

Preparation of Functional Pasta Supplemented with Amaranth Pregelatinized Extruded Flour

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Semolina pasta with improved nutritional properties is increasingly demanded by consumers. High protein content and low starch digestibility are features desired for a functional pasta. In this work, amaranth flour was used for 50% supplementation of semolina in pasta preparation. Raw amaranth flour increased protein from 11.7 g/100 g to about 14.0 g/100, dietary fiber from 6.8 g/100 g to about 8.0–10.0 g/100 g. However, raw amaranth flour deteriorated the texture of cooked pasta by increasing hardness from 15.7 to 52.8 N and reducing cohesiveness from 0.71 to 0.55. Pregelatinizations at 50 and 100% were explored to reduce the adverse effects of raw amaranth flour. The gelatinized amaranth flour mimicked the texture of semolina pasta, although cooking loss increased from 3.3 g/100 g to about 9.2 g/100 g, and the water absorption showed a marked reduction from 116.4% to about 80.0–84.0%. The *in vitro* starch digestibility was similar for pasta made with semolina and pasta containing pregelatinized starch, although digestibility decreased from 95 to 85% for raw amaranth flour. Overall, the results showed that amaranth flour offered advantages and drawbacks for the formulation of pasta with improved nutritional features.

Keywords: pasta, semolina, amaranth flour, starch gelatinization, pasta texture

INTRODUCTION

The formulation and characterization of functional foods have been a growing field in academic research and industrial production. An increasing number of consumers are demanding food products with health benefits. Functional ingredients in food products include dietary fiber, prebiotics and probiotics, antioxidant compounds and vitamins. The challenge to producing functional foods is to achieve sensory, texture, and cooking quality that are similar to traditional food (Alongi and Anese, 2021).

Pasta is consumed in practically all world regions, is easy to produce and prepare and relatively inexpensive. Pasta formulation offers flexibility for supplementation to increase protein content (Jayawardena et al., 2018), dietary fiber (Tudorică et al., 2002), or antioxidant capacity (Laus et al., 2017). Dehulled pulses (e.g., lentils and peas) are important sources of protein supplementation for pasta (Bresciani et al., 2021). Oat, wheat and rice brans are commonly used to improve soluble fiber content (Krishnan et al., 2012). Polyphenols from different sources are increasingly incorporated in commercial pasta formulations (Turco et al., 2019). However, the use of non-wheat ingredients

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commonly yields pasta with undesired characteristics, including disrupted texture and sensorial attributes. On the other hand, the surge of consumers with healthy (e.g., natural ingredients) and socio-ecological (non-animal ingredients) demands triggered the research on new pasta formulations. In particular, pulses offer certain economical and dietary advantages, since these flours have been explored for the management of diabetes (Ramdath et al., 2016). Reports have shown that pasta produced from amaranth flour exhibits decreased texture firmness and cooking time, while pasta from quinoa mainly shows an increased cooking loss (Schoenlechner et al., 2010). Amaranth has high protein with essential amino acids and waxy type starch with small granule size, which can confer specific characteristics to foods (pasta, bakery products, snacks, etc.) in the form of pregelatinized ingredients (Garcia-Valle et al., 2021). Green banana flour is a by-product that has been proposed for improving pasta quality and nutritional properties (Garcia-Valle et al., 2021). The high protein content of the chickpea flour (21.73 g/100 g) is advantageous to improve the nutrition characteristic of semolina pasta. The addition of chickpea flour in the formulation led to reduced in vitro starch digestibility as a consequence of the high protein and dietary fiber content. Interactions of dietary fiber with starch (Zheng et al., 2016) and inhibition of α -amylase by cellulose (Dhital et al., 2015) are potential mechanisms involved in decreased starch digestibility. Pregelatinization by extrusion was used for conditioning mango flour to improve texture and starch digestibility in pasta formulations (Garcia-Valle et al., 2021). Yellow lentil flour was subjected to extrusion-cooking to obtain pasta with improved texture (Bresciani et al., 2021).

Non-conventional flours offer nutritional advantages over traditional cereal flour. For instance, pulses are relatively inexpensive and commonly contain large amounts of proteins. However, the preparation of pasta incorporating nonconventional wheat flour commonly results in texture problems. Hydrocolloids (e.g., gums) have been used in combination with non-conventional flours to fix the texture requirements for pasta (Chauhan et al., 2017). However, hydrocolloids might increase pasta fabrication costs and add undesired sensorial attributes. Partial or total gelatinized flours are ad hoc alternatives for hydrocolloids (Bento et al., 2021). Pasta prepared with gelatinized flour exhibited optimal cooking time as well as high protein and dietary fiber contents. Motivated by the aforementioned reports, the present study aimed to evaluate the gelatinization degree (50 and 100%) of amaranth flour in the structuration of the pasta and its effect on the texture and in vitro starch digestion. Semolina pasta and pasta with raw amaranth flour were used as control.

MATERIALS AND METHODS

Materials

Wheat flour (*Triticum durum*) was obtained from San Blas Milling Co. (Puebla, Mexico). Amaranth grains were collected in an experimental crop field in Central Mexico (Texcoco, Estado de México, México). Amaranth grains were milled and sieved at 40 mesh. The different chemical reagents were purchased at JT Baker (Avantor Performance Materials, Center Valley, PA, Unite States), Sigma Aldrich (Sigma Chemical Co., St Louis MO, USA), and Fermont (Productos Químicos Monterrey, Monterrey, México).

Flour Extrusion

Amaranth flour was extruded with a single screw-extruded (Beutelspacher, CDMX, Mexico) at a constant rotation rate of 75 rpm. Before extrusion, flour was conditioned for 18 h with a water content of 35% (base of dry flour) in a plastic bag. Two constant temperatures were used in the three zones of the extruder (first zone of the barrel, blend zone, and end zone): 50 and 80 °C. Two batches were prepared for each sample. After extrusion, the pellets were milled and sieved (mesh 40, mean particle size 0.042 cm). Two degrees of flour gelatinization of 50% (E50) and 100% (E100) were used for pasta preparation.

Pasta Formulation and Processing

Pasta formulations consisted of blends of durum wheat semolina and amaranth flours: S100 denotes 100% semolina, S50-A50 denotes 50% semolina and 50% amaranth flour, S50-E50 denotes 50% semolina and 50% amaranth flour with 50% gelatinization, and S50-E100 denotes 50% semolina and 50% amaranth flour with 100% of gelatinization. Sheeting was carried out with a manual Pasta & Co. machine (Metaltex Molsheim, France). Afterward, the pasta formulations were dried at 45°C for 4 h in an oven under a relative humidity of about 40% (Biotecnica del Bajio S.A. de C.V., Celaya, Mexico). The pasta formulation was stored at 25 \pm 2°C in sealed plastic containers until further analysis.

Chemical Composition

Moisture, ash, fat and protein (N \times 6.25) contents were determined according to AACC Method 44–15, 08–01, 30–25, and 46–13, respectively (AACC, 2010). Total starch was measured according to AACC method 76.13 (AACC, 2010) using a total starch analysis kit from Megazyme (Wicklow, Ireland). Total dietary fiber (TDF) was assessed for cooked samples at the optimum cooking time, using two different methods: 1) Method AACC 2000 32–05, which quantifies the indigestible material after heat treatment of the food sample (i.e., non-starch polysaccharides), and 2) enzyme-based method proposed by McCleary (McCleary, 2007), which allows total resistant starch fractions to remain associated to the TDF (non-starch polysaccharides) residues, under conditions that resemble the physiological behavior. All analyses were performed in triplicate.

Thermal Properties

The gelatinization degree (DG) of raw pasta was performed with differential scanning calorimetry (DSC). Approx. 2.0 mg sample and 7 μ L of deionized water were equilibrated for 12 h at room temperature in a DSC pan, and subjected to the heating ramp for the range 20–120°C at a heating rate of 10°C/min in a DSC model 2010 (TA Instrument, New Castle, NJ). The temperature of the phase transition and enthalpy (area under the phase transition

curve) were calculated with the software of the equipment. The degree of gelatinization (DG) was calculated as:

$$DG = \frac{Entalpy of the raw pasta - Enthalpy of the cooked pasta}{Enthalpy of the raw pasta} \times 100$$
(1)

Cooking Quality

The cooking quality (optimum cooking time and cooking loss) of the pasta formulations was assessed according to AACC 2000 66–50 method (AACC, 2010). The water absorption was determined for 10 g of pasta sample (cut into 5 cm long pieces), and cooked in 300 ml at optimum cooking time. The pasta was then drained and rinsed with 20 ml of distilled water at room temperature for 2 min. The sample was weighed after reaching temperature. The water absorption was determined as follows:

$$Water absorption = \frac{Weight of cooked drained pasta - Weight of raw pasta}{Weight of raw pasta}$$
(2)

Texture Analysis

Two batches of each pasta formulation were cooked at boiling temperature for optimum cooking time. Five subsamples were used for texture profile analysis (TPA) using the texture analyzer (Brookfield CT3, Middleboro, MA 02346–1031) within 5 min after cooking. The measurements were carried out with a cylindrical probe (TA 4/1000, DD 38.1 mm, L 20 mm) and 10 kg load cell. The samples were compressed to 75% of their original height. The determining variables were pasta hardness, adhesiveness, cohesiveness, elasticity, and chewiness. TexturePro CT software was used to record the data and estimate the texture parameters.

In vitro Starch Digestibility

The *in vitro* starch digestibility was determined according to the procedure described by Bello-Pérez et al. (Bello-Perez et al., 2019), where starch digestion was carried out from the mouth to the small intestine. Results were presented as a percentage of digested on a dry basis.

Statistical Analysis

Results were reported as the mean \pm standard error. Data were analyzed by one-way analysis variance (ANOVA) (p < 0.05) followed by LSD multiple comparison procedure to determine significant differences among samples (SPSS v. 20 IBM, NY).

RESULTS AND DISCUSSION

Chemical Composition

Pasta made with semolina-amaranth flour blends contained reduced moisture contents (4.0-4.7 g/100 g) relative to the pasta made with semolina (6.6 g/100 g). The differences can be attributed to the reduced gluten content in pasta containing amaranth flour.

Gluten exhibits a marked capacity for water retention, providing pasta with enhanced elasticity (Almutawah et al., 2007). Protein, ash and fat contents increased for pasta supplemented with amaranth flour, and the increase was not dependent on whether the flour was gelatinized or not. The increase of such fractions is linked to the high contents in amaranth flour, posing this flour as a suitable ingredient to increase the nutritional characteristics of pasta. On the other hand, the total starch content showed an important decrease from about 65.0 g/100 g for semolina pasta to about 55.0 g/100 g for pasta with amaranth flour. Decreased starch content is an important issue given the metabolic syndrome problems related to high ingesta of digestible carbohydrates (Regalado-Rentería et al., 2020). Dietary fiber showed an important increase caused by the non-starch components of the amaranth flour (Tamsen et al., 2018). It should be recalled that the method AACC 200 32-05 determines total dietary fiber (non-starch polysaccharides), but resistant starch (fraction of the dietary fiber) is not quantified. The second method determines both non-starch polysaccharides and resistant starch. Hence, the results in Table 1 indicated that the pasta contains resistant starch that acted as dietary fiber.

Cooking Quality

The optimum cooking time of semolina pasta was about 16.6 min and decreased with the addition of the amaranth flour (**Table 2**). Gelatinization reduced the optimum cooking time to about 10-12 min, which was an expected result given the preconditioned structure of pasta with gelatinized starch. The optimum cooking time agrees well with the values reported by Yalcin and Basman (Yalcin and Basman, 2008) for gluten-free corn noodles prepared with different proportions of gelatinized starch. The cooking time reduction was accompanied by a negative effect as reflected by the increased levels of cooking loss, which increased from about 3.3 g/100 g for semolina pasta to about 9.2 g/100 g for pasta made with gelatinized flour.

The cooking loss parameter reflects the solids that were leached in the cooking process. It is an index of the resistance to disintegration during pasta cooking (Table 2). The cooking loss of semolina pasta was about 3.3 g/100 g, and increased to 9.2 g/100 g for pasta containing gelatinized amaranth flour. This pattern is related to short chains present in amaranth starch (Paredes-López et al., 1994) that were disorganized during the extrusion (partial or complete gelatinization) and during cooking. The absence of a gluten network to entrap the pasta components (e.g., proteins and starch) and the availability of amylose chains by the gelatinization process led to pasta with high losses during cooking (Marti and Pagani, 2013). These results agree with Larrosa et al. (Larrosa et al., 2016), who found that gluten-free pasta exhibited high cooking loss values as a consequence of the lack of a robust network to stabilize the pasta structure in the cooking process. The water absorption capacity was negatively correlated (Pearson correlation = -0.96, p < 0.03) with the cooking loss. Gluten is hydrated to form a stabilizing network with a high capacity for water retention (Roccia et al., 2009). The reduced or even lack of gluten fraction in a pasta formulation weakened the structure, leading to a poor capacity of water retention during cooking.

Sample	Moisture (g/100 g)	Protein (g/100 g)	Ash (g/100 g)	Fat (g/100 g)	Total starch (g/100 g)	Dietary fiber (AACC) (g/100 g)	Dietary fiber (McCleary, 16 h)* (g/100 g)
S100	6.6 ± 0.2^{a}	11.7 ± 0.2 ^a	1.1 ± 0.03 ^a	0.88 ± 0.02 ^a	64.73 ± 1.3 ^a	5.78 ± 0.04^{a}	6.56 ± 0.85^{a}
S50-A50	4.6 ± 0.1^{b}	13.7 ± 0.2 ^b	2.2 ± 0.03^{b}	2.80 ± 0.27^{b}	56.01 ± 0.72 ^b	7.43 ± 0.25^{b}	7.99 ± 0.13^{b}
S50-E50	4.7 ± 0.1^{b}	13.8 ± 0.1 ^b	2.3 ± 0.1^{b}	2.47 ± 0.12 ^b	$54.57 \pm 0.83^{\circ}$	$6.93 \pm 0.08^{\circ}$	$9.64 \pm 0.14^{\circ}$
S50- E100	$4.0 \pm 0.1^{\circ}$	13.9 ± 0.1 ^b	2.2 ± 0.01^{b}	$2.26 \pm 0.07^{\circ}$	55.27 ± 1.04 ^{b,c}	7.42 ± 0.21^{b}	$9.35 \pm 0.93^{\circ}$

Values are reported as means ± standard error. Column with different lower-case letters in columns indicate significant differences (p < 0.05). The letter "Sx" denote the percentage of semolina content, "Ax" denote the amaranth flour content, "Ex" denote the extruded amaranth flour content, * cooked pasta,

Sample	Optimum cooking time (min)	Cooking loss (g/100 g)	Water absorption (%)	Raw pasta dimensions (mm)	Cooked pasta dimensions (mm
S100	16.6 ± 0.6^{a}	3.3 ± 0.9^{a}	116. 4 ± 4.1 ^a	W: 6.2 ± 0.3^{b} T: 2.7 + 0.4 ^B	W:7.3 ± 0.4 ^a T: 3.0 ± 0.2 ^A
S50-A50	12.5 ± 0.4^{b}	6.9 ± 1.0^{b}	86.8 ± 0.3^{b}	$W: 5.9 \pm 0.2^{a}b$ T: 2.1 + 0.1 ^A	$W:7.2 \pm 0.1^{a}$ T: 2.9 + 0.2 ^A
S50-E50	$11.8 \pm 0.2^{\circ}$	$9.2 \pm 0.9^{\circ}$	$84.2 \pm 3.0^{\circ}$	W: 5.6 ± 0.4^{a} T: 2.5 $\pm 0.3A^{B}$	W: 7.1 ± 0.1 ^a T: 3.6 ± 0.8 ^B
S50-E100	10.0 ± 0.4^{d}	$9.2 \pm 0.1^{\circ}$	80.2 ± 0.6^{d}	W: 5.8 ± 0.2^{a} T: 2.4 ± 0.2^{A}	$W:7.1 \pm 0.5^{a}$ T: 3.4 ± 0.6 ^B

Values are reported as means ± standard error, Values are reported as means ± standard error, Column with different letters in columns indicate significant differences (p < 0.05). The letter "Sx" denote the percentage of semolina content. "Ax" denote the amaranth flour content. "Ex" denote the extruded amaranth flour content. W denote width, T denote thick of pastas.

The addition of amaranth flour produced pasta with reduced width and thickness dimensions. This effect is probably due to the reduced water retention capacity of pasta with lowered gluten content. However, the effect was not visible for the width of cooked pasta where the dimension differences were not statistically significant (p < 0.05). In contrast, the thickness was higher for pasta containing gelatinized flour. The amylose released in the gelatinization process produced a pasta structure with the capacity of retaining water during the cooking process.

Texture

Cooked semolina pasta (S100) and cooked pasta containing gelatinized flour (S50-E50 and S50-E100) exhibited similar hardness values (13.0-15.0 N), although the latter pasta with slightly smaller values (Table 3). In contrast, the hardness of cooked pasta containing non-gelatinized amaranth flour (S50-A50) was markedly high (52.8 N). Probably, the native starch granules of the amaranth flour (small size and high amylopectin content) induced the increased hardness value of the cooked pasta. This effect was reflected in adhesiveness, which was the smallest value for semolina pasta (0.07 N-s), increased for pasta containing gelatinized flour (0.43-0.62 N.s) and achieved the highest value (2.04 N.s) for pasta with non-gelatinized flour. Chewiness showed a similar trend. In contrast, cohesiveness and elasticity were only slightly affected by the addition of amaranth flour. In particular, the amaranth flour addition reduced the pasta elasticity, an effect that can be ascribed to the reduced gluten content. Schoenlechner et al. (Schoenlechner et al., 2010) reported that the reduced firmness of amaranth pasta may be attributed to the lack of gluten, which led to a weak

structural matrix. They used casein and egg protein to supplement gluten and improve pasta texture. The results in Table 3 showed that pregelatinized flour improved some texture features (e.g., hardness and elasticity), while maintaining the reduced gluten content.

Thermal Properties

Figure 1. a illustrates the thermal flow patterns for the different pasta formulations. Three endothermic peaks were obtained, which can be attributed as follows (Table 4). The first peak at the lowest peak temperatures of 63-65°C reflects the denaturation of the gluten (Falcão-Rodrigues et al., 2005) and the proteins contained in the amaranth flour (Martínez and Añón, 1996). The second peak located at about 74.5-77.0°C can be linked to starch gelatinization, whereas the peak at the highest temperature of 85.0-99.0°C to amylose-lipid complexes (Biliaderis et al., 1985). Figure 1. b shows the thermal flow pattern of the different pasta formulations. Except for the semolina pasta S100, the pasta formulations containing amaranth flour exhibited the three endothermic peaks illustrated in Figure 1. a. The onset and peak denaturation temperatures showed a slight increase with the addition of the amaranth flour. However, the denaturation enthalpy showed an important decrease from 7.9 J/g for S100 to 0.9 J/g for S50-E100. This suggests that gluten from semolina interacted with proteins to form a weak matrix with lowered ability to retain water (Table 2). As expected, the addition of gelatinized amaranth flour led to a decrease in the gelatinization enthalpy. Also, the gelatinized amaranth flour produced pasta with a reduced content of amylose-lipid inclusion complexes, as revealed by the reduced enthalpy of the thirst endothermic peak.

Sample	Hardness (N)	Adhesiveness (N.s)	Cohesiveness	Elasticity	Chewiness (N)	
S100	15.7 ± 1.4 ^a	0.07 ± 0.03^{a}	0.71 ± 0.05 ^a	0.73 ± 0.03 ^a	8.74 ± 1.5 ^a	
S50-A50	52.8 ± 1.9^{b}	2.04 ± 0.45^{b}	0.55 ± 0.02^{b}	0.61 ± 0.04^{a}	21.64 ± 1.1 ^b	
S50-E50	$13.2 \pm 1.7^{\circ}$	$0.62 \pm 0.18^{\circ}$	0.62 ± 0.10^{a}	$0.63 \pm 0.06^{b,a}$	5.23 ± 1.3°	
S50-E100	14.2 ± 2.1 ^{a,c}	$0.43 \pm 0.12^{\circ}$	0.61 ± 0.08^{a}	$0.62 \pm 0.06^{b,a}$	$6.14 \pm 2.0^{\circ}$	

Values are reported as means ± standard error. Column with different lower-case letters in columns indicate significant differences (p < 0.05). The letter "Sx" denote the percentage of semolina content. "Ax" denote the amaranth flour content. "Ex" denote the extruded amaranth flour content.



In vitro Digestibility

Figure 2 shows the digestograms of the cooked pasta. All cases presented an initial fast starch hydrolysis rate, achieving starch conversions of the order of 25–30%. For longer times, the hydrolysis curves evolved in different directions with an apparent slower hydrolysis rate. A log-of-slope (LOS) analysis of the digestograms was carried out to verify the presence of two

phases in the starch hydrolysis kinetics. Figure 2 also shows the LOS results where two-phase hydrolysis was exhibited. The slope of the LOS plots corresponds to the starch hydrolysis rate according to first-order kinetics. The enzymatic hydrolysis transited from a fast rate for small times, up to 100 min, to a slow rate for longer times. This means that the enzymatic hydrolysis of starch in the pasta formulations did not follow a uniform pattern, but rather it transited to a slower regimen at a certain time (Butterworth et al., 2012). The transition might be linked to changes in the pasta structure as a consequence of the enzymatic action. For relatively short times, starch is highly available for amylolytic enzyme binding. The advance in the starch hydrolysis disrupted the pasta structure, dispersing the starch chains in the aqueous dispersion and hardening the interaction between the enzyme and the substrate. The results in Figure 2 suggested that the digestograms can be fitted with a double exponential function of the form

$$X(t) = X_{\infty} + A_{fast} \exp(-k_{fast}t) + A_{slow} \exp(-k_{slow}t)$$
(3)

where X(t) is the starch conversion, X_{∞} is the limiting conversion (i.e., for long times), A_{fast} and A_{slow} are constants, and k_{fast} and k_{slow} are hydrolysis rates for the fast and slow phase, respectively. Since X(t = 0) = 0, the constants should satisfy the constraint $X_{\infty} + A_{fast} + A_{slow} = 0$. The continuous lines in Figure 2 depict the least-squares fitting of the experimental data by Eq. (Alongi and Anese, 2021). For all cases, the mean quadratic error is less than 5%. The difference between the fast and slow hydrolysis rate constants was one order of magnitude, showing a clear separation of the time scales of the starch hydrolysis (Table 5). That is, the fast hydrolysis takes place at time scales of the order of 80 min, whereas the slow hydrolysis rate at time scales of the order of 800 min. The pasta made with non-gelatinized amaranth (S50-A50) starch presented the lower values of both the fast and slow hydrolysis rates. In contrast, the pasta formulations containing gelatinized amaranth flour exhibited higher values of the estimated hydrolysis rates. These results could be expected since the gelatinized starch is easier to hydrolyze than granular starch (Biliaderis et al., 1985). The effect was also reflected in the limiting starch conversion. The lowest value was shown by the pasta S50-A50 (about 85.27%) and the highest value (100%) by the pasta with the highest content of gelatinized amaranth flour. Table 5 also shows the digestible starch fractions according to the classification by Englyst et al. (Englyst et al., 1992). The pasta made with non-gelatinized amaranth flour (S50-A50) presented the lowest value of the rapidlñy digestible starch (RDS) fraction, although the value was only slightly smaller than that presented by the pasta made only with semolina (S100). The pasta formulation containing



FIGURE 2 | LOS computations showing two hydrolysis time scales. (A) Cooked S100, (B) Cooked S50 A50. (C) S50-E50. (D) S50-E100.

Sample	First peak (protein denaturation)				Second peak (starch gelatinization)			Third peak (starch-lipid complex)				
	То (°С)	Тр (°С)	Tc (°C)	∆H (J/g)	То (°С)	Тр (°С)	Tc (°C)	∆H (J/g)	То (°С)	Тр (°С)	Tc (°C)	∆H (J/g)
S100	57.2 ± 0.4 ^a	63.5 ± 0.5ª	70.7 ± 0.2a	7.9 ± 0.3 ^a	N/D	N/D	N/D	N/D	N/D	N/D	N/D	N/D
S50-A50	59.7 ± 0.2 ^a	64.6 ± 0.1 ^a	68.9 ± 0.6 ^a	1.8 ± 0.7 ^a	72.0 ± 0.2 ^b	76.9 ± 1.2 ^b	81.0 ± 0.5 ^b	6.0 ± 0.7 ^b	91.9 ± 0.3 ^a	98.9 ± 0.1ª	106.7 ± 0.5ª	2.2 ± 0.5 ^a
S50-E50	58.8 ± 0.4 ^b	63.5 ± 0.1 ^b	67.9 ± 0.1 ^b	1.7 ± 0.4 ^a	69.0 ± 0.4 ^c	73.2 ± 0.2°	78.3 ± 0.6 ^c	3.8 ± 0.1°	89.9 ± 0.7 ^b	95.1 ± 1.3 ^b	105.9 ± 2.8ª	1.9 ± 0.3 ^a
S50-E100	61.2 ± 0.08 ^c	65.2 ± 0.2 ^c	70.9 ± 0.5 ^c	0.9 ± 0.03 ^b	73.0 ± 0.3 ^d	76.2 ± 0.5 ^b	80.7 ± 0.1 ^b	2.9 ± 0.01 ^d	81.0 ± 0.2 ^c	85.4 ± 0.3°	90.6 ± 1.2^{b}	0.6 ± 0.2 ^c
Amaranth Flour	N/D	N/D	N/D	N/D	69.1 ± 0.1°	74.4 ± 0.6 ^c	85.5 ± 1.1 ^d	5.2 ± 0.1 ^e	93.4 ± 1.5 ^a	99.1 ± 0.6 ^d	104.6 ± 0.2 ^a	1.4 ± 0.2 ^b

TABLE 4 | Thermal properties of flours and raw pastas made with wheat semolina and amaranth flours.

Values are reported as means \pm standard error. Column with different lower-case letters in columns indicate significant differences (p < 0.05). The letter "Sx" denote the percentage of semolina content. "Ax" denote the amaranth flour content. "Ex" denote the extruded amaranth flour content. T_o, onset temperature, T_p, peak temperature, T_o, conclusion temperature, ΔH , phase transition enthalpy.

Sample	$k_{\text{fast}} \times 10^{-2} \text{ (min}^{-1}\text{)}$	k _{slow} × 10 ^{−3} (min ^{−1})	X _∞	RDS (%)	SDS (%)	RS (%)
S100	1.2 ± 0.03^{a}	2.0 ± 0.04^{a}	93.03 ± 0.12 ^a	35.05 ± 0.34 ^a	36.54 ± 0.43^{a}	28.41 ± 0.67 ^e
S50-A50	1.1 ± 0.03 ^a	1.3 ± 0.02^{b}	85.27 ± 0.23 ^b	33.79 ± 0.28^{b}	30.12 ± 0.38^{b}	36.09 ± 0.56 ^k
S50-E50	1.2 ± 0.02^{a}	1.9 ± 0.02^{a}	95.37 ± 0.16 ^c	35.75 ± 0.31 ^a	36.56 ± 0.26 ^a	27.69 ± 0.43 ^e
S50-E100	1.6 ± 0.03^{b}	$2.5 \pm 0.04^{\circ}$	100.0 ± 0.08 ^d	37.68 ± 0.40 ^c	37.33 ± 0.18 ^a	24.99 ± 0.21°

TABLE 5 | Kinetics constants and digestibility fractions of cooked pasta made with wheat semolina and amaranth flours.

Values are reported as means ± standard error. Column with different lower-case letters in columns indicate significant differences (p < 0.05).). The letter "Sx" denote the percentage of semolina content. "Ax" denote the amaranth flour content. "Ex" denote the extruded amaranth flour content. KRDS denote kinetics constant for RDS, kSDS denote kinetics constant for SDS, RDS denote rapidly digestible starch content, SDS denote slowly digestible starch content, RS denote resistant starch content.

100% of gelatinized flour showed the highest value of the RDS, which can be attributed to the increased susceptibility of leached starch chains to amylolytic enzymes. In the same way, this pasta exhibited the lowest value of the resistant starch (RS) fraction, which is in line with the highest value of the limiting starch conversion.

DISCUSSION

The use of non-traditional flours as innovative ingredients for functional starchy food formulations has been a research and commercial trend in the recent decade (Nagash et al., 2017). The results of this work showed that amaranth flour is a visible complement to improving the nutritional quality of semolina pasta. Amaranth flour adds proteins and dietary fiber while reducing the total starch content of pasta, which are improvements over semolina pasta. However, the incorporation of amaranth flour degraded the pasta texture, and possibly the acceptability of consumers. Many studies have proposed to use of different polysaccharides to compensate for the degradation of pasta texture by non-conventional flours. For instance, Cai et al., (Cai et al., 2016) used xanthan gum for improving the cohesiveness of rice noodles. Feng et al. (Feng et al., 2020) studies chitosan, xanthan gum and sodium alginate as texture supplements for pasta based on wet sweet potato. The results showed that such polysaccharides improved the texture and cooking properties of the pasta. However, there is an increasing group of consumers preferring food products without additives. The notion that natural and traditional recipes are linked to a healthy food practice has motivated the exploration of other alternatives to improve pasta quality. The results of this work showed that gelatinized flour can play the role of a texture improver to mimic the characteristics of semolina pasta. Gelatinization had the drawback that the starch digestibility might be increased. Besides texture improvement, pasta additives should maintain or even reduce starch digestibility, a must that should be considered in the face of the so-called metabolic syndrome. Overall, the results reported in the present study are motivating, although more research is required to reduce the adverse effects of gelatinized starch on pasta digestibility. The combination of gelatinized flour with a minimal amount of polysaccharides might be a way to obtain pasta with desired physicochemical, textural and digestibility characteristics.

CONCLUSION

The addition of amaranth flour in the formulation of traditional pasta based on semolina increased the nutritional characteristics. Protein and dietary fiber increased by the supplementation of semolina with amaranth flour. However, native amaranth flour degraded the cooked pasta texture by increasing hardness and reducing elasticity. Gelatinized amaranth flour was used instead to alleviate the drawbacks of granular amaranth flour in texture, leading to pasta that mimics the textural characteristics of semolina pasta. However, gelatinized amaranth flour induced undesired effects, such as increased cooking loss and in vitro starch digestibility. Overall, gelatinized amaranth flour is a viable ingredient to improve the nutritional characteristics of semolina pasta, although its incorporation should be tuned to reduce the adverse effects on cooking quality and starch digestibility. The results of this work showed that amaranth flour with a partial gelatinization degree might be used to obtain some nutritional advantages while reducing adverse effects in texture and digestibility.

DATA AVAILABILITY STATEMENT

The raw data supporting the conclusions of this article will be made available by the authors, without undue reservation.

AUTHOR CONTRIBUTIONS

LAB-P conceived the study, analyzed, and interpreted the data. JAC-V performed the experiments. R-C-G, OP-R, and JA-R carried out the analysis and interpretation of the data. LAB-P and JA-R drafted the manuscript. All authors read and approved the final version of the paper.

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