

# Lightweight Concrete with an Agricultural Waste – Buckwheat Husk

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This paper presents a parametric experimental study which investigates the potential use of buckwheat husk wastes (as an aggregate in concrete) and opoka combination for producing a lightweight composite. The study was carried out to investigate the influence of the buckwheat husk extract on the setting and hardening of pastes in which cement was partially replaced by 0, 33, or 50 % of natural pozzolana. The kinetics of hydration up to 48 h was studied using isothermal conduction calorimetry. Some of the physical and mechanical properties of buckwheat husk concrete with various levels of cement replacement of opoka without and with sand (part of buckwheat husk was replaced in the same volume of sand) were investigated.

Experimental data on the compressive strength of concrete utilizing buckwheat husk with and without sand and cement at varying proportions are presented. The results suggest that buckwheat husk may be used as an aggregate, particularly in lightweight concrete, panel and blocks for walls at a relatively cheaper price.

*Key words:* waste; buckwheat husk; cement; pozzolana; isothermal calorimetry; lightweight concrete.

## 1. INTRODUCTION

Isolation and low-cost materials are important for non-load-bearing and load-bearing lightweight concrete masonry units [1], especially for agricultural constructions because, in general, farm buildings are of one or two floors and carry low loads. Therefore, the strength of constructed buildings is not as important as heat isolation and construction cost [2].

Lightweight aggregate concretes are broadly divided into three groups based upon their use and physical properties: structural, structural/insulating, and insulating. The bulk density, thermal conductivity, and compressive strength ranges, normally associated with each class of concrete, are summarized in Table 1. The first group is termed low strength, corresponding to low density, and is mostly used for insulation purposes. The second group is of moderate strength and is used for filling and block concrete. The third group is structural lightweight concrete and is used in reinforced concrete [3]. Organic lightweight aggregate concrete is produced by replacing normal aggregate with organic aggregate in concrete or mortar, either partially or fully, depending on the requirements of strength and density.

**Table 1.** Lightweight aggregate concrete classified according to use and physical properties [3]

Properties	Lightweight aggregate concrete		
	insulating	structural/ insulating	structural
Concrete density, kg/m <sup>3</sup>	240–800	800–1440	1440–1840
Compressive strength, MPa	0.7–(2.0)3.4	(2.0)3.4–17.0	> 17.0

To produce lightweight concrete, numerous methods can be used. The most popular method is the use of natural, synthetic or organic lightweight aggregates. Examples of lightweight aggregates are pumice, coal slag, flying ash, and various agricultural wastes and residues. It contains relatively large quantities of lignocellulosic materials available in the form of agricultural residues. The organic waste used in lightweight concrete is mainly of plant origin and includes rice and wheat husk, straw, sawdust, cork granules, coconut fibre and shell [4, 5].

A large quantity of lignocellulosic wastes is generated worldwide from various sources such as agriculture, construction, wood and furniture industries leading to environmental concerns. The use of these wastes in making cement-bonded construction materials can reduce the magnitude of the problems. However, in this effort, there are various restraints such as the compatibility of these wastes with cement and a limited composite strength [6].

In the course of growing and cleaning buckwheat grains, large amounts of wastes are formed (straw, husk, fine particles of grains). These agricultural wastes can be considered as valuable renewable raw materials for chemical, food, and pharmacological industries, suitable for the production of polysaccharides, cellulose, dyes and food additives, furfural, and drugs [7]. It has also been found that aqueous extracts of buckwheat straw and husk inhibit the corrosion of low-carbon steel [8]. However, until now, the main part of buckwheat wastes has not been used in chemical industry and is burnt in fields or used as a fuel for boilers, fillers for pillows, packing of fruits and fragile goods, and feed [9].

Much less data on these wastes are available as compared to the related products of rice processing. The buckwheat husk (BH) contains more organic matter and significantly less inorganic matter in comparison to rice husk widely used in cement-bonded composites [5].

Polysaccharides obtained by successive extraction with water, ammonium oxalate, and sodium hydroxide

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from buckwheat and sunflower husks contain not only polysaccharide fractions, but also protein–polysaccharide complexes and low-molecular-weight compounds [10].

The extractability of the hemicellulose component strongly depends on the alkali concentration. Increasing the alkali concentration increases the total yield of hemicelluloses [11].

Polysaccharides extractable with water from buckwheat husk have a relatively high glucose content. Polysaccharides obtained by alkaline extraction have a more complex monosaccharide composition which consist of residues of rhamnose, arabinose, xylose, mannose, glucose, and galactose [7, 10, 11].

The polysaccharide (PS) content depends on the buckwheat husk plant strain and recovery procedure. The total yield of PSs from buckwheat husk varies from 3.2 to 6.3 wt.% based on the dry raw material. The largest amount of polysaccharides is recovered by alkaline extraction and the smallest amount by aqueous extraction [12]. The yield of PSs from buckwheat wastes obtained in the extraction with water varies from 2.14 to 2.76 wt.% [7].

Samples of a dry extract (DE) were obtained by evaporation of liquid extracts on a water bath. The yield of a dry extract of raw material and water mixture in the ratio of 1 : 25, depending on the concentration of extractive substances in the solution, is no less than 2 % relative to the mass of the starting raw material. Such a low DE yield as compared with the amount of extractive substances in solution can be due to the presence of volatile components in the extract.

As Kuhne and Meier have reported [13], hemicelluloses cause the main inhibiting effects. The alkaline medium, which is caused by cement, decomposes hemicelluloses into the substances that negatively influence cement setting.

Colleparidi et al. found that the addition of glucose, gluconate, or lignosulfonate stabilized ettringite in the C<sub>3</sub>A – gypsum – lime system [14].

Buckwheat husk (BH) as an organic waste is a significant problem in buckwheat cultivating areas, because it is not used profitably and is generally burned after harvest, which causes environmental problems. The use of BH in concrete production may solve this environmental problem and provide an advantage in producing lightweight and low-cost concrete.

The aim of this study was to examine the effect of BH on the engineering properties of concrete.

An earlier work of one by us and colleagues [15] confirmed that in the production of lightweight concrete with sawdust or other fine organic agricultural additives the abundant usage of high-grade Portland cement is not rational, because the same amount of less active mineral binders or pozzolan mineral additive gives identical or even better results.

Opoka has been found [16] to be an efficient additive which reduces the harmful influence of hemicellulose on the hydration of Portland cement.

This paper gives a review of results of a recent research into the use of these wastes in making cement-bonded composites for building materials.

## 2. MATERIALS AND METHODS

Buckwheat hulls (Fig. 1), from which extracts were prepared, sifted through a sieve in order to separate them from dust and other impurities as well as crops seeds. A fraction with a particle size of no less than 2 mm was selected for experiments. Then the husk samples were washed with water and dried in an open air to constant weight. Their moisture was about 6.87 %. The apparent density of buckwheat husk was about 125.5 kg/m<sup>3</sup>.



Fig. 1. Buckwheat husk

The buckwheat husk was poured with water (water extracts) and with saturated Ca(OH)<sub>2</sub> solution (alkali extracts). A buckwheat husk sample was mixed with water at a ratio 1 : 10 by weight, the mixture was heated at 80–90 °C for 24 h, and then the formed extracts were filtered. The concentration of buckwheat husk water extract was 0.4 ± 0.04 %. Samples of a dry extract (DE) were obtained by evaporation of liquid extracts on a water bath. The yield of a dry extract at raw material was 4.4 ± 0.2 % relative to the mass of the starting raw material.

The presence of pigments in the soluble substances of buckwheat husk results in a different colour of extracts.

Lithuania contains abundant natural pozzolana called opoka, which consists of amorphous SiO<sub>2</sub> and finely dispersed CaCO<sub>3</sub> [17]. The composition of opoka (natural pozzolana) is presented in Table 2 and Table 3. The pozzolana was ground in a laboratory mill to a specific surface area of 1200 m<sup>2</sup>/kg (determined by the Blaine method). The hydraulic activity of pozzolana was 220 mg of CaO/g.

Table 2. Oxide composition of opoka

Component, wt.%						
SiO <sub>2</sub>	Al <sub>2</sub> O <sub>3</sub>	Fe <sub>2</sub> O <sub>3</sub>	CaO	MgO	Na <sub>2</sub> O	K <sub>2</sub> O
54.1	2.7	1.3	23.2	0.62	0.24	0.87

The quantitative chemical composition analysis of Portland cement clinker by X-ray fluorescence spectroscopy (XRF) on a Bruker X-ray S8 Tiger WD spectrometer equipped with a Rh tube with energy of up to 60 keV. Powder samples were measured in Helium atmosphere and data were analysed with SPECTRA Plus QUANT EXPRESS standard software. The accuracy of measurements – < 0.01 % (with standards).

The composition of the Ordinary Portland cement (OPC) clinker is shown in Table 4 and Table 5. The OPC was obtained by mixing OPC clinker and 4.5 % gypsum (specific surface area determined by the Blaine method was 320 m<sup>2</sup>/kg).

**Table 3.** Mineral composition of opoka

Component, wt. %							
Amorphous SiO <sub>2</sub> -nH <sub>2</sub> O	Ancerite Ca(Fe,Mg,Mn)(CO <sub>3</sub> ) <sub>2</sub>	Calcite CaCO <sub>3</sub>	Cristobalite SiO <sub>2</sub>	Muscovite KAl <sub>2</sub> (AlSi <sub>3</sub> O <sub>10</sub> )(F,OH) <sub>2</sub>	Plagioclase NaAlSi <sub>3</sub> O <sub>8</sub>	Quartz SiO <sub>2</sub>	Tridymite SiO <sub>2</sub>
22.2	1.3	36.1	19.2	4.1	1.1	12.0	4.0

**Table 4.** Oxide composition of Portland cement clinker

Component, wt. %						
SiO <sub>2</sub>	Al <sub>2</sub> O <sub>3</sub>	Fe <sub>2</sub> O <sub>3</sub>	CaO	MgO	Na <sub>2</sub> O	K <sub>2</sub> O
19.72	5.41	4.21	62.76	3.41	0.16	1.08

**Table 5.** Mineral composition of OPC clinker

Component, wt. %			
C <sub>3</sub> S	β-C <sub>2</sub> S	C <sub>3</sub> A	C <sub>4</sub> AF
63.19	8.87	7.21	12.81

Besides the OPC pastes for control purposes, two groups of pozzolana (opoka) incorporating specimens with cement replacement ratios of 33 and 50 % by weight were prepared and formed using the standard water and buckwheat husk extract (Table 6). The monitoring of consistency and setting time was directly carried out using a Vicat apparatus.

**Table 6.** Composition of specimens of cement pozzolana (B) paste mix with water and buckwheat husk extract

Mix no.	OPC, %	Opoka, %	W(E)/B
M1W	100	0	0.255
M2W	67	33	0.345
M3W	50	50	0.425
M1E	100	0	0.255*
M2E	67	33	0.345*
M3E	50	50	0.425*

\* Buckwheat husk extract concentration was 0.9 %.

Infrared spectra were recorded using a PerkinElmer Fourier transform-infrared (FT-IR) system Spectrum X spectrometer. Tablets were prepared from 1 mg of the sample and 200 mg of KBr. The spectral analysis was performed in the range of 4000–400 cm<sup>-1</sup> with a spectral resolution of 1 cm<sup>-1</sup>.

An eight channel TAM Air III isothermal calorimeter was used to investigate the heat evolution rate of the cement–opoka binding material (B). Glass ampoules (20 ml) each containing 3 g dry cementitious material were placed in the calorimeter and the injection units for each

ampoule filled with amounts of water (buckwheat husk extract) equivalent to a W(E)/B ratio of 0.5. After a steady temperature of 25 °C had been reached, the water was injected into the ampoules and mixed inside the calorimeter with the dry material for 20 s (frequency 2–3 s<sup>-1</sup>). The heat evolution rate was then measured over a period of 50 h. Repetition of the measurements showed deviations in total heat below 3 % for samples of similar type. Apart from the first minutes of water additive and mixing, the heat evolution rates were essentially identical. The accuracy of measurement – 0.03 W/g.

The buckwheat husk concrete specimens were formed using the OPC (Table 4 and Table 5) without and with sand (part of buckwheat husk was replaced by the same volume of sand). The size of the specimens used for all the tests was 40 × 40 × 160 mm. The concrete was then batched with the composition shown in Table 7. All concrete mixes had the total binder (cement plus opoka) content of 680 kg/m<sup>3</sup>.

For the first three days, the samples were kept in moulds at 20 ± 1 °C and 100 % humidity. Then the specimens were stored for 7, 28 days, and 3 months in a chamber at 20 °C and a 95 % humidity.

The physical properties of the pastes and the strength of mortars were determined in accordance with LST EN 196-3:2005+A1:2009 [18] and LST EN 196-1:2007 [19], respectively.

The moisture content of samples was determined on the wet basis  $W_{wb}$ :

$$W_{wb} = \frac{m - m_{dry}}{m} \cdot 100 \%, \quad (1)$$

where  $m$  is the mass of a wet sample, and  $m_{dry}$  is the mass of a dry specimen.

The X-ray diffraction (XRD) data were collected with a DRON-6 powder X-ray diffractometer with the Bragg–Brentan geometry, using Ni-filtered CuK $\alpha$  radiation operating at a voltage of 30 kV and an emission current of 20 mA. The step-scan covered the angular range of 5–60° (2 $\theta$ ) in steps of 2 $\theta$  = 0.02°.

**Table 7.** Actual mix proportions for 1 m<sup>3</sup> lightweight concrete

Concrete mixture	Material content (kg) for 1 m <sup>3</sup> of concrete mixture				
	Buckwheat husk	Cement	Opoka	Sand	Water
	Without sand				
CRM1	170	680	-	-	455
CRM2	170	453	227	-	455
CRM3	170	340	340	-	455
	With sand				
CRM1S	130	680	-	390	455
CRM2S	130	453	227	390	455
CRM3S	130	340	340	390	455

### 3. RESULTS AND DISCUSSIONS

#### 3.1. Investigation of the fresh buckwheat husk extracts

The husk was treated under two different conditions: neutral (water) and alkali (saturated  $\text{Ca}(\text{OH})_2$  solution). In order to find out the influence of the extraction conditions on the extract composition, an alkaline extract was prepared. It resulted in an alkaline medium during the preparation of buckwheat cement concrete and dissolved the materials present in the buckwheat husk. The alkaline buckwheat husk extract was prepared analogously to the aqueous one; in this case, the saturated calcium hydroxide was used. The results of this investigation are presented in Fig. 2 and Table 8.

Fig. 2 shows infrared spectra (IR) measured for two DE samples obtained from aqueous and alkali extracts of BH. The IR spectra of aqueous and alkali extracts of buckwheat husk contain groups of different intensity bands in the following frequency ranges ( $\text{cm}^{-1}$ ): 3600–2935, 1750–1516, 1450–1260, 1150–1020, and  $\sim 700$ –400, which are typical of polysaccharides of plant origin, including the buckwheat waste [8]. The transmission bands in the 1200–800  $\text{cm}^{-1}$  region give information about the polysaccharide types present [11]. The absorption bands were identified using the available structural-spectroscopic data on polysaccharides [12].

A comparison of the data of IR spectroscopy of dry materials existing in buckwheat husk with the results presented in the literature [10, 11] has shown that they are almost identical, implying the prevalence of various polysaccharides in aqueous and alkaline extracts of buckwheat husks. No more exact composition of buckwheat husk extracts has been determined, and data presented in the literature were used.

At the second stage of the experiment, we investigated the influence of the buckwheat husk extract on the setting and hardening of pastes in which cement was partially replaced by 0, 33, or 50 % of natural pozzolana. All replacements were made by the mass of cement.

**Table 8.** IR absorption maxima of dry buckwheat husk extract samples

Assignment [8, 9]*	Wave number $\nu$ , $\text{cm}^{-1}$	
	1 – DE from an aqueous extract of buckwheat husk	2 – DE from an alkali extract of buckwheat husk
1	3404	3384
2	2934	2335
3	1631	1619
4	1409, 1324, 1259	1420
5	1101, 1024	1125, 1052, 1026
6	790, 538	670, 618, 469
7	-	2496, 1420, 865

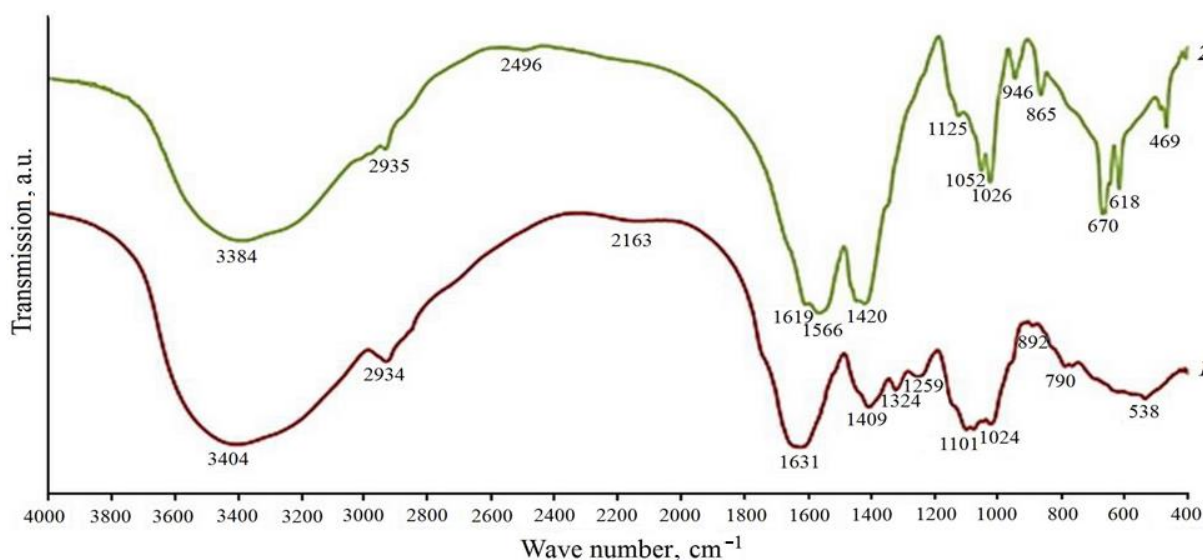
\* 1 – OH stretching vibrations in CHOH and  $\text{CH}_2\text{OH}$  groups; 2 – alkyl groups' stretching vibrations; 3 – OH bending vibrations, C=O stretching vibrations in carbonyl and carboxy groups; 4 – group of bands including C–H bending modes: in-plane, internal (scissor), and external (wagging); 5 – stretching vibrations of C–OH and C–O–C bonds in polysaccharides; 6 – group of bands including out-of-plane bending modes of OH groups and pulsation vibrations of pyranose rings [11, 12]; 7 –  $\text{CO}_3^{2-}$  groups' vibrations.

#### 3.2. Investigations of the initial and the final setting times

The hydration characteristics of buckwheat husk extract-modified cement specimens were determined by measuring the initial and the final setting times (Table 9).

The initial and the final setting times of the control cement were estimated to be 90 and 150 min, respectively. Both initial and final setting times are increased the cement paste mixed with the extract. The initial and the final setting times were 160 min and 330 min, respectively.

When part of cement was replaced with opoka and these mixes were prepared with water, the initial and the final setting times slightly changed as compared with the control cement paste. Thus, the initial and the final setting times varied within 95–80 min (initial) and 165–160 min (final), respectively, when the opoka content in the binder was 33 or 50 wt.%.



**Fig. 2.** The Fourier transform-infrared (FT-IR) spectrum of DE samples obtained from aqueous (1) and alkali (2) extracts of buckwheat husk



**Table 9.** Initial and final sets of cement pozzolana paste mix with water and buckwheat husk extract

Mix	Initial set time	Final set time
Mixing with water		
M1W	90 min	150 min
M2W	95 min	165 min
M3W	80 min	160 min
Mixing with extract		
M1E	160 min	330 min
M2E	75 min	205 min
M3E	60 min	195 min

However, when these binders with opoka were prepared with the buckwheat husk extract, their initial setting times significantly decreased (to 75 and 60 min, respectively). The initial setting time of these specimens with opoka is shorter than of analogous specimens prepared with water. Opoka also minimizes the influence of extracts on the cement final setting time to 205 and 195 min, respectively, but it remains longer as compared with specimens prepared with water.

### 3.3. Investigation of isothermal calorimetry

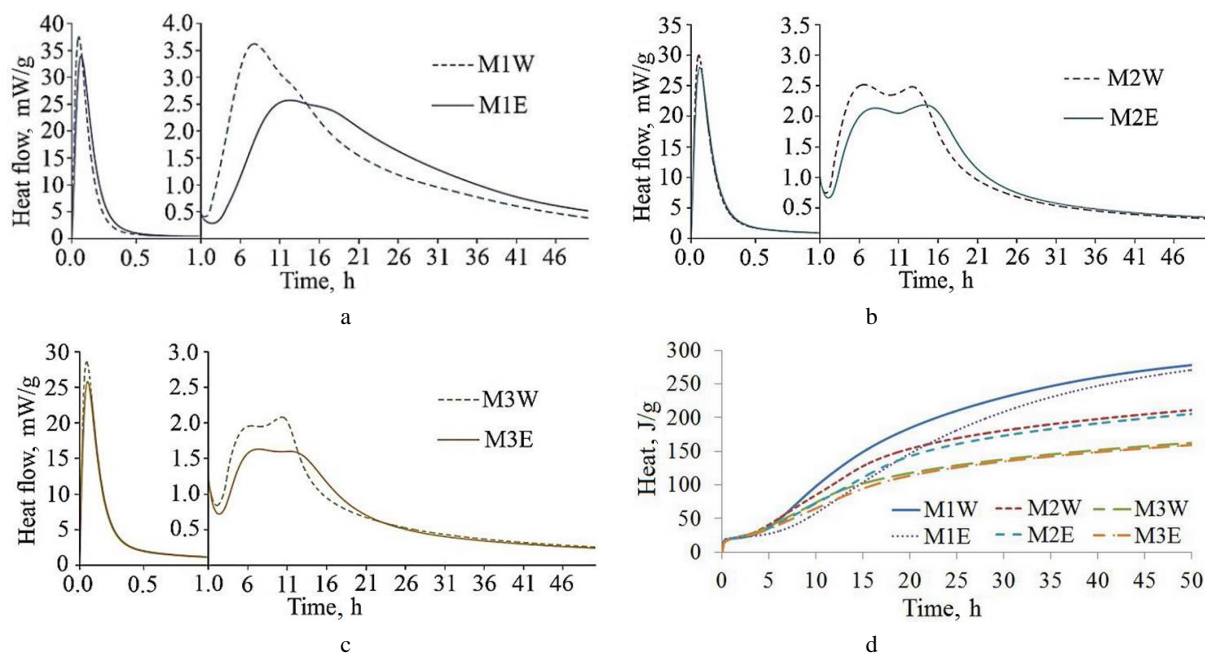
The process of the hydration of OPCs is commonly subdivided into several periods: initial (I), induction (II),

acceleration (III), and retardation (IV). It is possible to sum up the acceleration and the retardation periods into the “main period”. The calorimetric curve of many OPCs displays two significant heat flow maxima during the main period (Fig. 3).

The first is attained several hours after the beginning of the acceleration period, while the second one appears during the deceleration period.

The heat released during the first maximum of heat flow (called the “silicate reaction peak”) is mainly attributed to the heat release from the silicate reaction, which includes the dissolution of  $C_3S$  and the precipitation of portlandite and C-S-H phase. The second heat flow maximum (called the “sulfate depletion peak”) is characterized by heat release due to the dissolution of  $C_3A$  and the secondary formation of ettringite [20–22].

Three different periods of time were defined as proposed by Sauvat et al. [23], i.e.,  $t_1$ ,  $t_2$  and  $t_3$ , which correspond to the minimal dormant time, the first peak time, and the second peak time, respectively. The increased amount of pozzolana (from 33 % to 50 %) prolongs the dormant period to 1.69 and 2.05 hours in comparison with the control cement specimens without opoka, whose dormant period is 1.46 hours (Table 10).



**Fig. 3.** Calorimetric curves of thermal flow in Ordinary Portland cement pastes (a) and in pastes where cement was partially replaced by 33 % (b) or 50 % (c) of opoka prepared using standard water and buckwheat husk extract, and hydration heat in the samples (d)

**Table 10.** The influence of extracts on hydration characteristic times and enthalpy

Mix	First exothermal effect		Induction period $t_1$ , h	Second exothermal effect $t_2$		Third exothermal effect $t_3$		Heat of hydration, J/g	
	Peak time, h	Heat flow, mW/g		Peak time, h	Heat flow, mW/g	Peak time, h	Heat flow, mW/g	24 h	48 h
M1W	0.056	37.60	1.46	7.75	3.63	12.8	2.80	205.5	274.9
M1E	0.075	34.04	2.37	12.87	2.58	17.5	2.42	174.9	266.8
M2W	0.061	30.07	1.69	6.56	2.52	12.7	2.49	166.5	208.5
M2E	0.066	28.08	1.98	8.14	2.14	14.2	2.19	157.6	202.8
M3W	0.061	28.62	2.05	6.39	1.96	10.3	2.09	126.8	160.3
M3E	0.064	25.05	2.33	7.44	1.58	11.4	1.55	124.9	158.8

In general, natural pozzolana seems to accelerate the hydration of  $C_3S$  (the  $t_2$  peak time decreased from 7.75 h (control) to 6.56 – 6.39 h). The acceleration of the early hydration of  $C_3S$  caused by pozzolana has been ascribed to the fineness of pozzolana particles, which offer a greater surface for the precipitation of C-S-H from the pore solution. In other words, acceleration is not due to the pozzolanic reaction which can be observed, at the very best, after several days.

The general rate of heat evolution-time curve of Portland cement shows that the rate of retardation does not change significantly with pozzolana incorporation. However, one peculiarity of this period that should be noted is related to the shoulder or the peak that results from the renewed ettringite AFt formation. As the amount of pozzolana incorporation is increased, the peak becomes extraordinarily high. In fact, this may be the delayed  $C_3S$  hydration peak due to an excessive amount of pozzolana present in the cement rather than an AFt peak, or a combination of both. Another possible reason would be the amount of gypsum being higher than necessary. The gypsum content is so adjusted in cements that  $C_3A$  forms ettringite while the cement paste is still plastic. Therefore, the excess gypsum might have resulted in the conversion of monosulfate phase into ettringite. This conversion may be responsible for the accentuated second peak in the rate of the heat evolution curve.

The buckwheat husk extract particularly delays and inhibits the Portland cement hydration and extends the induction period from 1.46 h for pure cement to 2.37 h for paste prepared using the buckwheat husk extract. The  $t_2$  value increases from 7.75 to 12.87 h for the Portland cement pastes prepared with the buckwheat husk extract. This result confirms that the hydration of silicate phases is significantly delayed by buckwheat husk extractives.

Upon replacing part of the Portland cement with opoka, the influence of the buckwheat husk extract on the hydration of the binding material substantially changes. The calorimetric curves of Ordinary Portland cement replaced with 33 % (Fig. 3 b) and 50 % (Fig. 3 c) (by weight) of opoka pastes prepared using the standard water and buckwheat husk extract are similar (Fig. 3 b, c). Opoka additive minimized influence of buckwheat husk extract to cement hydration. In specimens with opoka are hydrating a less amount of  $C_3S$  (33 and 50 % (by weight) of cement replaced by opoka), but until 19 h and 13 h hydration (respectively) they emit more heat than pure cement prepared with the extract (Fig. 3 d).

### 3.4. Investigation of the buckwheat husk concrete properties

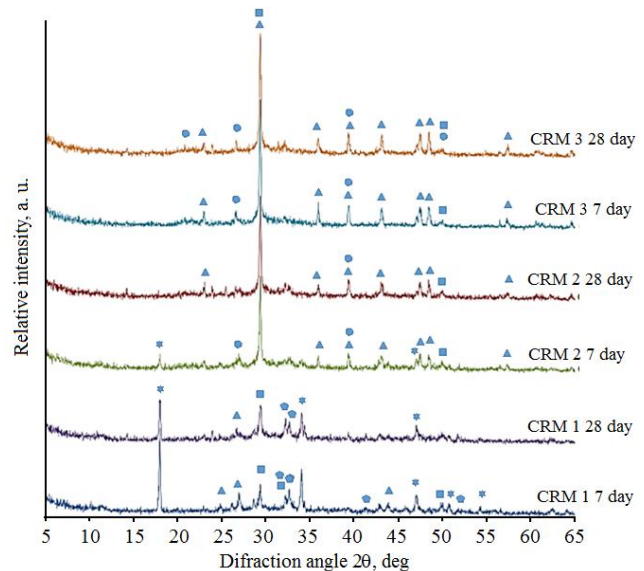
At a corresponding time, the compressive strength of the wet buckwheat husk concrete samples was measured. Crushed samples were weighed and then dried at a temperature of 80 °C until a constant mass.

Concrete samples without sand addition are relatively weak and brittle. The replacement of part of buckwheat husk by sand increases the density of the concrete and at the same time significantly increases the compressive strength of a sample. After the test, such sample deforms but not crumbles. A summary of the test results regarding

the compressive strength, moisture content and dry density of buckwheat husk concrete is given in Table 11.

**Table 11.** The test results of dry density and compressive strength of the wet buckwheat husk concrete samples

Concrete mixture	Dry density, kg/m <sup>3</sup>	Compressive strength, MPa		
		Moisture content of samples, %		
		3 day	7 day	28 day
CRM1	925	<u>1.58</u>	<u>2.51</u>	<u>3.01</u>
		22.83	15.88	16.90
CRM2	845	<u>1.68</u>	<u>2.53</u>	<u>3.08</u>
		21.37	14.82	14.38
CRM3	809	<u>1.33</u>	<u>2.16</u>	<u>2.91</u>
		22.06	16.02	16.30
		7 day	28 day	3 month
CRM1S	1410	<u>3.87</u>	<u>5.86</u>	<u>7.45</u>
		23.21	15.22	19.99
CRM2S	1309	<u>5.53</u>	<u>7.35</u>	<u>8.56</u>
		22.22	17.70	25.95
CRM3S	1263	<u>4.79</u>	<u>6.76</u>	<u>8.19</u>
		22.86	19.68	23.53



**Fig. 4.** X-Ray diffraction pattern of dried buckwheat husk concrete mixtures (CRM1, CRM2, CRM3) after 7 and 28 days of hydration. Indexes: ● – SiO<sub>2</sub>; ▲ – CaCO<sub>3</sub>; ■ – CSH; ★ – Ca(OH)<sub>2</sub>; ◆ – C<sub>3</sub>S

The experimental results show that after 28 days of hydration the compressive strength and dry bulk density of all specimens (without and with opoka replacement) of concrete without sand range within 2.91 – 3.08 MPa and 809 – 925 kg/m<sup>3</sup>, respectively. For concrete with an additive of sand, these values change within 5.86 – 7.35 MPa and 1263 – 1410 kg/m<sup>3</sup>, respectively. For all mixtures, there is satisfied the lower limit (2.0 – 17.0 MPa and 800 – 1440 kg/m<sup>3</sup>) to be used for structural/insulating purposes [3].

Results of investigations show that the properties of buckwheat husk concrete are better when not pure cement is used, but when 33 % of cement is replaced with opoka.

In order to make a more exact estimation of the hydration process, dry samples for X-Ray analysis were prepared without the sand that is usually used. Specimens were ground to a powder, and the fibrous fraction (fibres of buckwheat husk) was eliminated using a 50 μm sieve.

The X-Ray diffraction investigation data (Fig. 4) show that the hydration of cement with opoka replacement is more complete, and after 28 days no peaks typical of initial cement minerals in X-Ray diffraction pattern curves of dried buckwheat husk concrete mixtures are identified.

#### 4. CONCLUSIONS

A comparison of IR spectroscopy data of aqueous and alkaline buckwheat husk extracts' composition showed the prevalence of various polysaccharides. The dry extract yield in the buckwheat husk was  $4.4 \pm 0.2$  % relative to the mass of the starting raw material.

The buckwheat husk extract increases both the initial and the final setting time of OPC. The partial (33 % and 50 %) replacement of cement with opoka significantly decreases the influence of the extract on the initial setting time and minimizes the influence on the final setting time.

The thermogravimetric investigation has revealed a strong modification and delay of silicate phase hydration by buckwheat husk extractives.

The test results show that the buckwheat husk aggregate and cement-opoka binding material combinations have a potential to be used in the production of a new lightweight concrete. The compressive strength of buckwheat husk concrete ranges from 3.0 to 8.0 MPa, and its dry bulk density changes from 800 to 1400 kg/m<sup>3</sup>.

The results of the study show that better properties of buckwheat husk concrete are obtained when not pure cement is used, but when 33 % of cement is replaced with opoka.

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