

# Thermal, Pasting and Micro-Structural Properties of Starch and Flour of Tartary Buckwheat (*F. Tataricum*).

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**Abstract**— Tartary buckwheat (*F. tataricum*) is a rich source of bioactive components and is a major pseudo-cereal for processed functional foods. In buckwheat flour, starch is the main component which plays an important role in the functional and technological properties of end-use food products. In the present investigation starch was isolated; flour and starch isolates of buckwheat were studied for their pasting behaviour, thermal and microstructural characteristics. Granules of starch were oval, spherical and polygonal in shape and having noticeable flat surface areas. Scanned image of flour showed more agglomeration of granules. Results revealed that, presence of protein, lipids and other components affects the pasting and thermal properties of flour and starch. Pasting temperature of flour was higher than starch indicates the higher temperature requirement for gelatinization. RVA analysis showed higher peak, trough, and final and set back viscosities of starch than corresponding viscosities of flour. Slightly higher onset temperature was observed for flour than starch sample while conclusion temperature of starch was higher than flour. Difference in enthalpy change was noticed and starch showed higher values for enthalpy change. Understanding of these properties of buckwheat starch and flour aids in the selection of processing parameters in the food industry, and opens up new avenues of industrial applications.

**Keywords**—Starch; RVA; DSC; SEM.

## 1. INTRODUCTION

Starch is the major source of carbohydrate and the most abundant polysaccharide in plants existing naturally within the plant cells in the form of granules. The applications of starch in food industries in the form of thickener, bulking agent and gelling agent are mainly governed by its pasting, gelatinization, and solubility, swelling and morphological properties. Size of particles of starch granules contributes to specific functional properties such as texture, volume, consistency, and moisture and shelf stability (Raeker et al., 1998). In addition, the size of starch granules affects the temperatures of gelatinization, which in turn influences the properties of food (Soral-Smietana et al., 1984). Gelatinization of starch explains the disruption of the molecular order within the starch granules during heating in the presence of water. Indication of the loss of an organized structure includes irreversible swelling of starch granules, loss of birefringence and crystallinity. The process of gelatinization is energy absorbing than can be monitored through Differential Scanning Calorimetry (DSC) which has been used for analysis of starch (Freitas et al., 2004). In DSC thermogram, thermal stability of samples can be observed by the onset temperature or peak temperature, while the

transformed proportion is reflected by the area under the endothermic peak, representing the enthalpy change (Li-Chan and Ma, 2002). It is generally accepted that swelling of granules increase the viscosity during heating of starch in water. Breakdown refers the rigidity/fragility of the swollen granules. The properties of the swollen granules and the soluble materials leached out from the granules are cooperatively ruling the viscosity parameters during pasting (Doublie et al., 1987). Growing demand of starch has created interest in searching the non-conventional starch sources and buckwheat is a promising new starch source as it produces starch with diverse physicochemical, structural and thermal properties that give it a broad range of potential industrial applications. Tartary buckwheat (*F. tataricum*) is an underutilized crop and has been grown in several hundreds of hectares in Asia, Eastern and Central Europe, and in North America. In India, tartary buckwheat is grown in mountainous regions where the climate is cold, dry, and harsh, has a higher resistance to stress than common buckwheat (Lin et al., 1992). Because of the nutritious advantages of buckwheat, this crop has been considered as a raw material for functional food and medicine. Buckwheat is consumed in the form of flour for making noodles, pancakes, and pasta; and, as whole-seed groats for making porridge, salad, and tea. Minor crops are usually less investigated and hence the information on the ultra-structural details of buckwheat starch is limited. The size and shape of starch granules are species specific (Stark and Lynn, 1992). Kim et al. (1977) reported that buckwheat starch granules are polygonal but slightly larger than rice; have a 25% amylose content by iodine colorimetry; a gelatinization temperature of 61–65°C; and high cool paste viscosity in a Brabender ViscoAmylograph. Enhanced understanding of the structure and morphology of starch granules and analysis of thermal properties of starch can provide the important information to direct the processing and the utilization of starch. Even though lots of work has been done on starch systems, few studies have analysed starch of buckwheat. Thus, the present investigation focuses on pasting, thermal and micro-structural analysis of flour and starch of tartary buckwheat.

## 2. MATERIALS AND METHODS

### 2.1 Materials

Grains of tartary buckwheat (*F. tataricum*) of cultivar Shimla B-1 used in this study were procured from National Bureau of Plant Genetic Resources Regional Station, Shimla,

India. The grains were screened to remove foreign matter and stored in sealed container at room temperature till further use. The flour were prepared by grinding seeds on laboratory mill and stored in polyethylene bags at 10°C.

### 2.2 Isolation of starch

Starch was isolated from tartary buckwheat grain by following the alkaline steeping method (Choi et al., 2000). Grains were steeped in 0.25% aqueous NaOH solution for 18 h at room temperature and stirred three times during this period. Grains were washed 2-3 times with distilled water and ground in a blender at full speed for 2 min. Slurry was filtered step wise through 100 mesh (150µm) and 270 mesh (53µm) sieves. The starch was isolated from the filtrate by centrifugation at 25,000g for 20 min. The supernatant was discarded, and the top yellowish layer of protein was removed. This step was repeated to obtain a white starch layer. The starch layer was resuspended in distilled water, shaken and centrifuged as described above. Thereafter, the isolated starch was dried in hot air oven at below 40°C for 8-10 hours and stored at room temperature in sealed container.

### 2.3 Determination of amylose content

The amylose content of starch and flour was determined according to modified methods of Williams et al., (1970). The standard curve used for amylose was  $Y = 0.0089X + 0.0528$  ( $r = 0.99$ ), where  $X$  = amylose content (%), and  $Y$  = absorbance at 680 nm, based on fractionation of rice starch by Montgomery and Senti (1958).

### 2.4 Scanning electron microscopy

The granule morphology of flour and starch sample was observed in a scanning electron microscope (SEM). The sample was mounted on an aluminium stub using a double sided copper tape and coated with gold in sputter coater. The samples were examined at suitable magnification at an accelerated voltage of 10kV. The selected regions were photographed and saved.

### 2.5 Pasting properties

Pasting properties of starch were determined using a Rapid Visco Analyser (Perten Instruments, Australia). Starch (3 g, 14% moisture basis) was mixed with calculated amount of distilled water in the RVA sample canister. The slurry was manually homogenized using plastic paddle to avoid lump formation before RVA run. A programmed heating and cooling cycle was used where the samples were held at 50°C for 1 min, heated to 95 °C in 3.30 min, held at 95°C for 3 min before cooling to 50°C in 3.30 min and holding at 50°C for 2 min. The mixture was stirred at a constant speed of 160 rpm during the test. A RVA plot of viscosity (cP) versus time (s) was used to determine peak viscosity (PV), trough viscosity (T), breakdown viscosity(BD), final viscosity (FV), set back (SB), peak time (P time) and pasting temperature (P temp).

### 2.6 Thermal properties

Thermal properties of starch and flour of buckwheat were analysed using a Differential Scanning Calorimeter 2920 (TA Instruments, New Castle, DE, USA) according to Bao et al. (2004) with some modifications. Starch (6.0 mg, db) was weighed into an aluminium pan and 6µl distilled water was

added. The pan was hermetically sealed and equilibrated at room temperature for 1 h, then heated at the rate of 10°C/min from 30°C to 120°C with an empty sealed pan as a reference. Parameters such as onset (To), peak (Tp), conclusion (Tc) temperature and enthalpy ( $\Delta H$ ) of gelatinisation were determined by software provided in the system.

## 3. RESULTS AND DISCUSSION

Amylose content is an important factor affecting properties like swelling power, solubility, pasting and gelatinization properties of flour and starch. The fraction of amylose in buckwheat flour depicted in Table 2 was 14.99% which was lower than the range (19-28%) reported by Qin et al. (2010) for thirty nine varieties of buckwheat flour. Amylose content of starch (32.99%) was in consistent with the corresponding findings in earlier studies reported 22% to 33% amylose content in buckwhwat starch (Li et al., 1997; Pandey et al., 2015) , and comparable to the amylose content of cereal, root, tuber and legume starches. However, amylose content of buckwheat was also reported as high as 46.6% (Qian et al., 1998).

Scanned images of flour and starch were shown in Fig. 1. Morphological characteristics like size, shape and distribution of starch granules are attributed to the biological origin of the grain (Vandeputte and Delcour, 2004). The microstructure of the starch revealed that the shape of starch granules were polygonal, oval or spherical with noticeable flat areas due to compact packing in the grain endosperm. As regard to size, diameter of granules ranged from 5.5-10.7µm. No fissures were observed in the surface of starch granules. The size and shape of granules were in agreement with previous reports (Qian et al., 1998; Qian and Kuhn, 1999). Particle size of starch granules were in range of 2-14 µm, which were 1.6-2.4 time smaller than corn and wheat starch granules responsible for low starch recovery during wet extraction (Zheng et al., 1998). The smooth surface of granules indicated that the particles were not damaged during isolation of starch. Compared to starch, granules of buckwheat flour were more agglomerated, certainly due to the presence of proteins and lipids, which could be forming complexes with starch (Nierle, 1990).

Starch is widely applied in the food industry partially due to viscosity change of starch paste during heating and cooling. The pasting curve represents changes in behavior of paste viscosity of flour and starch with change in temperature and mainly varies with composition of flour and characteristics of starch. Pasting properties of starch and flour of buckwheat are depicted in Table 1. Increase in viscosity during heating may be attributed to the swelling of granules, as a result of loss of crystalline order and absorption of water (Bao and Bergman, 2004). Initiation temperature for gelatinization of buckwheat flour was 74.25°C which was slightly higher than the range of 71.3°C-72.1°C reported by Shevkani et al. (2014) for amaranth flour and 64°C for rice flour reported by Qui et al. (2015). The high pasting temperature of flour as compare to other pseudo- cereals and cereals flour indicates its higher resistance towards swelling. Pasting temperature of starch was 72.76°C that was higher

than that reported (62.8°C) by Li et al. (1997) for tartary buckwheat starch. Yoshimoto et al. (2004) studied eight kinds of buckwheat cultivars and reported 70°C as maximum pasting

Table 1 Pasting properties of buckwheat flour and starch

	Flour	Starch
Peak Viscosity (cP)	743.66±7.23	3101.66±30.82
Trough Viscosity (cP)	730±3	1389.66±47.35
Breakdown viscosity (cP)	16±3.6	1693.66±33.65
Final Viscosity (cP)	1472.66±3.05	3736.66±53.46
Setback Viscosity (cP)	742±6.2	2330.66±6.02
Peak Time (min)	6.47±0.13	3.13±0
Pasting temperature (°C)	74.25±0	76.76±0.02

Values expressed as mean ± SD (n=3).

temperature of starches. The point of maximum swelling of starch granules is indicated by peak viscosity. Peak viscosity of flour (743.66cP) was found to be lower than the starch (3101.66cP). Yoshimoto et al. (2004) reported peak viscosities of starches of various cultivars of buckwheat range from 2712cP to 3132cP. Break down viscosity, is the measure of resistance of gel to disintegrate at high temperature, noticed 1693.66cP for starch was in consistent with the findings of Lui et al. (2014) observed 1612cP for buckwheat starch. Lower breakdown viscosity of flour than starch represents greater resistance to shear thinning and high stability of paste. Final viscosity represents the ability of starch/flour to form a viscous paste, was found to be 1472.66cP and 3736.66cP for flour and starch respectively. Final viscosity of starch in present observation was lower than the reports of Lui et al. (2014) noticed 4208cP for buckwheat starch. Increase in final viscosity from peak viscosity might be due to the aggregation of amylose molecules (Miles et al., 1985). Set Back viscosity is the measure of syneresis upon cooling of cooked paste. Set back viscosity of starch was in range (2172cP -2712cP) registered by Yoshimoto et al. (2004) and comparatively lower set back viscosity was noticed by Lui et al. (2014) for buckwheat starches. Viscosity of flour was lower than starch at every point during RVA analysis, which might be due to presence of less percent of starch in flour than starch isolate, as the pasting properties depends mainly on starch present in sample. Hamaker and Griffin (1993) and Yang and Chang (1999) noticed that proteins in flour restrict starch granules swelling and reduce the amylogram viscosity. Protein removal was found to increase the paste viscosity of starch (Yang and Chang, 1999). The role of lipids on the pasting properties was observed as positive relation with pasting temperature while negative relation with viscosity (Shevkani et al., 2014; Singh et al., 2014), results of present investigation agreed with it.

Table 2 Thermal properties and amylose content of flour and starch of buckwheat

	Flour	Starch
Onset temperature (°C)	67.14±1.2	66.35±1.03
Peak temperature(°C)	77.5±0.8	78.2±0.2
Conclusion temperature(°C)	81.9±0.4	84.1±0.0
Enthalpy change(J/g)	12.5±1.3	16.2±0.5
Amylose content (%)	14.99±0.29	32.12±0.24

Values expressed as mean ± SD (n=3).

Transition temperatures ( $T_0$ ,  $T_P$ ,  $T_C$ ) and gelatinization enthalpy ( $\Delta H$ ) of starch and flour of buckwheat are shown in Table 2. The value of  $T_0$  represents the internal structure of the granule during its disintegrations, which results in the release of polysaccharide into the immediate medium (Singh et al., 2006). Differential scanning calorimetric studies of flour and starch of buckwheat showed that the gelatinization temperature of the flour (67.14°C) was slightly higher than that of the starch (66.35°C). The difference could be attributed to the presence of mucilage in the flour. In addition to protein, the mucilage also contains complex polysaccharides which could compete with starch for moisture and result in a higher onset starch gelatinization temperature in the flour (Taxi et al., 1972). Similar trend of transition temperatures were observed by Jane et al. (1992) for taro starches and flours. The peak temperature for flour (77.5°C) and starch (78.2°C) were slightly higher than paste temperature obtained by RVA (74.25°C, 76.6°C respectively). The results of present investigation agreed with the earlier studies (Li et al., 1997; Yoshimoto et al., 2004) reported  $T_0$  ranged from 59°C to 64°C for various cultivars of buckwheat. The peak temperature and conclusion temperature of starch was similar to the finding of Lui et al. (1997) reported  $T_P$  78°C and  $T_C$  83.9°C for tartary buckwheat starch. During DSC analysis the transformed proportion of starch is reflected by the area under the endothermic peak, representing the enthalpy change ( $\Delta H$ ). The enthalpy of gelatinization of starch (16.2 J/g) was comparable with the range registered in literature by other authors (Yoshimoto et al., 2004; Lui et al., 2014), however, it was higher than the findings of some other investigations (Li et al., 1997; Zheng et al., 1998; Qian and Kuhn, 1999) reported range of enthalpy change 2.14-12.7 J/g. The differences might be due to the different varieties and growing location of the observed buckwheats.

#### 4. CONCLUSION

In the present study, variations in the thermal and pasting properties were observed between flour and starch of tartary buckwheat. Flour showed lower viscosity than starch at every stage of RVA analysis. DSC analysis resulted in the higher onset temperature of flour than starch indicated that protein and mucilage in flour interfere with water absorption of starch granules and form basis for more energy requirement for occurrence of gelatinization. Granules of buckwheat starch were irregular in shape and having large flat surface areas indicating compact packing in endosperm. Size of granules was smaller than starch of cereal sources

which make its wide applications in food and non-food industries. Scanning electron microscopy revealed that granules of flour were more agglomerated than starch granules. Future work is necessary to characterise fine structures of tartary buckwheat starch and analyse their relationships with properties of starch.

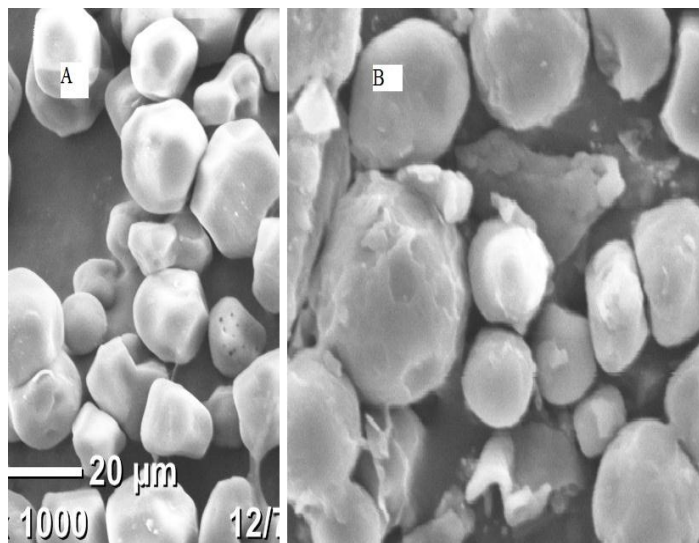


Fig. 1 Scanning electron micrographs of tartary buckwheat flour A (X2000) and starch B (X3000).

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