Effect of Phosphorylation and Copper(II) or Iron(II) Ions Enrichment on Some Physicochemical Properties of Spelt Starch

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ABSTRACT

This paper provides an assessment of the effect of saturation of spelt starch and monostarch phosphate with copper or iron ions on selected physicochemical properties of the resulting modified starches. Native and modified spelt starch samples were analyzed for selected mineral element content using Atomic Absorption Spectroscopy (AAS). Thermodynamic properties were measured using DSC, and pasting properties by RVA. Flow curves of 5% pastes were plotted and described using the Herschel-Bulkley model. The structure recovery ratio was measured. AAS analysis established the presence of iron(II) and copper(II) ions in the samples of modified starches and that potassium and magnesium ions had leached from them. In comparison to unfortified samples, enriching native starch with copper(II) ions decreases value of all temperatures of phase transformation about 1.3-2.7 °C, but in case of monostarch phosphates bigger changes (2.8-3.7 °C) were observed. Fortified native spelt starch with copper(II) ions caused increasing the final viscosity of paste from 362 to 429 mPa·s. However, presence iron(II) ions in samples caused reduced its final viscosity by 170 (spelt starch) and 103 mPa·s (monostarch phosphate). Furthermore, enriching monostarch phosphate contributed to reduce degree of structure recovery of pastes from 70.9% to 66.6% in case of copper(II) ions and to 59.9% in case of iron(II) ions.

Key words: chemically modified starch, enrichment, phosphorylation, spelt starch

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INTRODUCTION

Starch is one of the most widely used ingredients in the food industry, but its range of application extends to other industries, including pharmaceuticals. It is used for thickening, stabilization, texture smoothing, aroma encapsulation and fat substitution as well as affecting the moisture content of a product (Kaur et al. 2012). This polymer is easily modified by physical and chemical treatment, and therefore manufacturers often use starch to adjust the properties of their products. New functional, physical and chemical properties imparted to modified starches have brought new food products onto the market (Hung and Morita 2005). The use of various starch modification methods enables to obtain products with different properties; therefore, selecting the appropriate method of modification starch is a crucial part of the manufacturing process. The use of modified starch in food technology must impart the desired properties to food products, but should not hinder the production process. Consequently, selection of starch should take into account nutritional, sensorial, production, distribution and marketing requirements.

The chemical modification of starch is based on the reactivity of amylase and amyllopectin functional groups. For the food industry, starch phosphates are particularly important. They are created by incorporating acid substituents of orthophosphate into the starch. The method of enrichment the acid molecule determines whether a mono- or di-starch phosphate is obtained (Fortuna and Rożnowski 2002).

The food industry approves only of starch phosphates with a low substitution ratio so as not to alter the calcium to phosphorus ratio in food products (Fortuna and Rożnowski 2002). Monostarch phosphates are used as thickening agents in soups, sauces, dressings and, in particular, cold jelly desserts, as well as being ingredients in puddings, thermized cottage cheese, food concentrates, milk shakes and low calorie margarine. EU Commission Regulation 1129/2011 lists starch phosphate as an additive authorized for food use at quantum satis levels.

Corn, wheat, rice and potatoes are the major sources of starch used worldwide, but new sources of starch with new functional properties are constantly being sought. Although spelt starch has been known in the food industry for some time, this species of wheat is experiencing a renaissance. Spelt wheat belongs to the Gramineae Juss family (Campbell 1997); however, there are no definite data on the origins of spelt and there is some uncertainty about its classification. Most sources classify spelt as a subspecies of common wheat, Triticum aestivum, although some regard it as a separate species (Blatter et al. 2003; Campbell 1997; Kohajdová and Karovičová 2008). Modern life-styles are altering dietary habits: lack of time and constant hurry have increased the demand for quickly prepared convenience foods which have been heavily processed and frequently decreased of important nutrients. Therefore, the nutritional value of such products is generally lower than that of traditionally prepared meals. These and other factors have led to the growth of vitamin and mineral supplementation in foods (Anonymous 1999; Lozoff et al. 2012; Ma and Betts 2000). In order to meet the growing quality expectations of consumers, new convenient and safe methods of supplying minerals should be developed, not only preserving nutritional qualities but even enhancing them.

Anemia, the result of low levels of hemoglobin in the blood, was the most widespread blood disease at the end of 20th century. Iron deficiency, the main cause, is responsible for 80% of cases, although less commonly, vitamin B12 or folic acid deficiency may also be factors. Anemia often affects young people and women in reproductive age. Despite low iron intake in the diet and frequent cases of anemia, iron enrichment of food is limited to a narrow range of products (Cook 1995; Cook 2005; Zimmermann and Hurrell 2007).

Copper is another important microelement. As a component of cofactors, it is involved in oxidation and reduction processes, and affects the metabolism of iron and collagen. Infants with low birth weight are the major risk group for copper deficiency. Adults are rarely affected, although it may be triggered by zinc supplementation or high fructose consumption (Anonymous 2010; Festa and Thiele 2011; Hong et al. 2001).

This paper provides an assessment of the effect of the phosphorylation of spelt starch and the enrichment of spelt starch and monostarch phosphate with copper(II) and iron(II) ions on selected physical and chemical properties of the resulting starch samples. It also evaluates the level of ion incorporation during the modification treatment and seeks to ascertain whether changes in the properties of modified starches are significant when compared with natural starch.
MATERIAL AND METHODS

The research material was spelt starch, extracted from spelt flour (ECCO 700) under laboratory conditions. The flour, milled in 2010, was obtained from Młyń Gospodarczy, Radzyń Podlaski, Poland.

Spelt starch preparation

Spelt starch was isolated from spelt flour following the procedure described by Moreira et al. (2012), with minor changes. Spelt flour (1 kg) was soaked in 500 mL of distilled water at room temperature. The suspension was placed on a nylon sieve (100 μm mesh) and the kneaded dough was rinsed with distilled water. The dough was continuously kneaded and rinsed until pure gluten was obtained. The resulting starch milk was poured into 200 mL plastic centrifuge vessels and centrifuged for 20 minutes at 4000 rpm. After spinning, the supernatant was poured off and the layer of impurities was removed with a spatula. A new portion of starch milk was added and the whole was mixed and centrifuged again. The process was repeated three times, after which the supernatant and impurities were removed, distilled water was added, and the whole was mixed and centrifuged again. After the final spinning, the supernatant was poured off, the impurities was removed, and the sediment was transferred to plastic dishes, ground and air-dried at room temperature while being constantly stirred. After drying, the polymer was ground in a Retsch RM-200 mortar and filtered through a 125 μm Retsch AS-200 sieve. The obtained spelt starch contained 20.89% amylose, 0.27% protein and 9.4% moisture.

Phosphorylation

The spelt starch was used to make monostarch phosphates using Neukom’s modified method (Richter et al. 1968). Spelt starch (216 g d.m.) was rinsed with water in a 1G Schott funnel, and then transferred to a vessel, to which 12.28 g of sodium dihydrophosphate and 7.81 g of sodium hydrophosphate diluted in a small amount of water were added. The mixture was stirred for 5 minutes. The resulting starch suspension was dried in Petri dishes at room temperature to a moisture content of about 10%. The dried mixture was ground in a porcelain mortar, and then heated for 1 hour in an Erlenmeyer flask over an oil bath at 160°C while being constantly stirred. The cooled powder was added to 150 cm³ of 60% ethanol, and filtered through a 1G Schott funnel. This process was repeated four times with 96% ethanol used for the last run. The resulting sample was transferred to Petri dishes and dried at room temperature to air humidity. After drying, the resulting monostarch phosphates were ground in a Retsch, RM-200 mortar and filtered through a 125 μm Retsch AS-200 sieve. Both the spelt starch and the spelt monostarch phosphate were subjected to saturation with copper(II) and iron(II) ions.

Copper(II) and Iron(II) ions saturation

The native spelt starch and the spelt monostarch phosphate were enrichment with copper(II) ions according Przetaczek-Rożnowska et al. (2012) method and iron(II) ions according to procedure described by Rożnowski et al. (2014). The starch sample (100 g) were stirred for 5 minutes with deionized water (250 cm³) and filtered. Next, the starch was transferred to a G3 Schott funnel and stirred for 5 minutes at 330 rpm with 200 cm³ of salts solution (39.6 g CuSO₄·5H₂O in 1 dm³ of water or 18.31 g FeSO₄·5H₂O in 1 dm³ of water). The procedure was repeated twice. After completion, the sample was rinsed in deionized water until sulfide ions were no longer present (lack of white sediment during reaction with BaCl₂). The resulting starch was air dried on paper tissue to a moisture content of 10%, then ground in a Retsch RM-200 grinder. After grinding, the samples were filtered through a 125 μm Retsch AS-200 sieve.

Determination of mineral content

Spelt starch, monostarch phosphate and samples enriched with minerals were analyzed for potassium, magnesium, copper and iron content by AAS. The measurements were performed using an atomic absorption spectrometer (Avanta Sigma, GBC, Australia) following Antanasopoulos’ (2004) method, selecting wavelength λ for each of the elements: K – 769.9 nm, Mg – 285.2 nm, Ca – 422.7 nm and Fe –248.3 nm. Limit of detection (LOD) and limit of quantification (LOQ) were determined by measuring absorbance according to Doerffel (1989). The content of phosphorus was measured according to ISO 3946:1982.

Color analysis

The color of the starches under investigation was determined using an X-Rite Color i5.
spectrophotometer. \( D_{65} \) illuminants were applied for the d/8 geometry and 10 mm measurement slot. Color was determined in specular excluded mode for CIE 1964 additional observer. The CIE \( L^*, a^*, b^* \) coordinates were determined using X-Rite Color Master SW, while the saturation values (\( C^* \)) was computed according to Yam and Papadakis (2004).

**Thermodynamic properties of starch gelatinization**

The thermodynamic parameters of gelatinization were measured using a Netzsch DSC 204 F1 Phoenix differential scanning calorimeter (Netzsch GmbH, Germany). In order to determine the thermal properties of the samples, 3.5 mg of sample and 10.5 mg water were stored at room temperature in calorimetry pans, sealed hermetically and stored for 24 hours at room temperature. The temperature of the samples was increased from 20 to 110°C at a rate of 10°C/min. After analysis, the pans were stored at 4°C for 7 days, after which the thermodynamic properties of retrogradation were determined in a similar manner. An empty calorimetry pan was used as the reference. The resulting thermograms enabled the temperatures specific for gaseous transformations to be determined.

**Pasting properties (RVA)**

Pasting properties were determined using a Rapid Visco Analyzer (RVA). A 5% suspension of starch in distilled water was prepared in metal vessels. The RVA procedure was conducted as follows: samples were stirred at 960 rpm for 1 minute at 50°C, then the starch suspension was stirred at the same temperature for 1 minute with a speed 160 rpm., Next, the samples was heated from 50° to 95°C with a heating rate 4.5°C/min. The temperature 95°C was maintained for 5 minutes. Then the sample was cooled down to 50°C in the space of 10 min and kept at that temperature for 5 min, during which time was the function of viscosity against time constantly measured.

**Flow curves**

The flow curves of monostarch phosphate and spelt starch pastes as well as samples enriched with copper(II) or iron(II) ions were charted using a Rheolab MC1 rotational rheometer (Physica Meßtechnic GmbH, Germany), with Z3 DIN coaxial cylinders (25.00 mm internal, 27.12 mm external diameter) as the measurement system. The rheometer and thermostat (Viscotherm2) were controlled by US 200 software (Physica Meßtechnic GmbH, Germany). To determine flow curves, 5% w/w pastes of starch in distilled water were prepared. These samples were stirred at 300 rpm for 15 minutes at room temperature. Next, the vessel was heated for 30 minutes in a 95°C ± 1°C water bath, constantly stirred at 300 rpm. The paste was then immediately placed in the rheometer measurement vessel, and kept for 10 minutes at a constant temperature of 50°C ± 0.5°C.

The flow curves were charted for the increasing shear rate in the range of 1-500 s\(^{-1}\), over 3 minutes, for the constant 500 s\(^{-1}\)shear rate over 2 minutes, for the diminishing shear rate in the range of 500-1 s\(^{-1}\), over 3 minutes.

The curves were described according to the Herschel-Bulkley rheology model

\[
\tau = \tau_0 + K(\dot{\gamma})^n
\]

where: \( \tau \): shear stress (Pa), \( \tau_0 \): the yield strength (Pa), \( K \): consistency index (Pa·s\(^n\)), \( \dot{\gamma} \): shear rate (s\(^{-1}\)), \( n \): flow index. The parameters were computed with the use of the US 200 SW.

**Degree of structure recovery of starch paste Time - test**

The recovery ratio of 5% starch paste was measured in the manner described for determining flow curves. The samples were kept in the measurement vessel of the rotational rheometer for 10 minutes in 50°C ± 0.5°C and then analyzed using the modified Mezger (2002) and Pająk et al. (2012) procedure, with constant shear rates of: 1 s\(^{-1}\), for 3 minutes; 500 s\(^{-1}\), for 3 minutes; and 1 s\(^{-1}\), for 10 minutes.

The ratio of structure reconstruction (\( R \)) was computed as percentage of apparent viscosity (\( \eta_2 \)) obtained during the first minute of the third step based on the apparent viscosity value (\( \eta_1 \)) determined in the first 3 minutes of measurement:

\[
R = \left( \frac{\eta_2}{\eta_1} \right) \times 100\%
\]

**Statistical analysis**

All calculations and charts were carried out using Excel 2007. All treatments were repeated three times. The results were tested for \( \alpha = 0.05 \) (Tukey's HSD test) using Statistica 8.0 SW to identify homogeneous groups.
RESULTS AND DISCUSSION

Mineral components of native and modified starch

The analyses indicated that both phosphorylation and the enrichment with copper(II) or iron(II) ions of spelt starch and the resulting monostarch phosphates lead to decrease of potassium and magnesium naturally present in the starch (Table 1). Magnesium, although being the least abundant of the minerals present in the starch and monostarch phosphate examined in this investigation, was the most resistant to leaching. Similar results have been observed in the saturation of potato and corn starches (Różnowski et al. 2014). The fact that the leaching rate of calcium and magnesium from the starch and monostarch phosphate was lower than that of potassium demonstrates that bivalent ions are much more strongly bound by such features as "calcium bridges" (Pałasiński, 1968), which resist leaching much more effectively than univalent ions. Phosphorus is an important mineral component in starch grains; its content varies depending on the plant source (Swinkels 1985; Singh et al. 2003). Saturation with minerals resulted in a statistically significant loss of phosphorus (Table 1). Różnowski et al. (2014) also reported leached phosphorus from potato starch granules during modification with FeSO₄, whereas phosphorus content in corn starch after rinsing with ions solutions showed no statistically significant changes relative to control corn starch.

Table 1 - Mineral content in the starch samples examined (mg/100 g dry matter)

<table>
<thead>
<tr>
<th>Samples</th>
<th>K</th>
<th>Mg</th>
<th>Ca</th>
<th>Cu</th>
<th>Fe</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>ss</td>
<td>22.32±0.13</td>
<td>4.56±0.27</td>
<td>8.15±0.19</td>
<td>nd</td>
<td>nd</td>
<td>61.97±0.79³</td>
</tr>
<tr>
<td>ss_Cu</td>
<td>nd</td>
<td>nd</td>
<td>nd</td>
<td>11.17±0.32</td>
<td>nd</td>
<td>37.48±0.54⁴</td>
</tr>
<tr>
<td>ss_Fe</td>
<td>nd</td>
<td>0.67±0.37</td>
<td>nd</td>
<td>nd</td>
<td>29.93±0.75</td>
<td>51.67±0.64⁴</td>
</tr>
<tr>
<td>LOD</td>
<td>1.9</td>
<td>0.3</td>
<td>0.6</td>
<td>0.3</td>
<td>1.00</td>
<td>-</td>
</tr>
<tr>
<td>LOQ</td>
<td>3.3</td>
<td>0.5</td>
<td>1.2</td>
<td>0.5</td>
<td>2.01</td>
<td>-</td>
</tr>
<tr>
<td>ps</td>
<td>15.2±0.10</td>
<td>3.97±0.55</td>
<td>9.93±0.06</td>
<td>nd</td>
<td>nd</td>
<td>1011.56±1.02⁴</td>
</tr>
<tr>
<td>ps_Cu</td>
<td>nd</td>
<td>nq</td>
<td>3.43±0.06</td>
<td>503.6±4.01</td>
<td>nd</td>
<td>211.15±0.47⁵</td>
</tr>
<tr>
<td>ps_Fe</td>
<td>nq</td>
<td>3.57±0.15</td>
<td>1.20±0.10</td>
<td>nd</td>
<td>392.63±2.21</td>
<td>215.10±0.68⁸</td>
</tr>
<tr>
<td>LOD</td>
<td>0.6</td>
<td>0.1</td>
<td>0.5</td>
<td>0.3</td>
<td>7.7</td>
<td>-</td>
</tr>
<tr>
<td>LOQ</td>
<td>1.2</td>
<td>0.2</td>
<td>1.1</td>
<td>0.5</td>
<td>15.3</td>
<td>-</td>
</tr>
</tbody>
</table>

ss = spelt starch (native); ss_Cu = spelt starch enriched with copper(II) ions; ss_Fe = spelt starch enriched with iron(II) ions; ps = spelt monostarch phosphate; ps_Cu = spelt monostarch phosphate enriched with copper(II) ions; ps_Fe = spelt monostarch phosphate enriched with iron(II) ions; nd = below Limit of Detection (LOD); nq = between Limit of Detection (LOD) and Limit of Quantification (LOQ).

The phosphorus present in starch affects its ion exchange properties (Richter 1968; Jane et al. 1996; Lin and Czuchajowska 1998; Różnowski et al. 2014), which was confirmed in the present study: during saturation, spelt monostarch phosphate absorbed ten times more iron(II) ions than natural starch, and about forty five times more copper (Table 1). Natural starch absorbed iron better than copper, while the monostarch phosphate absorbed copper more effectively (Table 1). This shows that the effectiveness of mineral enrichment depends on both the nature of the starch and the mineral being enriched.

Color analysis

The color parameters were analyzed in order to investigate the influence of presence of copper(II) or iron(II) ions on color of modified starches, as a first factor in sensorial impression. The derived color parameters indicate that both enrichment of spelt starch and its phosphorylation affect color (Fig. 1). All colors parameter were compared to those in the unfortified starch samples. Mineral saturation decreased the lightness of the natural starch from 97.46 to 94.05 for starch enriched with copper(II) and to 92.01 for native starch fortified with iron(II) ions. As well the monostarch phosphate had distinctly lower L* values (by 12 units) than native spelt starch, which shows that starch modification, whatever the type, reduces lightness. Similar results have been observed in the
saturation of potato and corn monostarch phosphate (Rożnowski et al. 2015) and was also reported for phosphorylated spelt starch obtained during heating at 130 or 145 °C for 2 or 3 hours (Rożnowski et al. 2016a) or phosphorylated starch obtained from pumpkin starch during heating at 145 °C (Rożnowski et al. 2016b).

Figure 1- Value of color parameters of native and phosphorylated spelt starch and enriched with iron or copper ions (a) value of Lightness; (b) chroma of the color c*; (c) value of parameter a*; (d) value of parameter b*. Values expressed are mean ± standard error (n = 3). Values followed by the same letter in a figure are not significantly different at α = 0.05.

Values of a* or b* parameters examined for saturated sample with copper(II) ions were lower than values obtained for unfortified samples. On the other hand, the native spelt starch and monostarch phosphate enriched with iron(II) ions were characterized by 3 times higher value of a* parameter. Our research confirmed observation reported for phosphorylated potato or corn starch saturated with copper(II) or iron(II) ions (Rożnowski et al. 2015). In case the b* parameter there did not observed constant relationship. The value of this parameter for native starch saturated with iron(II) ions was over 4 times higher than value obtained for native spelt starch, meantime the monostarch phosphate fortified with iron(II) ions was characterized by lower value of b* parameter (12.54) than unfortified monostarch phosphate (18.03). Our research did not confirm observation reported for phosphorylated potato or corn starch (Rożnowski et al. 2015) were enrichment phosphorylated corn or potato starch with iron(II) ions caused increase value of b* parameter.

Unenriched monostarch phosphate had higher color saturation (C*) value in comparison with native spelt starch. Starch enriched with copper(II) had the lowest saturation (1.81) what confirmed observation reported for potato or corn phosphorylated starches (Rożnowski et al. 2015).

DSC of the thermodynamic properties

The value of gelatinization temperatures (T_o, T_p, and T_c) of analyzed spelt starch were similar to those found for wheat starch by Aggarwal and Dollimore (1998). However, there were lower than those published for wheat starch by Nakazawa and Wang (2003) and higher than those recorded also for wheat starch by Jacobs et al. (1998) or Gunaratne and Corke (2007). The gelatinization temperatures as T_o, T_p and T_c are indicators of the stability of starch granules or the starch crystal structure in water (Aparicio-Saguilan et al. 2005; Lehmann et al. 2002). Morikawa and Nishinari (2000) indicated that starch modification changes gelatinization and reorganizes the structure patterns in starch. Polymer esterification increases the peak temperature (T_p) (Sagar and Merrill 1995; Rajan et al. 2006), which was confirmed in the present work: spelt monostarch phosphate showed markedly higher T_p than natural spelt starch (Table 2). This is attributed to alterations of the crystal structure and change of the amorphous areas of starch granules (Tester and Debon 2000). The DSC of monostarch spelt phosphate also showed considerably higher end temperature (T_c), both in the initial measurement, and after 7 days' storage (Table 2). The enrichment of copper(II) into native spelt starch and monostarch phosphate and iron(II) ions into spelt starch decreased the T_o, T_p and T_c of gelatinization during the first day. This may indicate structural changes of both the starch and the monostarch phosphate in reaction to mineral incorporation.
However, our research did not confirm this, which suggests that pasting parameters depend not only on the presence of phosphorus, but also on the mineral enriched in the starch.

Table 2 - Gelatinization parameters

<table>
<thead>
<tr>
<th>Samples</th>
<th>Gelatinization</th>
<th>Retrogradation</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>T&lt;sub&gt;g1&lt;/sub&gt; (°C)</td>
<td>T&lt;sub&gt;g2&lt;/sub&gt; (°C)</td>
</tr>
<tr>
<td>ss_cu</td>
<td>51.5±0.15&lt;sup&gt;a&lt;/sup&gt;</td>
<td>50.4±1.69&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>ss_fe</td>
<td>54.0±1.00&lt;sup&gt;b&lt;/sup&gt;</td>
<td>60.1±0.12&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>ps_cu</td>
<td>51.2±0.12&lt;sup&gt;c&lt;/sup&gt;</td>
<td>70.2±0.15&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>ps_fe</td>
<td>56.5±0.26&lt;sup&gt;c&lt;/sup&gt;</td>
<td>64.0±0.06&lt;sup&gt;e&lt;/sup&gt;</td>
</tr>
</tbody>
</table>

 ss = spelt starch (native); ss_cu = spelt starch enriched with copper(II) ions; ss_fe = spelt starch enriched with iron(II) ions; ps = spelt monostarch phosphate; ps_cu = spelt monostarch phosphate enriched with copper(II) ions; ps_fe = spelt monostarch phosphate enriched with iron(II) ions. Values followed by the same letter in a column are not significantly different at α = 0.05 (Tukey's HSD test). Each value is the mean ± standard deviation of triplicate measurements.

Table 3 - Pasting properties

<table>
<thead>
<tr>
<th>Samples</th>
<th>T&lt;sub&gt;p1&lt;/sub&gt; (°C)</th>
<th>PV(mPas)</th>
<th>TV(mPas)</th>
<th>BD(mPas)</th>
<th>FV(mPas)</th>
<th>SB(mPas)</th>
</tr>
</thead>
<tbody>
<tr>
<td>ss</td>
<td>87.1±1.69&lt;sup&gt;a&lt;/sup&gt;</td>
<td>205±9&lt;sup&gt;b&lt;/sup&gt;</td>
<td>190±10&lt;sup&gt;b&lt;/sup&gt;</td>
<td>232±20&lt;sup&gt;a&lt;/sup&gt;</td>
<td>362±18&lt;sup&gt;b&lt;/sup&gt;</td>
<td>366±19&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
<tr>
<td>ss_cu</td>
<td>85.2±0.18&lt;sup&gt;a&lt;/sup&gt;</td>
<td>332±1&lt;sup&gt;b&lt;/sup&gt;</td>
<td>226±2&lt;sup&gt;a&lt;/sup&gt;</td>
<td>307±2&lt;sup&gt;c&lt;/sup&gt;</td>
<td>429±1&lt;sup&gt;a&lt;/sup&gt;</td>
<td>404±1&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>ss_fe</td>
<td>86.9±0.01&lt;sup&gt;a&lt;/sup&gt;</td>
<td>153±1&lt;sup&gt;d&lt;/sup&gt;</td>
<td>69±1&lt;sup&gt;c&lt;/sup&gt;</td>
<td>137±2&lt;sup&gt;a&lt;/sup&gt;</td>
<td>192±4&lt;sup&gt;d&lt;/sup&gt;</td>
<td>175±4&lt;sup&gt;c&lt;/sup&gt;</td>
</tr>
<tr>
<td>ps</td>
<td>78.1±0.42&lt;sup&gt;ab&lt;/sup&gt;</td>
<td>235±4&lt;sup&gt;b&lt;/sup&gt;</td>
<td>82±1&lt;sup&gt;c&lt;/sup&gt;</td>
<td>153±4&lt;sup&gt;d&lt;/sup&gt;</td>
<td>228±2&lt;sup&gt;e&lt;/sup&gt;</td>
<td>146±2&lt;sup&gt;c&lt;/sup&gt;</td>
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<tr>
<td>ps_cu</td>
<td>65.2±0.85&lt;sup&gt;c&lt;/sup&gt;</td>
<td>312±8&lt;sup&gt;e&lt;/sup&gt;</td>
<td>85±2&lt;sup&gt;c&lt;/sup&gt;</td>
<td>228±14&lt;sup&gt;a&lt;/sup&gt;</td>
<td>182±4&lt;sup&gt;d&lt;/sup&gt;</td>
<td>97±1&lt;sup&gt;d&lt;/sup&gt;</td>
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<td>ps_fe</td>
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<td>216±12&lt;sup&gt;b&lt;/sup&gt;</td>
<td>35±4&lt;sup&gt;d&lt;/sup&gt;</td>
<td>292±6&lt;sup&gt;a&lt;/sup&gt;</td>
<td>125±3&lt;sup&gt;c&lt;/sup&gt;</td>
<td>90±3&lt;sup&gt;d&lt;/sup&gt;</td>
</tr>
</tbody>
</table>

 ss = spelt starch (native); ss_cu = spelt starch enriched with copper(II) ions; ss_fe = spelt starch enriched with iron(II) ions; ps = spelt monostarch phosphate; ps_cu = spelt monostarch phosphate enriched with copper(II) ions; ps_fe = spelt monostarch phosphate enriched with iron(II) ions; T<sub>p</sub> = pasting temperature; PV = peak viscosity; TV = trough; BD = breakdown; FV = final viscosity; SB = setback.
Values followed by the same letter in a column are not significantly different at α = 0.05 (Tukey's HSD test). Each value is expressed as the mean ± standard deviation of triplicate measurements.

**Flow curves and structure recovery of starch paste**

The flow curves of 5% pastes of starch and the resulting monostarch phosphate, as well as of samples enriched with copper(II) and iron(II), are shown in Figure 2, while the Herschel-Bulkley parameters describing the curves are shown in Table 4.

**Figure 2** - Flow curves of 5% pastes (a) ss = native spelt starch; ss_Cu = spelt starch enriched with copper ions; ss_Fe = spelt starch enriched with iron ions; (b) ps = spelt monostarch phosphate; ps_Cu = spelt monostarch phosphate enriched with copper ions; ps_Fe = spelt monostarch phosphate enriched with iron ions.

**Table 4** - Herschel-Bulkley parameters for 5% pastes and structure recovery (R%) of starch paste

<table>
<thead>
<tr>
<th>Samples</th>
<th>Herschel-Bulkley parameters</th>
<th>Recovery (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>$\tau_0$ (Pa)</td>
<td>K (Pa · s$^n$)</td>
</tr>
<tr>
<td>ss</td>
<td>4.38±1.0$^a$</td>
<td>1.06±0.11$^b$</td>
</tr>
<tr>
<td>ss_Cu</td>
<td>3.17±0.4$^a$</td>
<td>1.52±0.08$^a$</td>
</tr>
<tr>
<td>ss_Fe</td>
<td>4.19±1.2$^a$</td>
<td>2.60±0.75$^a$</td>
</tr>
<tr>
<td>ps</td>
<td>4.03±0.3$^a$</td>
<td>0.43±0.03$^c$</td>
</tr>
<tr>
<td>ps_Cu</td>
<td>3.73±0.5$^a$</td>
<td>0.51±0.15$^c$</td>
</tr>
<tr>
<td>ps_Fe</td>
<td>4.30±1.0$^a$</td>
<td>0.34±0.06$^c$</td>
</tr>
</tbody>
</table>

ss = spelt starch (native); ss_Cu = spelt starch enriched with copper(II) ions; ss_Fe = spelt starch enriched with iron(II) ions; ps = spelt monostarch phosphate; ps_Cu = spelt monostarch phosphate enriched with copper(II) ions; ps_Fe = spelt monostarch phosphate enriched with iron(II) ions; $\tau_0 =$ shear stress; K = consistency index; n = flow index.

Values followed by the same letter in a column are not significantly different at α = 0.05 (Tukey's HSD test). Each value is expressed as the mean ± standard deviation of triplicate measurements.
The rheological measurements indicate that over the range of the shear rates applied, starches behaved as non-Newtonian shear-thinning fluids that are close to the yield strength (Fig. 2, Table 4). The shear stress of the 5% pastes was affected over the entire range of measurement by both the enriched mineral and the chemical modification (Fig. 2). Enrichment native spelt starch with copper(II) contributed to a downshifting of the flow curves, while iron(II) caused an increase in the shear stress. Spelt monostarch phosphates behaved differently. The presence of iron(II) ions led to a decrease in shear stress over the entire range of measurement, while copper(II) ions did not affect the shear stress compared with the pure monostarch phosphates. Similar correlations have been reported with respect to natural potato and corn starches enriched with iron (Rożnowski et al. 2014); for any given shear rate, enriched pastes had much lower shear stress than natural starch pastes. Neither phosphorylation nor mineral enrichment affected the flow index (n) (Table 4). Rożnowski et al. (2014) reported otherwise, finding that the enrichment of potato starch with iron considerably lowered the flow index, while similar treatment of corn starch increased this value. This shows that plant origin affects the rheology of enriched samples. It is probable that origin of starch determines how the incorporated minerals are bound, thus also affecting the rheology of the samples. This is confirmed by the neutral impact of phosphorylation and copper(II) or iron(II) enrichment on the yield strength (τ₀), (Table 4), and the strong impact on this parameter of iron enrichment in potato and corn starch.

Spelt starch saturated with copper(II) or iron(II) had considerably higher values for the consistency index (K), indicating that the viscosity of the enriched pastes was considerably higher than that of the natural starch pastes (Table 4). The native spelt starch enriched with ions did not indicate the difference in structure recovery of starch pastes compared with native spelt starch paste without ions (Table 4). Research carried out on the enrichment of iron(II) in potato and corn starch (Rożnowski et al. 2014) showed a considerable weakening of structure recovery. The presumption is that the ions incorporated in the starch bind differently in starches extracted from different plants, thus enabling new intermolecular bonds to be created in spelt starch but not in potato and corn starch. However, the combination of modification (esterification and enrichment with ions) contributed to weakening of structure recovery.

CONCLUSIONS

Natural spelt starch as well as spelt monostarch phosphate effectively absorbed iron(II) and copper(II) ions and can be used as carriers in mineral supplementation. However, the degree of mineral saturation of spelt starch or monostarch phosphate depends on both the type of the starch and the mineral incorporated in the polymer. Chemical modification of starch, both by phosphorylation and enrichment, leads to decrease of elements naturally present in this starch, including phosphorus, potassium and magnesium. DSC thermal analyses of gelatinization showed that starch enrichment had a significant effect on gelatinization parameters. The extent of the changes depended on the type of starch and the mineral enriched. The pasting properties showed that the presence of iron decreases the final viscosity of the pastes, while copper(II) increases the peak viscosity of both the natural starch and the spelt monostarch phosphate. The extent of rheological changes in spelt starch and spelt monostarch phosphate was affected by saturation with minerals. The presence of iron(II) and copper(II) ions in spelt starch pastes considerably increased the consistency index values, while the incorporation of iron(II) and copper(II) ions in the monostarch phosphate led to a decrease in the structure recovery index after the preset shear rate was decreased.

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