed an advantageous effect on CPB hardening which includes the amount of bleeding water (separation of water from the fresh backfill material) and a combination of drainage of part of the mixing water and the settlement of paste backfill after its placement in the cap.