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Effect of germination on the functional and moisture sorption properties of high pressure processed foxtail millet grain flour

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10 Abstract Foxtail millet is one of the commonly cultivated, nutritionally competitive source of protein, fibre, phytochemicals and other micronutrients, as compared to major 11 cereals like wheat and rice. Considering the potential of these grains, the high pressure 12 processed flours of germinated (GFMF) and non-germinated foxtail millet (NGFMF) grains 13 were studied for its functional, moisture sorption and thermodynamic properties. Germination 14 15 and high pressure processing of foxtail millet grains significantly improved the functional properties of the flour. Apart from this, the moisture sorption isotherms of both the flours 16 were determined at 10, 25 and 40 °C and the sorption data was fitted to Guggenheim-17 Anderson-De Boer (GAB) sorption model. The monolayer moisture content for NGFMF and 18 GFMF ranged between 3.235 - 2.364 g g⁻¹ and 2.987 - 2.063 g g⁻¹, respectively. The isosteric 19 heat of sorption ranged between -76.35 kJ mol⁻¹ to -38.23 kJ mol⁻¹ for NGFMF and 172.55 kJ 20 mol⁻¹ to -34.02 kJ mol⁻¹ for GFMF at a moisture range of 0 to 36%, whereas, the integral 21 entropy of sorption for NGFMF ranged between -0.404 to -0.120 kJ mol⁻¹ K⁻¹ and for GFMF 22 between -0.667 to -0.383 kJ mol⁻¹ K⁻¹. Along with the validation of the compensation theory, 23 the values of spreading pressures lied in the range of 0 - 0.078 J m⁻² for NGFMF and 0 -24 0.124 J m⁻² for GFMF, while, the glass transition temperatures ranged between 82.25 to 28.67 25 °C for NGFMF and from 51.11 to 11.83 °C for GFMF at all three temperatures. 26

Keywords Foxtail millet flour, germination, high pressure processing, functional
properties, moisture sorption isotherms, thermodynamic properties

30 Introduction

Water sorption characteristics are necessary for predicting the shelf life and determining the 31 32 moisture content and critical activity for acceptability of products that deteriorate, mainly due to moisture gain, and also for drying, packaging and storage (Katz and Labuza 1981). A food 33 moisture sorption isotherm describes the relationship between the moisture uptake in food 34 and the relative humidity of the air with which the food is in equilibrium at a constant 35 temperature (Lagoudaki et al. 1993). Thermodynamics has been reported as an approach to 36 understand the properties of water and calculate the energy requirements of heat and mass 37 transfer in biological systems (Rizvi and Benado 1983; Fasina et al. 1997; Fasina et al. 1999). 38 39 Properties such as isosteric heat of sorption, net integral enthalpy and entropy and spreading pressures are important for understanding the energy requirement during dehydration, the 40 food microstructure, physical phenomena on the food surfaces and sorption kinetic 41 parameters (Rizvi and Benado 1983). Also, the glass transition temperature is an important 42 43 physical parameter, which serves to explain the physical and chemical behaviour of food 44 systems (Bell and Labuza 2000), and it is defined as the temperature at which the material 45 changes from the glassy to the rubbery state for a given heating rate.

Foxtail millet (Setaria italica (L.) P. Beauv.) is the sixth highest yielding grain amongst all 46 47 the millet varieties and has been identified as a major millet in terms of worldwide production (Sharma et al. 2017). Foxtail millet ranks second in the total world millet production of 48 49 around 29 million tonnes (in 2015-16 season). It originates in the Yellow River Basin 50 country, which makes China a leading foxtail millet producer holding 80% of world production, followed by India with 10% of world production (Zhang et al. 2017). Foxtail 51 52 millet is one of the oldest cultivated cereal grains belonging to the Setaria genus, of Poaceae family and subfamily Panicoideae. Sharma et al. (2017) have discussed the versatility of 53 54 foxtail millet as a food source and its potential to address food and nutrition security. These 55 authors have reviewed the effects of various processing techniques on the properties of 56 foxtail millet grains.

It has been recognised that non-thermal processing techniques like high hydrostatic pressures (HHP), can potentially inactivate anti-nutritional factors like phytate and tannin content in cereal grains and preserve their constituents and quality (Estrada-Girón et al. 2005; Yu et al. 2015; Zhang et al. 2017). In addition to high pressure processing, germination, is one of the traditional approaches used to supplement the bio-accessibility of nutrients in plant-based 62 diets. The process of germination in cereal grains results in an increase in enzymatic activity, soluble protein content and break down of starch into simple sugars, resulting in the 63 formation of typical colours and flavours (Barreiro et al. 2003). Moisture sorption properties 64 have also been studied for various cereals and its products like various cereal grains and 65 legumes (Al-Mahasneh et al. 2014), milk-foxtail millet powder (Simha et al. 2016), two 66 millet grain varieties (Aviara et al. 2016), tef (Eragrostis tef (Zucc.) Trotter) grain flour 67 (Abebe and Ronda 2015), bulgar (Erbaş et al. 2015), Greek durum wheat semolina (Pollatos 68 et al. 2013), RTE breakfast cereal formulation (Sandoval et al. 2009), oat and rice flour (Brett 69 70 et al. 2009), maize meal and millet flours (Ikhu-Ornoregbe and Chen 2005). In general, very limited moisture sorption data is available for germinated cereals, especially for high pressure 71 processed and germinated millet grains. This study was therefore conducted to assess the 72 effect of germination on the functional, moisture sorption and thermodynamic properties of 73 high pressure processed foxtail millet grain flour. 74

75 Materials and methods

76 Sample collection

Foxtail millet (*Setaria italica* L.) grains were procured from authorized grain centres. All
procured grains were a month old since harvesting and were tested for viability using
tetrazolium chloride test (Lakon 1949). The moisture content of the native grains was found
to be 8.41±0.45%, following which the grains were stored in polythene bags at 4 °C until
further use. The chemicals used for analysis were procured from Sigma-Aldrich Chemical
Co., (St. Louis, Mo, USA).

83 Germination of foxtail millet grains

The foxtail millet grains were washed using 1.5% formaldehyde solution to prevent any 84 contamination. To remove the traces formaldehyde, the grains were then washed thoroughly 85 with distilled water and tested for any residues using the ferric chloride method described by 86 VICH Steering Committee (2002). Prior to germination, the foxtail millet grains were first 87 soaked in deionised water for 15h and then were subjected to germination in a controlled-88 environment growth cabinet. The temperature was maintained at 25±2 °C and the relative 89 humidity was controlled at 50%; these conditions were based on the work reported by 90 Sharma et al. (2015). Following the method described by Elkhalifa and Bernhardt (2010), the 91 92 activities of alpha amylase and protease were monitored at regular intervals of 12 hours, and the germination period of 72h represented the time taken for their activities to reach a maximum value., as. The maximum alpha amylase and protease activity was found to be attained after 72h (Fig. 1). The germinated foxtail millet grains were dried to a final moisture content of 7 - 8%, packed and stored in air-tight containers at 4 °C until further analysis. Non-germinated foxtail millet grains were used as a control.

98 High pressure soaking

Both germinated and non-germinated foxtail millet grains (25g each) were subjected to high 99 100 hydrostatic pressures in a temperature-controlled pressure vessel system (vessel dimensions: 37mm diameter and 246mm length, Food LAB 900, Stansted Fluid Power Ltd., Stansted, 101 102 UK). A blend of castor oil and ethanol in a ratio of 80:20 v/v was used as a medium to transmit pressure to the packed samples in the pressure vessel chamber. The foxtail millet 103 grains were treated at a pressure of 400 MPa for 60 mins at a temperature of 60 °C. For this, 104 25g foxtail millet grain samples were suspended in 150ml of deionised water and sealed with 105 virtually no headspace polyethylene pouches of dimensions $3 \text{cm} \times 20 \text{cm}$ by employing 106 CryovacTM. All the experiments were triplicated. 107

After high-pressure processing under specified conditions of pressure and temperature for a stipulated time, the water was completely drained from the pouches and the water occluding to the grains separated, was then blotted using a tissue paper. The processed grains were then subjected to freeze drying and were finally milled into fine flour passing through sieves of mesh sizes ranging between 100 and 200µm, using vibratory sieve shaker (Fritsch Analysette 3, Germany). The resulting flour was packed and stored in air-tight containers at 4 °C until further analysis.

115 Determination of functional properties

The bulk density, porosity and flowability of the optimized foxtail millet flour, as the angle of 116 repose (as a static measure for flowability), were determined by the method of Sjollema 117 (1963). The occluded air content was determined by the methods used by Jha et al. (2002). 118 The wettability was determined by the method given by Muers and House (1962). The 119 dispersibility of the optimized baby food was determined by the method described by the 120 American Dry Milk Institute (ADMI 1965). The insolubility index was determined by the 121 modified ADMI method described by Jha et al. (2002). The water, oil absorption capacity 122 123 and swelling capacity of the product were determined by the method described by Sodipo and

Fashakin (2011). The emulsion activity, emulsion stability, foaming capacity and foamingstability were determined using the methods described by Elkhalifa and Benhardt (2010).

Finally, the gelatinization temperatures were determined using Differential Scanning 126 Calorimetry, where a suspension of the resulting flour in deionised water was prepared in a 127 128 ratio of 1:3; equilibrated for 15-18 h at ambient temperature; and presented to the differential scanning calorimeter, equipped with a built-in software (Model Q 2000a TA Instruments, 129 New Castle, U.S.A.). A 10-12 mg suspension sample was weighed precisely in an aluminium 130 sample pan and hermetically sealed. A similar empty pan was used as a reference for the 131 analysis. A temperature range of 10 to 140 °C was used for scanning the sample in the pan at 132 133 a heating rate of 10 $^{\circ}$ C min⁻¹. The resulting thermogram gave the values of onset (T_o), peak (T_p) and conclusion temperatures (T_c) in ^oC. 134

135 Sorption studies

136 Gravimetric method described by Lang et al. (1981), was used to determine the sorption isotherms for processed foxtail millet flour. The saturated salt solutions of eight reagent-grade 137 138 salts, namely, lithium chloride, magnesium chloride, potassium carbonate, magnesium nitrate, sodium chloride, potassium chloride, potassium nitrate, and potassium sulfate, were chosen as 139 140 mentioned by Greenspan (1977) to cover the water activity range from 0.11 to 0.97. It is important to note that the water activity values for these salts can show slight variations at 141 142 different temperatures. Therefore, different water activity values of these salts at temperatures 10, 25 and 40 °C were used in this study. These saturated solutions were placed in air tight, 143 wide mouth glass jars. Approximately 1 g of sample was weighed into a glass beaker placed 144 on glass bead support inside the glass jar. The glass beakers were equilibrated for three days 145 with the humid atmosphere before placing the sample. Then the moisture sorption 146 characteristics were studied at 10, 25 and 40 °C (± 2 °C) by placing the sample containing 147 glass jars in incubators for equilibration. These conditions were chosen in this study, as they 148 represent the possible storage conditions for cereals like rice (Ondier et al. 2012) The samples 149 in bottles were weighed at a regular interval of 3 days till a constant weight was attained. 150 Equilibrium was considered to have accomplished when the weight difference between two 151 successive weighing was less than 1 mg. Experiment was conducted in triplicates for all the 152 samples at each relative humidity and temperature conditions. After the equilibrium has been 153 attained, the moisture content of the samples was determined and was expressed in gram per 154 155 litre (g/L). The average standard deviation between the replicates was calculated to be less than 1% of the mean of the three values. These average values of equilibrium moisture
content were then plotted against relative humidities or water activities to obtain moisture
sorption isotherms.

159 Sorption model

Various approaches have been used to describe sorption isotherms of foods. Amongst which, Guggenheim-Anderson-De Boer (GAB) (Pérez-Alonso et al. 2006) is mostly used to study the correlation between the equilibrium moisture content (M) and water activity (a_w) for various foods. The following linearized form of GAB model were used for evaluating the best fitted values of model constants using linear regression:

166
$$W = W_o \frac{Gka_w}{(1 - ka_w)(1 - ka_w + Gka_w)}$$
(1)

167

168 The GAB model could be rearranged into a second-degree polynomial equation as given 169 below:

170
$$\frac{a_w}{w} = aa_w^2 + ba_w + c$$
 (2)

171 where, *a*, *b* and *c* are model constants and significantly depend on type of regression analysis. 172 A nonlinear regression analysis of a_w/W v/s a_w yields a polynomial equation of the second 173 order. The coefficients *a*, *b* and *c* were thus obtained from this polynomial equation and then 174 substituted to obtain GAB constants W_o , *G* and *k*. W_o is the moisture content corresponding to 175 saturation of all primary adsorption sites by one water molecule, *G* is the Guggenheim 176 constant and *k* is the factor correcting for properties of multilayer molecule with respect to 177 the bulk liquid.

178 Thermodynamic properties

The integral enthalpy, also known as, net isosteric heat of sorption was determined usingClausius-Clayperon equation (Rizvi 1986; Bell and Labuza 2000):

181
$$\frac{\partial(\ln a_w)}{\partial(\frac{1}{T})} = \frac{Q_{st}}{R}$$
(3)

where, *T* is the absolute temperature (K), Q_{st} is the net isosteric heat of sorption (kJ mol⁻¹) and *R* is the universal gas constant (8.314 × 10⁻³ kJ mol⁻¹ K⁻¹).

The calculation procedure assumed that net heat of sorption was independent of temperature change. The slope of the plot of ln (a_w) versus 1/T at constant moisture content gave the net isosteric heat of sorption (Eq. 3). Adding the latent heat of vaporization of pure water at 10, 25 and 40 °C to the net isosteric heat of sorption (Q_{st}), gave the values of isosteric heat of sorption. A cubic spline interpolation was used to determine a_w values at various temperatures for a given moisture content (M).

Substituting free Gibbs energy in the Gibbs- Helmholtz equation (McMinn and Magee 2003;
Kaya and Kahyaoglu 2005), the differential entropy was calculated using the following
equation:

193
$$-\ln(a_w) = \frac{-(Q_{st}+\lambda)}{RT} - \frac{\Delta S}{R}$$
(4)

where, λ is the latent heat of vaporization of pure water (44.045 kJ mol⁻¹ at 25 °C), and ΔS is the integral entropy of sorption (kJ mol⁻¹ K⁻¹). The maximum storage stability point is when the water activity and temperature gives the minimum value of integral entropy (Alpizar-Reyes et al. 2017).

From the intercept of the plot of $\ln (a_w)$ versus 1/T at specified moisture levels of eq. (4), the integral entropy was evaluated.

200 Isokinetic theory

A linear relationship between the isosteric heat of sorption (differential enthalpy) and differential entropy established the enthalpy- entropy theory (McMinn and Magee et al. 2003; Arslan and Togrul 2006):

$$204 \qquad Q_{st} = T_{\beta} \Delta S_d + \alpha \tag{5}$$

where, Q_{st} is the isosteric heat of sorption (kJ mol⁻¹), T_{β} is the isokinetic temperature (K), and a is the constant or Gibb's free energy (kJ mol⁻¹).

The isokinetic temperature (T_{β}) is a characteristic property of the material surface (Aguerre et al. 1986), and it represents the slope of the enthalpy- entropy linear relationship (Eq. 5). Linear regression of Q_{st} with ΔS_d evaluated T_{β} . A statistical test proposed by Krug et al. (1976) verified the enthalpy- entropy compensation theory by defining the harmonic mean temperature (T_{hm}) as:

212
$$T_{hm} = \frac{n}{\sum_{1}^{n} \left(\frac{1}{T}\right)}$$
(6)

213 where, T_{hm} is the harmonic mean temperature (K), and n is the number of temperature levels. 214 Linear enthalpy- entropy compensation theory applies only when $T_{\beta} \neq T_{hm}$; the process is 215 enthalpy driven if $T_{\beta} > T_{hm}$ and entropy driven if $T_{\beta} < T_{hm}$ (Leffler and Grunwald 1963). Eq. 216 (6) shows that T_{hm} is a function of temperature levels selected for the experiment and is 217 independent of the material considered. Analyses were conducted with $\alpha = 0$ to evaluate the 218 effect of temperature on the sorption behaviour by introducing a temperature correction factor 219 (Aguerre et al. 1986) as:

220
$$-\ln(a_w) = \frac{Q_{st}}{R(\frac{1}{T} - \frac{1}{T_\beta})^{-1}}$$
(7)

221 Spreading pressures

Spreading pressure was evaluated based on the analytical procedure described by Iglesias etal. (1976) and Fasina et al. (1999). Spreading pressure is given by the following equation:

$$224 \qquad \pi = \frac{KT}{A_m} \int_0^{a_w} \frac{x}{x_m a_w} \tag{8}$$

where, π is the spreading pressure (J m⁻²), *K* is the Boltzman constant (1.380 × 10⁻²³ J K⁻¹) and A_m is the surface area of a water molecule (1.06 × 10⁻¹⁹ m²).

This integral becomes indeterminate at $a_w = 0.0$. Therefore, spreading pressure was evaluated by dividing the total limit (0.0 to a_w) into a very small interval of 0.0 to 0.05 and the remaining 0.05 to a_w . The integral in the first infinitesimal a_w range (0.0 to 0.05) was evaluated assuming a linear relationship (Henry's law) between m and a_w with usual notations (Fasina et al. 1999) as:

$$232 \qquad \pi = \frac{KTx}{A_m x_m} \tag{9}$$

The integral in the second a_w range (0.05 to a_w) was evaluated after fitting isotherm data (*m*, a_w) to the GAB model equation and the spreading pressure was obtained by integrating the equation in the second interval of 0.05 to a_w (Iglesias et al. 1976):

236
$$\pi = \frac{KT}{A_m} ln \left[\frac{1 + Cka_w - ka_w}{1 - ka_w} \right]_{0.05}^{a_w}$$
(10)

Adding the results of the two selected a_w intervals produced the spreading pressures at any studied temperature and water activity combination.

239 *Glass transition temperature*

240 Differential scanning calorimetry was used to determine the glass transition temperature. A suspension of processed foxtail millet flours and deionised water was prepared in a ratio of 241 1:3 and equilibrated for 15-18 h at ambient temperatures and were analysed using a TA 242 Instruments model Q-2000 differential scanning calorimeter, equipped with a built-in 243 244 software (TA Instruments, Newcastle, U.S.A.). For this, 10-12 mg of flour, equilibrated at a range of water activities (0.11 - 0.85) and temperatures $(10, 25 \text{ and } 40 \text{ }^\circ\text{C})$ were weighed in 245 246 an aluminium sample pan and hermetically sealed. A similar empty pan was used as a reference for the analysis. A temperature range of -50 to 150 °C was used for scanning the 247 sample in the pan with a heating rate of 2 °C min⁻¹ (Alpizar-Reyes et al. 2017). The resulting 248 thermogram gave the values of onset (T_o) , peak (T_p) and conclusion temperatures (T_c) in ^oC 249 and gelatinisation enthalpies (ΔH) in kJ kg⁻¹. The final values obtained were an average of 250 three replicates for each sample, with a maximum deviation of less than 1.5%. The midpoint 251 252 of the baseline shift obtained using DSC was taken as the glass transition temperature (T_{e}) . The measurements were made in triplicates. Further, to deduce the plasticizing effect of water 253 on T_g , the DSC data was put in the Gordon-Taylor equation: 254

255
$$T_g = \frac{W_1 T_{g1} + K W_2 T_{g2}}{W_1 + K W_2}$$
(11)

Where, W_1 and W_2 are the mass fractions of the flours and of water, respectively, T_{g1} is the T_g value of flour at zero moisture content and T_{g2} is the T_g value of water (-135 °C), and *K* is a constant.

259

261 Statistical analysis

The differences amongst the samples were determined using one-way analysis of variance (ANOVA) using Minitab 17.0 software. Using the Duncan's test the differences were considered statistically significant at p-value ≤ 0.05 . The degree of fitness of the model was evaluated using determination coefficient (R²).

266 **Results and discussion**

267 Functional properties of the flours

268 The functional properties of the processed germinated foxtail millet flour (GFMF) as well as non-germinated foxtail millet flour (NGFMF) are shown in Table 1. Germination of foxtail 269 270 millet increased the values of porosity, occluded air content, water and oil absorption capacity, dispersibility, swelling capacity, wettability, flowability, gelatinization enthalpy, 271 272 emulsification activity and stability; while decreased the values of bulk density, insolubility index, foaming capacity and stability. Albarracín et al. (2016) reported that the water 273 274 absorption and water solubility values were higher for germinated brown rice samples as compared to the control brown rice samples. Similar results were also stated by Chinma et al. 275 (2009) for varieties of germinated tigernut flour; Devisetti et al. (2014) for millet flours and 276 Elkhalifa and Bernhardt (2010) for germinated sorghum flour. 277

278 Moisture sorption characteristics

The effect of germination was studied for the stability of the high pressure processed foxtailmillet flours.

281 *Equilibrium moisture content- water activity*

The relationship between the EMC and water activity (a_w) at a constant temperature forms the 282 MSI. Therefore, foods with different moisture content have different water activity, thus they 283 form their own sorption isotherm (Alpizar-Reyes et al. 2017). Fig. 2 (a, b) shows the 284 experimental isotherms for NGFMF and GFMF at 10, 25 and 40 °C, respectively, with which 285 the thermodynamic functions related to moisture sorption in porridge flours were determined. 286 Both figures depict that the EMC increased with increasing water activity ranging from 0.11 287 to 0.97 at a constant temperature. A constant linear increase in EMC at lower water activities 288 ranging from 0.11 to 0.55 was observed for NGFMF and from 0.11 to 0.76 for GFMF, which 289

290 could be due to the ability of the flour to lower the vapour pressure, with the decrease in the relative humidity. While, a rapid increase in EMC at higher water activities from 0.55 to 0.97 291 for NGFMF (Fig. 2(a)) and an impulsive increase (Fig. 2(b)) in the EMC from 0.76 to 0.97 292 for GFPF could be attributed to the high sorptive capacity of hydrophilic starch in foxtail 293 294 millet flours consisting of a large number of free hydroxyl sites (Pollatos et al. 2013). This impulsive increase in the case of GFMF could be because of elevated enzyme activity and 295 breakdown of complex sugars into simple sugars during germination (Wu et al. 2013). 296 Barreiro et al. (2003) also found similar results for malted barley flours. Therefore, it could 297 298 be concluded from the results that GFMF would require improved storage conditions with relative humidity not exceeding 76% and NGFMF not exceeding 55%. 299

300 The increase in temperatures did not have a significant effect on the moisture sorption behaviour of both the flours. Though it was observed that for NGFMF, the EMC decreased 301 with the increase in temperatures for a given water activity, showing a less hygroscopic 302 303 behaviour of NGFMF at higher temperatures. This is because high temperatures increase the kinetic energy of the water molecules, thus increasing the distance between them and causing 304 easy binding of these water molecules to the suface of the flours (Pollatos et al. 2013; 305 306 Alpizar-Reyes et al. 2017). On the other hand, an opposite trend was depicted by GFMF, which could be attributed to the higher content of sugars due to germination, thus increasing 307 the active sites for the sorption of water by proteins (Barreiro et al. 2003; Pollatos et al. 308 2013). In addition to this, the MSI's of both NGFMF and GFMF were of the characteristic 309 310 sigmoidal shape (BET II type), which is likely found in all types of dry foods rich in starch 311 and sugars.

312 Sorption models

The experimental sorption data for both the flours were fitted using GAB model, with R^2 313 value of more than 0.998 at all temperatures and RMSE value of less than 0.051. The model 314 parameters were determined using non-linear regression method. Table 2 and 3 shows the 315 316 model parameters for NGFMF and GFMF, respectively. MSIs of both the flours can be divided into an: (1) initial monolayer region (a_w less than 0.2), where the water molecules are 317 318 strongly absorbed as critical moisture content that starts the chemical reactions in foods. It is 319 also a measure of the availability of active sorption sites; (2) an intermediate multilayer region (a_w in the range of 0.2-0.6), that causes lipid oxidation and physical changes in foods, 320 and (3) a final condensed water region (a_w greater than 0.6), which ascertains the microbial 321

growth (Labuza and Altunakar 2007). The values of monolayer moisture content (W_o) for 322 both NGFMF and GFMF decreased with the increase in temperatures from 10 to 40 °C, as 323 expected by the theory of physical sorption (Pollatos et al. 2013). The values for W_{ρ} for 324 NGFMF and GFMF ranged between 3.235 - 2.364 g g⁻¹ and 2.987 - 2.063 g g⁻¹, respectively. 325 326 The composition and nature of the product was responsible for the difference in the values of the monolayer moisture content in germinated and non-germinated flours. Thus, NGFMF 327 being a starch rich flour was found to be more hygroscopic and GFMF being a sugar rich 328 flour due to germination, was found to be less hygroscopic. 329

330 The constant G is related to the monolayer sorption heat which associates the chemical potential difference with the superior layers of moisture in the flour structure. It has been 331 332 established that low temperatures assist the formation of strong adsorbent and adsorbate interactions, thus causing an increase in the G values with the increase in temperature 333 (Alpizar-Reyes et al. 2017). While, this study reported a decrease in the G values on 334 increasing the temperature from 10 to 25 °C and then an increase in the G values on further 335 increasing the temperature from 25 to 40 °C, for both NGFMF and GFMF. Thus, it suggested 336 that the flour-water interactions in both NGFMF and GFMF decreased as follows: $10 \text{ }^{\circ}\text{C} > 40$ 337 $^{\circ}C > 25 \ ^{\circ}C$. These results agreed with Pollatos et al. (2013) and Simha et al. (2016). 338

Another constant *k* is related to the enthalpy of water multilayer sorption indicating a measure of the interactions between multilayer molecules and adsorbent. When, *k* values are in the range of 0.24 to 1.0, GAB model best describes the sigmoid shaped isotherms (Lewicki 1997). For NGFMF and GFMF, the *k* values were in the range of 0.780 - 0.881 and 0.852 -0.944, respectively. Since the values $k \ll 1$, so it was inferred that the flours comprised of a structured state of adsorbate in the adjacent layers to the monolyer (Alpizar-Reyes et al. 2017).

346 *Thermodynamic properties*

Dried foods like the flours that have been studied here, undergo various changes during
storage due to the processes like crystallization, dissolution, diffusion, swelling, etc.
Therefore, various thermodynamic properties like integral enthalpy and entropy determines
the condition of maximum food stability.

351

The degree of bound water in the flour estimates the variation in the integral enthalpy which 353 is also known as net isosteric heat of sorption (Q_{st}) . It indicates the interaction of water 354 vapour with solid substrates in the flours, thus explaining the endothermic or exothermic 355 nature of the interaction (Pushpadass et al. 2014). Fig. 3 (a, b) shows the relationship between 356 Q_{st} and the moisture content at constant temperature for both NGFMF and GFMF. The 357 negative values of Q_{st} in the graph indicated the exothermic nature of sorption for both flours. 358 Strong water-substrate interactions were suggested at low moisture contents, as the water 359 binding capacity of the flours increased with the increase in the negative values of Q_{st} . The 360 values of Q_{st} ranged between -76.35 kJ mol⁻¹ to -38.23 kJ mol⁻¹ for NGFMF and 172.55 kJ 361 mol⁻¹ to -34.02 kJ mol⁻¹ for GFMF at a moisture range of 0 to 36%. At the higher moisture 362 contents, the value of Q_{st} became close to the values of heat of sorption of pure water, thus 363 indicating the excess moisture bounded to the flours or condensation in the capillaries of flour 364 particulates. The difference in the Q_{st} of both flours at low moisture contents indicated that 365 the degree of binding of water molecules was higher for GFMF as compared to the NGFMF. 366 This could be because of the availability of the higher energy polar sites on the surface in the 367 initial stages of sorption of GFMF, causing water molecules to form a monomolecular layer. 368 In addition to this, as the moisture content further increased for both the flours, the 369 availability of these sites decreased and sorption occurred at less active sites, yielding lower 370 heats of sorption (Lagoudaki and Demertzis 1994). Similar findings were observed for 371 372 chestnut and wheat flours (Moreira et al. 2010); Japanese durum wheat flour (Chuma et al. 2012); whole wheat flour (Martín- Santos et al. 2012) and Greek durum wheat semolina 373 (Pollatos et al. 2013). 374

375 Integral entropy

The integral entropy of sorption (ΔS) for NGFMF ranged between -0.404 to -0.120 376 kJ mol⁻¹ K⁻¹, with a minimum value at 4% moisture content and 40 °C temperature and for 377 GFMF between -0.667 to -0.383 kJ mol⁻¹ K⁻¹, with a minimum value at 8% moisture content 378 and 10 °C temperature. Fig. 4 (a, b) shows the integral entropy as a function of moisture 379 content at 10, 25 and 40 °C for both flours. These findings were in accordance with Simha et 380 al. (2016). The minimum ΔS value was observed for GFMF as compared to NGFMF, thus 381 interpreting that GFMF is more stable than NGFMF at a temperature range of 10 to 40 °C. 382 This stability is because the water molecules in the flour forms more ordered arrangement, 383

bond between adsorbate and adsorbent being the strongest. Therefore, less moisture is 384 available for spoilage reactions. The subsequent increase in the values of ΔS with the increase 385 in moisture content for both the flours, reflected the formation of multi-layers and freely held 386 molecules of water (Rizvi 1986). The increasing temperatures also had a great impact on the 387 integral entropy of both the flours. With the increase in temperature from 10 to 40 °C, the 388 values of ΔS decreased for NGFMF and increased for GFMF, suggesting that NGFMF was 389 more stable at higher temperatures, while GFMF was more stable at lower temperatures. This 390 391 could be attributed to the composition of the flours.

392 *Enthalpy-entropy compensation theory*

The isokinetic theory or the enthalpy-entropy compensation theory determines the existence 393 of true compensation by comparing the difference between the harmonic temperature (T_{hm}) 394 and isokinetic temperature (T_{β}) $(T_{hm} \neq T_{\beta})$. This theory helps in evaluating the sorption 395 reactions, confirming if the reactions are enthalpy driven or entropy driven. According to 396 397 Leffler and Grunwald (1963), if $T_{\beta} > T_{hm}$ the process is enthalpy driven, while if $T_{\beta} < T_{hm}$ the process is considered to be entropy controlled. The T_{hm} values were found to be 297 K and 398 303 K at all three temperatures for NGFMF and GFMF, respectively. The results of this study 399 showed that the T_{β} values were greater than T_{hm} at all three temperatures, therefore it was 400 concluded that the sorption reactions for both NGFMF and GFMF were enthalpy-controlled 401 402 processes in the studied moisture content range. This further confirmed the predominance of mesopores in the interactions between water-flour (Azuara and Beristain 2006). These values 403 were calculated by fitting the data in Eq. 7. Fig. 5 (a, b) depicts the enthalpy-entropy 404 compensation obtained by plotting the integral properties of both flours at constant 405 406 temperatures of 10, 25 and 40 °C. The linear relationship between the two thermodynamic properties confirmed the existence of compensation theory, which has been already justified 407 408 for various foods by Fontan et al. (1982).

409 Spreading pressures

The spreading pressures (π) represents the excess of free energy at the surface of the flours, and provides an indication of the increase in surface tension at free sorption sites due to adsorbed molecules (Fasina et al. 1999). The spreading pressure isotherms for both NGFMF and GFMF were plotted against the water activity at 10, 25 and 40 °C (Fig. 6 (a, b)). The values of π lied in the range of 0 – 0.078 J m⁻² for NGFMF and 0 – 0.124 J m⁻² for GFMF, indicating higher surface tension on the surface of GFMF at high water activities. The graph showed that the value of π increased with increasing water activity and decreased with increasing temperatures. While, at water activities above 0.76 the values of π significantly increased with increasing temperatures. Similar trend was observed for cassava flour (Ayala-Aponte 2016) and milk-foxtail millet powder (Simha et al. 2016).

420 *Glass transition temperatures*

DSC thermograms obtained for both NGFMF and GFMF at a range of water activities 421 between 0.11 to 0.97 at 10, 25 and 40 °C, gave the values of glass transition temperatures 422 (T_g) . T_g was obtained from the midpoint of the typical heat capacity change in the 423 thermogram line over a range of temperatures (Sandoval et al. 2009). Increasing temperatures 424 did not have any significant effect on the T_g values of both flours. Fig. 7 (a, b) shows the 425 relationship between T_g and water activity at constant temperature. It was observed that as the 426 water activity increased the values of T_g decreased linearly for both the flours, ranging 427 428 between 82.25 to 28.67 °C for NGFMF and from 51.11 to 11.83 °C for GFMF at all three temperatures. The lower values of GFMF could be attributed to the conversion of starch into 429 sugars and other compounds due to germination of foxtail millet flours. The low molecular 430 431 weight of sugars and other compounds formed of germination, have been reported to cause plasticizing effect of small molecules on starchy products (Chen and Yeh 2000). While, the 432 decrease in T_g values with increasing temperatures and water activities could be explained by 433 the plasticizing effect of water. Increasing temperatures causes decrease in the attractive 434 435 forces, thereby causing slow motion of water molecules, that gets easily adsorbed on the flour surfaces, thus decreasing the T_g values. Whereas, increasing water activities, increases the 436 437 moisture content in the flours that acts as plasticizers, thus changing the glassy material into 438 rubbery state (Alpizar-Reyes et al. 2017).

The experimental values of T_g for NGFMF and GFMF stored at different water activities and temperatures were fitted in Eq. 3.8.12 with a R² value of 0.991. The estimated values of T_g were found to be 124.36 °C for NGFMF and 101.82 °C for GFMF. This difference in the result could be due to erratic decrease in the experimental T_g values with increasing water activity, predominantly in NGFMF as compared to GFMF. Similar justification was provided for lower T_g values of dry amorphous materials in comparison to pure starch materials by Sandoval et al. (2009).

446 Conclusion

The germinated and non-germinated flours obtained from high pressure processed foxtail 447 millet grains were studied for their moisture sorption characteristics to understand the 448 interdependence between the water and the foxtail millet grain components. Thus, the 449 450 moisture sorption isotherms provided information regarding the stability of these flours in any environment. Apart from this, the thermodynamic properties were also studied that gave 451 information on the changes that took place in the flours during storage. Sorption moisture 452 isotherms showed a sigmoidal type II shape. Experimental data for these isotherms at 10, 25 453 and 40 °C fitted satisfactorily to the GAB model. Sorption properties of processed 454 455 germinated and non-germinated foxtail millet flour indicated that the percent of bound water decreased for NGFPF and increased for GFPF, as the temperature increased, confirming that 456 457 the availability of active sites decreased due to a reduction in the total sorption ability for NGFPF and vice-versa for GFPF. This work also suggested that the moisture sorption, 458 459 thermodynamic properties, spreading pressures and glass transition temperatures can be related to obtain conditions guaranteeing maximum stability for storage of the flours. 460

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TABLES

Property	GFMF	NGFMF
Bulk density (g l ⁻¹)	592±12.2 ^b	732±10.4 ^a
Porosity (% volume)	67.8±0.5ª	53.4 ± 0.6^{b}
Occluded air content (ml 100ml ⁻¹)	105.3±2.0 ^a	94.6 ± 1.8^{b}
Water absorption capacity (g 100g ⁻¹)	$157.4{\pm}1.8^{a}$	148.2±2.3 ^b
Oil absorption capacity (g 100g ⁻¹)	98.0±0.5ª	$80.4{\pm}1.1^{b}$
Dispersibility (%)	82.2±0.5ª	75.4 ± 0.3^{b}
Insolubility index (ml)	3.1 ± 0.2^{b}	4.7±0.1ª
Swelling capacity (ml g ⁻¹)	5.2 ± 0.2^{a}	4.0 ± 0.3^{b}
Wettability (s)	58.0±1.0ª	$52.0{\pm}1.0^{b}$
Flowability (θ)	70.1±0.4ª	62.3±0.9 ^b
Emulsion activity (%)	64.5±0.1ª	48.2 ± 0.0^{b}
Emulsion stability (%)	82.4±0.2 ^a	66.5±0.1 ^b
Foaming capacity (%)	5.2 ± 0.4^{b}	$8.8{\pm}0.5^{a}$
Foaming stability (%)	57.8 ± 0.7^{b}	78.8 ± 0.9^{a}
Gelatinization properties:		
1. Onset temperature (°C)	70.05 ± 0.92^{a}	70.05 ± 1.03^{a}
2. Peak temperature (°C)	73.16±1.11 ^a	$73.01{\pm}1.13^{a}$
3. Conclusion temperature (°C)	77.25 ± 0.88^{a}	77.13 ± 0.98^{a}
4. Gelatinization enthalpy (J/g)	2.306±0.110 ^a	1.563 ± 0.099^{b}

Values are mean of three replications (n=3) and expressed as mean \pm standard deviation. Means with different superscript within the same row are significantly different at p \leq 0.05.

Table 2 Estimated parameters of GAB sorption model for NGFMF

<i>T</i> (°C)	W _o (%)	G	k	R ²
10	3.235±0.100 ^a	43.171±0.150 ^a	0.780±0.020 ^a	0.99
25	$2.662{\pm}0.150^{b}$	28.260 ± 0.200^{b}	$0.820{\pm}0.020^{a}$	0.99
40	2.364 ± 0.090^{b}	32.720±0.190 ^c	$0.881 {\pm} 0.025^{a}$	0.99

Values are mean of three replications (n=3) and expressed as mean \pm standard deviation.

Table 3Estimated parameters of GAB sorption model for GFMF

<i>T</i> (°C)	Wo (%)	G	k	R ²
10	$2.987{\pm}0.090^{a}$	42.701 ± 0.220^{a}	$0.852{\pm}0.025^{a}$	0.99
25	$2.476{\pm}0.095^{a}$	28.912 ± 0.150^{b}	0.896 ± 0.030^{a}	0.98
40	2.063 ± 0.090^{a}	33.661±0.150 ^c	0.944 ± 0.010^{a}	0.99

652 Values are mean of three replications (n=3) and expressed as mean \pm standard deviation.

FIGURES



669	Fig. 1	Protease and amylase activity variation during the germination of foxtail millet
670		grains (Temperature= 25±2 °C; RH=80%)
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672		
673		
674		
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676		





Fig. 2 Moisture sorption isotherms for (a) NGFMF and (b) GFMF at 10, 25 and 40 °C
obtained by static gravimetric method





Fig. 3

Net isosteric heat of sorption (Q_{st}) as a function of moisture content for (a) NGFMF and (b) GFMF. The isoteric heat of sorption at any temperature can be obtained by adding the latent heat at that temperature to the Qst values.







699 Fig. 4Integral entropy (ΔS) as a function of moisture content of (a) NGFMF and (b)700GFMF





Integral enthalpy-integral entropy compensation for (a) NGFMF and (b) GFMF

Fig. 5







Fig. 6 Spreading pressure isotherms as a function of water activity of (a) NGFMF and(b) GFMF





Fig. 7 Effect of water activity on glass transition temperatures of (a) NGFMF and (b) GFMF