

EFFECT OF PROCESSING ON THE STARCH AND GLYCEMIC
PROPERTIES OF DICHPRIA SPP.

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**Effect of Processing on the Starch and Glycemic
Properties of *Digitaria spp.***

by

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Thesis presented in partial fulfilment for the degree of:

Master of Technology (Food Technology)

Department of Food Technology

Faculty of Applied Sciences

Cape Peninsula University of Technology

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DECLARATION

I, Michelle Jordaan, hereby declare that the work contained in this proposal is my own original work and that it has not previously, in its entirety or in part been submitted at any other university for a degree.



Michelle Bernitta Jordaan

13. Oct. 2013.

Date

Abstract

Acha starch was isolated and purified from clean and milled acha grain. Functional, thermal and physicochemical properties of acha starch were analysed using appropriate methods. Wheat starch was used as the reference standard. Acha bread from acha grain was baked and the consumer sensory acceptability was evaluated and white wheat bread was used as the reference standard. The effect of baking, boiling, steaming and microwaving on the starch and glycemic properties of the acha starch was evaluated. With regard to thermal properties, gelatinisation temperature of acha and iburu starches typifies that of waxy starch. Acha starch has similar retrogradation temperature profiles as that of wheat. There were however significant differences in some of the functional properties (pasting and turbidity) and physico-chemical properties (*in vitro* starch digestibility), but no significant difference in the texture profile analysis (TPA) and water binding capacity (WBC). WBC of both acha varieties was higher than that for wheat starch. Due to its high break down viscosity, white acha starch can be included in foods that are subjected to high temperature processing. This indicates that both acha starch varieties can be used for hot and cold desserts as well as for soft jelly like sweets and confectionery toppings. A pre-screening exercise using carboxymethyl cellulose (CMC), Xanthan gum, yeast and acha starch as the variables was successful in concluding a recipe which rendered acha bread with the optimum specific loaf volume for both white and black acha bread. The optimum recipe consisted 8.0 % acha starch, 2.0 % xanthan gum, 2.0 % CMC and 1.0 % yeast. The majority of the consumer panellists found the crust colour, taste and aroma to be moderately desirable. This implies that most consumers find acha bread to have the potential to be marketed as wheat free bread. The different processing methods baking, boiling, microwaving and steaming, affected the black and white acha starch hydrolysis. The amount of starch hydrolysed for the different processing methods was in the following order: baking > boiling > microwaving > steaming. It can thus be concluded that different processing methods affects the micro structure and physical properties of the acha and wheat samples

which thus influence their starch hydrolysis. The equilibrium percentage of starch hydrolysed after 180 min incubation was affected differently for the various starches, black acha, white acha and wheat starch by the different processing methods and times. In the case of baking black acha starch and wheat bread were affected similarly. However, this was not the case for microwaving, steaming and boiling, where both acha starch varieties and wheat starch were affected in the same way. The rate of starch hydrolysis for both acha varieties and wheat grain for the different processing methods, steaming, boiling, microwaving and baking was affected to the same degree respectively.

Language and style used in this thesis are in accordance with the requirements of the International Journal of Food Science and Technology. This thesis represents a compilation of manuscripts where each chapter is an individual entity and some repetition between chapters has, therefore, been unavoidable.

TABLE OF CONTENTS

Chapter	Page
Declaration	ii
Abstract	iii
Table of Contents	v
List of Figures	ix
List of Tables	xiii
Acknowledgment	xiv
Dedication	xiv
1 Motivation and design of the study	
1.1 Introduction	1
1.2 Statement of the Research Problem	2
1.3 Broad Objectives	3
1.3.1 Specific objectives	3
1.4 Research Hypothesis	3
1.5 Research Delimitation	3
1.6 Significance of the Research	4
1.7 Expected Outcomes, Results and Contributions of the Research	5
References	5
2 Literature Review	
2.1 Overview of <i>Digitaria spp.</i>	7
2.2 Uses of Acha	7
2.3 Nutritional Composition of Acha	10
2.4 Physical Properties of Acha Starch	10
2.5 Chemical Properties of Cereal Starch	12
2.6 Functional Properties of Cereal Starch	15
2.7 Overview of Starch in relation to Digestion	18
2.8 Glycemic Properties of Starch	21
2.9 Measurement Techniques Resistant Starch	23
2.10 Effect of Processing on Functional and Glycemic Properties of Starch	23

References	27
3 Physicochemical, Functional and Thermal Properties of Acha Starch	
Abstract	32
3.1 Introduction	32
3.2 Materials and Methods	34
3.2.1 Source of acha grain and materials	34
3.2.2 Cleaning of acha grain	34
3.2.3 Isolation and purification of starch from acha grain	34
3.3 Physicochemical Properties of Acha Starch	34
3.3.1 Starch characteristics of acha starch	34
3.3.2 Total starch analysis for acha and wheat starch	34
3.3.3 Resistant starch analysis of acha and wheat starch	36
3.3.4 Digestible starch	39
3.3.5 Determination of rapidly and slowly digestible starch by in vitro rate of starch digestion	39
3.3.6 Determination of total amylose content of acha starch	40
3.3.7 Proximate composition of acha starch	41
3.4 Functional Properties of Acha Starch	41
3.4.1 Textural properties of starch gels	41
3.4.2 Turbidity of starch suspensions	42
3.4.3 Water binding capacity	42
3.4.4 Pasting properties	42
3.5 Thermal Properties of Acha Starch	43
3.6 Results and Discussion	44
3.6.1 Proximate composition of acha starch	44
3.6.2 Amylose content of acha starch	46
3.6.3 In vitro starch digestibility of acha and wheat starch	47
3.6.4 Functional properties of acha Starch	49
3.6.5 Thermal properties of acha starch	57
3.7 Conclusion	62
References	63

4	Production of Acha Bread and its Sensory Characteristics	
	Abstract	69
4.1	Introduction	69
4.2	Materials and Methods	70
	4.2.1 Source of acha grain and materials	70
	4.2.2 Cleaning of acha grain and production of acha flour	70
	4.2.3 Experimental design for acha bread	70
	4.2.4 Production of acha bread	71
	4.2.5 Physical properties of acha bread	71
	4.2.6 Sensory properties of acha bread	74
	4.2.7 Data analysis	74
4.3	Results and Discussion	74
	4.3.1 Effects of ingredients on acha bread loaf volume	74
	4.3.2 Sensory properties of acha bread	76
	4.3.3 Sensory attributes of interest to the consumer panellists	81
4.4	Conclusion	86
	References	87
5	Effect of Processing on the Starch and Glycemic Properties of Acha starch	89
	Abstract	89
5.1	Introduction	90
5.2	Materials and Methods	90
	5.2.1 Source of acha grain and materials	90
	5.2.2 Cleaning of acha grain	91
	5.2.3 Steaming of acha	91
	5.2.4 Microwaving of acha	91
	5.2.5 Production of acha bread	91
	5.2.6 Analysis of total starch of processed acha and wheat products	93
	5.2.7 Resistant starch analysis of processed acha and wheat products	93

	5.2.8 Digestible starch of processed acha and wheat products	94
5.2.9	Determination of rapidly and slowly digestible starch by <i>In vitro</i> rate of starch digestion	94
5.3	Statistical analysis	95
5.4	Results and Discussion	
	5.4.1 Effect of processing on starch hydrolysis	96
	5.4.2 <i>In-vitro</i> starch hydrolysis rate and hydrolysis index	106
	5.4.3 Modelling the effect of boiling on acha starch hydrolysis	108
	5.4.4 Modelling the effect of steaming on acha starch hydrolysis.	111
	5.4.5 Modelling the effect of microwaving on acha starch hydrolysis.	114
5.5	Total, Resistant and Digestible Starch in Acha and Wheat Bread	117
5.6	Total, Resistant and Digestible Starch in Acha and Wheat Grain	119
	References	
5.7	Total, Resistant and Digestible Starch in Processed Acha and Wheat Grain.	121
5.8	Estimated Glycemic Index (EGI) of Processed Acha, Wheat Grain and Bread.	125
5.9	Conclusions and Recommendations	129
	References	130
6	Summary, Conclusion and Recommendation	134

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LIST OF FIGURES

Figure		Page
2.1	<i>Digitaria iburua</i> (A), <i>Digitaria exilis</i> (B), <i>Digitaria</i> spp grain size	8
2.2	Taxonomic relationship of grasses	9
2.3	Scanning electron micrograph of fonio (<i>Digitaria exilis</i> Staph) starch (a). Koulli variety (1310 x)(b).	13
2.4	Amylose and amylopectin structure	14
2.5	Categories of starch on the basis of enzyme digestion	20
3.1	Experimental outlines for first research chapter	35
3.2	Extraction of starch from Acha grain	37
4.1	Production of acha bread.	73
4.2	Specific loaf volume of acha bread	77
4.3	Mean sensory scores for acha bread	79
4.4	Breads of different grains	80
4.5	Cluster count per age group	84
5.1	Experimental outline for the effect of processing on acha	92
5.2	Effect of incubation time on starch hydrolysis from acha and wheat bread.	98
5.3	Effect of incubation time on starch hydrolysis from microwaved acha and wheat.	100
5.4	Effect of incubation time on starch hydrolysis from boiled acha and wheat	102
5.5	Effect of incubation time on starch hydrolysis from steamed acha and wheat.	104
5.6	Predicted hydrolysed starch from black acha, white acha and wheat bread	107
5.7	Predicted hydrolysed starch from black acha, white acha and wheat after boiling for (A) 15 min; (B) 25 min; (C) 35 min	110
5.8	Predicted hydrolysed starch from black acha, white acha and wheat after steaming for (A) 15 min; (B) 25 min; (C) 35 min	113
5.9	Predicted hydrolysed starch from black acha, white acha and	116

- 5.10 Rapidly digestible starch (RDS), and slowly digestible starch (SDS) of milled and wheat bread 122
- 5.11 Rapidly digestible starch (RDS), and slowly digestible starch (SDS), of microwaved acha and wheat grain. 122
- 5.12 Rapidly digestible starch (RDS), and slowly digestible starch (SDS), of boiled acha and wheat grain. 123
- 5.13 Rapidly digestible starch (RDS), and slowly digestible starch (SDS), of steamed acha and wheat grain. 123
- 5.14 Estimated glycemic Index (EGI) of black acha bread, white acha bread and wheat bread. 126
- 5.15 Estimated glycemic Index (EGI) of acha and wheat grain boiled for 15,25 and 35 minutes 128
- 5.16 Estimated glycemic Index (EGI) of acha and wheat grain steamed for 15,25 and 35 minutes 128
- 5.17 Estimated glycemic Index (EGI) of acha and wheat grain microwaved for 3,6 and 9 minutes 128

LIST OF TABLES

Table	Page	
2.1	Nutritional composition of acha compared to other cereals	11
2.2	Essential amino acid content of fonio compared to maize	11
2.3	Factors affecting the glycemic index of foods	22
3.1	Proximate and amylose composition of acha starch	45
3.2	The percentage of starch hydrolyzed within 0 -180 min of starch from acha and wheat grain.	48
3.3	Analysis of variance (ANOVA) for the effect of time and type of starch grain on hydrolysed starch.	48
3.4	Textural characteristics of acha starch gel	50
3.5	Pasting properties of Acha of Starch	52
3.6	Turbidity and WBC of acha starch ¹	56
3.7	Gelatinization of acha starch	58
3.8	Retrogradation of cereal starch	61
4.1	Ingredients and levels used for acha bread production.	72
4.2	Fractional (2^{4-1}) factorial design with resolution IV for acha bread production.	72
4.3	Analysis of variance for black acha specific loaf volume	72
4.4	Analysis of variance for white acha specific loaf volume	72
4.5	Demography of respondents (N = 30)	78
4.6	Pattern/structure for coefficients obtained using varimax rotation of three factor solution for sensory attributes of acha bread	83
4.7	Gender * Cluster Number of Case Cross tabulation	85
4.8	Race * Cluster Number of Case Cross tabulation	85
4.9	Acha varieties * Cluster Number of Case Cross tabulation	89
5.1	Analysis of variance for the effect of processing, processing time, incubation time and sample on starch hydrolysis of acha and wheat.	97
5.2	Tests of between-subjects effects of baking on starch hydrolysis from acha and wheat bread	98
5.3	Analysis of variance for effect of microwave on starch hydrolysis	101

5.4	Analysis of variance for the effect of boiling on starch hydrolysis	103
5.5	Analysis of variance for the effect of steaming on starch hydrolysed from acha and wheat	105
5.6	Model parameters for bread from black acha, white acha and wheat grain ^{1, 2}	106
5.7	Model parameters for black acha, white acha and wheat grain as affected by boiling ^{1, 2}	109
5.8	Model parameters for black acha, white acha and wheat grain as affected by steaming ^{1, 2}	112
5.9	Model parameters for black acha, white acha and wheat grain as affected by microwave	115
5.10	Total starch, resistant starch and digestible starch in bread samples.	118
5.11	Total starch, resistance starch and digestible starch of processed grain samples	120

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Dedication

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CHAPTER 1

MOTIVATION AND DESIGN OF THE STUDY

1.1 Introduction

Digitaria spp. belongs to the family Graminae and the tribe Poaceae (Dalziel, 1937; Jideani, 1999) and includes 230 species that are widely distributed in the tropics and the subtropics (Adoukonou-Sagbadja *et al.*, 2006). Two types exists namely *Digitaria exilis* (acha, fonio or hungry rice) and *Digitaria iburua* (iburu, black fonio or petit mil) [Jideani, 1999; Philip and Itodo, 2006]. In this study the name Acha will be used to indicate the *Digitaria sp.* Acha is by large the oldest African cereal, it has been cultivated for thousands of years in West Africa (Mali, Burkina Faso, Guinea and Nigeria) and Dominican Republic (Morales-Payán *et al.*, 2003; Jideani *et al.*, 2008) where it is used as a staple food (National Research Council, 1996). *Digitaria exilis* grows under varying conditions from poor dry upland soils to hydro-morphic valleys. Acha was once classified as one of the “lost crops of Africa”, but is now “rediscovered” and considered for improvement as cultivated species (Ibrahim, 2001; Morales-Payán *et al.*, 2003). It is valued because of its unique taste and nutritional value (Philip and Itodo, 2006).

Acha grains contain about 7-9% crude protein with high leucine (9.8%), methionine (5.6%) and valine (5.8%). It has been reported that its methionine content is twice as high as those of egg proteins (Temple and Bassa, 1991; National Research Council, 1996; Ghana Grains Development Project, 1993). Hence, it is the most nutritious grain, containing high essential amino acids that are deficient in today's major cereals such as maize, sorghum, barley, wheat and rye. Consequently, acha has important potential as a complement for standard diets (National Research Council, 1996).

Food products where acha grains have been used include porridge, couscous and it is also mixed with other flours to make bread, pastries and is even oven popped (Jideani, 1999; Philip and Itodo, 2006) as well as traditional beverages such as beer (Philip and Itodo, 2006). Jideani *et al.* (2007, 2008) demonstrated the possibility of producing gluten free loaf from acha flour. In Togo acha is a staple food crop that is cultivated for home consumption and

used during religious and cultural festivals (Adoukonou–Sagbadja *et al.*, 2006). Acha grains prepared similar to a bean type meal, were reported as the most popular acha-based diet in Togo- mostly prepared at special occasions for chiefs, dignitaries and guests. The more common acha meals were acha couscous, rice, porridge and paste. In some regions acha is used solely or mixed with sorghum, pearl millet and maize to prepare local beer named Tchoukouto. Togolese farmers also consider acha as having medicinal value in being useful for those suffering from diabetes among others (Adoukonou- Sagbadja *et al.*, 2006). Similar medicinal use has also been associated with acha in Nigeria (Jideani, 1999).

Starch is the main storage carbohydrate of plants. It is deposited as insoluble, semi-crystalline granules in storage tissues (grains, tubers, roots) and to lesser extent in most vegetative tissues of plants (Copeland *et al.*, 2009). Energy (50-70%) in the human diet is obtained from starch, providing a direct source of glucose- an essential substrate in brain and red blood cells for generating metabolic energy (Copeland *et al.*, 2009). Starch is a food ingredient with a wide applications- thickener, gelling agent, bulking agent and water retention agent (Pomeranz, 1991; Singh *et al.*, 2003), making them an excellent ingredient for the manufacture of foods such as custards, porridges, puddings, cookies and sausages. There is a growing demand for starches for the modern food industry creating interest for new sources of this polysaccharide. Perry *et al.* (2003) reported that the availability of a reliable source of starch from agriculture is considered to have an important factor in human development. This is important especially as the glycemic response to excessive consumption of rapidly digesting starch may be a factor in some diet-related illnesses.

Starchy foods like acha usually are not eaten raw and must undergo one type of heat processing or the other for palatability and bioavailability. Different processing methods include boiling, microwave, baking, drying, extrusion, steaming, drum-drying, popping, pressure-cooking and others. Therefore, processing methods may modify starch in various ways consequently affecting digestion and nutritional value. It is thus of interest to investigate the physicochemical, functional and thermal properties of acha grain as well as the effect of processing on the acha starch.

1.2 Statement of the Research Problem

The properties of acha have received some attention. De Lumen *et al.* (1993) reported acha as a promising underutilised African cereal because of its high content of sulphur-amino acids. Jideani *et al.* (1994) studied the proteins of acha. Lasekan (1994) studied the effect of germination on alpha-amylase activities of acha. The carbohydrates of acha have been investigated (Jideani *et al.*, 1999). Jideani (2007, 2008) studied the potential of acha flour in non-gluten bread. However, the effect of processing on the starch component and glycemic properties has not received attention from scientists.

1.3 Broad Objectives

The aim of the research was to determine the effect of different processing methods on the starch and glycemic properties of acha with a view to establishing relationship between form and functional properties of starch constituent of acha.

1.3.1 Specific objectives

The objectives of the research were to:

1. Establish some functional and thermal properties of acha starch.
2. Establish the physicochemical properties of acha starch.
3. Evaluate the changes in glycemic properties of acha due to different processing methods.

1.4 Research Hypothesis

The hypotheses in this study comprised of

1. The starches from the two acha cultivars (*D. exilis* and *D. iburu*) differ in their functional and thermal properties.
2. Different processing methods differ in their effect on the starch and glycemic properties of acha.
3. The effect of processing on the starch and glycemic properties of acha differs among the cultivars.

1.5 Research Delineation

Two cultivars (*D. exilis* and *D. iburua*) were used in the study.

1.6 Significance of the Research

This study on the effects of different processing techniques on the starch and glyceemic properties of acha will benefit the following sectors; research and development, patients suffering from diet dependent illnesses, health food manufacturers, fast moving consumer goods manufacturers, industrial food manufacturers and the economy.

1.6.1 *Research and development*

1. It will put focus on acha as a crop that can be utilized for other uses apart from being a staple crop i.e. low GI snack foods especially in this era of global obesity, or as a partial substitute for wheat flour based products i.e. bread
2. Research and development departments of manufacturing companies can assist in optimizing procurement by making use of locally procured acha grain instead of opting for imported foreign grains.

1.6.2 *Health Foods*

It will draw attention to the benefits of acha as a dietary supplement for diabetic patients, the development of products aimed at weaning of toddlers, the old and frail as well as immune-compromised individuals. It will aid in controlling and preventing many metabolic diseases such as cardiovascular diseases and patients suffering from hyperlipidaemia. It will be beneficial for use by patients with celiac disease as well as for promoting and maintaining large bowel health.

1.6.3 *Manufacturing*

It will create a spin off in the technological development specifically with the aim of improving mechanised processing technique for acha grain.

1.6.4 *Economy*

It will draw closer attention to this indigenous underutilized crop and its

usefulness, especially in this time of increasing food prices and world-wide food shortages.

1.7 Expected outcomes, results and contributions of the research

1. Identification of the processing techniques that slows down the rate of acha starch digestion.
2. New research data on the effect of processing on acha starch properties.
3. New starch source made available.
4. Novel use of this indigenous African cereal as a starch based material.
5. Non wheat bread from acha with nutritional and health benefits.
6. At least one scientific article will be published from the work and the information will be presented at a scientific conference.

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CHAPTER 2

LITERATURE REVIEW

2.1 Overview of *Digitaria* spp.

Digitaria exilis (white) and *Digitaria iburua* (black) are two species of acha grains (Figure 2.1). Other common names are fonio or hungry rice. It is indigenous to Africa particularly regions of Mali, Burkina Faso, Guinea and Nigeria (Ayo, 2003) and the Dominican Republic (Morales-Payán *et al.*, 2003; Jideani *et al.*, 2008). Acha falls under the grass family Gramineae (Poaceae) and belongs to the same sub-family as maize, sorghum, pearl millet and barley. This is illustrated in Figure 2.2.

Acha has a unique small size of 0.4 to 0.5 mm (Adoukonou- Sagbadja *et al.*, 2006). Gomez & Gupta (1993) reported acha grains to be very small with an individual grain weight of 0.5 to 0.6 mg. Ayo (2004) recorded this tiny grain weight to be about 0.5 mg with a diameter of between 500 to 700 nm. Acha's kernel is much smaller than the average pearl millet, sorghum and maize kernel (Phillip & Itodo, 2006). Irving & Jideani (1997) reported that acha can be used for the same technological processes as cereals similar to its anatomical structure. It is thus of interest for this research to determine the exact physical characteristics of these two *Digitaria* spp.

2.2 Uses of Acha

Acha is essentially cultivated as a staple food crop for most ethnic groups of West Africa (Phillip & Itodo 2006). Acha can be used in a variety of meals like breakfast and supper. It can be made into a porridge and couscous, or it can be ground and mixed with other flours to make bread, pastries, oven popped and even brewed into a beer.

In Togo acha is known for its medicinal values hence it is used to treat blood clots, chronic diarrhoea, loss of appetite, stomach aches and as a useful diet for individuals suffering from diabetes (Adoukouno-Saghadja *et al.*, 2006). Although classified as a lost crop of Africa it is one of the world's best palatable cereal. Assessing the effects of processing on acha starch properties and glycemic index can lead to increased focus on the potential of



A



B



C

Figure 2.1 *Digitaria iburua* (A), *Digitaria exilis* (B), *Digitaria iburua* grain size (C).

Family

Poaceae

Tribe

Oryzeae

Eragrostideae

Paniceae

Andropogoneae

Aveneae

Triticeae

Species

Eleusine coracana
(Finger millet)

E. tef
(tef)

Digitaria spp.
(Acha & iburu)

Zea mays
(maize)

Sorghum bicolor
(Sorghum)

Avena sativa
(Oats)

T. aestivum
(Wheat)

H. vulgare
(Barley)

Oryza sativum
(Rice)

Figure 2.2 Taxonomic relationship of grasses (Clayton and Renvoize, 1986)

this crop in terms of developing food products that will be beneficial to those suffering from diabetes and related illnesses.

2.3 Nutritional Composition of Acha

Phillip & Itodo (2006) reported that acha contains about 8-11% proteins and its digestibility is better than that of sorghum or millet. Moisture content of acha grain is 7.2% compared to millet of 6.5% (Ayo, 2004). This moisture content is lower than that reported by Chukwu (2008). The total lipid content (Table 2.1) for acha is higher than that of brown rice, wheat and barley but lower than that for maize and millet (CIRAD, 2004). The protein content of hulled acha has been reported to be lower than that of brown rice, wheat and barley, slightly higher than that of maize and millet (CIRAD, 2004).

With a high essential amino acid, leucine 9.7%, methionine 5.6% and valine 5.8% acha is considered as one of the world's most nutritious grain (Phillip & Itodo, 2006). Essential amino acids content of acha compared to Obatanpa (quality protein maize) and Okomasa (full season maize) as depicted in Table 2.2, indicates that acha contain significantly higher methionine, valine and leucine values than the all season maize and the high quality protein maize (National Research Council, 1996 & Ghana Grains Development Project, 1993).

According to Ayo (2003), the high water absorption capacity of acha is a characteristic that is linked to high amount of pentosans; acha contains 33 g/kg pentosans. The high amount of pentosans in acha will be beneficial during baking due the consequent increase in water absorption capacity during the baking process. Ayo & Nkama (2004) reported an increase in bread moisture content with an increase in acha flour to the recipe.

Acha residue protein has been shown to contain high levels of glycine, glutamate/glutamine and leucine (Jideani, 1999). The structural role of glycine in cereal proteins depends on the amino acid sequence of the glycine-containing peptides, which is responsible for the elastic properties of gluten necessary in bread making and which only occurs in wheat gluten (Jideani, 1999).

Table 2.1 Nutritional composition of acha compared with other cereals

Nutrient	Cereal					
	Hulled acha	Brown rice	Sorghum	Millet	Maize	Wheat
Sugar (%)	85.0	86	82.5	80.0	82.0	82.0
Protein (%)	10.0	9.0	12.0	13.0	11.0	14.0
Fat (%)	4.0	2.5	4.0	5.0	5.5.0	2.0
Ash (%)	1.0	1.5	1.5	2.0	1.5	2.0

Source: CIRAD (2004)

Table 2.2 Essential amino acid content of acha compared to Obatanpa and Okomasa maize on percentage dry matter¹

Amino Acid	Fonio	Okomasa (Full season maize)	Obatanpa (Quality protein maize)
Threonine	0.37	0.24	0.34
Cystine	0.25	0.19	0.26
Valine	0.55	0.33	0.48
Methionine	0.45	0.17	0.15
Isoleucine	0.4	0.23	0.30
Leucine	1.05	0.77	0.88
Phenylalanine	0.57	0.31	0.39
Lysine	0.25	0.23	0.36
Tryptophan	1.60	0.06	0.10

Source: NRC (1996), GHANA GRAINS DEVELOPMENT PROJECT (1993)

¹ Amino acids expressed as % of protein

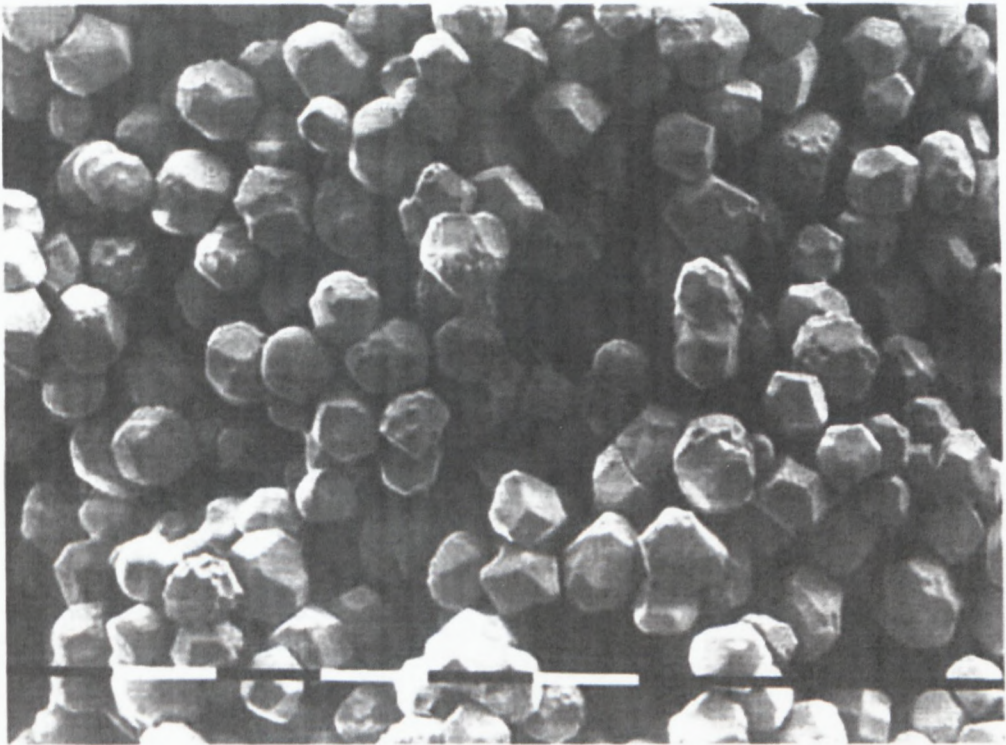
2.4 Physical Properties of Acha Starch

Starch is laid down in the tissue of many plants and according to Kearnsley & According to Dsiedzia (1995) the commercial reality of starch recovery is limited to mainly wheat, maize, potato, tapioca and rice. Starch is found in the granules and its shape and size is relative to its source. The size of starch granules varies from 2-150 micron in size; from round to polygonal and truncated in shape. Rice starch is the smallest of the cereal starches, and also generally the most uniform in size.

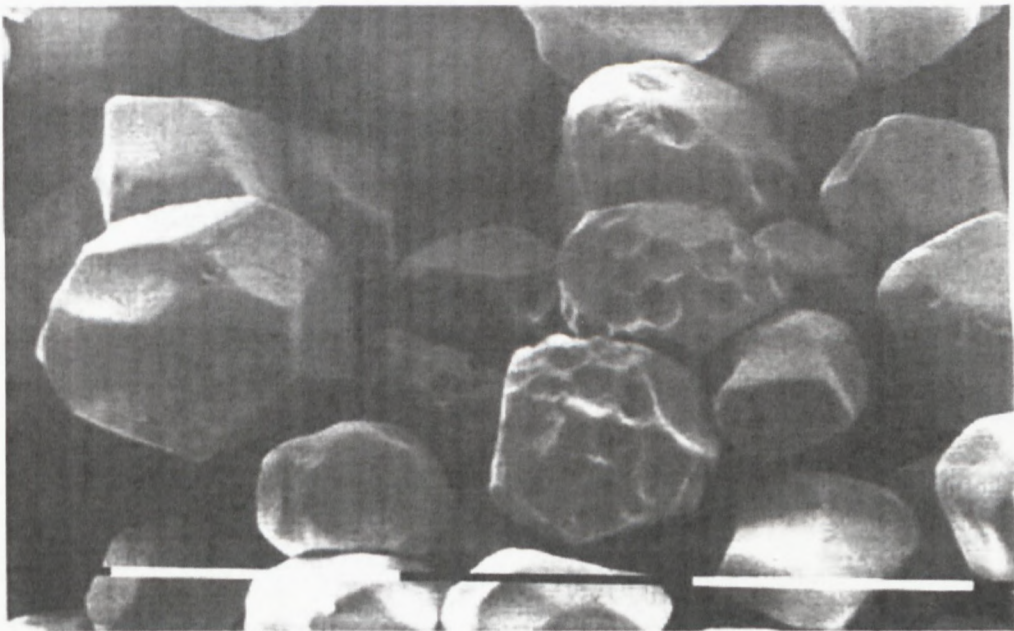
Jideani (1999) reported that white and black acha have not been used to produce starch for industrial applications, but that the starch from these two grains are similar in structure and physiochemical properties to starch from conventional cereal grains. Electron photomicrographs of dry acha starch granules from two varieties (Hothia and Koulli) of *Digitaria exilis* grown in Ethiopia, at different magnifications, are presented in Figures 2.3 (Carcea & Acquistuicci, 1997). Microscopic examination conducted by Carcea & Acquistuicci (1997) revealed that starch granules from these two varieties are morphologically very similar, being in both cases, small in size and mostly polygonal. In both varieties the average diameter of the granules was 8 μm but Koulli appeared to be more uniform. Carcea & Acquistuicci (1997) also observed that acha starch has a morphology (small size, angular, irregular shape) which is common to other millet starches. Starch granule dimension is a characteristic peculiar to each plant species. Due to the potential of small-granule starches to be used in many ways such as fat substitutes, as stabilizers in baking powder, as laundry stiffening as well as in the manufacture of degradable plastic films, it has received some investigations by Daniel & Whistler (1990) as well as by Muir & O'Dea (1992). Polygonal starch granule contains deep indentations on their surfaces (Figure 2.3). These indentations could be due to the dense packing of the endosperm and especially the dense packing of the adjacent protein bodies on the developing starch granules (Carcea & Acquistuicci, 1997).

2.5 Chemical Properties of Cereal Starch

Kearnsley & Dsiedzia (1995) reported that starch granules consist of two glucose polymer structures, namely amylose and amylopectin (Figure 2.4), as



(a)



(b)

Figure 2.3 Scanning electron micrograph of fonio (*Digitaria exilis* Staph) starch. (a) Koulli variety (1310x); (b) Hothia variety (2100x). (Carcea & Acquistucci, 1997)

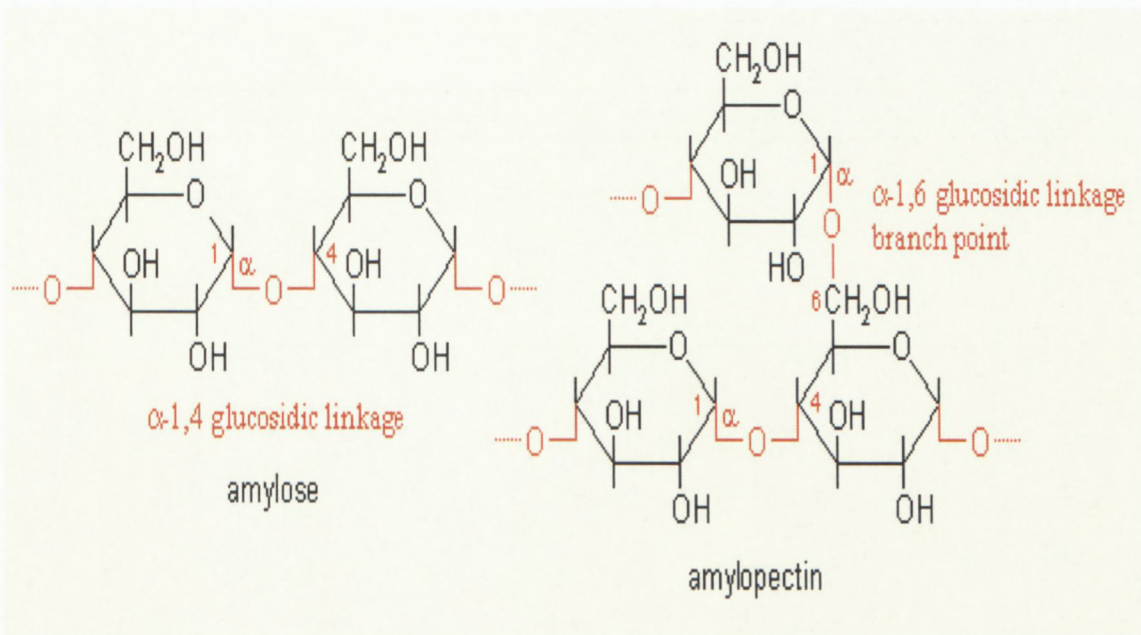


Figure 2.4 Amylose and Amylopectin structure (Ophardt, 2003)

well as moisture, lipids, proteins and mineral ions. The backbone of this molecule is the α 1-4 linked anhydro-glucose unit but with branching points occurring at the carbon 6 primary alcohol groups, every 10 to 12 glucose units. The side chain of the main backbone is 20 to 30 glucose units in length. Amylose has traditionally been considered to be a linear polymer composed of glucopyranose units linked by α -D-(1-4) glycosidic linkages, while amylopectin is a branched polymer (Karim *et al.*, 2000).

The molecular structures and physicochemical properties of starches from waxy barley and two low-amylose cultivars of barley were examined by Yoshimoto *et al.* (2002) and compared with those of a normal barley cultivar. The waxy barley starch had no detectable amylose. The amylose content was 22% and 11.4% in the two low-amylose barley starches, respectively, being much lower than that in the normal barley cultivar starch 25.4%. The two low-amylose barley starches contained the branched amylose with a lower number of chains (6.1 to 10.4) than normal barley starch cultivar (Yoshimoto *et al.*, 2002).

Acha starches were reported to be of the non-waxy type, and there were significant differences in amylose between the two acha varieties Hothio (22.6%) and Koulli (26.1%) (Carcea & Acquistucci, 1997). The amylose in the Koulli starch was higher than that of Hothio and wheat (25%) starch, but was much higher than that of pearl millet starch reported by Badi *et al.* (1976). It is unknown at this stage if the same differences exist between *Digitaria exilis* and *Digitaria iburua*. Cereal starches normally have a moisture content of 13 to 14% while other starches have a moisture content of 18 to 20%. The moisture content of Hothio variety has been reported to be higher than that of Koulli and wheat starch (Carcea & Acquistucci, 1997).

2.6 Functional Properties of Acha Starch

Starch is an important ingredient for the food industry and is extensively used as a binding agent, thickener, gelling agent, bulking agent and water retaining agent (Sandhu & Singh, 2007). A brief review of the functionality of starch in general and acha will follow.

Starch swelling characteristics

Starch granules swell when heated in excess water and their volume fraction and morphology plays an important role in the rheological behaviour of starch dispersions (Sandhu & Singh, 2007). Pylar (1988) reported that when an aqueous starch suspension is heated, the starch granules imbibe water at 50% of its weight, without showing any change in its appearance. At a critical temperature of 60°C the weaker bonds are dissociated and the granules begin to swell progressively and start to lose some of the higher opacity and birefringence. With continued increase in temperature to the gelatinization range, there is a more extensive disruption of the hydrogen bonds and continues swelling of the granules. According to Pylar (1988) this process continues until the swollen granules have taken up all the surrounding free water, to form a starch paste or gel of various clarity and viscosity.

Various starches consist of a birefringence pattern; the centre of this pattern can either be eccentric or concentric and will correspond to the hilum of the starch granule. The different birefringence patterns of different starches indicate different degrees of crystallinity in individual's starches (Kearnsley & Dsiedzig, 1995). The intra- and inter- molecular hydrogen bonding, responsible for the structural integrity of the starch granule, accounts for its insolubility in water. When a starch suspension is heated in water, heat overcomes the hydrogen bonding forces which holds the granules together, allowing swelling or gelatinization of the starch granule to occur. Acid and enzymes have little hydrolyzing effect on the un-gelatinized starch (Kearnsley & Dsiedzig, 1995). Thermal gelatinization of starch in the presence of alkali, using different temperature regimes, is therefore, likely to result in wide variations in the properties of gelatinized starch such as colour, paste viscosity, gel properties and retrogradation tendency (Karim *et al.*, 2000).

Gonza'lez-Reyes *et al.* (2002) reported the swelling pattern of *Okenia* (*Okenia hypogaea*) and corn starches with 5% (w/v) total solids. *Okenia* starch presented a slight increase in swelling when heated from 30 to 60°C. However, the swelling of corn starch was constant in a similar temperature range. When increasing the temperature, a gradual increase in swelling of both starches was observed with increase in temperature up to 95°C. This

was due to the breaking of the molecular bonds when increasing the temperature of the starch granules and the subsequent incorporation of water molecules within their structure (Gonzales & Perez, 2002).

Cereal starches exhibit a two stage swelling and solubility pattern that is indicative of two sets of internal bonding forces. Whereas waxy starches swells more readily than do the normal starches, presumably because of the lack of reinforcement of the internal network provided by the linear fractions. Potato starch expands at a markedly lower temperature of 75°C and expands more rapidly and to a greater extent, so that the outline of the granules become vague. The swelling power of acha and wheat starch over a temperature range of 50-90°C was investigated by Carcea & Acquistuicci (1997). Swelling in the early stages indicated disintegration of the granular structure and significant difference ($p \leq 0.05$) were observed in the swelling power. Acha starch swelled from 60°C and up to 65°C and had lower swelling power than wheat starch. This is indicative of the degree of bonding forces within the acha starch granules.

Acha solubility characteristics

Although a significant difference ($p \leq 0.05$) was observed between fonio Hothio and fonio Koulli during exposure to the entire temperature range, the two fonio varieties behaved similarly for both swelling and solubility power compared to wheat starch. These stronger bonding forces within the fonio granules can be observed in the solubility of the two fonio types compared to wheat starch. Both fonio types showed lower solubility values than wheat for temperature of 50°C - 65°C, thus less solute were released from fonio than wheat starch.

In the case of solubility for Okenia and corn starch at 5% solids, over a range of 30°C - 60°C, the solubility increased slightly, increasing significantly from 70°C (Gonzales *et al.*, 2003). At this temperature, the granular structure is lost and solubilisation of the constituent polymers was promoted.

Starch paste viscosity characteristics

Holm *et al.* (1988), reported that hot paste viscosity at 95°C and final cooked

paste viscosity (after cooling to 25°C or 50°C) of extruded flour and starches are closely related in terms of their molecular size distribution. Carcea & Acquistucci (1997) investigated the pasting properties of 5% wheat and fonio starch by means of a viscoamylograph and observed that wheat starch showed the typical two step pasting curve whereas both fonio varieties showed a single step pasting curve. These studies indicated that in both fonio slurries, the onset of gelatinization, indicated by an increase in viscosity is apparent for fonio at higher temperatures (> 50°C) than for wheat starch. The Hothia variety showed the highest peak viscosity followed by wheat starch. Koulli starch gave the lowest viscosity values and this can be attributed to the higher power of Koulli starch at temperatures higher than 70°C. As reported by Carcea & Acquistucci (1997), there might be negative correlation between amylose content and peak viscosity. Hothia gave the lowest amylose content (22.6%) and showed the highest peak viscosity, whereas Koulli gave the highest amylose (26.1%) and the lowest peak viscosity.

Starch textural characteristics

Understanding gel textural properties of starch is imperative for selecting the appropriate starch for end use suitability. Textural properties of corn starch varieties differ significantly from each other (Sandhu & Singh, 2007). The gel firmness of corn starch is mainly caused by retrogradation of starch which in turn is associated to syneresis of water and crystallization of amylopectin, leading to harder gels. Starches that exhibit harder gels tend to have higher amylose content and longer amylopectin chains. The mechanical properties of starch gels depends on various factors including the rheological characteristics of the amylose matrix, the volume fraction and the rigidity of the gelatinized starch granules as well as the interaction between the dispersed and continuous phases of the gel. These factors in turn depend on the amylose content and the structure of the amylopectin (Sandhu & Singh, 2007).

2.7 Overview of Starch in relation to Digestion

Starches contribute as much as 70-80% of the total carbohydrate in normal

diet and can thus be classified on the basis of the rate at which digestive enzymes hydrolyse starch into three categories (Figure 2.5) namely, rapidly digestible starch (RDS), slowly digestible starch, (SDS) and resistant starch (RS) [Chung *et al.*, 2008a]. RDS and SDS is measure of the glucose released after 20 and 100 min of incubation, respectively. RS is the starch not hydrolysed after 120 min incubation (Englyst *et al.*, 1992). However, RS is commonly defined as the sum of starch and products of starch degradation not absorbed in the small intestine of healthy humans. RS escapes digestion and is fermented in the colon (Asp, 1992). Resistant starch is starch that escapes digestion in the small intestine of healthy individuals and passes through to the large intestine where it acts as dietary fibre. There are four types of resistant starch (Figure 2.5). RS1 is physically inaccessible or digestible e.g. unprocessed whole grain and those found in legumes. RS2 occur in its natural form such as in uncooked potato and green banana flour. RS3 are formed when starch containing foods are cooked and cooled as in bread, cornflakes and cooked and chilled potatoes. RS4 are formed in starches that have been chemically modified to resist digestion (Asp *et al.*, 1996). Different processing techniques and type of raw materials thus affects the resistant starch content and type within any given food stuff (Hyun-Jung *et al.*, 2008). Chewing can also affect the amount of starch escaping digestion, that is, decrease in RS is relative with increase in chewing (Muir & O'Dea, 1992).

Resistant starch has major health benefits in areas such as individuals colonic health, their energy and glycemic management. Research on dietary fibre has demonstrated the importance of fermentation of indigestible carbohydrates in the human colon. It provides a substrate for microbial fermentation in the large intestine, and therefore as a dietary fibre is beneficial in areas such as diabetes, cardiovascular disease and bowel cancer (Buttriss, 2002). Jideani, (1999) reported that acha is usually consumed as whole grain. It is known that whole grain consumption reduces the risk of certain cancers, stroke, diabetes and cardiovascular disease. Jideani *et al.* (2008) developed non wheat acha bread where a deliberate attempt was made to remove the sugar from this product and as such is believed to serve as a good alternative to wheat bread for diabetic patients and those allergic to

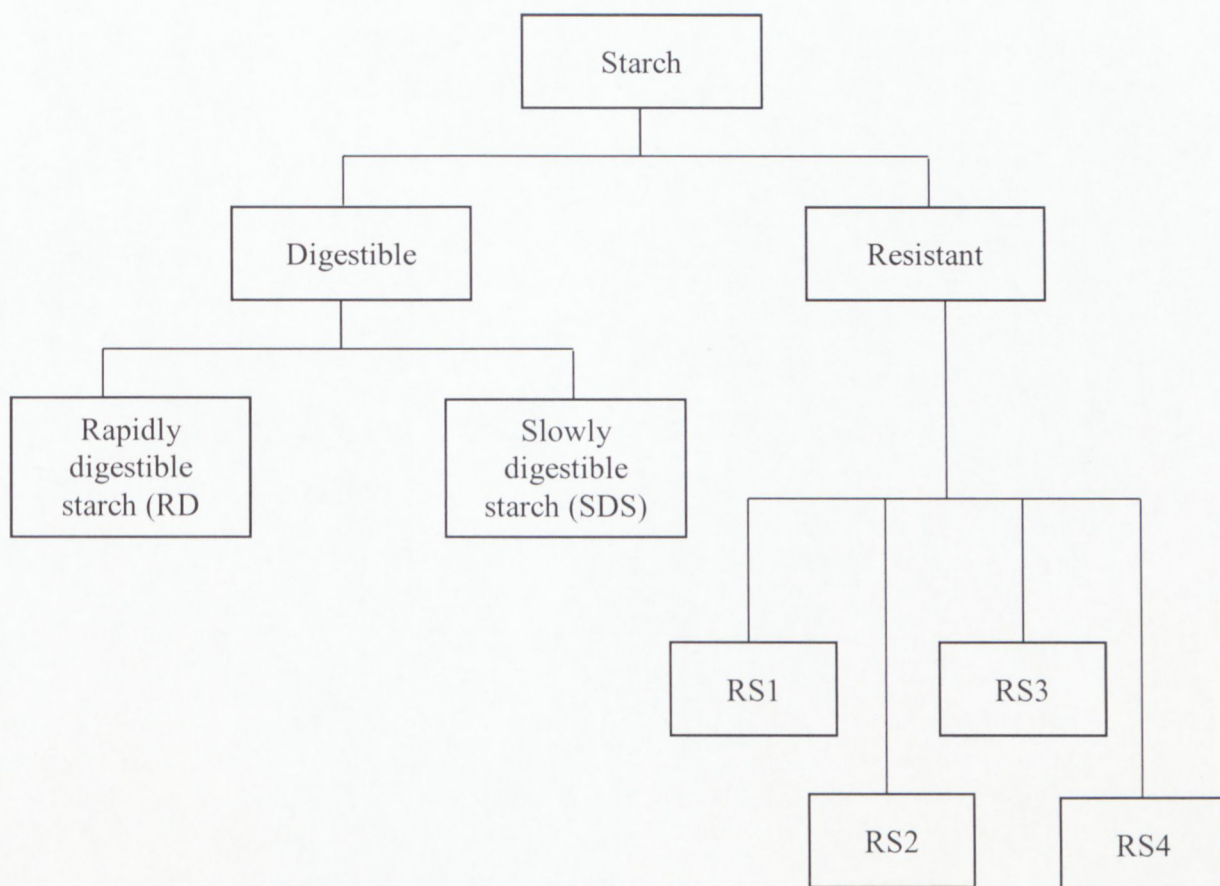


Figure 2.5 Categories of starch on the basis of enzyme digestion

gluten. Acha has been identified as a major food for diabetic patients in Nigeria (Jideani, 1999).

2.8 Glycemic Properties of Starch

The glycemic index is the rate of increase in blood sugar level relative to the glucose taken as 100. The rate of increase in blood sugar level is the area under the curve for two hours following ingestion of 50 g of available carbohydrates (Bird *et al.*, 2000). Glycemic index (GI) of food is the degree to which the food specifically carbohydrates increases blood glucose in proportion to glucose itself. GI is used as a measure of the effects of carbohydrates on blood sugar levels. High GI indicates that the carbohydrate will be broken down rapidly releasing glucose rapidly into the blood stream. Low GI on the other hand is indicative of a slow break down of the carbohydrates thus releasing glucose slowly and gradually into the bloodstream (Buttriss, 2002).

The GI plays a major role in blood sugar control of diabetics since a lower GI is relative to a lower insulin demand thus a better long term blood glucose control and a reduction in blood lipids (Buttriss, 2002). Following a low GI diet thus reduces the risk of individual developing type-2 diabetes. However, a high GI food which releases glucose rapidly in the bloodstream is suitable for athletes who need quick energy release during a fast event where speed and spurge of energy is required as well as diabetic individuals undergoing hypoglycaemia (Buttriss, 2002). On the other hand, endurance athletes also consume low GI food before and during an endurance event to ensure a slow release of energy during long distance running (Buttriss, 2002).

Pawlak *et al.* (2002) reported a 71% increase in body fat in male rats fed with high GI diet compared to those fed with a low GI diet. GI values can be interpreted as percentages and are commonly categorised as: Low GI 55% or less, Medium GI 56-79%, high GI 70-99% and with straight glucose as 100% (Pawlak *et al.*, 2002). The GI of foods depends on a number of variables e.g. the type of starch i.e. amylose vs. amylopectin, the chemical and physical nature of the food as well as the combination of the meal e.g. protein and fat content (Buttriss, 2002). The presence of protein and fat in food may lower the GI of the food; this however is not advisable due to risk

Table 2.3 Factors affecting glycemic index of foods

Factor	Comment
Particle size	Intact grains such as whole wheat, barley, whole corn and whole rye have much lower GI values than flours (tiny particles) made from grains.
Processing	Milling, beating, grinding, mixing, mashing and refining foods raise the GI index of that foods
Protein and Fat	The presence of protein and fat in food may lower the GI.
Cooking	Cooking usually decreases the digestibility of the food and would have an effect of raising the GI of that food.
Resistant starch	When starches are cooked and then cooled, the crystalline structure within the food changes to resistant starch which is more difficult to digest. Therefore cold cooked starch has a lower GI. This is especially true for mealy meal. The cooked cold maize has a lower GI than the hot freshly prepared porridge.

Source: (Steenkamp, 2008)

associated with high fat content meals (Steenkamp, 2008). Other factors affecting the glycemic properties of foods are outlined in Table 2.3 and include particle size, processing, protein and fat, cooking, resistant starch (Steenkamp, 2008).

The GI and resistant starch content are two important indicators of starch digestibility (Hu *et al.*, 2004). RS and SDS are highly desirable forms of dietary starch as they are known to have low glycemic index (GI) (Eyaru *et al.*, 2008). The amount of resistant starch and digestibility rate of acha starch is not known. As stated by Hu *et al.* (2004) resistant starch is slowly digestible and may be used as a means of slow release of glucose.

2.9 Measurement Techniques for Resistant Starch

Englyst *et al.* (1992) devised a scheme for the classification and measurement of nutritionally important starch fractions including the three main forms of RS. The method is based upon analysis of starch fractions in foods as eaten, which is based on the recovery of even the physically enclosed RS1. There are however various *in vitro* assay systems and *in vivo* methods to measure the amount of resistant starch escaping digestion in the small intestine.

An *in vitro* assay system was developed that mimics the physiological conditions for starch digestion. This assay is useful in predicting which foods and processing techniques results in high amounts of starch escaping digestion in the small intestine (Muir & O 'Dea, 1992). Another *in vitro* method that includes the measurement and classification of starch into rapidly available glucose (RAG), slowly available glucose (SAG) and starch fraction, involves the measurement by means of HPLC of the glucose released from a test food during incubation with digestive enzymes under controlled conditions (Englyst,1999). In this work, extracted acha starch was analyzed for rapidly digestible starch (RDS), slowly digestible starch (SDS), resistant starch (RS) utilizing the modified Goñi *et al.* (1997) methods.

2.10 Effect of Processing on Functional and Glycemic Properties of Starch

Starch can be modified by various chemical and physical processing techniques (Gonzales & Perez, 2002). For this reason different processing

techniques are currently in use for the production of pre-cooked, modified and instant starches, which has led to an increase in ready-to-eat cereals, ready-to-eat meals, infant formulae and snack foods (Gonzales & Perez, 2002). Examples of different processing techniques are baking, soaking, microwaving and cooking.

Effect of milling on cereal starch

Various grain or dry bran fractionation processes have been developed to produce safe and healthy whole wheat products (Hemery *et al.*, 2007). Some of the biochemical markers were starch, aleurone cell walls, aleurone content and endosperm. Three different grain or bran dry fractionation processes, namely: conventional milling, de-branning process and the production of aleurone-rich fractions from coarse bran were tested. This led to the development of new processes in order to exploit all the nutritional benefits of whole grain as well as to produce new wheat foods and wheat-based ingredients with improved nutritional quality (Hemery *et al.*, 2007). Various whole grain products with improved nutritional benefits in terms of low GI foods can be found on the market, namely whole grain low GI bread and whole grain low GI porridge/cereal (Eyaru *et al.*, 2008). No investigation could be found where the nutritional benefit of whole acha grain flour has been compared to whole wheat grain flour.

Effect of baking on cereal starch

Collado-Fernandez (2003) reported three stages during the baking process as oven spring; gelatinization of starches (70 to 90°C) and browning and aroma formation above 100°C. During the oven spring stage, yeast activity continues until the dough reached about 50°C and enzymatic activity continues at about 60°C where conversion of starch to sugar continues. Enzymatic activity increases to until about 80°C. The enzymatic activity plays a major role in the quality of the bread with regard to structure, loaf volume and crumb structure (Collado-Fernandez, 2003). Inadequate enzyme activity leads to a small loaf volume due to the starch that becomes too rigid too soon, whereas too much activity on the other hand leads to weak structure, which

collapses completely. Gelatinization of starch is promoted at temperatures of about 50 to 60°C and therefore denaturation of proteins with the coagulation of gluten (Collado-Fernandez, 2003). During baking, dehydration of gluten occurs and water is moved from the gluten to the starch, swelling occurs as the starch gelatinizes. When gluten collapses, the starch not only supports the structure of the dough but also contributes to the firmness of the bread when the starch crystallizes upon cooling of the bread.

Investigating the microstructure of starch in dough, fresh bread and aged bread by means of light microscopy, Hu *et al.* (2004) observed a partial segregation of starch from the protein phase. On baking, the starch gelatinization yielded a continuous starch network which itself was homogenous and consisted of swollen and connected starch granules. The same studies also revealed that two polymers, namely amylose and amylopectin were phased separately and the amylose was accumulated in the centre of the starch granule. The baking quality rendered by various wheat and non-wheat flour is attributable to factors such as starch structure, namely the crystalline organisation of the starch granules as well as the chemical structure of amylopectin and amylose (Cauvain, 2003).

Effect of soaking on cereal starch

Native starch does not absorb water at room temperature and its viscosity is nearly zero, whereas extruded starches absorb water rapidly to form a paste at room temperature. This paste is formed by solubilised macromolecules but also includes particles swollen with water (gel) (Colonna *et al.*, 1981). These two properties are affected by the particle size, with smaller particles showing increased rates and levels of solubilisation. Water activity index correlates well with cold-paste viscosity because only damaged starch granules absorb water at room temperature and swell, creating increased viscosity. Miao *et al.* (2009) reported that swelling waxy maize starch were characterised by an initial phase of slight swelling, a second phase of rapid swelling and a final stage of maximum swelling. A similar trend has been observed for waxy rice starch (Miao *et al.*, 2009). No evidence could be found of investigations into a trend of this nature for acha starch granules.

Soaking red kidney beans in water significantly reduced the RDS, TS, RS and SDS as compared to raw beans. Similarly, both varieties of peas (yellow and green) showed a high degree of loss in RDS. The loss in RDS was accompanied by a gain in SDS. Eyaru *et al.* (2008) further reported that the soaking of seeds in plain water produces swelling of the tissues and water uptake without cell separation. This could have been the cause of increase in solubility of the starch granules during soaking which might have subjected the RS to enzymatic hydrolysis, therefore, yielding more SDS (Eyaru *et al.*, 2008). The effect of soaking on the glycemic properties of acha starch has not been investigated.

Effect of microwaves cooking on cereal starch

Microwaves are non-ionising energy that can generate heat deep inside the penetrated medium by the molecular friction in an alternating electromagnetic field. Microwave irradiation appears to be applicable to starch processing, but has not been used on a commercial scale. It causes several changes in the functional, rheological and morphological characteristics of lentil starches (Gonzales & Perez, 2002). Microwave irradiation also reduced the retrogradation tendency of lentil starch. The effect of microwave irradiation on fonio starch properties have not been an object of research.

Effect of boiling on cereal starch

According to Frame (1994), this is the simplest type of cooking process and one with the least mechanical action and least starch damage. The glycemic responses of boiled products are less than the corresponding extruded products (Mercier & Conterrali, 1989).

Eyaru *et al.* (2008) reported that boiling of previously soaked legumes brought changes in starch fractions that were different to that of unsoaked-boiled legumes. Boiling soaked red kidney beans caused a lesser increase in RDS (232%) compared to the unsoaked beans (339%) but the SDS contents were almost the same for both cooked beans. There was a higher decrease in RS content during boiling of soaked samples which meant that soaking of beans prior to cooking made them more digestible than cooking raw beans (Eyaru *et al.*, 2008).

Processed rice seeds must be boiled or steamed before consumption. Studies conducted by Frei *et al.* (2003) to investigate the *in vitro* starch digestibility and the GI of six different indigenous rice cultivars from Philippines indicated that substantial differences exist in the estimated GI between rice cultivars. Values ranged between 68 and 109 for cooked rice and between 64 and 87 for stored rice containing retrograded starch (Frei *et al.*, 2003). Waxy rice starches exhibit a similar trend as starch granules when starch (100 mg) was soaked in water (5 ml) and heated in a water bath at the required temperature (60 to 80°C) for 30 minutes with constant shaking (Miao *et al.*, 2009). Acha grain is also consumed as rice or couscous (Phillip & Itodo, 2006). The effect on the glycemic properties of processing acha grain into rice or couscous has not been investigated.

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CHAPTER 3

Physicochemical, Functional and Thermal Properties of Acha Starch

Abstract

The thermal, functional and physicochemical properties of two acha cultivars (*Digitaria exilis*, white and *Digitaria iburua*, black) were investigated. Wheat was used as reference. The acha starch was extracted from cleaned and dried acha grains. The thermal, functional and physicochemical properties of both acha starches and wheat starch as reference were investigated. There was no significant difference ($p > 0.05$) between white acha, black acha and wheat starches. Wheat starch had the highest gelatinization temperature 34.75°C and 74.56°C for T_o and T_p . White and black acha recorded 25.05, 26.7°C and 68.20, 70, 54°C for T_o and T_p respectively. The gelatinization temperature range ($T_c - T_o$) for white and black acha were between 62 and 66°C whereas that of wheat starch was 61°C. No significant ($p > 0.05$) difference existed between, white and black acha gels in hardness, resilience and springiness. There were significant differences ($p < 0.05$) in peak viscosity and final viscosity between the black and white acha and wheat starch. There was a significant difference ($p < 0.05$) in the water binding capacity between wheat starch (0.083 g/100g) and both acha cultivars, black acha starch (0.83 g/100g) and white acha starch (1.33 g/100g). However there was no significant ($p > 0.05$) difference between black acha and white acha starch. There was similarity in the nutritional composition and amylose content of white acha starch and black acha starch. There was however a significant difference ($p < 0.05$) in the nutritional composition with regard to the protein, fat and carbohydrate content between black, white acha and wheat starch. Black and white acha starch had a higher fat and protein content than wheat starch, whereas wheat starch had higher carbohydrate content.

3.1 Introduction

Digitaria spp. belongs to the family Graminae and the tribe Poaceae (Dalziel, 1937; Jideani, 1999) and includes 230 species that are widely distributed in

the tropics and the subtropics (Adoukonou-Sagbadja *et al.*, 2004). Two types exist namely *Digitaria exilis* (acha, fonio or hungry rice) and *Digitaria iburua* (iburua, black fonio or petit millet) (Jideani, 1999; Philip & Itodo, 2006). Acha is by large the oldest African cereal, it has been cultivated for hundreds of years in West Africa (Mali, Burkina Faso, Guinea and Nigeria) and the Dominican Republic (Morales-Payán *et al.*, 2003; Jideani *et al.*, 2008). Acha grows under varying conditions from poor dry upland soils to hydro-morphic valleys (National Research Council, 1996). Acha was once classified as one of the “lost crops of Africa”, but is now “rediscovered” and considered for improvement as cultivated species (Ibrahim, 2001; Morales-Payán *et al.*, 2003). It is valued because of its unique taste and nutritional value (Philip & Itodo, 2006).

Acha grains contain about 7 to 9% crude protein with high leucine (9.8%), methionine (5.6%) and valine (5.8%). It has been reported that its methionine content is twice as high as those of egg proteins (Ghana Grains Development Project, 1993; National Research Council, 1996; Temple & Bassa, 1991). Hence, it is the most nutritious grain, containing high levels of essential amino acids that are deficient in today's major cereals such as maize, sorghum, barley, wheat and rye. Consequently, acha has potential as a complement for standard diets (National Research Council, 1996).

Food products where acha grains have been used include porridge, couscous and it is also mixed with other flours to make bread, pastries and is even oven-popped (Jideani, 1999; Philip & Itodo, 2006). It is also used in traditional beverages such as beer (Philip & Itodo, 2006). Jideani *et al.* (2007, 2008) demonstrated the possibility of producing gluten free loaf from acha flour.

Starch is the main storage carbohydrate of plants. It is deposited as insoluble, semi-crystalline granules in storage tissues (grains, tubers, roots) and to a lesser extent in most vegetative tissues of plants (Copeland & Blazek, 2009). Energy (50 to 70%) in the human diet is obtained from starch, providing a direct source of glucose, an essential substrate in brain and red blood cells for generating metabolic energy (Copeland & Blazek, 2009). Starch is a food ingredient with wide applications such as thickener, gelling agent, bulking agent and water retention agent (Pomeranz, 1991; Singh *et al.*,

2003), making it an excellent ingredient for the manufacture of foods such as: custards, porridges, puddings, cookies, and sausages. There is a growing demand for starches for the modern food industry creating interest for new sources of this polysaccharide. Perry & Colman (2003) reported that the availability of a reliable source of starch from agriculture is considered to have an important factor in human development. This is important; especially as the glycemic response to excessive consumption of rapidly digested starch may be a factor in some diet-related illnesses. Togolese farmers also consider acha as having medicinal value in being useful for those suffering from diabetes among others (Adoukonou-Sagbadja *et al.*, 2006). Similar medicinal use has also been associated with acha in Nigeria (Jideani, 1999). Currently there is however a lack of information on the physicochemical, thermal and functional properties of acha starch. The objective of this chapter was to investigate the physicochemical, thermal properties as well as the functional properties of acha starch.

3.2 Materials and Methods

The experimental outline for this chapter is detailed in Figure 3.1. Description of each step in the study will follow.

3.2.1 Source of acha grain and materials

Two acha cultivars namely, *Digitaria exilis* (white) and *Digitaria iburua* (black) were purchased from Grace Africa, Salt River, Cape Town, South Africa.

All equipment and materials were either obtained from the Department of Food Technology, Cape Peninsula University of Technology (CPUT) or Pioneer Foods Laboratories (Bokomo Foods pilot plant and SASKO technical laboratory), Cape Town, South Africa.

All chemicals were obtained from Laboratory and Scientific Ltd., Maitland, Cape Town, Republic of South Africa and the Department of Food Technology CPUT.

3.2.2 Cleaning of acha grain

The cleaning procedure entailed screening the grains for 3 minutes through a 1000, 750, 500 and 125 micron screens to remove all foreign matter such as

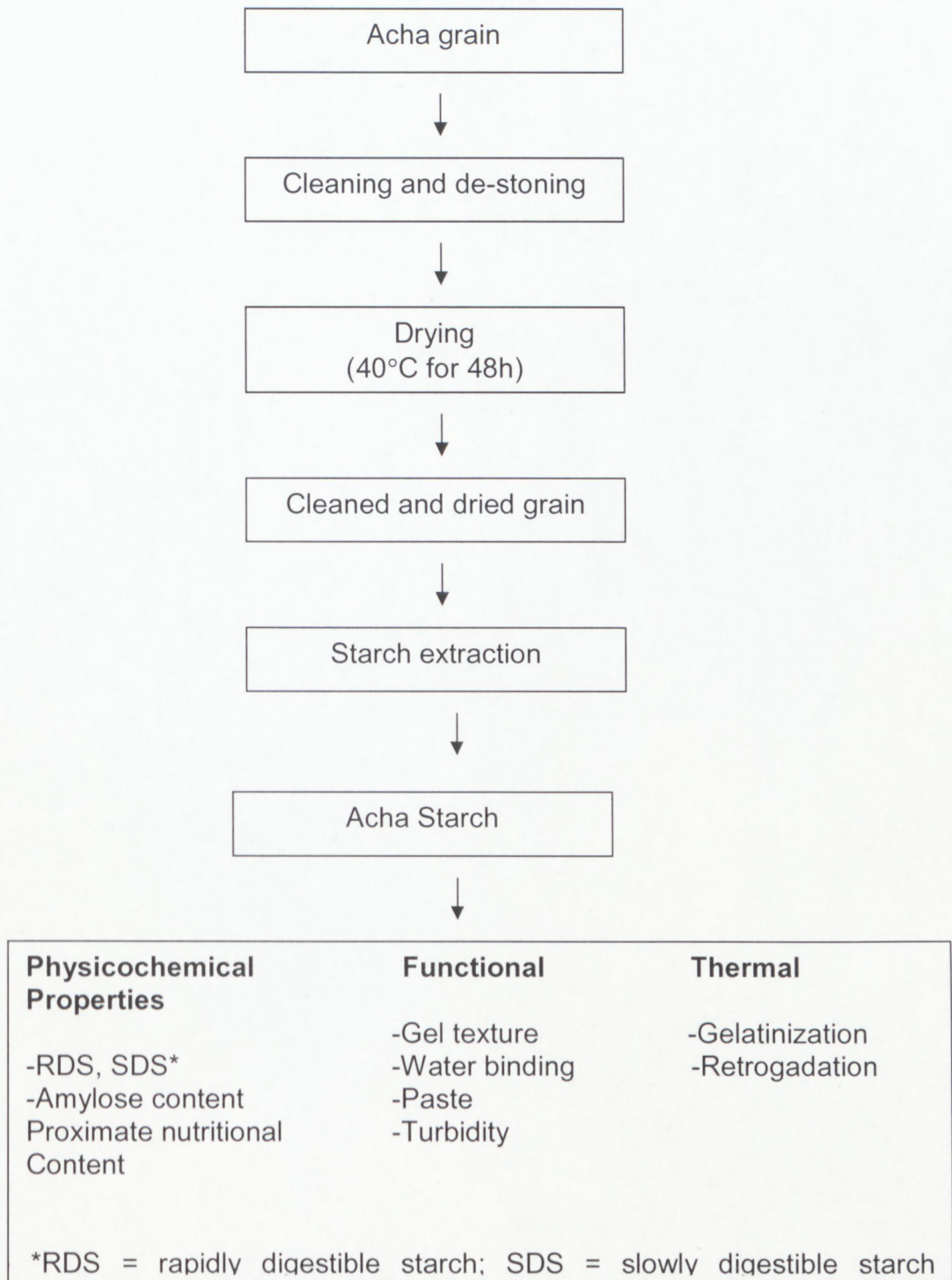


Figure 3.1 Experimental outlines for first research chapter

stones, foreign grain and small sticks. The process of dry cleaning was then followed by rinsing the grains three times with water and drying at 40°C for 48 h (Jideani & Podgorski, 2010). The cleaned and dried samples were then placed in sealed polyethylene bags and stored in a refrigerator at a temperature of $3 \pm 2^\circ\text{C}$ until required.

3.2.3 Isolation and purification of starch from acha grain

The method described by Betancur & Torruco-Uco (2001) as outlined in Figure 3.2 was used to extract starch from milled acha grain. Cleaned, washed and dried acha grain were milled and sieved through a 20 mesh screen. The acha flour was then dispersed in distilled water at a ratio of 1:8 w/v. The pH was then adjusted to 11 with 1 N NaOH and the solution was stirred for 1 h. The suspension was then passed through an 80 and 100 mesh screens to separate the fibre solids from the liquid containing the protein and the starch. The suspension was allowed to sediment for 30 min to recover the starch, and then the solubilised protein was separated. The starch was washed 3 times with distilled water, and centrifuged at 4250 rpm for 10 min during the last wash to recover the starch. The product was then dried at 60°C in a drying oven (Defy, Gemini Ltd., Germiston) for 1 h then milled and sieved through a 20 mesh screen. The isolated starch was then analysed for physicochemical, functional and thermal properties.

3.3 Physicochemical Properties of Acha Starch

3.3.1 Starch characteristics of acha starch

Extracted acha starch was analyzed for rapidly digestible starch (RDS), slowly digestible starch (SDS), resistant starch (RS), amylose and total starch utilizing the modified Goñi *et al.* (1997) methods.

3.3.2 Total starch analysis for acha and wheat starch

Total starch was determined enzymatically according to the modified Goñi *et al.* (1997) method. Triplicate samples of both acha starches and wheat starch were analyzed separately. Starch (50 mg) was dispersed in 50 ml centrifuge tubes with 6 ml of KOH. The samples were shaken for 30 min at room

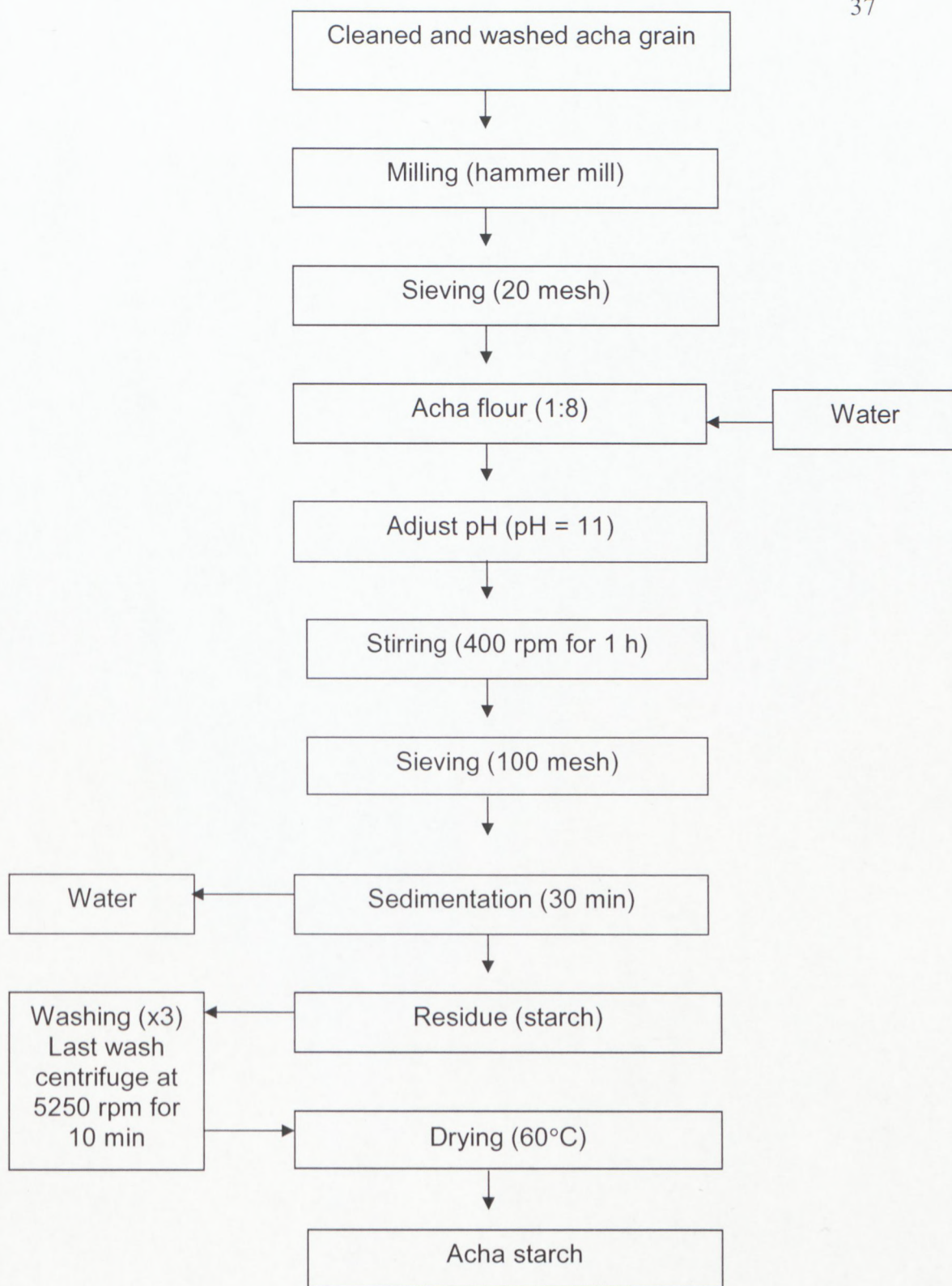


Figure 3.2 Extraction of starch from Acha grain (Betancur & Torruco-Uco, 2001)

temperature. The solubilized starch was then hydrolyzed by the addition of 3 ml 0.4 M sodium acetate buffer (pH 4.8) and 60 μ l of amyloglucosidase (A7420 Sigma Aldrich) from *Aspergillus niger* (67.4 U/mg 50 mg AMG in 50 ml buffer solution). The samples were transferred into Erlenmeyer flasks and then incubated at 60°C for 45 min in a shaking water bath. The samples were then centrifuged for 10 min at 3000 rpm. After centrifugation, the glucose concentration was determined in the supernatant using a glucose oxidase-peroxidase (GAGO-20 Sigma Aldrich) kit. Sample aliquots (1 ml) of the supernatant were measured into test tubes and to that 2 ml of the assay reagent was added. The sample were then incubated at 37°C for 30 min, 2 ml H₂SO₄, was carefully added to the test tubes after incubation and were then thoroughly mixed. Colour absorption was measured against a reagent blank at a wavelength of 540 nm. The glucose concentration was converted into starch by applying the factor 0.9.

3.3.3 Resistant starch analysis of acha and wheat starch

Resistant starch (RS) content in acha starch and wheat starch were determined according to the Goñi *et al.* (1997) method. Dry starch (100 mg) was weighed into a 50 ml centrifuge tube. An aliquot (10 ml) of KCl-HCl buffer (pH 1.5) and 0.2 ml of pepsin (P7000 Sigma-Aldrich) solution (1 g pepsin /10 ml buffer KCl-HCl) were added. The solution was mixed well and was placed in a shaking water bath for 60 min at 40° C. After cooling samples to room temperature, 9 ml of 0.1M Tris - maleate buffer (pH 6.9) was added and was followed by adding 1 ml of α -amylase (A3176 Sigma-Aldrich) solution (40 mg α -amylase/ml Tris-maleate buffer). The samples were then mixed well and incubated for 16 h in a water bath at 37°C with constant shaking. The samples were centrifuged for 15 min at 3000 rpm, and the supernatants were discarded. The residues were moistened with 3 ml distilled water and 3 ml of 4 M KOH then mixed well for 30 min at room temperature with constant shaking. Thereafter, 5.5 ml of 2 M HCL and 3 ml of 0.4 M sodium acetate buffer (pH 4.75) were added, followed by the addition of 80 μ l of amyglolucosidase. The solution was then mixed well and was placed in a water bath at 60°C for 45 min with constant shaking. The solution was then centrifuged at 3000 rpm for 15 min and the supernatants were collected and

saved in a 50 ml volumetric flask. To 1 ml of the sample in a test tube was added 2 ml of the assay reagent. The samples were then incubated at 37°C for 30 min. Sulphuric acid (2 ml) was carefully added to the test tubes after incubation and were then thoroughly mixed. Colour absorption was measured against the reagent blank at a wavelength of 540 nm and the glucose concentration was converted into starch content by multiplying by the factor 0.9.

3.3.4 Digestible starch

DS was calculated as the difference between TS and RS (Goñi *et al.*, 1997).

3.3.5 Determination of rapidly and slowly digestible starch by *in vitro* rate of starch digestion

The procedure and model established by Goñi *et al.* (1997) was used to measure the *in vitro* starch hydrolysis. Triplicate samples of 50 mg of starch were dispersed in 10 ml of KCl-HCl buffer. Pepsin solution (0.2 ml) containing 1 g of pepsin in 10 ml HCl-KCl buffer (pH 1.5) was added to the samples. The samples were incubated at 40°C for 60 min in a shaking water bath. The volume was raised to 25 ml by adding 15 ml Tris-maleate buffer (pH 6.9) and adjusting the pH carefully. Starch hydrolysis was started by adding another 5 ml of Tris- maleate buffer containing 2.6 UI α -amylase to each sample.

The flasks were placed in a shaking water bath at 37°C with moderate agitation. Sample (1 ml) was taken from each flask every 30 min from 0-3 h. These aliquots were placed in a test tube at 100°C and were energetically shaken for 5 min to inactivate the α -amylase. The 2 ml of 0.4 M sodium acetate buffer (pH 4.75) was added to each aliquot and 60 μ l of amyloglucosidase was used to hydrolyze the digested starch into glucose after 45 min at 60°C in a shaking water bath. The volume was adjusted to 5 ml with distilled water and 0.5 ml was incubated with the glucose oxidase-peroxidase kit. The samples were analyzed in triplicate. The glucose was finally converted into starch by multiplying by 0.9. The rate of starch digestion was expressed as percentage of starch hydrolyzed at different times (30, 60, 90, 120, 150 and 180 min). RDS and SDS were determined according to Englyst *et al.* (1992). The RDS was defined as the percentage of starch

digested at 30 min and the SDS as the percentage of starch digested at 120 min. The digestion curve was modeled with the non-linear equation (Eq 1) established by Goñi *et al.* (1997) to describe the kinetics of starch hydrolysis.

$$C = C_{\infty} (1 - e^{-kt}) \quad \dots(1)$$

Where C is the percentage of starch hydrolyzed at time t (min); C_{∞} is the equilibrium percentage of starch hydrolyzed after 180 min, and k is the kinetic constant. The parameters C_{∞} and k were estimated for each sample using SPSS for Windows 19.0 non-linear regression.

3.3.6 Determination of total amylose content of acha starch

The method by Hoover & Ratnayake, (2001) was used to determine the amylose content of starch. Defatting of starch samples were conducted by weighing 5 g of starch accurately into a cellulose extraction thimble and covering the mouth with cotton wool. Lipids were then extracted with 120 ml of 75% n-Propanol at 85°C with a heating mantle for 7 h in a Soxhlet extractor. The lipid free starch was then air dried for 12 h and removed from the thimble and oven dried for 24 h at 30°C. Lipid-free starch (20 mg) was weighed into a round bottom screw-cap tube fitted with a Teflon-faced rubber liner in the cap. A series of mixtures of pure potato amylose and amylopectin (0%, 10%, 20%, 40%, 50% amylose) were prepared after which 20 mg of each was weighed into round-bottom tubes with caps. Eight millilitres of 90% DMSO was added to each round-bottom tube and vigorously mixed for 2 min using a vortex mixer. The tubes were then heated in a water bath at 85°C for 15 min with intermittent mixing. The tubes were then allowed to cool to room temperature (45 min). The samples were then diluted to 25 ml with water in a volumetric flask.

Determination of absorbance of the dispersed starch solution was then conducted after completion of the following steps. Diluted solution (1 ml) and 40 ml of distilled water were added into a 50 ml volumetric flask. Iodine solution (5 ml) was added and mixed vigorously. The volume was adjusted to 50 ml with distilled water and mixed vigorously. The colour was then allowed to develop for 15 min. The contents of the flask were then vigorously mixed by hand. The absorbance of the samples and each of the standard mixtures

were then measured at 600 nm against a reagent blank as the reference. The reagent blank contained all reagents in the same amounts without the sample containing starch.

Following the above procedure a standard curve was then plotted. The regression equation for the standard curve was then measured and used to calculate the total amylose content of the sample.

3.3.7 Proximate composition of acha starch

The moisture, fat, protein, fiber and ash content of the acha and wheat starch were determined in accordance with the standard methods of AOAC (1990). Carbohydrates were calculated by difference.

3.4. Functional properties of the Starches

3.4.1 Textural properties of starch gels

Texture profile analysis was conducted as described by Sandhu & Narpinder (2007) on an Instron apparatus (model 2519, UK, 3300 series, 2000 N capacity) to evaluate the textural properties of acha and wheat starch gels. A suspension of 45 g of starch in 455 ml of cold water was prepared. The suspension was then boiled on a hot plate (Model Z341 Ohaus Instr. US) with stirring until a gel was formed. The hot starch gel formed was then poured into small aluminium dishes (10 cm in diameter) and stored at 4°C. The gels were evaluated for their textural properties by means of texture profile analysis.

Each dish was placed on the food support plate (200 x 145 mm) of the Instron, the gel was compressed at a speed of 0.5 mm/s to a distance of 10 mm (30% of the original sample height) with a cylindrical plunger (5 mm diameter). The compressive force was then removed and the sample was re-compressed. Such a compressive sequence represents two bites (Karim *et al.*, 2000). During the test, compressive force was recorded as a function of the amount of compression (distance). Thus two, forces against distance, TPA curves could be derived from which hardness (height of first peak) and springiness (ratio between recovered height after the first compression and the original gel height) were determined. Chewiness was obtained by multiplying gumminess and springiness (Sandhu & Narpinder, 2007). These

were calculated automatically with the Blue-Hill 2 software.

3.4.2 Turbidity of starch suspensions

The turbidity of the acha starch suspension was measured as described by Perera & Hoover (1999). A 1% aqueous suspension of acha starch as well as the reference wheat starch was heated separately, in a water bath at 90°C for 1 h with constant stirring. The samples were then stored for 5 days at 4°C and turbidity was determined after 5 days by measuring the absorbance at 640 nm (UV-VIS Split Beam 8 Auto Cell Model UVS-2800 spectrophotometer Shanghai Jingke Scientific Instruments Co. Ltd. China) against a water blank. The turbidity was measured in Nephelometric Turbidity Units (NTU). NTU measures how much light is scattered in the suspension. The greater the scattering, the higher the NTU; thus the lower the clarity of the suspension.

3.4.3 Water binding capacity

Water binding capacity (WBC) of both the acha starch as well as the reference starch was measured as described by Medcalf & Gilles (1965). A suspension of 5 g starch in 75 ml distilled water was agitated for 1 h and centrifuged at 3000 rpm for 10 min. The free water was drained for 10 min and the drained wet starch was then weighed. The water binding capacity was then expressed as grams per hundred grams.

3.4.4 Pasting properties

The method of Holm *et al.* (1988) was used to determine the pasting properties of acha starch as well as the reference sample using a rapid visco analyzer (RVA 4 from Newport Scientific, Warriewood, Australia). The viscosity profiles of white acha, black acha and wheat starch were recorded using starch suspensions of 9% (w/w, dry basis) and 500 g total sample weight. The starch suspensions were prepared and then transferred to a rapid visco analyzer (RVA4 from Newport Scientific, Warriewood, Australia). The capacity of the RVA was 50 – 50 000 cP at 80 rpm. A programmed heating and cooling cycle were used where the samples were held at 25°C for 1 min and heated to 95°C at 1.5°C/min, a holding at 95°C for 30 min before

cooling from 95° to 50°C, 1.5°C/min and holding at 50°C for 2 min. Parameters recorded were peak viscosity, trough viscosity, final viscosity, breakdown viscosity and setback viscosity. Analysis was conducted in triplicate.

3.5. Thermal properties of the starches

Differential scanning calorimeter (DSC, 20 to 120°C, Perkin-Elmer instrument) was used to measure the thermal characteristics of the isolated starch. Both the acha starches and wheat starch (3.5 mg, dry weight) was separately loaded into a 40 µl capacity aluminium pan (Mettler ME-27331) and distilled water added using a Hamilton micro syringe to achieve a 70% starch–water suspension. Samples were sealed and allowed to stand for 5 h at room temperature before it was heated in the DSC. The DSC analyzer was calibrated using indium, and an empty aluminium pan was used as a reference (Sandhu and Narpinder, 2007). Sample pans were heated at a rate of 10°C/min from 20°C to 100°C. Thermal transition of starch samples was defined as T_o (onset temperature), T_p (peak of gelatinization), T_c (conclusion temperature) and ΔH_{gel} the enthalpy of gelatinization. Enthalpies were calculated on a starch dry weight basis automatically. The gelatinization temperature range (R) and peak height index (PHI) were calculated as described in equations 1 and 2.

$$R = 2(T_p - T_o) \quad \dots\dots 1$$

$$PHI = \Delta H / (T_p - T_o) \quad \dots\dots 2$$

After conducting thermal analysis, the samples were stored at 4°C for 7 days, for retrogradation studies. The aluminium sample pans containing the starches were reheated at the rate of 10°C/min from 25°C to 100°C to measure retrogradation. The enthalpies of retrogradation (ΔH_{gel}) T_o , onset temperature; T_p , peak temperature; H_1 , peak height; H, height; ΔH , enthalpy of gelatinization; T_c , end (conclusion) temperature, ΔT_r gelatinization temperature range ($T_c - T_o$), were calculated automatically by the Pyris

R9.1.software.

3.6 Results and Discussion

3.6.1 Proximate composition of acha starch

Table 3.1 shows the proximate content of two acha starch varieties and wheat starch. The moisture content of wheat, black and white acha starches was 11.0, 11.9 and 10.7% respectively. There was no significant difference between the white acha starch and wheat starch. The moisture content recorded for white acha starch and black acha starch were slightly lower than that reported for two acha cultivars (Fonio Hothia and F. Koilli), 15 and 13% respectively. The moisture value for white acha starch (10.7%) was in the same range of that reported for corn starch 10.1% (Gonzales-Reyes *et al.*, 2003).

The protein content of white (1.7%) and black (1.2%) acha differed significantly from that of wheat starch (0.5%). There was also a significant difference in the protein content between white (1.7%) and black (1.2%) acha starch). Protein content of acha starch reported in literature ranged between 0.3 and 0.7% for Hothia and Koulli respectively (Carcea and Acquistucci, 1997). Classical studies at the University of Illinois demonstrated that the variability of various chemical compounds such as the protein content in maize is of genetic, environment and cultural origin and that chemical composition can thus be changed through appropriate manipulation (FAO, 2012). The protein content of starch is one of the critical characteristics when starch is used in the manufacture of glucose syrup. The protein content of corn starch from Quinoa Corporation investigated by Lindeboom *et al.* (2005) was reported to be 1.2%; this is similar to the protein of black acha starch (1.2%) in this study. The lower the protein contents the better the chances of avoiding Maillard reaction (Gonzales-Reyes *et al.*, 2003). Although the protein content of acha starch (1.2% for black and 1.7% for white acha starch) being significantly higher than that of wheat starch, the protein content is within the acceptable range in starches that would possibly avoid Maillard reaction.

There was no significant difference ($p > 0.05$) between wheat and black acha starch in terms of dietary fibre, 2.2% and 1.5% respectively. There was

Table 3.1 Proximate and amylose composition of acha starch^{1,2}

Sample	Moisture	Fat	Carbohydrate	Protein	Amylose content
Wheat	11.0 ± 0.00 ^a	0.5 ± 0.00 ^a	88.0 ± 0.00 ^a	0.47 ± 0.47 ^a	45.8 ± 16.56 ^a
Black acha	11.9 ± 0.25 ^b	0.8 ± 0.00 ^a	86.0 ± 0.00 ^a	1.23 ± 1.23 ^b	42.1 ± 16.35 ^a
White acha	10.7 ± 0.26 ^a	1.3 ± 0.00 ^a	86.0 ± 0.00 ^a	1.73 ± 1.73 ^c	61.3 ± 15.46 ^a

¹Values are mean ± standard deviation.

²Any two means with different superscript in each column differ significantly (p < 0.05).

however a significant difference between black acha (1.5%) and white acha starch (0.45%). The difference in dietary fibre could be attributed to genetic difference between white acha and black acha starch. Dietary fibre recorded for white acha starch (0.45%) is higher than that reported in literature for lentil starches (0.21%) (Gonzales *et al.*, 2002).

The fat content of white acha starch and black acha differed significantly from that of wheat starch. This difference could be due to their genetic differences. The fat content of black acha (2.9%), white acha (2.0%) and wheat starch (1.8%) are much higher than that reported for corn and Okenia starch, 0.17% and 0.1% respectively (Gonzales-Reyes *et al.*, 2003). Higher values could affect gelatinization due to the formation of amylose-lipid complexes (Gonzales-Reyes *et al.*, 2003).

The carbohydrate content of white acha, black acha and wheat starch were 86.0, 86.0 and 88.0 g/100 g respectively. The carbohydrate content for wheat starch and the two acha varieties did not differ significantly ($p > 0.05$).

There were no significant differences between either the white or the black acha starches and wheat starch with regard to moisture content, dietary fibre and carbohydrate content. There was thus no significant difference in the nutritional composition of white acha starch and black acha starch with regard to carbohydrate, moisture and fat content. There was however significant differences in the nutritional composition with regard to the protein content and dietary fibre of black and white acha starches. This supports the hypothesis that there is significant difference between the two acha starch varieties.

3.6.2 Amylose content of acha starch

The amylose content of acha and wheat starches is detailed in Table 3.1. There was no significant difference in amylose content between black acha (42.1%), white acha (61.4%) and wheat (45.8%) starches. Lower amylose values were reported in literature for two other varieties of acha (Hothia and Koulli) 22.6 and 26.1% (Carcea and Acquistucci, 1997). In this instance the amylose content was measured in the presence of lipids which may complex with amylose and reduce

its iodine binding capacity (Morrison, 1988). Taylor *et al.* (1997) however reported amylose content of starch, extracted from sorghum cultivars, to be in the region of 35.7 to 36.9% for high pasting peak viscosity sorghum starch and 27.1 to 47.3% for low peak viscosity sorghum starch that received supplementary irrigation. Literature revealed that environmental effects may exert more influence on amylose content than genetic differences (Taylor *et al.*, 1997). There was a negative correlation between glycemic index, resistance starch and amylose content of starch (Chung *et al.*, 2008a). Higher amylose content may contribute to a lower hydrolysis index (HI) and estimated glycemic index (eGI). The results do not support the hypothesis that there is significant difference in the amylose content between black and white acha starches. Hence, amylose contents of the two acha starches are similar.

3.6.3 *In vitro* starch digestibility of acha and wheat starches

The starch hydrolysis of starch isolated from black and white acha and wheat grain with respect to incubation time are presented in Table 3.2. The analysis of variance (ANOVA) for the effect of time and type of starch on starch hydrolysis for the two acha variants and wheat grain are presented in Table 3.3. The amount of hydrolyzed starch was significantly ($p < 0.05$) affected by both time and starch type in the *in-vitro* starch digestibility. There was no significant ($p > 0.05$) difference in the amount of starch hydrolysed within 30, 60, 120 and 150 min for all types of starches. There was no significant difference in the total amount of starch hydrolysed after 180 min for wheat (56.79%) and white acha starch (57.13%). However, there was a significant difference ($p < 0.05$) between white acha starch (56.79%) and black acha starch (39.02%) in the amount of starch hydrolysed within 180 min. The values for white acha starch and wheat starch although slightly higher, still in the same range as those reported by Zhang *et al.* (2006), for waxy maize (46.6%), maize (53.0%) and wheat (50.0). The amount of starch hydrolysed in black acha starch was however much lower than the value reported by Zhang *et al.* (2006).

The two acha starch cultivars had fairly low RDS contents, white acha

Table 3.2 The percentage of starch hydrolyzed within 0 -180 min of starch from acha and wheat grain^{1,2,3}

Grain	Time (Min)						Total
	30	60	90	120	150	180	
Starch							
Wheat	87.4 ± 1.1	83.2 ± 1.4	82.9 ± 1.1	54.5 ± 1.6	66.9 ± 16.0	22.7 ± 5.3	56.8 ± 32.4 ^a
White acha	75.0 ± 4.3	72.8 ± 17.7	69.9 ± 27.2	67.7 ± 50.7	66.7 ± 28.6	47.8 ± 2.87	57.1 ± 33.0 ^a
Black acha	63.8 ± 1.9	62.1 ± 0.8	40.1 ± 0.9	38.8 ± 4.8	36.7 ± 2.9	31.7 ± 0.2	39.0 ± 20.3 ^b
Total	75.4 ± 10.5 ^a	72.7 ± 12.8 ^a	64.3 ± 23.4 ^a	56.7 ± 22.3 ^b	53.7 ± 28.4 ^b	34.1 ± 11.4 ^c	

¹Values are mean ± standard deviation.

²Any two means with different superscript in the total column differ significantly ($p < 0.05$).

³Any two means with different superscript in the total row differ significantly ($p < 0.05$).

Table 3.3 Analysis of variance (ANOVA) for the effect of time and type of starch grain on hydrolysed starch

Source	Type III Sum of				F	Sig.
	Squares	df	Mean Square			
Corrected Model	45977.356 ^a	20	2298.868	10.057	.000	
Intercept	163747.985	1	163747.985	716.325	.000	
Starch type	4509.892	2	2254.946	9.864	.000	
time	37542.264	6	6257.044	27.372	.000	
Starch type * time	3925.199	12	327.100	1.431	.190	
Error	9600.966	42	228.594			
Total	219326.307	63				
Corrected Total	55578.322	62				

^aR Squared = 0.827 (Adjusted R Squared = 0.745)

(75.0%) and black acha starch (63.8%) compared to wheat starch (87.42%) whereas the SDS for wheat (54.47%) and white acha starch (67.67%) was higher than that of black acha starch (38.79%). No literature values for RDS and SDS for both acha starch cultivars could be found. Chung *et al.*, (2008b), reported that isolated bean starch (63-65%) contained a substantially larger amount of SDS than those reported for cereal starches, maize (53.0%), waxy maize (47.6%), wheat (50.0%). The SDS for white acha starch (67.7%) is higher than the SDS reported by Chung *et al.*, (2008b) for maize and wheat. RDS content of acha starch was substantially higher than that reported for corn (24.4%), wheat (40.1%) and rice (32.4%) starches (Zhang *et al.*, 2006). On the contrary, the amounts of starch hydrolysed for white acha and wheat starch is much lower than that reported for bean starch which ranged from 63.1 – 65.8% (Chung *et al.*, 2008a). In another study by Chung *et al.* (2008b), unmodified starches showed a higher level of hydrolysis than did the modified starches after 180 min. After 180 min of digestion the amount of starch hydrolysed for white acha starch was much higher than the amount of starch hydrolysed from black acha starch. This difference supports the hypothesis that a significant difference exist in the in-vitro starch digestibility between black and white acha starches. White acha starch appears to have a faster rate of digestibility than black acha starch.

3.6.4 Functional Properties of Acha Starch

Textural characteristics of the starch gels

The textural characteristics of the acha and wheat starches are indicated in Table 3.4. The hardness for white acha starch, black acha starch and wheat starch were 0.9367 N, 0.9467 N and 0.9500 N, respectively. No significant difference in gel hardness was observed between black acha starch gel and white acha starch gel or between the wheat starch and the two acha starch varieties. These low hardness values indicate that both acha and wheat starches produced soft gels. Acha starch would thus be suitable for manufacturing of foodstuff where soft gels are required. Sandhu & Singh (2007) reported that starches which exhibit harder gels tend to have higher amylose content. They also reported that the gel

Table 3.4 Textural characteristics of acha and wheat starch gels^{1,2}

Cereal starch	Hardness(N)	Resilience(ratio)	Springiness (mm)
White acha	0.936 ± 0.041 ^a	1.199 ± 0.090 ^a	0.709 ± 0.112 ^a
Black acha	0.946 ± 0.015 ^a	1.239 ± 0.087 ^a	0.709 ± 0.132 ^a
Wheat	0.950 ± 0.010 ^a	1.240 ± 0.033 ^a	0.374 ± 0.273 ^a

¹Values are mean ± standard deviation.

²Any two means with different superscript in each column differ significantly (p < 0.05).

firmness is mainly caused by retrogradation which is associated with crystallization of amylopectin, leading to harder gels.

The resilience value is the first force before the maximum force (F) which is the capability of the gel to return to an original shape or position after having been compressed. The resilience for black and white acha and wheat starch gels were 1.1994, 1.2399 and 1.2402, respectively. There was no significant difference ($p > 0.05$) in resilience between the gels. Springiness for black acha gel was 0.7090 mm, white acha gel was 0.7090 mm and wheat gel was 0.3746 mm. There was no significant difference ($p > 0.05$) in the springiness of both acha starch gels and wheat gels. Springiness refer to the degree of gel rubberiness in the mouth (Lau *et al.*, 2000). A high degree of springiness is depicted by gel structure that breaks into large pieces during the initial compression; whereas a low degree of springiness is represented by a gel that breaks into smaller pieces during the initial compression test (Huang *et al.*, 2007). The starch gels broke into large pieces during compression. Hence, both acha and wheat gels are highly springy. Highly springy gels would not break down easily during mastication (Lau *et al.*, 2000). Acha starch gels were springy and resilient. This indicates that both acha starch varieties can be used for hot and cold desserts as well as for soft jelly like sweets and confectionery toppings.

Since there was no significant differences in the hardness, resilience and springiness between the two acha starch gels, the hypothesis that the starch from the two acha cultivars (*D. exilis* and *D. iburu*) differ in textural properties is rejected.

Pasting properties of acha starch

The pasting properties of wheat starch, black acha starch as well as white acha starch are summarized in Table 3.5. Peak viscosity is the ability of starch to swell before their physical breakdown, an indication of the water binding capacity of starch (Ikegwu *et al.*, 2010). The peak viscosity was 3506 cP for wheat starch, 3994 cP for black acha starch and 4936 cP for white acha gel. There was a

Table 3.5 Pasting properties of Acha of Starch^{1,2}

Sample	Peak1 (cp)	Trough1 (cp)	Breakdown viscosity (cp)	Final viscosity (cp)	Setback viscosity(cp)	Peak Time (min)	Pasting Temp. (°C)
Wheat	3506 ± 16 ^a	1960 ± 41 ^a	369 ± 21 ^a	4436 ± 82 ^a	2004 ± 62 ^a	4.18 ± 0.04 ^a	78.03 ± 0.46 ^a
Black acha	3994 ± 36 ^b	3125 ± 15 ^b	869 ± 34 ^b	5129 ± 60 ^b	2332 ± 54 ^b	5.78 ± 0.10 ^b	80.15 ± 0.44 ^b
White acha	4936 ± 58 ^c	3137 ± 6 ^b	2976 ± 95 ^c	5470 ± 50 ^c	2476 ± 190 ^b	6.69 ± 0.03 ^c	85.88 ± 0.49 ^c

¹Values are mean ± standard deviation.

²Any two means with different superscript in each column differ significantly (p < 0.05)

significant difference ($p < 0.05$) in the peak viscosity between white acha starch and black acha starch. Acha starch has a higher peak viscosity than that reported in literature for corn starch (Sandhu & Singh, 2007). The peak viscosity of various corn starches ranged between 804 and 1552 cP. The higher peak viscosity of acha starch is indicative of their higher water binding capacity and thus higher degree of starch swelling, an indication that acha starch may be good for products requiring high gel strength and elasticity. Trough viscosity is the measurement of the holding strength of the starch paste before it breaks down and viscosity decreases. This depends on the temperature and degree of mixing or shear stress. The trough viscosity for two acha starch varieties were similar, 3125 cP and 3137 cP for black and white acha starch. There was however a significant difference between the two acha starch varieties and wheat starch trough viscosity, 1960, 3125, 3137 cP for wheat starch, black acha and white acha, respectively. This significant difference ($p < 0.05$) in trough viscosity implies a difference in the paste holding strength of wheat starch and the two acha starch varieties. The holding strength of black acha and white acha starch is similar while the holding strength of wheat starch paste is much lower than the two acha starch paste varieties. This signifies that acha starch gels may be the better option for products requiring high holding strength without breaking down and thus lead to a decrease in viscosity. The trough viscosity are much higher than the values recorded for African Tall (662 cP) and Ageti (652 cP) corn starch (Sandhu & Singh, 2007).

The break down viscosity for wheat, black acha and white acha starches were 369, 869, 2976 cP respectively. The break down viscosity for white acha starch was significantly higher ($p < 0.05$) than those of wheat and black acha starches. Breakdown viscosity is the measure of disintegration of cooked starch. It is the difference between the peak viscosity and the trough viscosity. The higher the breakdown viscosity; the lower the ability of the sample to withstand heating and shear stress during cooking (Adebowale, 2004). This indicates that white acha starch being higher in break down viscosity, will have a lower ability to withstand heating and shear stress during cooking.

The final viscosity for black acha starch was 5129 cP, white acha starch 5470 cP and wheat starch 4436 cP. The final viscosity, which is the measure of the ability of starch to form a viscous paste for white starch was significantly higher ($p < 0.05$) than those of black acha starch and wheat starch. This difference could be due to the difference detected in the peak time and pasting temperatures of all three starches, white acha starch 6.69 min at 85.88°C, black acha starch 5.7 min at 80.15°C and wheat starch 4.18 min at 78.03°C. The higher pasting temperatures for acha starches indicated the higher resistance of their starches to swelling. The increase in viscosity with temperature may be contributed by the removal of water from the exuded amylose as the starch granules swell (Sandhu & Singh, 2007).

The setback viscosity is the measure of the degree of syneresis of starch upon cooling of the cooked starch paste, for white acha, black acha and wheat starch were 2476 cP, 2332 cP and 2004 cP respectively. There was no significant difference between the acha starches in setback viscosity. However, the wheat starch was significantly lower ($p < 0.05$) in setback viscosity compared to the acha starches. The high setback values for acha starches make them unsuitable for use where low syneresis rate is required, such as in frozen or refrigerated foods. The setback values are also indicative of the retrogradation tendency of starch gels. The higher the setback viscosity the lower the retrogradation during cooling of the products made from the flour (Ikegwu *et al*, 2010). This implies a significant difference in retrogradation tendency between wheat starch and the two acha starch varieties. Acha starches have lower retrogradation tendencies during cooling compared with wheat starch, suggesting that the degree of re-association of the wheat starch molecules was higher than that of acha starch molecules, upon cooling. Sandhu & Singh (2007) reported that the pasting properties of starch depend upon various factors such as the rigidity of starch granules, which in turn affects the degree of swelling of the starch granules. The significant differences in peak, break down, and final viscosity support the hypotheses that there are differences in the pasting properties between white and black acha starches.

Turbidity of acha starch

The turbidity for wheat, black acha and white acha starches are reported in Table 3.6 the values were 0.2097 NTU, 0.0003 NTU and 0.0193 NTU respectively for wheat, black and white acha starches. There was significant difference ($p < 0.05$) in the turbidity of the starches, with wheat starch being the highest followed by white acha starch and lastly black acha starch. The turbidity of starch in foods is important when used as thickener in sauces, as a carrier of flavours in beverages or as a suspending agent in liquid foods. The higher the NTU value the lower the clarity of the suspension. This signifies that both acha starch varieties are clearer than wheat starch, and would be the better option to use in products where starch clarity is required. Sandhu & Singh (2007) reported that turbidity development of starches is affected by inter-related factors such as starch granule swelling, leached amylose and amylopectin, amylose and amylopectin chain length. The substantial difference in turbidity between white acha starch and black acha starch supports the hypotheses that the white acha starch and black acha starch will differ in their functional properties.

Water binding capacity of acha starch

The water binding capacity (WBC) for acha and wheat starches are detailed in Table 3.6. The water binding for wheat, black and white acha starches were 0.83, 1.33 and 1.36 g/100g respectively. The WBC for both acha starches were significantly higher ($p < 0.05$) than for wheat starch. This difference in terms of WBC can be due to structural differences in starch and proteins of the starches (Celik *et al.*, 2005). Different proportions of crystalline and amorphous regions within the granules may be the result of the variations in WBC. Thus weakly bonded amorphous starch granules will imbibe less water (Carcea & Acquistucci, 1997). It appears that acha starch granules possess stronger bonded amorphous granules compared to wheat starch. There was however no

Table 3.6 Turbidity and WBC of acha starch^{1,2}

Starch	Turbidity(NTU) ³	Water binding capacity (g/100 g)
Black acha	0.0003 ± 0.001 ^a	1.333 ± 0.06 ^b
White acha	0.0193 ± 0.002 ^b	1.367 ± 0.06 ^b
Wheat	0.2097 ± 0.009 ^c	0.833 ± 0.23 ^a

¹Values are mean ± standard deviation.

²Any two means with different superscript in each column differ significantly (p < 0.05).

³NTU, nephelometric turbidity units

significant difference ($p > 0.05$) in WBC between black acha starch 1.33 g/100g and white acha starch 1.36 g/100g. This similarity does not support the hypotheses that the white acha starch and black acha starch will differ in their functional properties. Water binding capacity (WBC) is considered very important in foods such as sauces, batters, dough(s) and baked products.

The WBC of the starch affects important physical attributes such as the viscosity of sauces and batters and the texture of baked products. Sandhu & Singh (2007) reported WBC for starches from different corn varieties in the range of 0.82 to 0.97 g/100 g. The water binding capacity for acha starch is higher than the values reported by Sandhu & Singh (2007). This is also true in practice. Acha is known to swell far more than other cereals (Carcea & Acquistucci, 1997). Acha starch can thus be included as part of the ingredients in the manufacturing of sauces, batters, and dough(s) and baked products. It can be used as thickener, bulking agent and most importantly due to its high WBC as water retention agent.

3.6.5 Thermal properties of starches

Gelatinization of acha starch

The onset, peak and conclusion temperatures (T_o , T_p , and T_c) and enthalpy (ΔH) of wheat and acha starches are shown in Table 3.7. There was no significant difference ($p > 0.05$) in peak and onset temperature between the acha and wheat starches. Wheat starch had the highest onset (T_o) and peak (T_p) temperatures 34.8°C and 74.6°C, although it was not significant. White and black acha starches recorded (25.1° & 26.8°C) and (68.2° & 70.8°C) for (T_o) and (T_p), respectively.

Acha starches and the wheat starch differed significantly ($p < 0.05$) in conclusion temperature T_c and peak T_p temperatures. For wheat starch it was 96.0°C, black acha starch was 87.8°C and for white acha starch was 93.7°C. Miao *et al.* (2009) reported gelatinization temperatures for waxy maize starch of 59.9° (T_o), 69.1° (T_p) & 78.1°C (T_c). The onset temperature range of acha starch is much lower than that of waxy maize starch. This means that less energy is

Table 3.7 Gelatinization of acha starch^{1,2}

Starch	T _o (°C)	T _p (°C)	T _c (°C)	ΔH _{gel} J/kg	ΔT _r = (T _c - T _o)
Wheat	34.7 ± 25.20 ^a	74.56 ± 0.97 ^a	96.03 ± 0.49 ^b	-213.21 ± 169 ^a	58.33 ± 24.71 ^a
White acha	25.05 ± 12.12 ^a	68.20 ± 1.64 ^a	93.67 ± 0.57 ^b	188.42 ± 250 ^b	66.92 ± 11.55 ^{ab}
Black acha	26.75 ± 7.56 ^a	70.54 ± 6.07 ^a	87.85 ± 2.15 ^a	691.04 ± 106 ^c	62.8 ± 5.41 ^a

¹Values are mean ± standard deviation

²Any two means with different superscript in each column differ significantly ($p < 0.05$). T_o, onset temperature; T_p, peak temperature; H₁, peak height; ΔH, enthalpy of gelatinization; T_c, end (conclusion) temperature, ΔT_r gelatinization temperature range (T_c - T_o)

needed to induce gelatinization of acha starches. The peak gelatinization temperature of acha starch is the same as that reported in literature (Miao et al., 2009). Tester & Morrison (1990) reported that due to the structural differences in amylopectin, starches with the low gelatinization temperature has less crystallinity than the high gelatinization temperature starches, this was attributed to structural differences in their amylopectins. Hence, acha starches may have lower degree of crystallinity compared to wheat starch. Ezekiel *et al.* (2007) reported that low transition temperatures of between 50°-67°C for Kufri and Chandramukhi potato starch was due to the lower crystallinity, meaning that the higher the transition temperatures the higher the degree of crystallinity. This would lead to structural stability causing the starch granules to have a higher degree of resistance to gelatinization. Acha starches have low transition temperature which implies a low crystallinity and high amorphous regions (Miao *et al.*, 2009). The gelatinization temperature range ($T_c - T_o$) for white and black acha was 66.9° and 62.8°C respectively, whereas that of wheat starch was 58.3°C. The similarity of the gelatinization temperature range amongst the starches could be attributed to possible similarity in protein content and starch structure.

The melting enthalpy (ΔH) of gelatinization of black acha was 188.41 J/kg and white acha was 691.6 J/kg whereas that of wheat starch was 213.21 J/kg. The melting enthalpy values for white acha starch and black acha starch differed significantly ($p < 0.05$). The melting enthalpy for both acha starch varieties as well as wheat starch was higher than that reported for waxy maize starch (Chung *et al.*, 2008b). The higher melting enthalpy could be due to the difference in the alignments of the hydrogen bonds in the starch molecules. This is caused by the difference in the bonding forces between the double helices that form the amylopectin crystallites (Sandhu & Singh, 2007). Sandhu & Singh (2007) reported melting enthalpy values between 11200 and 12700 J/kg for various corn starches. This is much higher than the melting enthalpy of acha starch.

The similarity of the gelatinization and gelatinization temperature range of white and black acha starches do not support the hypotheses that there is a

significant difference between white acha and black acha starch in thermal properties. However, the significant difference in peak height and melting enthalpy substantiate the hypotheses.

Retrogradation properties of acha starch

The retrogradation of acha starch compared to that of wheat starch is summarised in Table 3.8. Retrogradation is the hydrogen bonding between starch chains that occurs after cooling of gelatinized starch paste (Hoover & Zhou, 2003). The transition temperatures of retrogradation for both acha starches and wheat starch are much lower than the transition temperatures of gelatinization. The onset temperature for wheat, white acha and black acha starches were 19.97°, 17.58° & 19.05°C respectively. Wheat starch had the highest onset temperature followed by black and white acha starch. However, the differences were not significant ($p > 0.05$). Sandhu & Singh (2007) reported onset temperatures for retrogradation in the range between 41.50° & 43.10°C for two corn starch varieties. These values are much higher than that recorded for white acha and black acha starches. According to Sandhu & Singh (2007) this could be due to a less organized manner of recrystallization of the amylopectin branched chains of the gels. The lower onset temperature for the acha starches could be due to weaker molecular bonding of the starch chains.

Values for peak temperature of retrogradation were 6°4.13, 67.05° & 64.84°C for wheat, white acha and black acha starches, respectively. There was no significant difference ($p > 0.05$) between the starches in peak temperatures. Literature values for different corn varieties varied between 52.40° & 54.50°C for African Tall corn starch and Partap corn starch respectively. These values were much lower than that recorded for white acha and black acha starch. Higher peak temperatures of acha starches could be indicative of the need for higher thermal energy for retrogradation.

Retrogradation enthalpy is an indication of the disentanglement and melting of the double helices formed during cooling (Adebowale *et al.*, 2004). The retrogradation enthalpy (ΔH) for white acha starch was 37.61 J/kg and for

Table 3.8 Retrogradation of cereal starch^{1,2}

Starch	T _o (°C)	T _p (°C)	T _c (°C)	ΔH _{retro} J/kg	ΔT _r = (T _c - T _o)
Wheat	17.97 ± 0.02 ^a	64.13 ± 5.35 ^a	94.73 ± 1.78 ^a	56.87 ± 32.62 ^{ab}	76.76 ± 1.76 ^a
White acha	17.58 ± 0.48 ^a	67.06 ± 4.58 ^a	94.83 ± 1.48 ^a	37.61 ± 12.88 ^a	77.25 ± 1.00 ^a
Black acha	19.05 ± 1.30 ^a	64.84 ± 0.53 ^a	95.95 ± 0.52 ^a	118.00 ± 40.58 ^b	76.90 ± 0.78 ^a

¹Values are mean ± standard deviation.

²Any two means with different superscript in each column differ significantly ($p < 0.05$). T_o, onset temperature; T_p, peak temperature; H₁, peak height; H, ΔH, enthalpy of gelatinization; ΔH_f, T_c, end (conclusion) temperature

black acha starch was 118.00 J/kg. There was a significant difference between black and white acha starches, with black acha starch being higher than that of white acha starch. No significant difference was observed for retrogradation enthalpy between wheat starch and both acha starch varieties. Wheat starch had a recorded value of 56.87 J/kg, white acha 37.61 J/Kg and black acha starch 118.00 J/Kg. The differences in the retrogradation enthalpy could be due to the difference in the degree of starch crystallinity of the retrograded starches (Sasaki *et al.*, 2000).

The conclusion melting temperature values for wheat, white acha and black acha starches were 94.73°C, 94.82°C and 95.95°C respectively. There was no significant difference in conclusion temperature of retrogradation between white acha and black acha starches. Whenever a material undergoes a change in physical state for example melting, or transforms from one crystalline form to another, or when it reacts chemically, heat is either absorbed (endothermic) or liberated (exothermic) (Karim *et al.*, 2000). Wheat and both acha starch cultivars presented an endothermic transition (retrogradation). Retrogradation has been identified as the dissociation of the amylopectin crystallites (Gonzales *et al.*, 2003). The retrogradation phenomenon is a process of re-crystallization of the starch polymers (Gonzales-Reyes *et al.*, 2003). In the case of retrograded starch, ΔH (J/kg) provides a quantitative measure of the energy transformation that occurs during the melting of recrystallized amylopectin as well as precise measurement of the transition temperatures (onset, T_o ; peak, T_p and conclusion (end), T_c) (Karim *et al.*, 2000). There is a significant difference in the ΔH and peak height values between white and black acha starches, supporting the hypotheses that there is a significant difference in the thermal properties between white and black acha starch. There were however no significant difference amongst the retrogradation temperature profile (T_o , T_p and T_c) for both white and black acha starch, this opposes the hypotheses.

3.7 Conclusion

The physicochemical, functional and thermal properties of acha starches

compared to wheat starch are reported in this chapter. The following conclusion can be made:

1. The two acha starch cultivars were low in RDS compared to wheat starch. SDS for wheat and white acha starch was higher than that of black acha starch.
2. Although there was no significant difference in the nutritional composition of white acha starch and black acha starch with regard to carbohydrate, moisture and fat content, there was however significant differences with regard to the protein content and dietary fibre between black and white acha starches.
3. There was no significant difference in amylose content between black acha (42.1%), white acha (61.4%) and wheat (45.8%) starches.
4. The peak time and pasting temperatures for acha starches, white acha starch 6.69 min at 85.88°C, black acha starch 5.7 min at 80.15°C is higher than that of wheat starch 4.18 min at 78.03°C. The higher pasting temperatures for acha starches indicated the higher resistance of their starches to swelling. Consequently, the water absorption capacity (WBC) of both acha varieties was higher than that for wheat starch. This indicates that both acha starch varieties can be used for hot and cold desserts as well as for soft jelly like sweets and confectionery toppings.
5. Acha starch cultivars and wheat starch are similar in thermal properties. The observed gelatinisation temperature of acha and iburua starches typifies that of non-waxy starch. Acha starch has similar retrogradation temperature profiles as that of wheat starch.
6. Acha starches may not be suitable for use where low syneresis rate is required, such as in frozen or refrigerated foods.
7. White acha has a lower ability to withstand heating and shear stress during cooking.

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CHAPTER 4

PRODUCTION OF ACHA BREAD AND ITS SENSORY CHARACTERISTICS

Abstract

Bread was baked with two acha cultivars (*Digitaria exilis*, white and *Digitaria iburua*, black) and their specific loaf volume as well as consumer acceptability were investigated and compared against that of white wheat bread. The sensory properties investigated were appearance, crust colour, crumb colour, aroma, taste, firmness, mouthfeel and overall acceptability on a 5-point hedonic scale (1 = undesirable, 5 = desirable). Results indicated that the majority of panel members found both varieties of acha bread, white acha and black acha to be moderately desirable 83.3 % and 81.7% respectively in terms of crust colour, taste and aroma.

4.1 Introduction

Acha is by large the oldest African cereal, it has been cultivated for years in West Africa (Mali, Burkina Faso, Guinea and Nigeria) and the Dominican Republic (Morales-Payán *et al.*, 2003; Jideani *et al.*, 2008) where it is used as a staple food (National Research Council, 1996). Food products where acha grains have been used are porridge, couscous mixed with other flours to make bread, pastries and oven-popped (Jideani, 1999; Philip & Itodo, 2006). It is valued because of its unique taste and nutritional value (Philip & Itodo, 2006). Togolese farmers also consider acha as having medicinal value in being useful for those suffering from diabetes among others (Adoukonou-Sagbadja *et al.*, 2004). Similar medicinal use has also been associated with acha in Nigeria (Jideani, 1999). Being gluten free, food products from acha, like acha bread has the potential to be consumed by individuals suffering from wheat intolerance and gluten allergies namely celiac disease. Jideani *et al.* (2007, 2008) demonstrated

the possibility of producing gluten free loaf from acha flour. However, nothing is known about the effect of processing on the starch and glycemic property of such bread. The objective of this study was thus to produce bread from whole white and black acha flour, assess its consumer acceptability and study the rate of starch hydrolysis of the loaf. The latter objective will be discussed in Chapter 5.

4.2 Materials and Methods

4.2.1 Source of acha grain and materials

Two acha cultivars namely: *Digitaria exilis* (white) and *Digitaria exilis* (black) were purchased from Grace Africa, Salt River, Cape Town, South Africa. All equipment and materials were obtained from the Department of Food Technology Cape Peninsula University of Technology (CPUT) or Pioneer foods laboratories (Bokomo Foods pilot plant and SASKO technical laboratory), Cape Town, and South Africa. All ingredients were obtained from Pick and Pay retail store in Brackenfell, Cape Town.

4.2.2 Cleaning of acha grain and production of acha flour

The cleaning procedure entailed screening the grains through a 1000, 750, 500 and 125 micron screens to remove all foreign matter such as stones, foreign grain and small sticks. The process of dry cleaning was then followed by washing three times by rinsing with water and drying at 40°C for 48 h (Jideani & Podgorski, 2010). The cleaned and dried grain samples were then placed in sealed polyethylene bags and stored in a refrigerator at a temperature of 3 to 5°C until it was required. Cleaned and dried acha grain was milled through a Hammer mill (9FQ-500 series, ZMA manufacturing Co.Ltd. China) and screened through a 20 mesh sieve to produce acha flour.

4.2.3 Experimental design for acha bread

Screening experiment (2^{4-1}) with design resolution IV was used to identify the ingredients (acha starch, xanthan gum, CMC and yeast) with significant effect on the loaf volume. The variables and their proportions as well as the runs are

detailed in Tables 4.1 and 4.2. There was ten design points in total with each design point conducted in triplicate. The ingredients that gave the optimal loaf volume were used to bake the optimal loaf following the procedure described in section 4.2.4. The baked bread was assessed for consumer acceptability as described in section 4.2.6.

4.2.4 Production of acha bread

The method of Jideani *et al.* (2007) was modified and used to produce acha bread. The standard recipe consisted of 100% flour, 1% salt, vegetable fat 2.5% and water (52%). Lite apple juice (7%) was used as a source of sugar for yeast fermentation. The quantities for acha starch, xanthan gum, CMC and yeast were added according to the factorial design (Tables 4.1 and 4.2).

The method used for production of acha flour and acha bread is explained in Figure 4.1. The batter was prepared by pre-mixing the dry ingredients together for 5 minutes at low speed in a Hobart mixer (model 1/6-H.P.). The wet ingredients were added and mixed at high speed for 10 minutes. The batter was divided into equal weights and poured into greased baking pans (10 x 20 cm). The batter in the pans was placed in the proof oven (Macadam's, Johannesburg) for 10 minutes at 40°C. The proofed batter was then baked at 180°C for 55 minutes in a rotary air oven (Macadam's, Johannesburg), in triplicate. After baking, the bread was de-panned and allowed to cool for 20 minutes at room temperature.

4.2.5 Physical properties of acha bread

The physical quality attributes that were analysed on the baked bread were loaf height, loaf weight and loaf volume. Loaf weight was weighed on a Mettler Toledo scale (RSA) with an accuracy of 0.01 g and maximum capacity of 2 kg. The loaf volume was measured by means of seed displacement method using a rectangular plastic container. The container was filled with acha grain, levelled and poured out and noted as the volume that filled the container. The acha bread was then placed in the same container and filled with the same acha grain

Table 4.1 Ingredients and levels used for acha bread production^{1,2}

Variable (%)	Symbol	x_i (%)		
		-1	0	+1
Acha starch	X_1	4	6	8
Xanthan gum	X_2	1.2	1.6	2.0
CMC	X_3	1.2	1.6	2.0
Yeast	X_4	1.2	2.0	2.8

¹Transformation of coded variable x_i to un-coded variable X_i levels could be obtained from $X_1 = 2x_1 + 6$, $X_2 = 0.4x_2 + 1.6$, $X_3 = 0.4x_3 + 1.6$, $X_4 = 0.4x_4 + 2$

²CMC: carboxyl methyl cellulose

Table 4.2 Fractional (2^{4-1}) factorial design with resolution IV for acha bread production

Design point	Acha starch	xanthan	CMC	Yeast
1	-1	-1	-1	-1
2	+1	-1	-1	+1
3	-1	+1	-1	+1
4	+1	+1	+1	-1
5	-1	-1	+1	+1
6	+1	-1	+1	-1
7	+1	-1	+1	+1
8	+1	+1	+1	+1
9	0	0	0	0
10	0	0	0	0

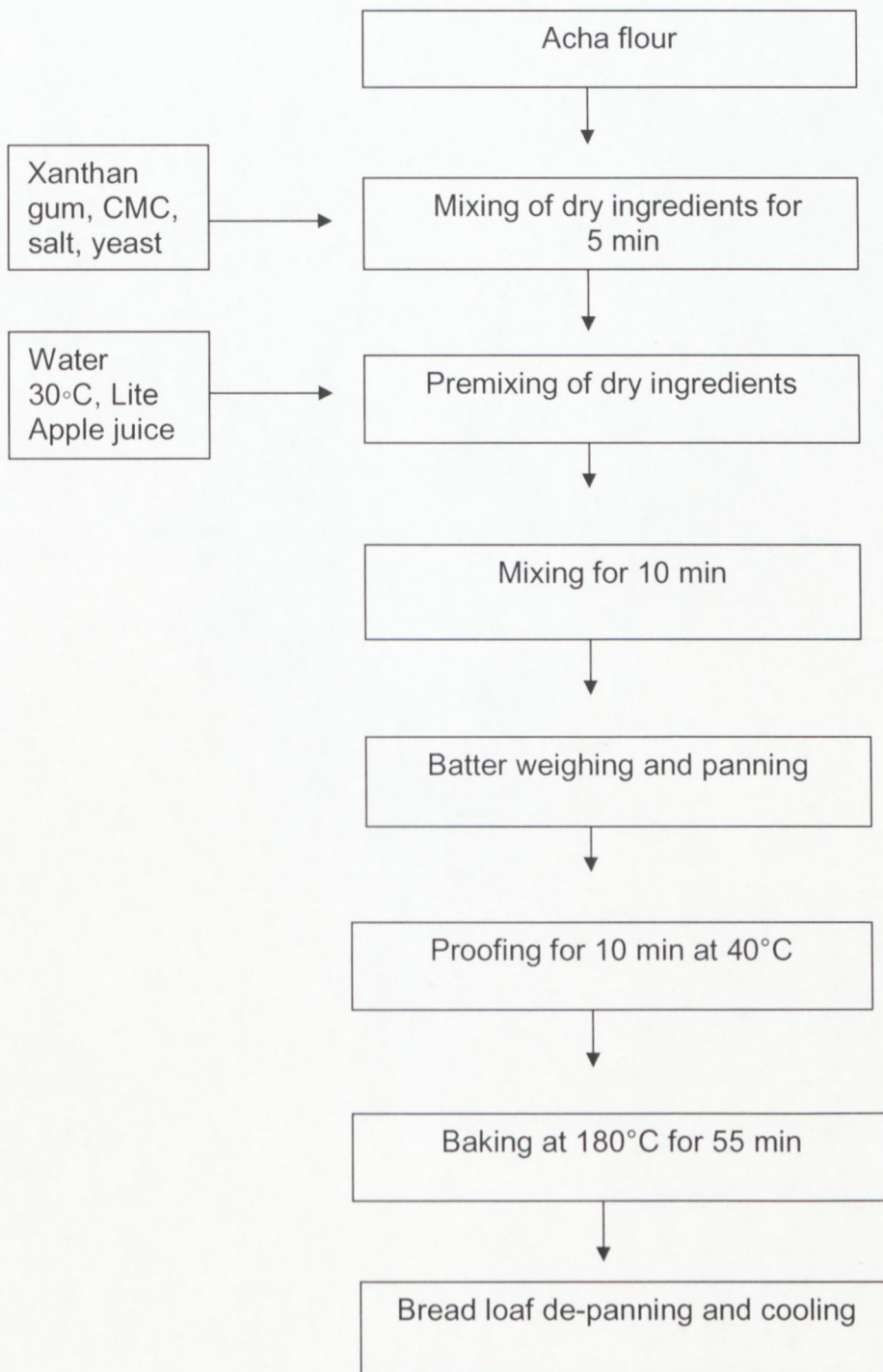


Figure 4.1

Production of acha bread

and levelled. The volume of the spilled acha grains was taken as the volume of the loaf (Ayo and Nkama, 2004; Jideani and Onwubali, 2009). The specific loaf volume was estimated as loaf volume divided by the loaf weight and expressed in ml/g.

4.2.6 Sensory properties of acha bread

The sensory evaluation method as described by Jideani *et al.* (2008) was used to conduct the consumer acceptability test on the optimal acha bread loaf. An evaluation panel of 30 members were used to evaluate the acha bread samples. A slice (3 mm in thickness) of white acha, black acha and wheat bread were served on separate white paper plates (15 cm). Random numbers were used for sample identification. Panellist were required to taste the sample and rate their preference for appearance, taste, aroma, crust colour, crumb colour, firmness, mouth feel and overall acceptability using a 5-point hedonic scale (1, undesirable; 2, undesirable; 3, neither desirable nor undesirable; 4, moderately desirable; 5, desirable).

4.2.7 Data analysis

Analysis of variance (ANOVA) was used to establish significant differences among treatments. Duncan's multiple range tests was used to separate means where significant difference existed. Sensory attributes as judged by the consumers were subjected to principal component analysis (PCA) in order to obtain three factors that will explain at least 80% of the variability in the panellist preference for acha bread. Hierarchical cluster analysis was used to determine the clusters inherent in the data. K-means cluster analysis was performed to classify the bread samples into the inherent cluster on the basis of these attributes or parameters (IBM SPSS, 2010).

4.3 Results and Discussion

4.3.1 Effects of ingredients on acha bread loaf volume

Tables 4.3 indicates the analysis of variance for the effect of ingredients on

Table 4.3 Analysis of variance for black acha specific loaf volume

Source ¹	Sum of Squares	df	Mean Square	F Value	p-value
Model	2.01	3	0.67	6.96	0.0222
Acha starch	0.034	1	0.034	0.36	0.5728
CMC	0.96	1	0.96	9.93	0.0198
Acha starch * CMC	1.02	1	1.02	10.59	0.0174
Residual	0.58	6	0.096		
Lack of Fit	0.56	5	0.11	8.02	0.2615
Pure Error	0.014	1	0.014		
Cor Total	2.59	9			

¹CMC = carboxymethyl cellulose

Table 4.4 Analysis of variance for white acha specific loaf volume

Source	Sum of Squares	df	Mean Square	F Value	p-value
Model	0.56	2	0.28	7.75	0.0217
Acha starch	6.355E-003	1	6.355E-003	0.18	0.6884
CMC	0.52	1	0.52	14.40	0.0090
Residual	0.22	6	0.036		
Lack of Fit	0.20	5	0.040	2.41	0.4519
Pure Error	0.016	1	0.016		
Cor Total	0.77	8			

specific loaf volume (SLV) of black acha bread. Carboxyl methyl cellulose (CMC) and the interaction between it and the acha starch significantly affected the specific loaf volume. A significantly greater effect (39.4%) was from the interaction between CMC and acha starch and 36.9% from CMC. Hence, the effectiveness of the CMC depends on the level of acha starch. A 6.8% of the effect came from the yeast and 0.23% from xanthan. Table 4.4 details the analysis of variance for the effect of ingredients on white acha SLV. CMC significantly ($p < 0.05$) affected the specific loaf volume contributing to 66.9% of the overall effect. The effects from acha starch, xanthan and yeast were 0.8%, 0.01% and 4.0% respectively. Figure 4.2 details the mean of SLV for the black and white acha bread. The bread produced with 8% acha starch, 1.2% xanthan gum, 2.0% CMC and 1.0% yeast had the lowest SLV (1.4 ml/g) for both acha varieties (run 6). The bread produced with 8.0% acha starch, 2.0% xanthan gum, 2.0% CMC and 1.0% yeast had the highest specific loaf volume for both black (3.1 ml/g) and white acha (2.2 ml/g) bread (run 4). This recipe was taken as the optimum loaf recipe for acha bread. The SLV differed significantly ($p < 0.05$) among the varieties, with black acha higher in SLV.

4.3.2 Sensory properties of acha bread

Table 4.5 details the demography of the respondents. A total of 30 consumers participated in the sensory evaluation of the acha breads. Of the 30 panel members 63.3% were female and 36.7% males. In terms of occupation 13.3% were students and 86.7% were employed. Black panel members were 10%, white 13.3% and coloured 76.7%. The age group younger than 20 was 3.3%, 16.7% between 20 and 24, 13.3% between 25 and 29 years, 20% between 30 and 34 and 46.7% between 35 and 39 years of age.

The mean scores of the consumer panellist rating are shown in Figure 4.3. Multivariate analysis of variance indicated that the panellist differed significantly ($p < 0.05$) in their preference for the acha bread and wheat bread. However, the panellists could not detect any difference between the black acha bread and that from the white acha bread. There was a significant interaction between the

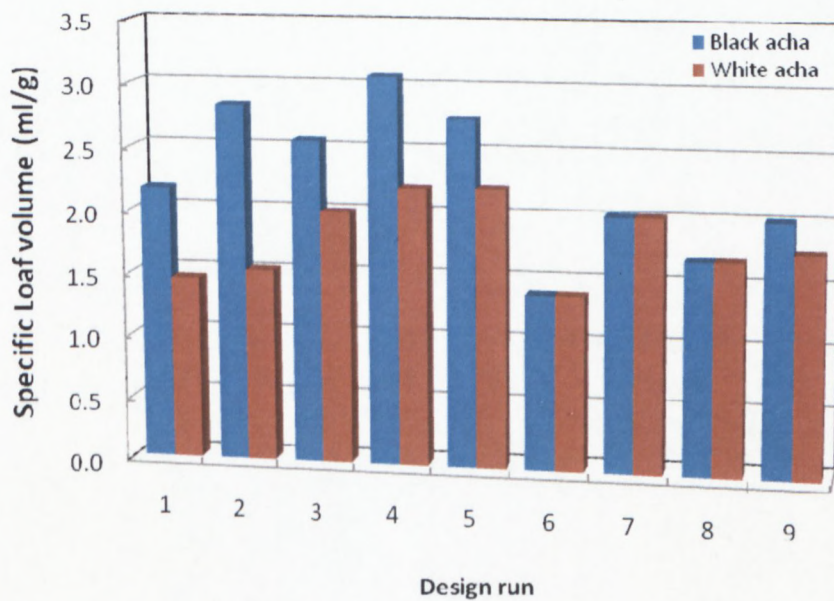


Figure 4.2 Specific loaf volume of acha bread

1 = acha starch (4%), xanthan (1.2%), CMC (1.2%), yeast (1.2%);

2 = acha starch (8%), xanthan (1.2%), CMC (1.2%), yeast (2.8%);

3 = acha starch (4%), xanthan (2%), CMC (1.2%), yeast (2.8%);

4 = acha starch (8%), xanthan (2%), CMC (2.0%), yeast (1.2%);

5 = acha starch (4%), xanthan (1.2%), CMC (2%), yeast (2.8%);

6 = acha starch (8%), xanthan (1.2%), CMC (2%), yeast (2.8%);

7 = acha starch (8%), xanthan (1.2%), CMC (2.0%), yeast (2.8%);

8 = acha starch (8%), xanthan (2%), CMC (2%), yeast (2.8%);

9 = acha starch (6%), xanthan (1.6%), CMC (1.6%), yeast (2.0%);

10 = acha starch (6%), xanthan (1.6%), CMC (1.6%), yeast (2.0%).

Table 4.5 Demography of respondents (N = 30) for acha and wheat bread consumer acceptability.

Item	Frequency (%)
Gender	
Female	19 (63.3)
Male	11 (36.7)
Occupation	
Student	4 (13.3)
Employed	26 (86.7)
Race	
Black	3 (10.0)
White	4 (13.3)
Coloured	23 (76.7)
Age	
Less than 20	1 (3.3)
20-24	5 (16.7)
25-29	4 (13.3)
30-34	6 (20)
35-39	14 (46.7)

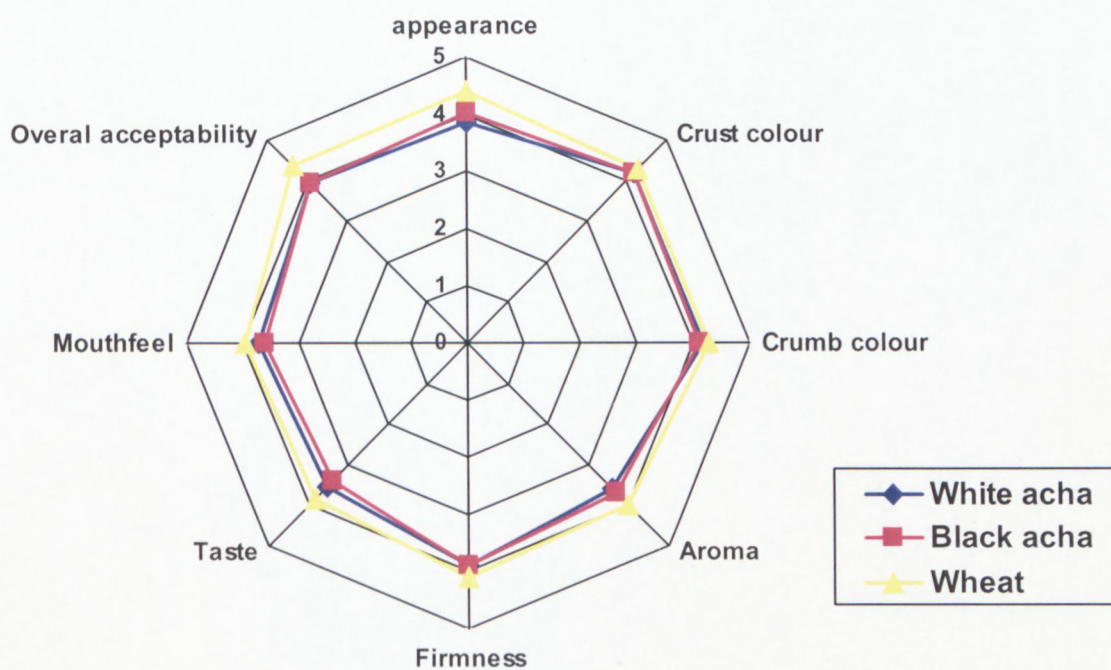


Figure 4.3 Mean sensory scores for acha bread on a 5-point hedonic scale
 1 = very undesirable; 2 = moderately undesirable; 3 = neither desirable nor undesirable; 4 = moderately desirable; 5 = very desirable



A

B

C

Figure 4.4 Bread from different grains (A) wheat bread; (B) white acha and (C) black acha bread

panellists and type of bread. This implies that the significant difference noted earlier among the panellists was in bread type as panellist score differed according to the bread. For appearance black acha scored higher than white acha bread while wheat bread scored higher than black and white acha bread. There was no difference in preference between the bread samples in crust and crumb colour. White acha bread was rated 4.2 and 4.2 respectively, for crust and crumb colour. Black acha bread 4.0 and 4.2 and wheat bread 4.3 and 4.3 respectively, for crust and crumb colour. For aroma wheat bread (4.1) was more preferred followed by black acha bread (3.7) which was more preferred than white acha bread (3.6). The acha bread and wheat bread were similar in firmness, however, they differed significantly ($p < 0.05$) in taste and mouthfeel with high preference for wheat bread. The bread samples differed significantly ($p < 0.05$) in overall acceptability, 4.4 for wheat bread and 3.9 for both acha breads. The high preference for wheat bread in some of the sensory attributes is expected as consumers are generally familiar with wheat bread. However, the high rating for acha bread indicating moderating acceptance in most of the sensory attributes as well as the similarity of acha bread to wheat bread in crust and crumb colour indicates the potential of acha bread to make it into the food basket of Africa.

Lack of differences ($p > 0.05$) between white and black acha loaves for aroma, taste, mouthfeel and overall acceptability does not support the hypotheses that there will be a significant difference in sensory parameters between black and white acha loaves.

4.3.3 Sensory attributes of interest to the consumer panellists

The suitability of the sensory data for factor analysis was assessed prior to performing principal component analysis (PCA). Inspection of the correlation matrix revealed the presence of many coefficients of 0.4 and above. The Kaiser-Meyer-Olkin value (0.898) exceeded the recommended value of 0.6. The Barlett's test of sphericity was significant ($P < 0.05$). Hence, factorability of the correlation matrix was supported. PCA revealed that the variation in the sensory

attributes of acha bread could be explained by three components with eigen values exceeding 1. Much of the variation (35.7%) in the data is explained by component 1, component 2 (32.2%) and component 3 (17.0%), with a cumulative variation of 84.9% (Table 4.6).

Varimax rotation was performed to aid in the interpretation of these three components. Table 4.6 indicates pattern or structure for coefficients obtained using varimax rotation of three factor solution for sensory attributes of acha bread. Crust colour and appearance loaded strongly on component 1, taste and mouthfeel on component 2 while aroma loaded strongly on component 3. The implication is that acha bread can be categorised into groups on the basis of crust colour, taste and aroma. K-means cluster analysis was performed to classify the bread samples into two groups (inherent in the data as indicated by a hierarchical cluster analysis) on the basis of these attributes or parameters.

Cluster 1 was the case for consumers who neither desired nor undesired (3) acha bread in crust colour and the samples were moderately undesirable in taste and aroma (17.5%, 21 out of 120 cases). Cluster 2 was the case where the bread samples were moderately desirable in crust colour, taste and aroma (82.5%, 99 out of 120 cases). This implies that most consumers find acha bread to have the potential to be marketed amongst other types of bread. It also implies that with small improvements in the taste and aroma the remaining 17.5% of consumers will switch to desirable.

Majority of the consumers in all age group moderately desired acha bread in crust colour, taste and aroma (Figure 4.5). The potential of acceptance for acha bread is high across all ages. This implies that acha bread has the potential to be consumed as an alternative to other wheat or non-wheat bread across all age groups. Majority of the cases for males (92.3%) and females (76.3%) were in cluster 2- moderately desirable in crust colour, taste and aroma (82.5% cases) (Table 4.7). Majority (53.7%) of the black panel members were in cluster 1, implying that they neither desired nor undesired (3) acha bread in crust colour (Table 4.8). Majority of the cases for white (81.3%) and coloured (88%)

Table 4.6 Pattern/structure for coefficients obtained using varimax rotation of three factor solution for sensory attributes of acha bread

	Component*		
	1	2	3
Appearance	.415	-.122	-.115
Crust colour	.512	-.198	-.179
Crumb colour	.308	-.044	-.056
Aroma	-.222	-.272	1.111
Firmness	.135	-.134	.387
Taste	-.235	.572	-.132
Mouth feel	-.133	.523	-.192
Overall acceptability	.047	.364	-.219
% of variance explained	35.7	32.182	17.0

*Extracted method: Principle Component Analysis. Rotation Method: Varimax with Kaiser Normalization

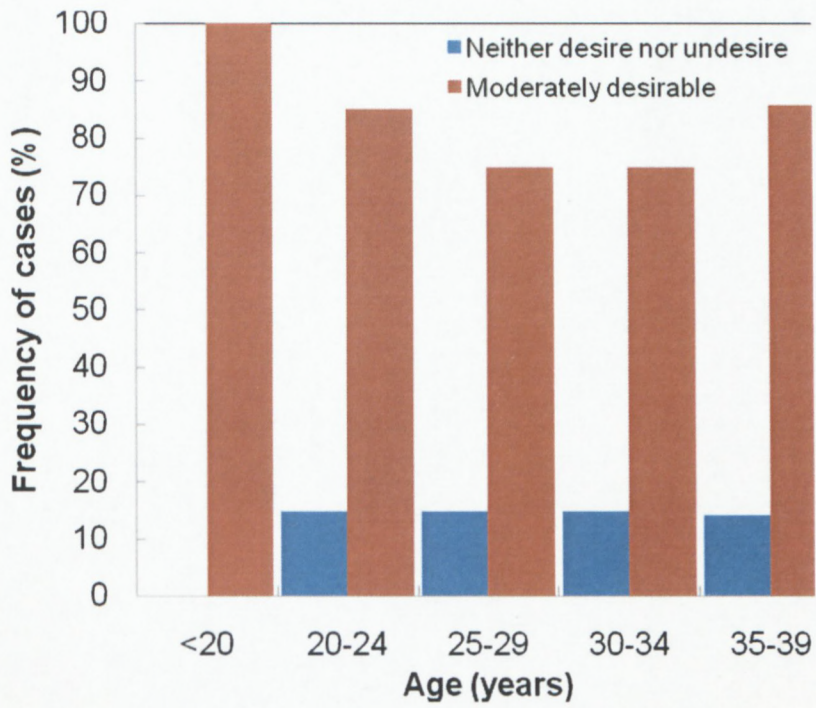


Figure 4.5. Cluster count per age group

Table 4.7 Gender * Cluster Number of Case Cross tabulation

Gender	Cluster Number of Case*		
	1	2	Total
Female	18(23.7%)	58(76.3%)	76(100%)
Male	3(6.8%)	41(93.2%)	44(100%)
Total	21(17.5%)	99(82.5%)	120(100%)

*Cluster 1 = consumers who neither desired nor undesired (3) acha bread in crust colour and who moderately undesired acha bread in taste and aroma; Cluster 2 = consumers who moderately desired the acha bread in crust colour, taste and aroma (82.5%, 99 out of 120 cases).

Table 4.8 Race * Cluster Number of Case Cross tabulation

Race	Cluster Number of Case*		
	1	2	Total
Black	7(58.3%)	5(41.7%)	12(100%)
White	3(18.8%)	13(81.3%)	16(100%)
Coloured	11(12%)	81(88%)	92(100%)
Total	21(17.5%)	99(82.5%)	120(100%)

*Cluster 1 = consumers who neither desired nor undesired (3) acha bread in crust colour and who moderately undesired acha bread in taste and aroma; Cluster 2 = is the case where the consumers who bread samples were moderately desired the acha bread in crust colour, taste and aroma (82.5%, 99 out of 120 cases).

panel members were in cluster 2- the case where the bread samples were moderately desirable in crust colour, taste and aroma. Acha bread has the potential to serve a niche market amongst white and coloured consumers.

Majority of the cases with regards to variety [black acha (83.3%), white acha (81.7%)] belong to cluster 2 indicating moderately desirable in crust colour, taste and aroma (Table 4.09). Hence, either of the acha varieties could be used in the production of acha bread. Acha bread thus has the potential to be consumed as alternative wheat free bread. The overall acceptability ratings for all consumer quality characteristics were the same for both white and black acha bread. The objective to produce black and white acha bread that is acceptable to the consumer and thus be used as a wheat free bread for individuals suffering from diabetes and celiac disease have been met.

Table 4.09 Acha varieties * Cluster Number of Case Cross tabulation

Acha variety	Cluster Number of Case*		
	1	2	Total
White	10(16.7%)	50(83.3%)	60(100%)
Black	11(18.3%)	49(81.7%)	60(100%)
Total	21(17.5%)	99(82.5%)	120(100%)

*Cluster 1 = consumers who neither desired nor undesired (3) acha bread in crust colour and who moderately undesired acha bread in taste and aroma; Cluster 2 = is the case where the consumers who moderately desired the acha bread in crust colour, taste and aroma (82.5%, 99 out of 120 cases).

4.4 Conclusion

Optimum acha bread was baked using 8.0 % acha starch, 2% xanthan gum, 2.0% CMC and 1% yeast. Majority of the panellist found the crust colour, taste and aroma to be moderately desirable. This implies that acha bread have the potential to be marketed as wheat free bread. Male and female as well as consumers of all ages found acha bread to be moderately desirable in crust colour, taste and aroma.

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CHAPTER 5

EFFECT OF PROCESSING ON THE STARCH AND GLYCEMIC PROPERTIES OF ACHA

Abstract

The effect of baking, steaming, boiling and microwaving on the starch and glyceemic properties of two acha cultivars (*Digitaria exilis*, white and *Digitaria iburua*, black) is reported. Processing method, sample and incubation time significantly ($p < 0.05$) affected starch hydrolysis. However, the processing times did not affect the starch hydrolysis significantly. Significantly ($p < 0.05$) more starch was released from baked products (49.4%) compared to steaming (7.2%), microwaving (10.7%) and boiling (19.4%). Significantly more starch was released from black acha (19.8%) compared to white acha (16.3%) and wheat (12.4%). The amount of starch hydrolysed for the different processing methods was in the following order: baking > boiling > microwaving > steaming. For baking, the wheat and both acha bread samples did not differ significantly in their response to hydrolysis. On the effect of microwaving on starch hydrolysis samples, incubation time, interaction between sample and incubation time affected starch hydrolysis significantly ($p < 0.05$) of wheat and black and white acha. During boiling, processing time, cereals, incubation time, interaction between processing time * incubation time and sample * incubation time significantly influenced the hydrolysed starch. The cereals differed significantly in the amount of starch released with high starch (28.1%) for white acha compared to black acha (23.8%) and wheat (6.3%). During steaming the processing time, cereals, incubation time as well as the two-way and three-way interaction all affected starch hydrolysis significantly ($p > 0.05$). Slowly digestible starch (SDS) and rapidly digestible starch (RDS) analysis results indicate that black acha grain (*Digitaria iburua*) cultivar is the better cultivar to use than white acha grain (*Digitaria exilis*) since it consisted of higher slowly digestible starch (SDS) and lower rapidly digestible starch content after baking, microwaving, boiling and steaming. Whereas white wheat bread, microwaved, boiled and steamed wheat

grain had a higher RDS and SDS compared to black acha. Acha white bread and processed (boiled, steamed and microwaved) acha and wheat grain consisted of low GI values and is thus suitable as a food supplement or meal replacer for those individuals suffering from diabetes and celiac disease.

5.1 Introduction

Digitaria exilis (white) and *Digitaria iburua* (black) are two species of acha grains. Other common names are fonio or hungry rice. It is indigenous to most African countries particularly regions of Mali, Burkina Faso, Guinea and Nigeria (Ayo, 2003) and the Dominican Republic (Morales-Payán *et al.*, 2003; Jideani *et al.*, 2008). Starchy foods like acha are not usually eaten raw and must undergo one type of heat processing or the other for palatability and bioavailability. Different processing methods include boiling, microwave, baking, drying, extrusion, steaming, drum-drying, popping, pressure-cooking and others. Therefore, processing methods may modify starch in various ways, consequently affecting digestion and nutritional value. De Lumen *et al.* (1993) reported acha as a promising underutilised African cereal because of its high content of sulphur-amino acids. Acha is recommended as a cereal suitable for the management of diabetes mellitus in West Africa (Jideani *et al.*, 1994). However, not much is known about the starch and glycemic properties of this grain. It is thus of interest to investigate the effect of processing specifically baking, boiling, steaming and microwaving on the starch and glycemic properties of acha in order to ascertain its potential in low glycemic food products. The outline of the investigation can be seen in Figure 5.1.

5.2 Materials and Methods

5.2.1 Source of acha grain and materials

Two acha cultivars *Digitaria exilis* (white) and *Digitaria exilis* (black) were purchased from Grace Africa, Salt River, Cape Town, South Africa.

All equipment and materials were either obtained from the Department of Food Technology Cape Peninsula University of Technology (CPUT) or Pioneer

foods laboratories (Bokomo Foods pilot plant and SASKO technical laboratory), Cape Town, South Africa.

All chemicals were obtained from the Department of Food Technology CPUT and Laboratory and Scientific Ltd. in Maitland, Cape Town, Republic of South Africa.

5.2.2 Cleaning of acha grain

The cleaning procedure entailed screening the grains through a 1000, 750, 500 and 125 micron screens to remove all foreign matter such as stones, foreign grain and small sticks. The process of dry cleaning was then followed by washing three times by rinsing with water and drying at 40°C for 48 h (Jideani & Podgorski, 2010). The cleaned and dried samples were then placed in sealed polyethylene bags and stored in a refrigerator at a temperature of 3 to 5°C until required.

5.2.3 Steaming of acha

Cleaned and dried acha grains (20 g) were rinsed in 500 ml distilled water. The water was drained through a sieve for 5 min. The drained grain was steamed using a steamer and three samples were drawn at 10 min interval for 15 to 35 minutes. The samples were cooled immediately on crushed ice for 10 min and thereafter were assayed for SDS, RDS, RS, TS and glycemic index as outlined in section 5.2.6 to 5.2.9

5.2.4 Microwaving of acha

Cleaned and dried acha grains (20 g) in 250 ml distilled water were processed by microwaving at medium for 3, 6 and 9 minutes, respectively. The samples were cooled immediately on crushed ice for 10 min and thereafter were assayed for SDS, RDS, RS, TS and glycemic index. See section 5.2.6 to 5.2.9.

5.2.5. Production of acha bread

The method of Jideani *et al.* (2007) was modified and used to produce acha

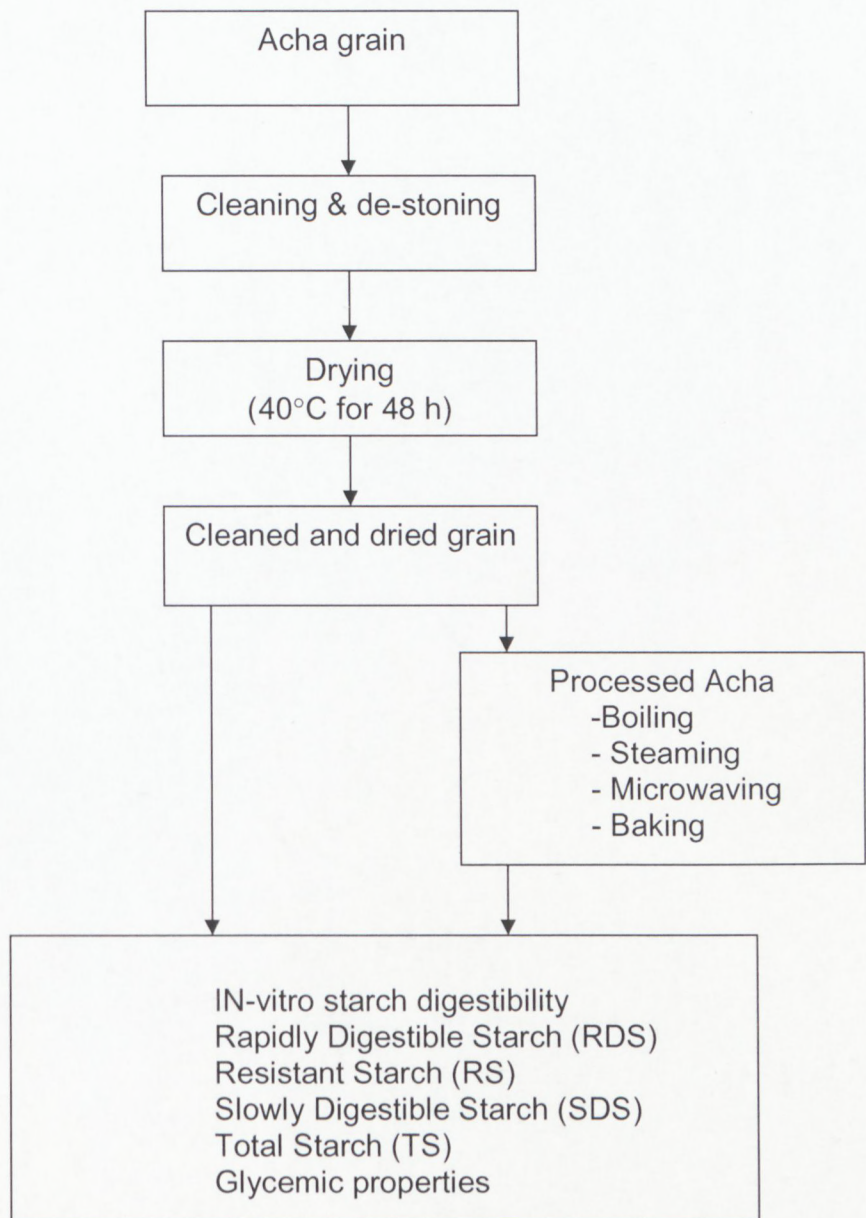


Figure 5.1 Experimental outlines for the effect of processing on acha

bread as described in Chapter 4. Bread samples (50 mg) were assayed for SDS, RDS, RS, TS and glycemic index following the method described in section 5.2.6 – 5.2.9 of this chapter.

5.2.6 Analysis of total starch of processed acha and wheat products

Total starch was determined enzymatically according to the modified method of Goñi *et al.* (1997). Triplicate samples of both processed acha and wheat products were analyzed for total starch. 50 mg of processed sample was dispersed in 50 ml centrifuge tubes with 6 ml of KOH. The samples were shaken for 30 min at room temperature. The samples were then hydrolyzed by the addition of 3 ml 0.4M Sodium acetate buffer (pH 4.8) and 60 µl of amyloglucosidase (A7420 Sigma Aldrich) from *Aspergillus niger* (67.4 U/mg 50 mg AMG in 50 ml buffer solution). The samples were transferred into Erlenmeyer flasks and then incubated at 60°C for 45 min in a shaking water bath. The samples were then centrifuged for 10 min at 3000 rpm. After centrifugation, the glucose concentration was determined in the supernatant using a glucose oxidase-peroxidase (GAGO-20 Sigma Aldrich) kit. 1 ml sample aliquots of the supernatant were measured out into test tubes and to that 2 ml of the assay reagent was added. The samples were then incubated at 37°C for 30 min, 2 ml H₂SO₄, were carefully added to the test tubes after incubation and were then thoroughly mixed. Colour absorption was measured against a reagent black at a wavelength of 540 nm and the glucose concentration was converted into starch by applying the factor 0.9.

5.2.7 Resistant starch analysis of processed acha and wheat products

RS content in acha grain bread and wheat grain and bread were determined according to the method of Gõni *et al.* (1997). Dry grain and processed grain samples (100 mg) was weighed into a 50 ml centrifuge tube. An aliquot (10 ml) of KCL-HCL buffer (pH 1.5) and 0.2 ml of pepsin (P7000 Sigma-Aldricht) solution (1 g pepsin /10 ml buffer KCL-HCL) were added. The solution was mixed well and was placed in a shaking water bath for 60 min at 40° C. After cooling

samples to room temperature, 9 ml of 0.1M Tris- maleate buffer (pH 6.9) was added and was followed by adding 1 ml of α -amylase (A3176 Sigma-Aldrich) solution (40 mg α -amylase / ml Tris-maleate buffer). The samples were then mixed well and incubated for 16 h in a water bath at 37°C with constant shaking. The samples were centrifuged for 15 min at 3000 rpm, and the supernatants were discarded. The residues were moistened with 3 ml distilled water and 3 ml of 4 M KOH then mixed well for 30 min at room temperature with constant shaking. Thereafter, 5.5 ml of 2 M HCL and 3 ml of 0.4 M sodium acetate buffer (pH 4.75) were added, followed by the addition of 80 μ l of amyglolucosidase. The solution was then mixed well and was placed in a water bath at 60°C for 45 min with constant shaking. Thereafter the solution was centrifuged at 3000 rpm for 15 min and the supernatants were collected and saved in a 50 ml volumetric flask. 1 ml of the sample was measured into test tubes and to that 2 ml of the assay reagent added. The samples were then incubated at 37°C for 30 min. Sulphuric acid (2 ml) was carefully added to the test tubes after incubation and were then thoroughly mixed. Colour absorption was measured against the reagent blank at a wavelength of 540 nm and the glucose concentration was converted into starch content by multiplying by the factor 0.9.

5.2.8 Digestible starch of processed acha and wheat products

Digestible starch (DS) was calculated as the difference between TS and RS (Goñi *et al.*, 1997).

5.2.9 Determination of rapidly and slowly digestible starch by *In vitro* rate of starch digestion

The procedure and model established by Goñi *et al.* (1997) was used to measure the *in vitro* starch hydrolysis. Triplicate samples of 50 mg of unprocessed grain, processed grain and acha and wheat bread were dispersed in 10 ml of KCL-HCL buffer. 0.2 ml pepsin solution containing 1 g of pepsin in 10 ml HCL-KCL buffer (pH 1.5) was added to the samples. The samples were incubated at 40°C for 60

min in a shaking water bath. The volume was raised to 25 ml by adding 15 ml Tris-maleate buffer (pH 6.9) and adjusting the pH carefully. Starch hydrolysis was started by adding to each sample another 5 ml of Tris- maleate buffer containing 2.6 UI α -amylase.

The flasks were placed in a shaking water bath at 37°C with moderate agitation. Aliquot samples (1 ml) were taken from each flask every 30 min for 3 h. These aliquots were placed in a test tube at 100°C and were energetically shaken for 5 min to inactivate the α -amylase. The 2 ml of 0.4 M sodium acetate buffer (pH 4.75) was added to each aliquot and 60 μ l of amyloglucosidase was used to hydrolyze the digested starch into glucose after 45 min at 60°C in a shaking water bath. The volume was adjusted to 5 ml with distilled water and 0.5 ml was incubated with the glucose oxidase-peroxidase kit. The samples were analyzed in triplicate. The glucose was finally converted into starch by multiplying with 0.9. The rate of starch digestion was expressed as percentage of starch hydrolyzed at different times (30, 60, 90, 120, 150 and 180 min). RDS and SDS were determined according to Englyst *et al.* (1992). The RDS was defined as the percentage of starch digested at 30 min and the SDS as the percentage of starch digested at 180 min. The digestion curve was modeled with the non-linear equation (Eq1) established by Goñi *et al.* (1997) to describe the kinetics of starch hydrolysis.

$$C = C_{\infty} (1 - e^{-kt}) \quad \dots\dots\dots (1)$$

where C is the percentage of starch hydrolyzed at time t (min); C_{∞} is the equilibrium percentage of starch hydrolyzed after 180 min, and k is the kinetic constant. The parameters C_{∞} and k were estimated for each sample using IBM SPSS for Windows 19.0 non-linear regression.

5.3. Statistical analysis

Analysis of variance (ANOVA) was used to determine differences among treatments. Duncan's multiple range tests was used to separate means where significant difference existed (IBM SPSS, 2010).

5.4 Results and Discussion

5.4.1 Effect of processing on starch hydrolysis

Table 5.1 indicates the effect of incubation time on the hydrolysed starch from baking, microwave, boiling and steaming of acha and wheat. Processing method, sample and incubation time significantly ($p < 0.05$) affected starch hydrolysis. However, the processing times did not affect the starch hydrolysis significantly. Significant interaction effect exists between processing * processing time, processing time * incubation time, processing * sample, processing * incubation time, sample * incubation time, processing * processing time * sample, processing * sample * incubation time, and processing * processing time * sample * incubation time.

There was a significant difference ($p < 0.05$) in the amount of starch released during baking, steaming, microwaving and boiling. More starch was released from baked products (49.4%) compared to steaming (7.2%), microwaving (10.7%) and boiling (19.4%) from wheat and both acha variants. Steaming resulted in significantly ($p < 0.05$) less hydrolysed starch compared to the others. Significantly more starch was released from black acha (19.8%) compared to white acha (16.3%) and wheat (12.4%). This variation is in line with other investigations. Eyaru *et al.* (2009) reported that starch digestibility of three legumes varied with the different processing techniques, pressure cooking raw grain, boiling raw grain, pressure cooking soaked grain, and boiling soaked grain. Increase in incubation time of acha and wheat grain resulted to increase in hydrolysed starch. Lee *et al.* (2005) reported that the hydrolysis rate of corn samples treated with microwaves were marginally lower than other samples treated with other cooking methods. There was a correlation between changes in starch hydrolysis rates and the differences in disruption of the crystalline regions due to the various processing methods. Different processing methods affects the microstructure of the samples which thus influences their physical properties and starch hydrolysis (Lee *et al.*, 2005). Effect of baking on starch hydrolysis is depicted in Figure 5.2 and Table 5.2. The bread samples did not differ

Table 5.1 Analysis of variance for the effect of processing, processing time, incubation time and sample on starch hydrolysis of acha and wheat

Source	Sum of Squares	Df	Mean Square	F	Sig.
Corrected Model	249448.8 ^a	208	1199.3	13.4	0.000
Intercept	205816.2	1	205816.2	2292.8	0.000
Processing	13033.3	1	13033.3	145.2	0.000
Processing time	561.5	4	140.4	1.6	0.183
Sample	4160.6	2	2080.3	23.2	0.000
Incubation time	57118.5	6	9519.6	106.1	0.000
Processing * Processing time	804.4	2	402.1	4.5	0.012
Processing time * sample	1180.7	8	147.6	1.6	0.110
Processing time * Incubation time	3306.98	24	137.7	1.5	0.052
Processing * sample	6856.0	2	3428.0	38.2	0.000
Processing * Incubation time	12505.3	6	2084.2	23.2	0.000
Sample * Incubation time	4263.9	12	355.3	3.9	0.000
Processing * Processing time * sample	348.6	4	87.1	1.0	0.423
Processing * Processing time * Incubation time	2308.1	12	192.3	2.1	0.014
Processing time * sample * Incubation time	5632.0	48	117.3	1.3	0.090
Processing * sample * Incubation time	4879.5	12	406.6	4.5	0.000
Processing * Processing time * sample * Incubation time	5365.3	23	233.3	2.6	0.000
Error	37432.3	417	89.8		
Total	450359.9	626			
Corrected Total	286881.1	625			

^aR Squared = 0.870 (Adjusted R Squared = 0.804)

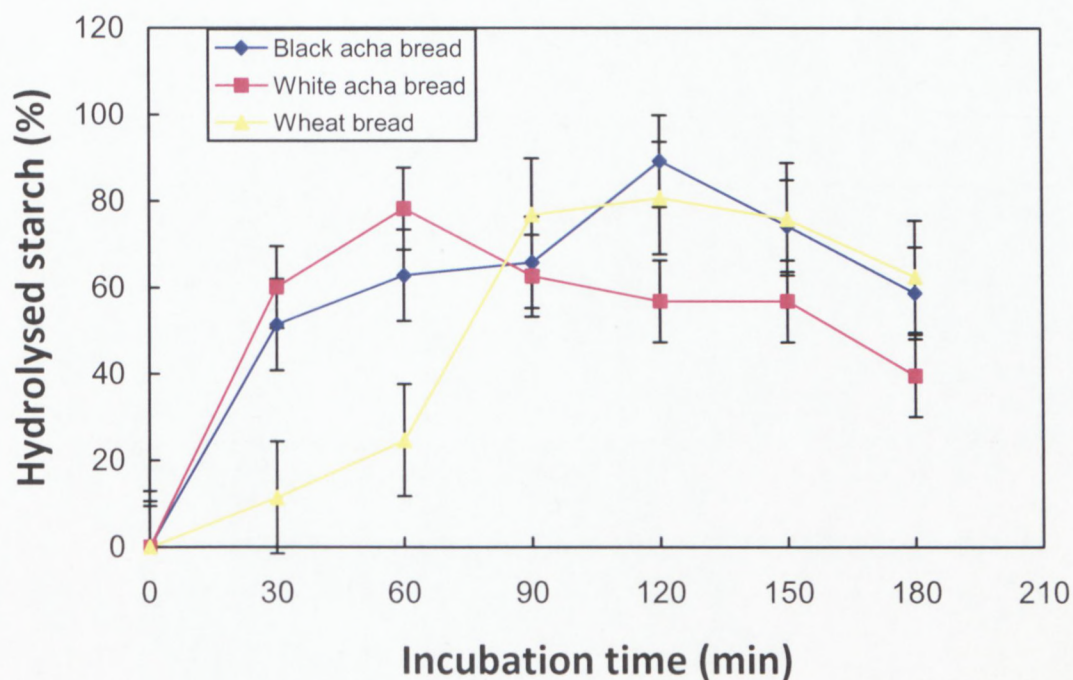


Figure 5.2 Effect of incubation time on starch hydrolysis from acha and wheat bread

Table 5.2 Tests of between-subjects effects of baking on starch hydrolysis from acha and wheat bread

Source	Type III Sum of Squares	df	Mean Square	F	Sig.
Corrected Model	50724.7 ^a	20	2536.2	8.5	0.000
Intercept	153926.5	1	153926.5	517.8	0.000
sample	131.5	2	65.8	0.221	0.803
Time Incubation	32709.6	6	5451.6	18.4	0.000
sample * time	17883.6	12	1490.3	5.0	0.000
Incubation					
Error	12486.1	42	297.3		
Total	217137.3	63			
Corrected Total	63210.8	62			

^aR² = .802 (Adjusted R Squared = 0.708)

significantly in their response to hydrolysis. This is in agreement with the report of Sayed et al. (2008) indicating that the differences in starch hydrolysis of spelt and wheat grain disappeared after baking. However, incubation time and interaction between the samples and the incubation time had significant effect on the starch hydrolysis. Hydrolysis of starch increased with increase in incubation time peaking at 120 min. Bread normally has an open cell structure where the enzyme solution is exposed to a greater protein-starch complex surface area (Van der Merwe *et al.*, 2001). This means that there is a higher accessibility of starch to enzymatic action and would thus lead to a higher starch digestibility rate in the bread than within the rest of the processed (steaming, microwaved and boiled) samples. Goni *et al.* (1996) found 76.1% starch digestion for wheat bread after 180 minutes, this however is still higher than the results in this study 49.4%.

Effect of microwave on starch hydrolysis is detailed in Figure 5.3 and Table 5.3. Samples, incubation time and interaction between sample and incubation time significantly ($p < 0.05$) affected starch hydrolysis. Significantly more starch was released from black acha (14.8%) and wheat (13.9%) compared to white acha (3.4%). There was no significant difference between the starch released from wheat and that of black acha. Significantly more starch was released after 90, 120 and 150 min of incubation. Processing time did not have significant effect on starch hydrolysis.

Effect of boiling on starch hydrolysis is detailed in Figure 5.4 and Table 5.4. Processing time, cereals, incubation time, interaction between processing time * incubation time and sample * incubation time significantly influenced the hydrolysed starch. Boiling for 15 min resulted in significant amount (22.7%) of hydrolysed starch. A significant decrease in starch hydrolysis resulted when boiled for 25 min (16.7%) and 35 min (18.9%). The cereals differed significantly in the amount of starch released with higher starch (28.1%) for white acha compared to black acha (23.8%) and wheat (6.3%). Significantly more starch was release after 90, 120 and 150 min of incubation.

Effect of steaming on starch hydrolysis is detailed in Figure 5.5 and Table 5.5. Processing time, cereals, incubation time as well as the two-way and three-

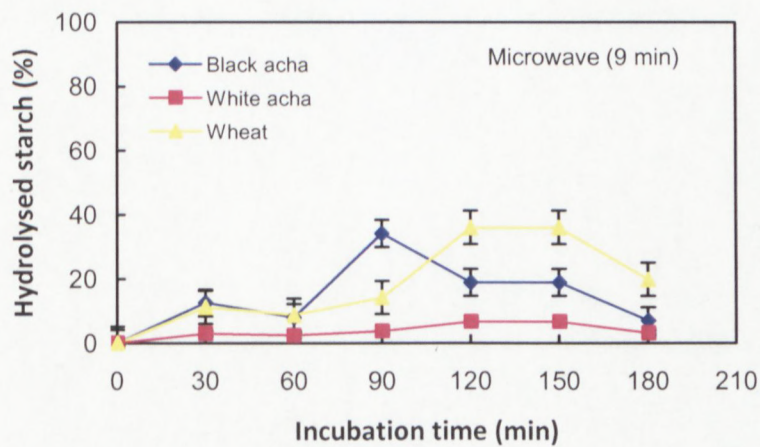
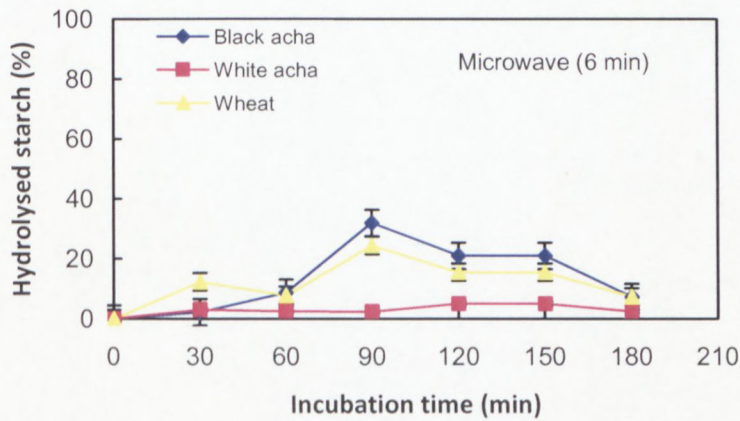
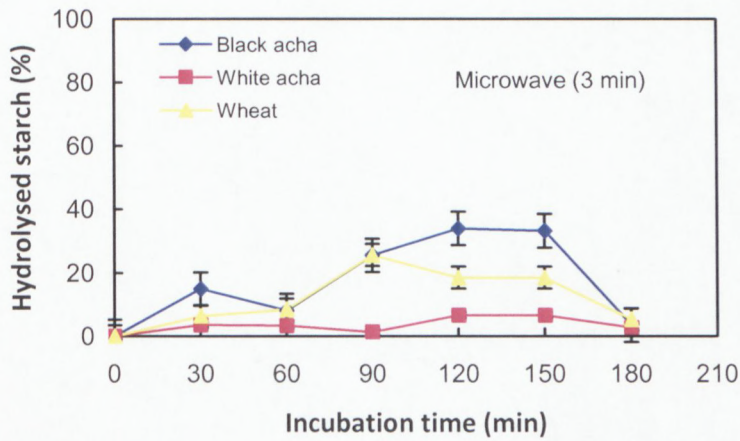


Figure 5.3. Effect of incubation time on starch hydrolysis from microwaved acha and wheat

Table 5.3. Analysis of variance for effect of microwave on starch hydrolysis

Source	Type III Sum of Squares	df	Mean Square	F	Sig.
Corrected Model	20248.1 ^a	62	326.6	4.7	0.000
Intercept	21616.5	1	21616.5	312.8	0.000
Processing time	230.1	2	115.0	1.7	0.193
Sample	5092.5	2	2546.2	36.8	0.000
Incubation time	8664.3	6	1444.0	20.9	0.000
Processing time * sample	488.3	4	122.1	1.8	0.140
Processing time * Incubation time	504.0	12	42.0	0.6	0.832
sample * Incubation time	3140.3	12	261.7	3.8	0.000
Processing time * sample * Incubation time	2128.5	24	88.7	1.3	0.189
Error	8707.4	126	69.1		
Total	50572.0	189			
Corrected Total	28955.5	188			

^aR² = 0.699 (Adjusted R² = 0.551)

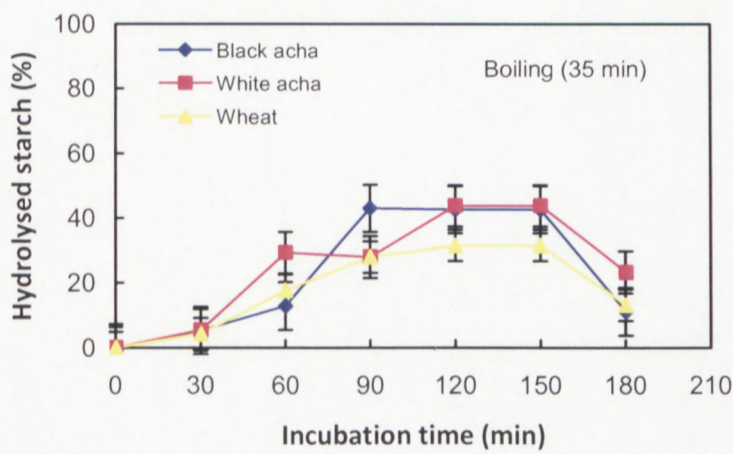
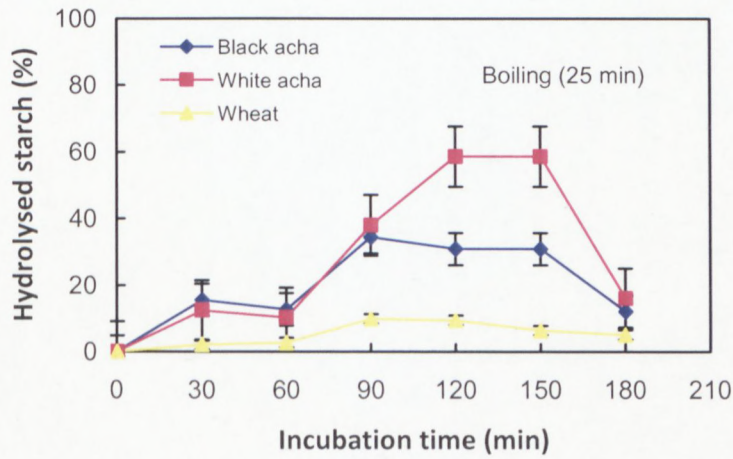
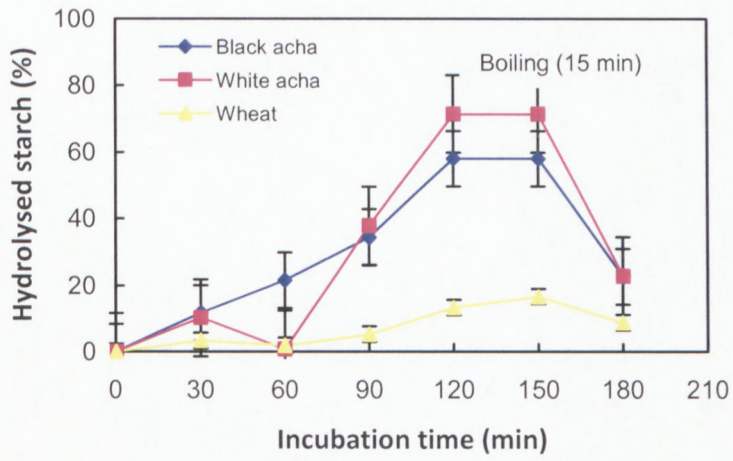


Figure 5.4. Effect of incubation time on starch hydrolysis from boiled acha and wheat

Table 5.4. Analysis of variance for the effect of boiling on starch hydrolysis of acha and wheat

Source	Type III Sum of Squares	df	Mean Square	F	Sig.
Corrected Model	70069.5 ^a	61	1148.7	12.0	0.000
Intercept	69233.8	1	69233.8	724.4	0.000
Processing time	925.0	2	462.5	4.8	0.009
Sample	16134.7	2	8067.4	84.4	0.000
Incubation time	34410.4	6	5735.1	60.0	0.000
Processing time * sample	603.9	4	151.0	1.6	0.184
Processing time * Incubation time	3100.9	12	258.4	2.7	0.003
Sample * Incubation time	11155.9	12	929.6	9.7	0.000
Processing time * sample * incubation time	2543.8	23	110.6	1.2	0.296
Error	12041.8	126	95.6		
Total	152743.0	188			
Corrected Total	82111.3	187			

^aR² = 0.853 (Adjusted R² = 0.782)

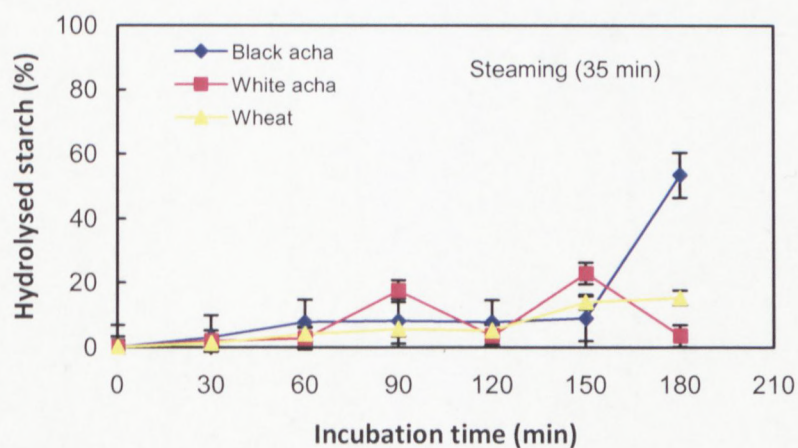
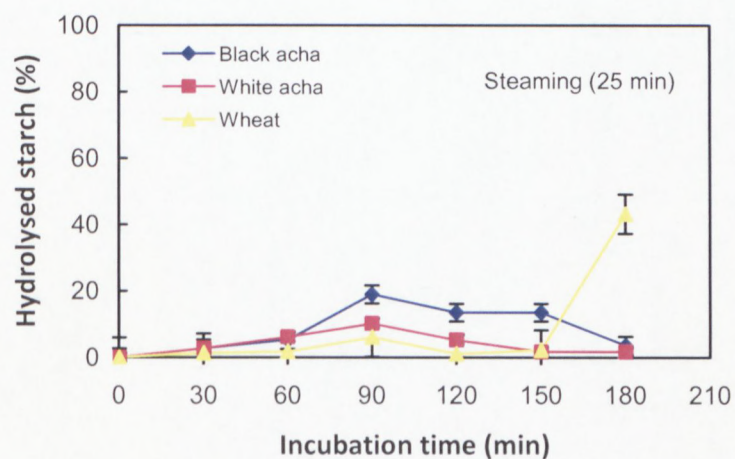
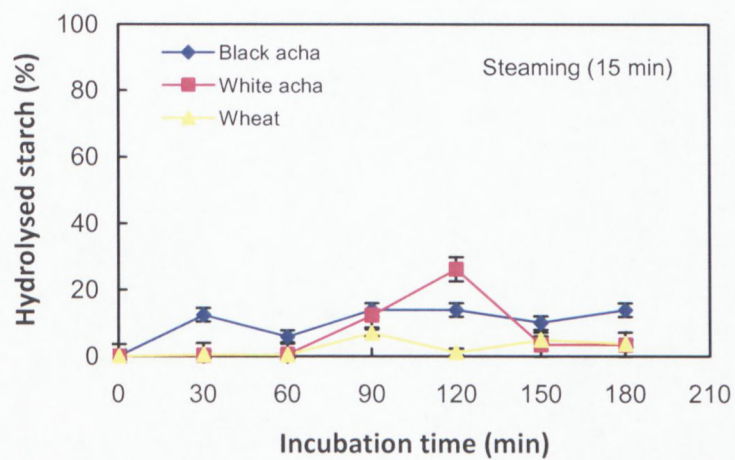


Figure 5.5. Effect of incubation time on starch hydrolysis from steamed acha and wheat

Table 5.5 Analysis of variance for the effect of steaming on starch hydrolysed from acha and wheat

Source	Type III Sum of Squares	df	Mean Square	F	Sig.
Corrected Model	16101.8 ^a	62	259.7	7.6	0.000
Intercept	9972.7	1	9972.7	292.3	0.000
Processing time	222.3	2	111.2	3.3	0.042
Sample	820.3	2	410.1	12.0	0.000
Incubation time	4585.8	6	764.3	22.4	0.000
Processing time * sample	431.5	4	107.9	3.2	0.016
Processing time * Incubation time	2002.2	12	166.9	4.9	0.000
Sample * Incubation time	2471.4	12	206.0	6.0	0.000
Processing time * sample * Incubation time	6039.4	24	251.6	7.4	0.000
Error	4197.0	123	34.1		
Total	29907.6	186			
Corrected Total	20298.8	185			

^aR² = 0.793 (Adjusted R² = 0.689)

way interaction all affected starch hydrolysis significantly ($p > 0.05$). There was no significant difference in the amount of starch released after 15 min (6.2%) and 25 min (6.4 %) of steaming. However, steaming for 35 min (8.9%) significantly increased the amount of hydrolysed starch. The increase in hydrolysed starch with increase in steaming time could be explained by the disruption of complexes such as starch-lipid complexes, which renders the weaker bonds to hydrolysis (Jideani & Scott, 2011). Wheat and white acha did not differ significantly in the amount of starch released during steaming. However, black acha released higher amount of starch (10.4%) compared to white acha and wheat. Grain sizes may have also contributed to the different amount of starch hydrolysed. Given that the acha grains are very small compared to wheat, they may be subjected to enzyme digestion faster thereby resulting in comparatively more starch hydrolysed. Incubation time increased significantly the amount of starch released with more starch released from 90 to 180 min of steaming.

The different processing methods, microwaving, boiling and steaming as well as the different processing times, affects the structure of the cereals and thus influenced the amount of starch hydrolysed in the different cereal samples. Van der Merwe *et al.* (2001) reported that different cooking methods of bread and cereal porridge and the way that cooking affects the structure of the food could greatly contribute to the differences in starch hydrolysis. Shorter processing time will lead to less disrupted starch granules and therefore less prone to enzyme digestion, whereas longer processing time would lead to increased starch granule disruption and increase in the formation of retrograded amylose during the cooling period.

5.4.2 *In vitro* starch hydrolysis rate

Modelling the effect of baking on *in vitro* acha starch hydrolysis

Table 5.6 and Figure 5.6 detail the model parameters for bread prepared from black acha, white acha and wheat. The equilibrium percentage of starch hydrolysed after 180 min did not differ significantly ($p > 0.05$) among the cereals. However, the rate of starch hydrolysis (k) differed significantly ($p < 0.05$) among

Table 5.6 Model parameters for bread from black acha, white acha and wheat bread^{1,2}

Bread	C_{∞}	k	R^2
Black acha	77.5 ± 5.7^a	0.039 ± 0.015^a	0.713
White acha	59.1 ± 5.5^a	1.394 ± 0.015^b	0.461
Wheat	91.4 ± 21.7^a	0.011 ± 0.005^a	0.763

¹ C_{∞} = equilibrium percentage of starch hydrolyzed after 180 min; k = kinetic constant; $R^2 = 1 - (\text{Residual Sum of Squares}) / (\text{Corrected Sum of Squares})$

²Values with the different superscript in a column are significantly different ($p < 0.05$).

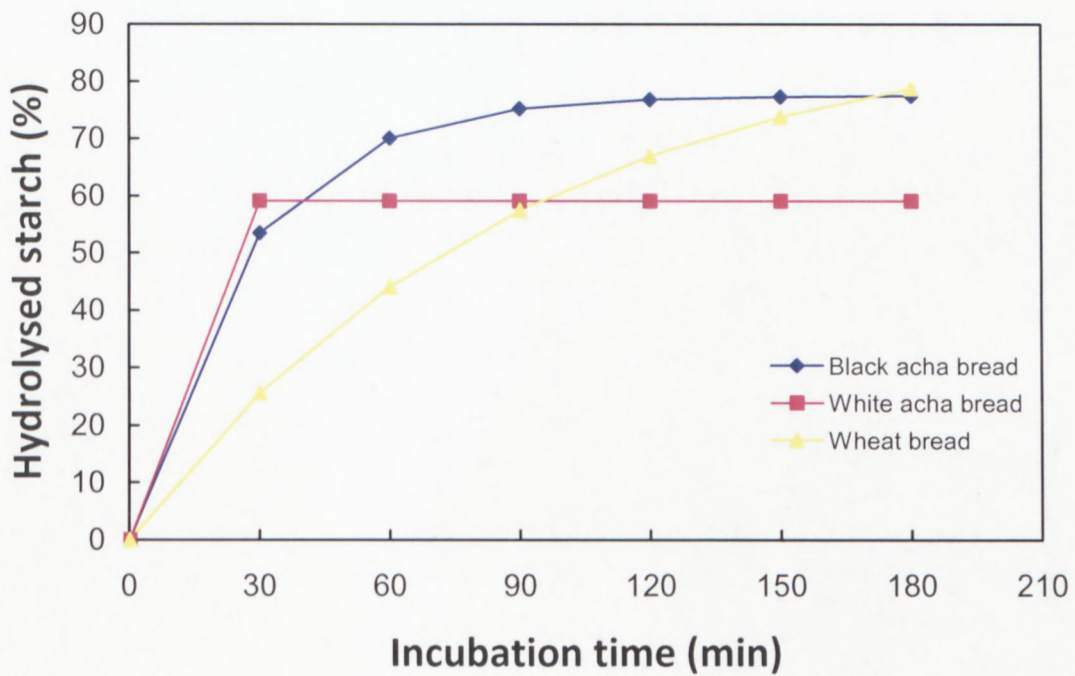


Figure 5.6. Predicted hydrolysed starch from black acha, white acha and wheat bread

the cereals. White acha showed a higher rate of hydrolysis compared to black acha and wheat. Wheat and black acha did not differ significantly in their rate of the cereals. White acha showed a higher rate of hydrolysis compared to black acha and wheat. Wheat and black acha did not differ significantly in their rate of hydrolysis. Sayed *et al.* (2008) reported a similar difference in starch hydrolysis of spelt whole grain bread and common wheat bread. The same results were reported by Skrabanja *et al.* (2001); spelt white bread had a significantly higher starch hydrolysis index than common wheat white bread. These differences in starch hydrolysis rate and extent could be attributed to difference in the chemical and physicochemical properties due to the genetic differences. Although there was no significant difference, the equilibrium percentage of starch hydrolyzed from black acha and wheat bread were higher compared to white acha bread. This does not support the hypothesis that white and black acha bread will differ in the equilibrium percentage of starch hydrolyzed after 180 min.

5.4.3 Modelling the effect of boiling on acha starch hydrolysis

The model parameters for black acha, white acha and wheat grain subjected to different boiling times are detailed in Table 5.7 and Figure 5.7. The equilibrium percentage of starch hydrolysed after 180 min did not differ significantly among the cereals for the same boiling time but differed significantly ($p < 0.05$) amongst the different boiling times (15, 25, 35 min). This difference in the equilibrium starch hydrolysis can be attributed to the disruption of the crystalline regions due to the various boiling times. Lee *et al.* (2005) reported similar results for different corn samples subjected to various processing methods and times. A positive correlation could be drawn for the changes in the starch hydrolysis and the different cooking times (Lee *et al.*, 2005). The equilibrium percentage of starch hydrolyzed for cereals boiled for 15 min was the highest for white acha 62.2% followed by black acha 48.1% and the lowest for wheat at 7.7%. The difference in the equilibrium percentage of starch hydrolyzed after 180 min for the different cereal samples could possibly be due to genetic and grain sizes differences.

Table 5.7 Model parameters for black acha, white acha and wheat grain as affected by boiling^{1,2}

Cereal	Boiling time (min)	C_{∞}	k	R^2
Black acha	15	48.1 ± 10.5 ^a	0.015 ± 0.009 ^a	0.603
White acha	15	62.2 ± 31.9 ^a	0.011 ± 0.011 ^a	0.448
Wheat	15	27.1 ± 40.1 ^a	0.004 ± 0.007 ^a	0.535
Black acha	25	25.9 ± 4.0 ^b	0.029 ± 0.019 ^b	0.431
White acha	25	46.4 ± 17.3 ^b	0.014 ± 0.013 ^b	0.420
Wheat	25	7.7 ± 1.7 ^b	0.019 ± 0.013 ^b	0.454
Black acha	35	34.6 ± 9.1 ^c	0.018 ± 0.014 ^c	0.423
White acha	35	38.7 ± 10.7 ^c	0.017 ± 0.013 ^c	0.267
Wheat	35	8.8 ± 1.5 ^c	0.036 ± 0.029 ^c	0.373

¹Values are mean ± standard error. C_{∞} = equilibrium percentage of starch hydrolyzed after 180 min; k = kinetic constant;

$R^2 = 1 - (\text{Residual Sum of Squares}) / (\text{Corrected Sum of Squares})$

²Values with the different superscript in a column (within boiling time) are significantly different ($p < 0.05$).

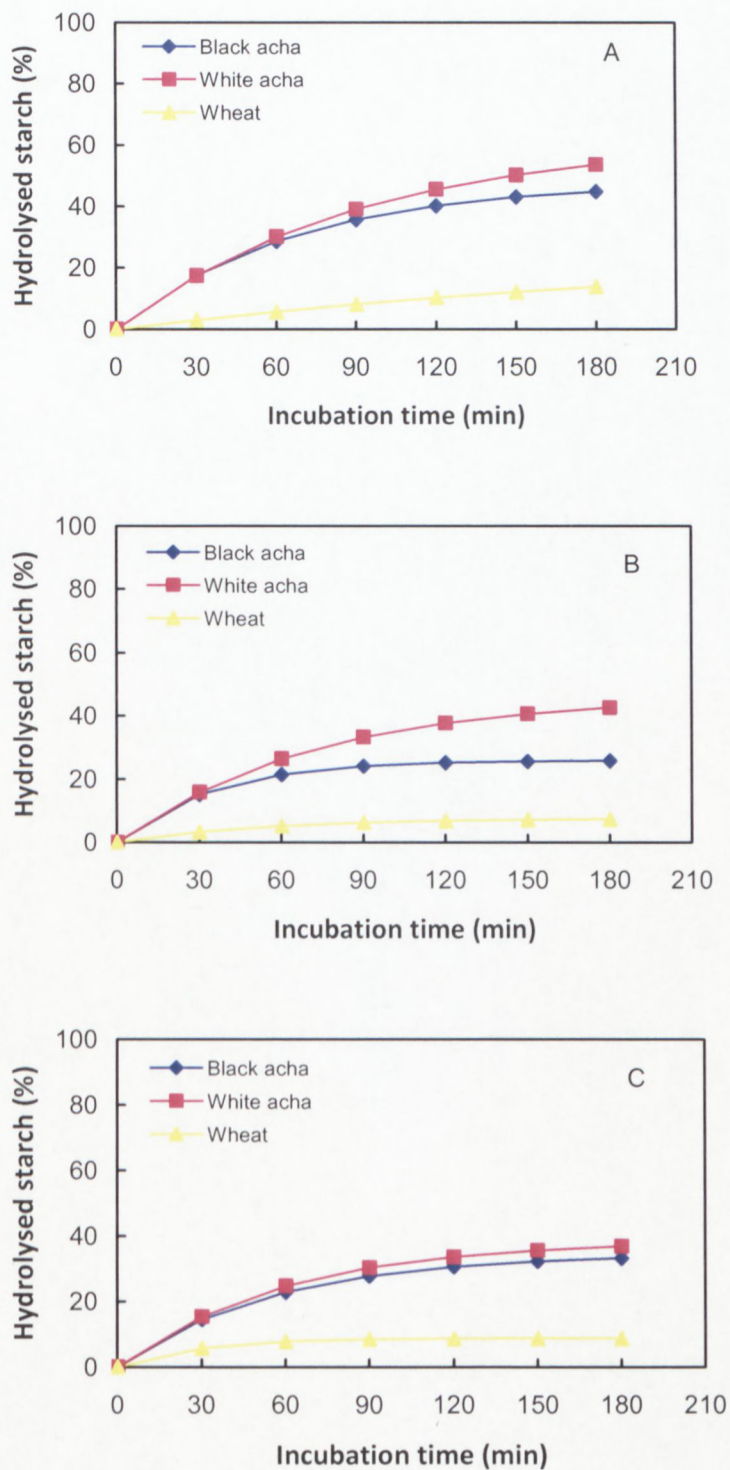


Figure 5.7. Predicted hydrolysed starch from black acha, white acha and wheat after boiling for (A) 15 min; (B) 25 min; (C) 35 min

amongst the cereals. The equilibrium percentage of starch hydrolyzed after 180 min for samples boiled for 25 min was 25.9% for white acha and 27.7% for wheat and black acha, 46.1%. Chung *et al.* (2008) reported similar differences in starch hydrolysis and physicochemical properties among pulse from the same species.

The rate of starch hydrolysis (k) did not differ significantly ($p > 0.05$) among the cereals for the same processing times. However the rate of starch hydrolysis increased with increase in boiling time of the different cereal grains. The highest rate of starch hydrolysis (k) was for cereal samples boiled for 35 min, with wheat 0.036 followed by black acha 0.018 and white acha 0.017. The increase in hydrolysis rate (k) was influenced by the difference in their microstructures caused by longer boiling times. Lee *et al.* (2005) demonstrated that the highest kinetic constant was experienced Table 5.8 and Figure 5.8 detail with the longest method (autoclaving) followed by stone pot cooking, electric pot cooking and lastly microwaving. Englyst *et al.* (1992) reported that the variation on rate and extent of starch digestion in foods depends on several intrinsic and extrinsic factors such as physical form of the food and its composition. There are significant differences ($p < 0.05$) in the rate of starch hydrolysis (k) and the equilibrium percentage of starch hydrolysed after 180 min of incubation for white acha, black acha and wheat grain for the different boiling times (15, 25 and 35 min). This supports the hypothesis that white acha and black acha will differ in their hydrolysis rate and kinetic constant.

5.4.4 Modelling the effect of steaming on the acha starch hydrolysis

The model parameters for samples from black acha, white acha and wheat grain subjected to different steaming times is detailed in Table 5.8 and Figure 5.8. Apart from wheat and white acha steamed for 25 min, the equilibrium percentage of starch hydrolysed after 180 min did not differ significantly ($p > 0.05$) among the cereals for the different steaming times (15, 25, 35 min). This could be due to minimal disruption of the starch crystalline regions at the various steaming times. The amount of moisture to which the cereals were subjected to, played a role in the degree of disruption of the cereals starch crystallinity during steaming. Less

Table 5.8 Model parameters for black acha, white acha and wheat grain as affected by steaming^{1,2}

Cereal	Steaming		C _∞	k	R ²
	time (min)				
Black acha	15	12.3 ± 2.3 ^a	12.3 ± 2.3 ^a	0.558 ± 0.017 ^a	0.201
White acha	15	15.4 ± 13.2 ^a	15.4 ± 13.2 ^a	0.012 ± 0.024 ^b	0.167
Wheat	15	3.2 ± 1.3 ^a	3.2 ± 1.3 ^a	0.025 ± 0.038 ^b	0.145
Black acha	25	11.7 ± 2.4 ^a	11.7 ± 2.4 ^a	0.024 ± 0.017 ^a	0.413
White acha	25	4.9 ± 1.0 ^b	4.9 ± 1.0 ^b	0.055 ± 0.078 ^a	0.168
Wheat	25	2.5 ± 0.6 ^b	2.5 ± 0.6 ^b	0.043 ± 0.054 ^a	0.210
Black acha	35	8.6 ± 0.7 ^a	8.6 ± 0.7 ^a	0.024 ± 0.007 ^a	0.817
White acha	35	10.6 ± 3.5 ^a	10.6 ± 3.5 ^a	0.024 ± 0.028 ^a	0.220
Wheat	35	12.9 ± 6.5 ^a	12.9 ± 6.5 ^a	0.010 ± 0.010 ^a	0.444

¹Values are mean ± standard error. C_∞ = equilibrium percentage of starch hydrolyzed after 180 min;

k = kinetic constant; R² = 1 - (Residual Sum of Squares) / (Corrected Sum of Squares)

²Values with the different superscript in a column (within steaming time) are significantly different (p < 0.05).

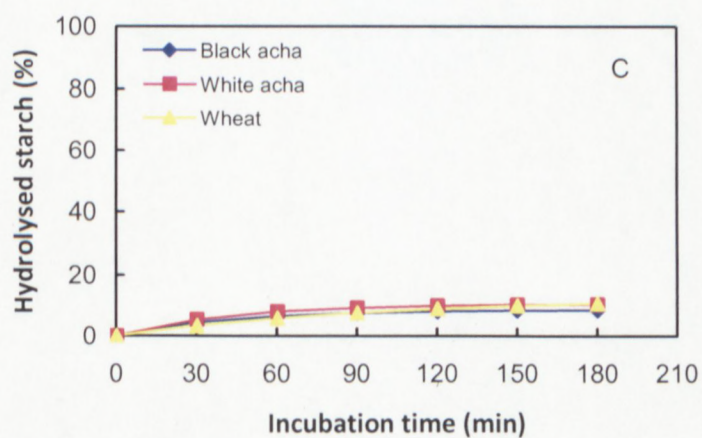
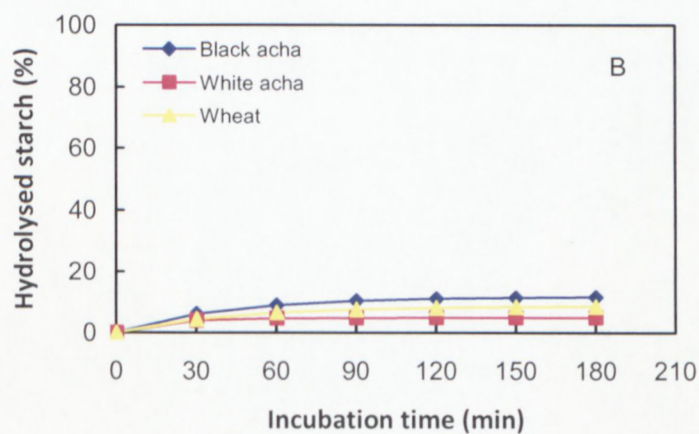
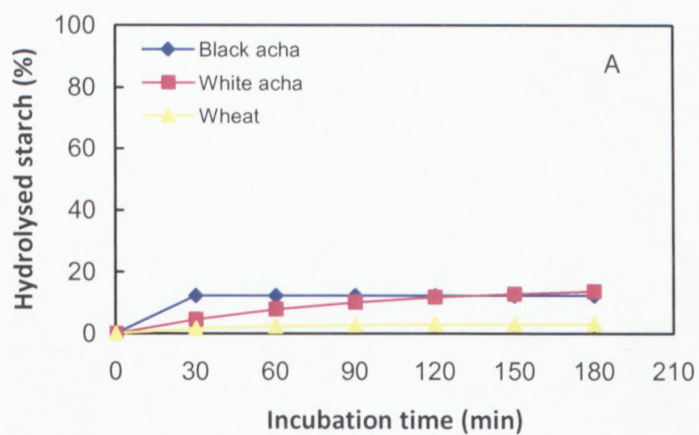


Figure 5.8. Predicted hydrolysed starch from black acha, white acha and wheat after steaming for (A) 15 min; (B) 25 min; (C) 35 min

disruption of the crystalline regions leads to less water infiltration and hydration of the starch granules (Sargum & Argot, 2000). A slight increase in the equilibrium percentage starch hydrolysis from the shortest steaming time of 15 min to the longest steaming time of 35 min was observed. This is in agreement with reports by Sargum & Argot (2000) who reported an increase in the starch hydrolysis with increase in gelatinization and thus an increase in the breakdown of the crystallinity of the starch molecules. However, the equilibrium percentage of starch hydrolyzed for wheat (2.5%) and white acha (4.9%) did not differ significantly ($p < 0.05$) after 25 min steaming, but differed significantly ($p < 0.05$) from black acha (11.7%) at the same steaming time. Van der Merwe *et al.* (2001) reported that the way the cooking affects the structure of the different samples would greatly affect the starch hydrolysis. Chung *et al.* (2008) reported similar differences in extent of starch hydrolysis for pulse species of the same cultivar and specie. They attributed these differences in extent of starch hydrolysis to differences in protein content, amylose content and melting temperatures.

The rate of starch hydrolysis (k) differed significantly ($p < 0.05$) for wheat and white acha after 15 min of steaming and black acha at the same steaming time. The rate of starch hydrolysis was the highest for black acha after 15 min, followed by wheat and white acha. There was however no significant differences ($p < 0.05$) in the rate of starch hydrolysis between black acha, white acha and wheat at the rest of the different processing times (25, 35 min). This does not support the hypothesis that white acha grain and black acha grain will differ in the hydrolysis rates when subjected to different processing times.

5.4.5 Modelling the effect of microwaving

Table 5.9 and figure 5.9 depicts the model parameters for samples from black acha, white acha and wheat grain subjected to different microwaving times. There were no significant difference ($p > 0.05$) in the equilibrium amount of starch hydrolyzed after 180 min for black acha microwaved for 3 min and black acha, white acha and wheat microwaved for different microwaving times (6, 9 min),

Table 5.9 Model parameters for black acha, white acha and wheat grain as affected by microwave^{1, 2, 3}

Cereal	Microwave				R ²
	time (min)	C _∞	k		
Black acha	3	22.9 ± 5.5 ^a	0.027 ± 0.026 ^a	0.260	
White acha	3	4.7 ± 0.8 ^b	0.027 ± 0.017 ^a	0.403	
Wheat	3	16.0 ± 3.5 ^a	0.028 ± 0.025 ^a	0.304	
Black acha	6	19.2 ± 5.6 ^a	0.022 ± 0.022 ^a	0.296	
White acha	6	3.8 ± 0.5 ^a	0.031 ± 0.017 ^a	0.504	
Wheat	6	14.5 ± 2.5 ^a	0.049 ± 0.053 ^a	0.268	
Black acha	9	18.1 ± 3.8 ^a	0.038 ± 0.041 ^a	0.237	
White acha	9	5.4 ± 0.9 ^a	0.019 ± 0.010 ^a	0.561	
Wheat	9	37.2 ± 26.8 ^a	0.009 ± 0.012 ^a	0.320	

¹Values are mean ± standard error. C_∞ = equilibrium percentage of starch hydrolyzed after 180 min;

k = kinetic constant; R² = 1 - (Residual Sum of Squares) / (Corrected Sum of Squares)

²Values with the different superscript in a column (within microwave time) are significantly different (p < 0.05).

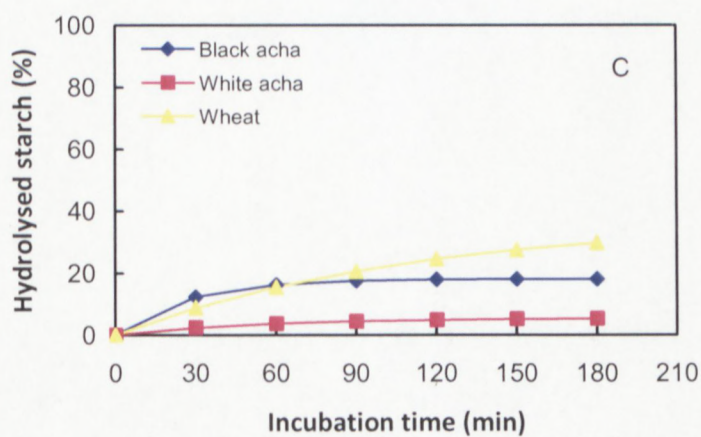
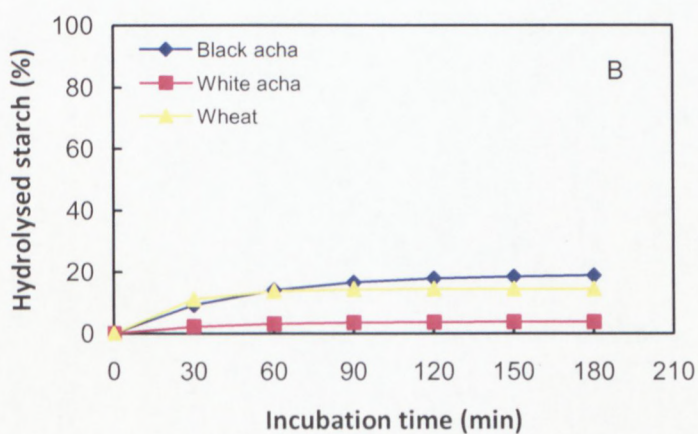
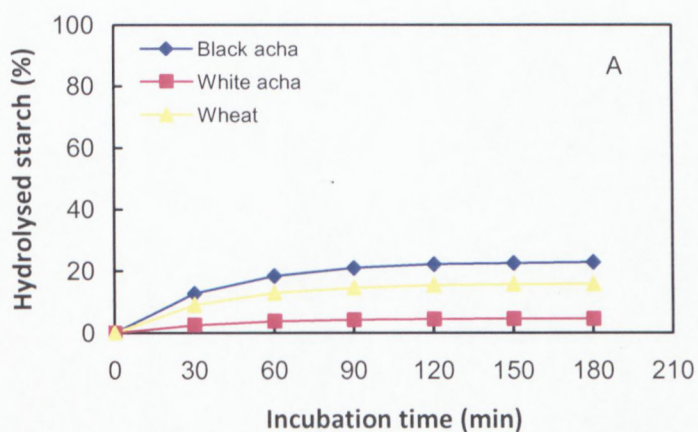


Figure 5.9. Predicted hydrolysed starch from black acha, white acha and wheat after microwave for (A) 3 min; (B) 6 min; (C) 9 min

respectively. This similarity indicates a similar degree of disruption of the cellular structure and therefore degree of susceptibility of starch to amylolytic enzymes after 3, 6, and 9 min of microwaving. There was however significant difference in the equilibrium percentage amount of starch hydrolyzed for white acha (4.7%) and the equilibrium amount of starch hydrolyzed for black acha (22.9%) and wheat (16.0%) after 3 min of microwaving. This difference in the equilibrium percentage hydrolysed starch could be due to genetic differences and therefore difference in the degree of disruption of the crystalline region of the starch molecules of the different cereals after 3 min of microwaving. Lee *et al.* (2005) reported that the overall starch hydrolysis of cooked rice was influenced by the differences in their microstructure, which were caused by the different processing methods. Similar results were reported by Eyaru *et al.* (2009). They highlighted that cooking increases starch digestibility and decreases enzyme resistant starch. There were no significant differences in the rate (k) of starch hydrolysis amongst the different samples or the different processing times. There was very little literature available on the effect of microwave processing on the rate of starch hydrolysis. However this similarity in the rate (k) of starch hydrolysis amongst the different samples or the different processing times does not support the hypothesis that white and black acha grain will differ in their rate of starch hydrolysis at different micro waving times.

5.5 Total, Resistant and Digestible starch in Acha and Wheat bread

The amount of total, resistant and digestible starch in white acha, black acha and wheat bread is shown in Table 5.10. There are significant difference ($p < 0.05$) in the total percentage starch between wheat bread (WB), white acha bread (WAB) and black acha bread (BAB). Total starch in WB, WAB and BAB were 6.6%, 42.0% and 52.3%, respectively. Hallstrom *et al.* (2011) reported that the starch content of white wheat bread to be 40.7%. Hence, the result is in agreement with the value of total WB reported in this work. The significant difference in the total starch between the acha bread varieties and the wheat bread could be due to the fact that the acha bread recipe consisted of added

Table 5.10 Total starch, resistant starch and digestible starch in bread samples^{1,2}

Bread	Starch (%)		
	Total	Resistant	Digestible
Wheat	6.6 ± 4.2 ^a	0.94 ± 0.09 ^a	5.7 ± 4.2 ^a
White acha	42.0 ± 2.5 ^b	0.74 ± 0.03 ^b	41.2 ± 2.5 ^b
Black acha	52.3 ± 4.1 ^c	0.93 ± 0.01 ^a	51.3 ± 4.1 ^c

¹Values are mean ± standard error.

²Values with the different superscript in a column are significantly different ($p < 0.05$).

acha starch, whereas the wheat bread did contain added starch in the recipe. There was no significant difference ($p > 0.05$) in the resistant starch between BAB (0.93%) and WB (0.94%). There was however significant difference ($p < 0.05$) in resistant starch between WAB (0.74%) and WB as well as between the two acha cultivars, WAB (0.74) and BAB (0.93). The results in this study were higher than that reported by Elmstahl (2002) for Hallakaka wheat bread (0.60%). Elmstahl (2002) however also reported wholemeal bread to contain 1.5% resistant starch. This contrast in quantity of resistant starch could be due to part of the wholemeal to consist of intact cereal grain which keeps the starch encapsulated in the grain structure. The difference in the resistant starch content between WAB and BAB could be due to genetic difference of the two acha grain varieties.

In terms of digestible starch, a significant difference ($p < 0.05$) was detected between the two acha bread variants and WB. The digestible starch was 5.7%, 41.2% and 51.3%, respectively for WB, WAB and BAB. The higher percentage of digestible starch in WAB (41.2%) and BAB (51.3%) could be attributed to the added acha starch as well as the lite apple juice in the bread recipe. Hence, addition of acha starch and lite apple juice to acha bread recipe is discouraged as it may add to the glycemic load. This suggests effect of ingredients on GI of a food product is additive. Consequently, combining low GI ingredients in a recipe could result in high glycemic load.

The significant difference in the total starch, resistant starch and digestible starch supports the hypothesis that significant differences exists in the starch of the two acha varieties.

5.6 Total, Resistant and Digestible Starch in Acha and Wheat Grain

The percentage of total, resistant and digestible starch of acha and wheat grain is show in Table 5.11. There are no significant difference ($p > 0.05$) in the percentage total starch between wheat and white acha grain. There was however significant difference ($p < 0.05$) between black acha grain and wheat grain in terms of total starch content as well as between white and black acha

Table 5.11 Total starch, resistant starch and digestible starch of grain samples^{1,2}

Grain	Starch (%)		
	Total	Resistant	Digestible
Wheat	15.02±0.28 ^a	5.89±0.04 ^a	9.14±0.28 ^a
White acha	16.23±0.97 ^a	6.07±0.82 ^a	10.17±1.71 ^a
Black acha	12.16±1.94 ^b	6.14±0.37 ^a	6.03±2.07 ^b

¹Values are mean ± standard error.

²Values with the different superscript in a column are significantly different (p < 0.05).

grain. The difference in the percentage total starch between white acha and black acha grain could be due to differences in their genetic structure. A resistant starch value of 3.1% was however reported by Hallstrom *et al.* (2011) for elevated amylose wheat grain. Resistant starch has many health benefits such as provision of dietary fibre, prebiotic benefits to the gut as well as overall health benefit. The higher resistant starch reported in the study could be due to the grains being whole and intact. Whole and intact grains are good sources of resistant starch, since the starch is encapsulated in the intact grain.

The resistant starch for wheat grain is 5.9%, for white acha starch grain, 6.07% and for black acha starch grain is 6.1%. There were no significant differences ($p > 0.05$) in resistant starch amongst the different grains. Hallstrom *et al.* (2011) reported a resistant starch value of 2.9% for whole grain wheat. This is much lower than the values reported in this study, wheat grain was 5.9%, white acha grain, 6.1% and black acha grain 6.1%. The digestible starch for white acha grain was highest at 10.2%, followed by wheat grain at 9.1% and black acha grain at 6.0%. There was no significant difference in the percentage digestible starch between white acha grain and wheat grain. There was however significant difference ($p < 0.05$) in digestible starch between white and black acha grain. The difference in total starch and digestible starch between white and black acha grain supports the hypothesis that a difference exist in the two acha grain varieties.

5.7 Total, resistant and digestible starch in processed acha and wheat grain.

Rapidly digestible starch (RDS) and slowly digestible starch (SDS) contents of acha and wheat bread are shown in Figures 5.10, microwaved acha and wheat grain in Figure 5.11 and boiled acha and wheat grain in Figure 5.12, steamed acha and wheat grain in Figure 5.13. Table 5.13 shows the starch hydrolysis of acha and wheat grain with respect to incubation time.

The RDS for baked bread ranged from 12% for wheat bread (WB), 51% for black acha bread (BAB) and 60% for white acha bread (WAB) (Figure 5.10).

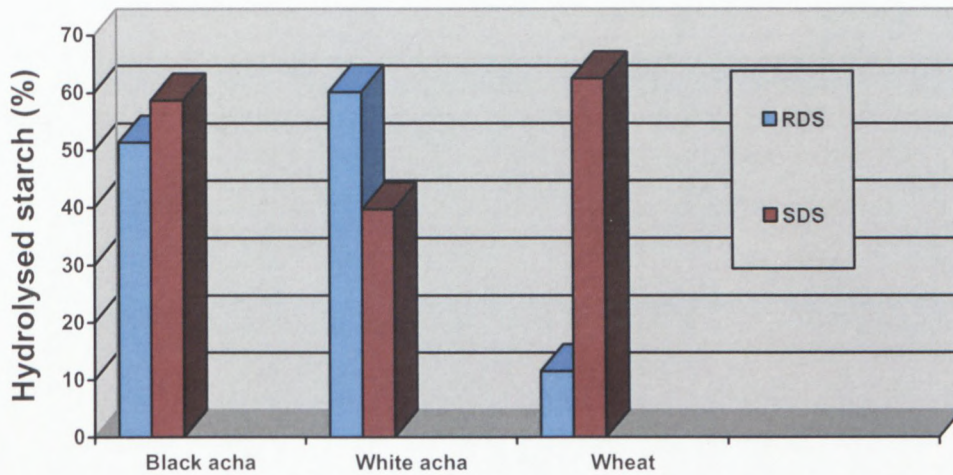


Figure 5.10 Rapidly digestible starch (RDS), and slowly digestible starch (SDS) of milled acha and wheat bread.

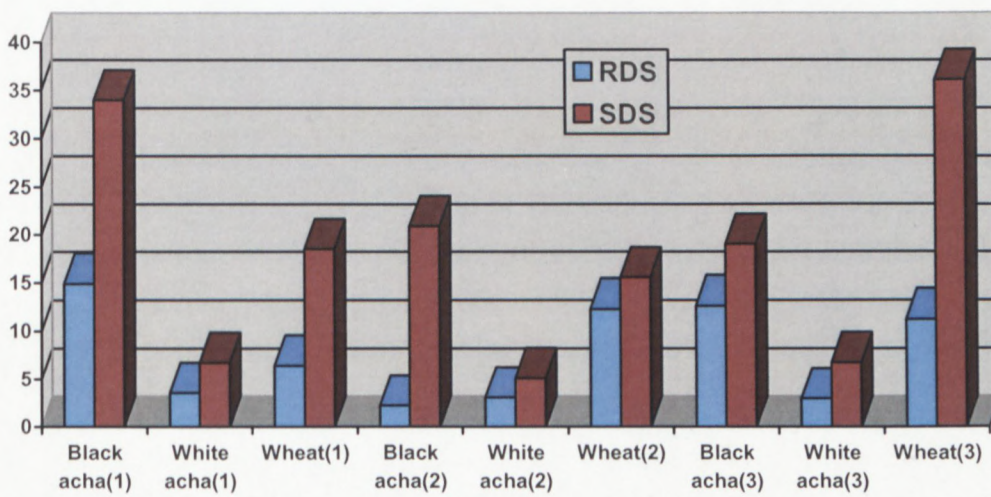


Figure 5.11 Rapidly digestible (RDS), and slowly digestible starch (SDS) of microwaved acha and wheat grain. 1: acha and wheat grain microwaved for 3 minutes, 2: acha and wheat grain microwaved for 6 minutes, 3: acha and wheat grain microwaved for 9 minutes.

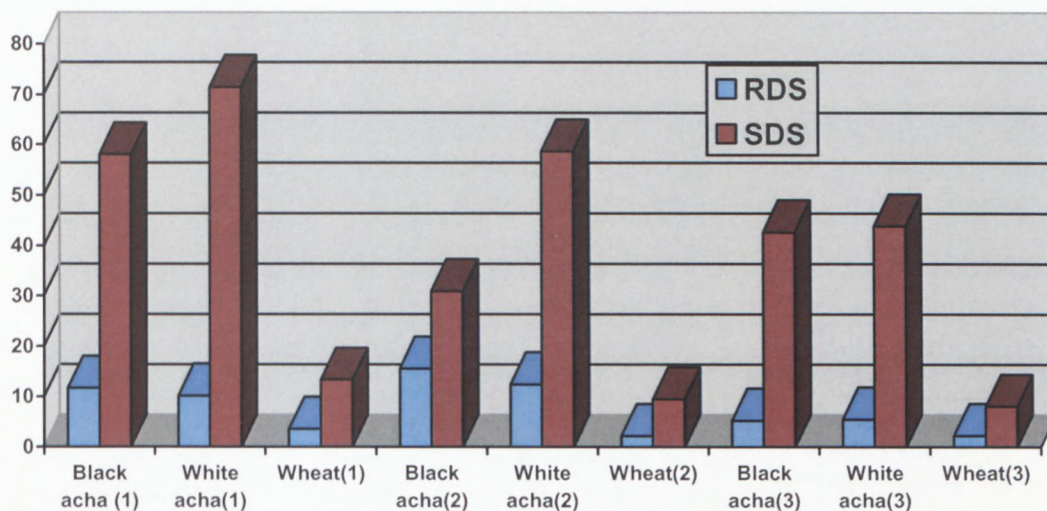


Figure 5.12 Rapidly digestible (RDS), and slowly digestible starch (SDS) Of boiled acha and wheat grain. 1: acha and wheat grain boiled for 15 minutes, 2: acha and wheat grain boiled for 25 minutes, 3: acha and wheat grain boiled for 35 minutes.

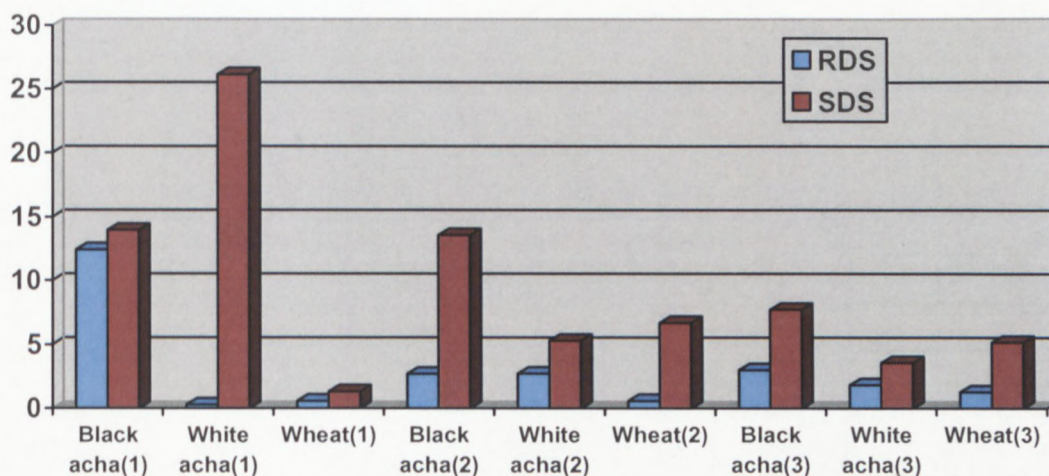


Figure 5.13 Rapidly digestible (RDS), and slowly digestible starch (SDS) of steamed acha and wheat grain. 1: acha and wheat grain steamed for 15 minutes, 2: acha and wheat grain microwaved for 25 minutes, 3: acha and wheat grain microwaved for 35 minutes.

The SDS content for BAB was the highest at 89% followed by WB, 62% and WAB at 56%. Goñi *et al.* (1997) reported a RDS in the region of 20% and SDS of about 25% for wheat bread. This SDS for wheat bread is much lower than that reported in this study (62%) and the RDS reported by Goñi *et al.* (1997) is much higher than the RDS results in this study (12%). Whereas Sayed *et al.* (2008) reported RDS of 41.9% and SDS of 5.8% for common wheat bread and a RDS of 42.3% and SDS of 6.5% for spelt bread. This difference might be due to differences in genetics and composition of the wheat cultivars, baking formulas and baking conditions. The higher RDS of acha breads compared to the wheat bread could be due to the difference in the recipe, since the acha bread recipe consisted of additional acha starch whereas the white bread recipe did not contain added starch.

The SDS for microwaved black acha grain (MBA) was higher than that of microwaved wheat grain (MWG) at 3 and 6 minutes of microwaving (Figure 5.11). For MBA the SDS was 34.0% for 3 minutes and 20.9% for 6 minutes of microwaving. Whereas for MWG, it was 18.5% at 3 minutes of microwaving and 15.5% at 6 minutes of microwaving. The SDS for microwaved white acha (MWA) was much lower than that of MBA as well as MW. The SDS for MWA was 6.69% for 3 minutes, 5.07% for 6 minutes and 6.72% for 9 minutes of microwaving. The SDS after 9 minutes of microwaving for MBA was 19.0% and for MW was 36.10%. The RDS for all microwaved grain samples were much lower than the SDS microwaved grain samples for the different processing times.

The SDS content of 15, 25 and 35 minutes boiled black acha (BBA), boiled white acha (BWA) and boiled wheat grain (BW) grain were significantly higher than the RDS for BBA, BWA and BW grain.

The SDS for steamed black acha (SBA), steamed white acha (SWA) and steamed wheat grain (SW) after 15 minutes of steaming was 13.94% (SBA) and 26.15% (SWA) and 1.32% (SW). The RDS for steamed acha and wheat grain was significantly higher than SDS for 15, 25 and 35 minutes of steaming. The SDS for SBA after 15, 25 and 35 minutes of steaming was higher than the SDS

for SWA and SW at the same steaming times.

A study RDS and SDS of whole grain acha porridge were conducted by Podgorski (2009) showed a RDS of 13.73% for white acha and 16.26% for black acha porridge. The RDS reported for white and black acha porridge was in the same range as the results for RDS in this study for boiling after 15 and 25 minutes, but much higher than the results for boiling of 35 minutes. The SDS content for white acha porridge was 18.47% and black acha porridge was 39.97%. This is lower than that reported in this study for BWA, 71.4% after 15 minutes and 43.8% after 35 minutes, but higher than the 9.45% after 25 minutes of boiling. The SDS for black acha porridge reported by Jideani & Podgorski (2010) is much lower than that reported in this study for 15 minutes of boiling (57.98%), 25 minutes (58.62%) and 35 minutes (42.62%). This difference in SDS and RDS can be attributed to the difference in boiling processing method with regard to sample size and boiling times. No evidence could be found of investigation of the effect of steam and microwave processing on acha and wheat grains.

According to Englyst *et al.* (1999), the nutrients in minimally processed cereal grains are encapsulated within the cell walls which retard the release and thus digestion of starch which thus lead to low RDS. High RDS foods are harmful for health, whereas high SDS foods are beneficial for reducing chronic diseases (FAO/WHO, 1997).

Results indicate that black acha bread as well as microwaved, boiled, steamed black acha grain is the better alternative in the dietary management of metabolic disorders such as diabetes and hyperlipidemia due to its lower RDS and higher SDS compared to the higher RDS and SDS for white wheat bread, microwaved, boiled and steamed wheat grain.

5.8 Estimated Glycemic index (EGI) of processed Acha and Wheat grain and bread.

Figure 5.14 shows the GI of acha and wheat breads. Black acha and wheat bread the GI differed significantly. For white wheat bread and white acha bread it

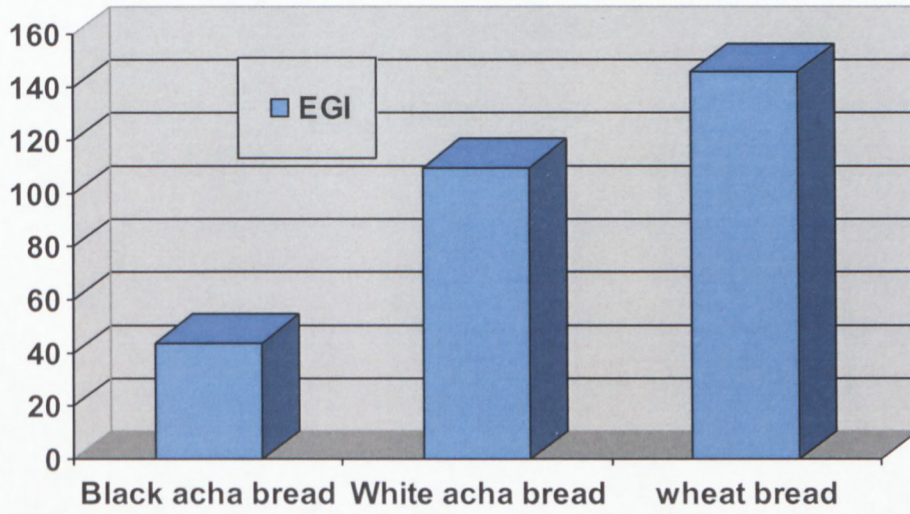


Figure 5.14 Estimated Glycemic Index (EGI) of black acha bread, white acha bread and wheat bread.

was 146 and 110 respectively whereas for black acha bread it was 44. Englyst (1999) reported that white wheat bread, corn flakes, and spaghetti are all examples of highly processed foods. The starch in white wheat bread is fully gelatinized and thus likely to be rapidly digested and absorbed and white wheat breads thus have high GI values. Glycemic index categories are as follows, low GI (0-55), intermediate (56 -69) and high GI (>70). Based on this, the black acha can be categorised as low GI whereas white wheat bread and white acha bread as high GI. Shangumam *et al.* (2007) however reported that the average GI values, calculated, taking white bread as a control (GI =100), for wheat-based, millet-based, expanded rice-based and popped rice based foods were 55.4, 93.4, 105 and 109 respectively. White acha bread in this study was thus in line with GI results reported for millet and popped rice based breads reported by Shanmugam *et al.* 2007. The difference in the GI between white acha, black acha and wheat bread could be attributed to the nature of available as well as non-available carbohydrates since both acha bread recipes consisted of added starch whereas the white wheat bread did not consist of added starch. The difference in GI values between white and black acha bread could be due to genetic differences. Englyst *et al.* (2003) reported that white wheat bread, corn flakes, and spaghetti is all examples of highly processed foods. The starches in white wheat breads are fully gelatinized and thus likely to be rapidly digested and absorbed and thus have high GI values.

After boiling for 15, 25 and 35 minutes, both acha varieties and wheat grain were categorised as low GI according to their GI index results (Figure 5.15). The GI ranged between 40 and 46 for both acha varieties and wheat grain, with wheat grain (46) the highest followed by white acha grain (45) after 35 minutes of boiling. Black acha grain had the lowest GI at all processing times, 40 after 15 minutes boiling and 41 after 25 and 35 minutes of boiling respectively.

For steaming the highest GI was for black acha grain (58) after 15 minutes of steaming and wheat grain with a GI of 56 after 35 minutes of steaming (Figure 5.16). This would categorise black acha grain steamed for 15 minutes and

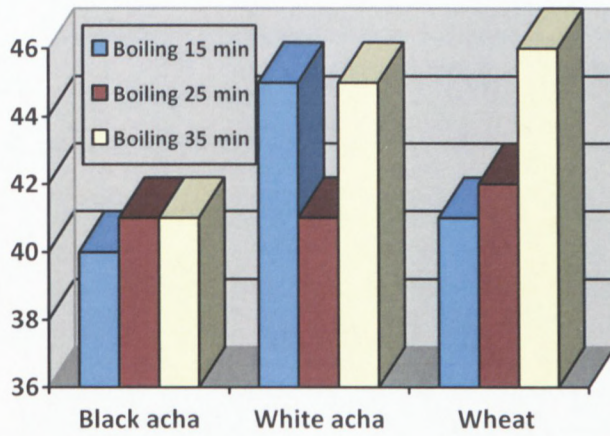


Figure 5.15 EGI of acha and wheat grain boiled for 15, 25 and 35 minutes.

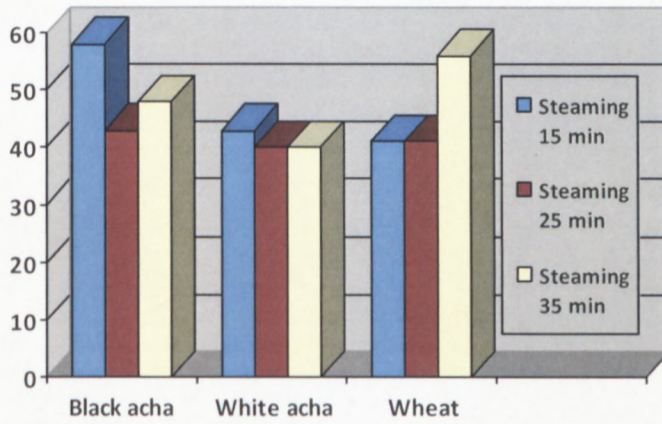


Figure 5.16 EGI of acha and wheat grain steamed for 15, 25 and 35 minutes.

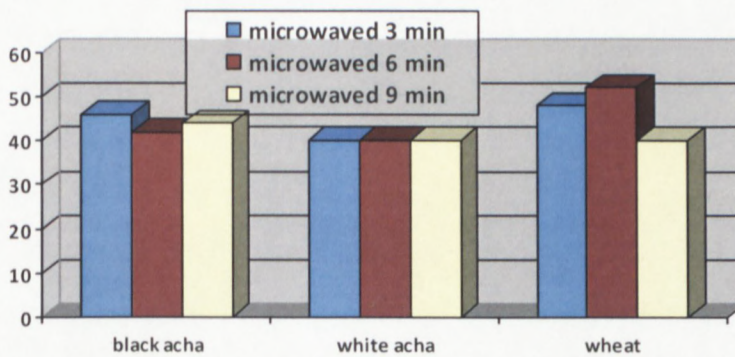


Figure 5.17 EGI of acha and wheat grain microwaved for 3, 6 and 9 minutes.

wheat grain steamed for 35 minutes as intermediary GI products. Steenkamp (2008) reported that the longer the processing, time the higher the GI value. White acha grain however consisted of GI value of 43 after 15 minutes steaming and 40 after 25 and 35 minutes steaming respectively and would thus be categorised as low GI. The average GI for both acha varieties and wheat grain were the lowest after 9 minutes of microwaving (Figure 5.17) and did not differ significantly, ranging between 40 for both wheat and white acha and 44 for black acha grain. Black acha and wheat grain had the highest GI value after 3 and 6 minutes of microwaving, with black acha consisting of 46 and wheat grain 48 after 3 minutes of microwaving and 42 and 52 after 6 minutes of microwaving respectively. White acha grain remained a constant 40 after 3, 6 and 9 minutes of microwaving. Both acha varieties and wheat grain have been categorised as low GI in this study and thus offer the possibility of being used for diabetic dietary management.

The physical form of food and the degree of processing imparted to the foods also influences the glycemic response of the food (Shanmugam *et al.*, 2007). Since the majority of the foods under this study was intact grain and has undergone minimum disruption, the low GI values for boiled, steamed and microwaved acha and wheat grain were to be expected. In many plant foods, such as legumes and minimally processed cereal grains (e.g., pearled barley), nutrients are encapsulated within cell walls (dietary fibre), which retard the release and hence digestion and absorption of starch and sugars, and these foods thus have low GI values (Englyst *et al.*, 1999).

5.8 Conclusion and Recommendation

The effect of processing on the starch and glycemic properties of acha is reported in this chapter. The following conclusion can be drawn:

1. The different processing methods baking, boiling, microwaving and steaming, affected the black and white acha starch hydrolysis. The amount of starch hydrolysed for the different processing methods was in the following order: baking > boiling > microwaving > steaming.

2. The equilibrium percentage of starch hydrolysed was affected differently for the various starches, black acha, white acha and wheat starch by the different processing methods and times. In the case of baking black acha and wheat bread was affected similarly. However, this was not the case for microwaving, steaming and boiling, where both acha starch varieties and wheat starch were affected in the same way.
3. The rate of starch hydrolysis for both acha varieties and wheat grain for the different processing methods, steaming, boiling, microwaving and baking was affected to the same degree.
4. SDS and RDS analysis results indicate that black acha grain (*Digitaria iburua*) cultivar is the better cultivar to use than white acha grain (*Digitaria exilis*) for the treatment and prevention of chronic disease due to its higher slowly digestible starch (SDS) and lower rapidly digestible starch content after baking, microwaving, boiling and steaming. White wheat bread, microwaved, boiled and steamed wheat grain had a higher RDS and SDS compared to black acha.
5. This study indicated that the acha white bread and processed (boiled, steamed and microwaved) acha and wheat grain consisted of low GI values and is thus suitable as a food supplement or meal replacer for those individuals suffering from diabetes and celiac disease.

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CHAPTER 6

Summary, Conclusion and Recommendation

The physicochemical, thermal and functional properties of two acha cultivars (*Digitaria exilis*, white and *Digitaria iburua*, black) were investigated. Wheat was used as reference. The acha starch was extracted from cleaned and dried acha grain. The physicochemical properties investigated were, proximate nutritional content, amylose content, SDS and RDS. The thermal properties investigated were gelatinization and retrogradation. The functional properties investigated were texture profile analysis, turbidity, pasting properties and water binding capacity. Bread was baked with two acha cultivars (*Digitaria exilis*, white and *Digitaria iburua*, black) and the sensory and physical properties investigated and compared against that of white wheat bread. The physical properties that were investigated were specific loaf volume as well as consumer acceptability. The sensory properties investigated were appearance, crust colour, crumb colour, aroma, taste, firmness, mouthfeel and overall acceptability. The effect of baking, steaming, boiling and microwaving on the starch and glycemic properties of two acha cultivars (*Digitaria exilis*, white and *Digitaria iburua*, black) was investigated. The aim of this research to determine the effect of different processing methods; baking, boiling, steaming and microwaving on the starch and glycemic properties of acha were met. The specific objective to evaluate the changes in glycemic properties of acha due to different processing methods (baking, boiling, steaming and microwaving) was also met.

The following conclusions were made.

1. Although there was no significant difference in the nutritional composition of white acha starch and black acha starch with regard to carbohydrate, moisture and fat content, there was however significant differences with regard to the protein content and dietary fibre between black and white acha starches. Wheat and white acha starch had no significant difference in the nutritional composition with regard to Moisture, fat and carbohydrates. There was no significant difference in amylose content between black acha, white acha and wheat starches.

2. Significant differences in some of the functional properties with regard to pasting and turbidity were observed, but no significant difference in the TPA and WBC. WBC of both acha varieties was higher than that for wheat starch. This indicates that both acha starch varieties can be used for hot and cold desserts as well as for soft jelly like sweets and confectionery toppings.
3. No significant differences with regard to the thermal properties of the two acha starch cultivars and wheat starch were observed. The observed gelatinisation temperature of acha and iburua starches typifies that of non-waxy starch. Acha starch has similar retrogradation temperature profiles as that of wheat starch.
4. Optimum acha bread could be baked using 8% acha starch, 2% xanthan gum, 2% CMC and 1.2% yeast.
5. Majority of the panellists found the crust colour, taste and aroma to be moderately desirable. Hence, acha bread has the potential to be marketed as wheat free bread.
6. Few differences in *in vitro* starch digestibility and physicochemical properties were observed among two acha cultivars and wheat starch. Starch hydrolysis was significantly ($P < 0.05$) affected by both time and starch grain. There was no significant difference in the amount of hydrolyzed starch between wheat and white acha starch after 180 min. The two acha starch cultivars had fairly low RDS contents compared to wheat starch whereas the SDS for wheat and white acha starch was higher than that of black acha starch.
7. The different processing methods baking, boiling, microwaving and steaming, affected the black and white acha starch hydrolysis. The amount of starch hydrolysed for the different processing methods was in the following order: baking > boiling > microwaving > steaming.
8. The rate of starch hydrolysis for both acha varieties and wheat grain for the different processing methods, steaming, boiling, microwaving and baking was affected to the same degree.
9. On the basis of slowly digestible starch (SDS) and rapidly digestible starch (RDS), black acha grain (*Digitaria iburua*) cultivar is the better cultivar to use than white acha grain (*Digitaria exilis*) for the treatment

and prevention of chronic disease due to its higher SDS and lower RDS after baking, microwaving, boiling and steaming.

10. This study indicated that the acha white bread and processed (boiled, steamed and microwaved) acha and wheat grain consisted of low GI values and is thus suitable as a food supplement or meal replacer for those individuals suffering from diabetes and celiac disease.

Some of the hypotheses stated below were supported while others were not supported by the test results in this study:

1. For the hypothesis that starches from the two acha cultivars (*D. exilis* and *D. iburu*) differed in their functional, physicochemical and thermal properties, test results indicated that only some functional and physicochemical properties supported this hypotheses while results from thermal analysis did not support the hypothesis.
2. The hypothesis that different processing methods differed in their effect on the starch and glycemic properties of acha was not supported. The rate of starch hydrolysis for both acha varieties and wheat grain for the different processing methods, steaming, boiling, microwaving and baking was affected to the same degree.
3. The effect of processing on the starch and glycemic properties of acha differed among the cultivars. Results of the baked samples supported this hypothesis whereas processing such as boiling, steaming and microwaving did not support this hypothesis. All processed acha grain could be categorised as low GI products.

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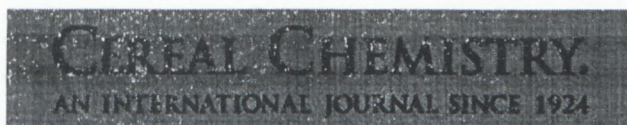
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**Physicochemical, Thermal and Functional Properties of Acha
(Digitaria spp.) Starch**

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1 **Physicochemical, Thermal and Functional Properties of Acha (*Digitaria* spp.) Starch**

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25 **Key words**

26 Gelatinization, Retrogradation, viscosity, starch.

27

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28

Abstract

29 Acha [*Digitaria exilis*, white (WTAS) and *Digitaria iburua*, black (BLAS)], consumed as
30 whole grain, has been reported to be high in resistant and slowly digestible starches but low
31 in rapidly digestible starch. It is a grain of choice in the management of diabetes in Africa.
32 Our objective was to investigate the physicochemical, thermal and functional properties of
33 starches from the two acha cultivars compared to wheat starch (WHTS). There were no
34 significant differences in amylose between BLAS (42.1%), WTAS (61.4%) and WHTS
35 (45.8%). WHTS had highest onset (T_o) and gelatinization (T_p) temperatures 34.7°C and
36 74.5°C, respectively, although not significant. BLAS and WTAS were 25.0, 26.7°C and
37 68.2, 70.5°C, respectively. The gelatinization temperature ranges did not differ significantly
38 between the varieties. Acha starches with lower transition temperature imply a low
39 crystallinity with high amorphous regions. Temperature of retrogradation was 64.1, 67.0 and
40 64.8°C for WHTS, WTAS and BKAS, respectively. The starches were similar ($p > 0.05$) in
41 gelatinization and retrogradation temperatures. There were significant differences ($p < 0.05$)
42 in peak viscosity and final viscosity between wheat and acha starches. However, no
43 significant difference ($p > 0.05$) existed between WTAS (3125 cP) and BLAS (3137 cP) for
44 trough viscosity and setback viscosity 2332 cP and 2476 cP respectively. Gels produced
45 from the three starches did not differ significantly in hardness, resilience and springiness. All
46 the starches produced highly springy soft gels. Water binding capacity differed significantly
47 between WHTS (0.83 g/100g) and both acha cultivars 1.33 g/100g for WTAS and 1.36
48 g/100g for BLAS. However, the acha cultivars did not differ significantly in WBC. Acha
49 starches can thus be utilized as an alternative to wheat starch when required in application
50 where thermal properties such as gelatinization are important.

51

Introduction

53 *Digitaria* spp. belongs to the family Graminae and the tribe Poaceae (Jideani 1999) and
54 includes 230 species that are widely distributed in the tropics and the subtropics. Acha is by
55 large the oldest African cereal, it has been cultivated for thousands of years in West Africa
56 (Mali, Burkina Faso, Guinea and Nigeria) and Dominican Republic (Jideani et al. 2008).
57 Two types exist namely *Digitaria exilis* (acha, fonio or hungry rice) and *Digitaria iburua*
58 (iburua, black fonio or petit millet), (Philip and Itodo 2006). It consists of a unique small size
59 of 0.4 to 0.5 mm seeds. Acha is high in essential amino acids, leucine 9.7%, methionine
60 5.6% and valine 5.8%. It is considered as one of the world's most nutritious grains (Philip
61 and Itodo 2006). It is also used for the management of diabetes in West Africa. Not much is

62 known about the thermal and functional properties of acha starch. Starch is a food ingredient
63 with wide applications such as thickener, gelling agent, bulking agent and water retention
64 agent, making them an excellent ingredient for the manufacture of foods such as: custards,
65 porridges, puddings, cookies, and sausages. There is a growing demand for starches for the
66 modern food industry creating interest for new sources of this polysaccharide. The
67 availability of a reliable source of starch from agriculture is considered to have an important
68 factor in human development (Asp et al. 1996). Our objective was to investigate the
69 physicochemical, thermal and functional properties of acha starch.

70

71

MATERIALS AND METHODS

72

Materials

74 Two acha cultivars namely, *Digitaria exilis* (white) and *Digitaria iburua* (black) were
75 purchased from Grace Africa, Salt River, Cape Town, South Africa.

76 All equipment and materials were either obtained from the Department of Food
77 Technology, Cape Peninsula University of Technology (CPUT) or Pioneer Foods
78 Laboratories (Bokomo Foods pilot plant and SASKO technical laboratory), Cape Town,
79 South Africa.

80 All chemicals were obtained from Laboratory and Scientific Ltd., Maitland, Cape
81 Town, Republic of South Africa and the Department of Food Technology CPUT.

82

Cleaning of acha grain

84 The cleaning procedure entailed screening the grains for 3 minutes through a 1000, 750, 500
85 and 125 micron screens to remove all foreign matter such as stones, foreign grain and small
86 sticks. The grains were washed three times with water, drained and dried at 40°C for 48 h.
87 The cleaned and dried samples were then placed in sealed polyethylene bags and stored in a
88 refrigerator at a temperature of 5°C until required.

89

Isolation and purification of starch from acha grain

91 The method described by Betancur and Torruco-Uco (2001) was used to extract starch from
92 milled acha grain. Wheat starch (S2760) bought at Sigma Aldrich was used as the reference.
93 Cleaned, washed and dried acha grain were milled and sieved through a 20 mesh screen. The
94 acha flour was then dispersed in distilled water at a ratio of 1:8 w/v. The pH was then

95 adjusted to 11 with 1 N NaOH and the solution was stirred for 1 h. The suspension was then
96 passed through an 80 and 100 mesh screens to separate the fibre solids from the liquid
97 containing the protein and the starch. The suspension was allowed to sediment for 30 min to
98 recover the starch, and then the solubilised protein was separated. The starch was washed 3
99 times with distilled water, and centrifuged at 4250 rpm for 10 min during the last wash to
100 recover the starch. The product was then dried at 60°C in a drying oven (Defy, Gemini Ltd.,
101 Germiston) for 1 h then milled and sieved through a 20 mesh screen. Starch from white acha
102 is denoted as WTAS and that from black acha as BLAS. The isolated acha starches and
103 wheat starch (WHTS) were then analysed for physicochemical, functional and thermal
104 properties.

106 **Physicochemical Properties of Acha Starch**

107 **Proximate composition and total amylose content of the starches**

108 The moisture, fat, protein, fiber and ash content of the acha and wheat starch were determined
109 in accordance with the standard methods of AOAC (1990). Carbohydrates were calculated
110 by difference.

111 The method by Hoover and Ratnayake (2001) was used to determine the amylose
112 content of starches. Defatting of starch samples were conducted by weighing 5 g of starch
113 accurately into a cellulose extraction thimble and covering the mouth with cotton wool.
114 Lipids were then extracted with 120 ml of 75% n-Propanol at 85°C with a heating mantle for
115 7 h in a Soxhlet extractor. The lipid free starch was then air dried for 12 h and removed from
116 the thimble and oven dried for 24 h at 30°C. Lipid-free starch (20 mg) was weighed into a
117 round bottom screw-cap tube fitted with a Teflon-faced rubber liner in the cap. A series of
118 mixtures of pure potato amylose (0%, 10%, 20%, 40%, 50% amylose) were prepared after
119 which 20 mg of each was weighed into round-bottom tubes with caps. Eight millilitres of
120 90% DMSO was added to each round-bottom tube and vigorously mixed for 2 min using a
121 vortex mixer. The tubes were then heated in a water bath at 85°C for 15 min with
122 intermittent mixing. The tubes were then allowed to cool to room temperature (45 min). The
123 samples were then diluted to 25 ml with water in a volumetric flask.

124 Determination of absorbance of the dispersed starch solution was then conducted after
125 completion of the following steps. Diluted solution (1 ml) and 40 ml of distilled water were
126 added into a 50 ml volumetric flask. Iodine solution (5 ml) was added and mixed vigorously.
127 The volume was adjusted to 50 ml with distilled water and mixed vigorously. The colour

128 was then allowed to develop for 15 min. The contents of the flask were then vigorously
129 mixed by hand. The absorbance of the samples and each of the standard mixtures were then
130 measured at 600 nm against a reagent blank as the reference. The reagent blank contained all
131 reagents in the same amounts without the sample containing starch.

132 Following the above procedure a standard curve was then plotted. The regression
133 equation for the standard curve was then measured and used to calculate the total amylose
134 content of the sample.

135

136 **Thermal properties of the starches**

137 Differential scanning calorimeter (DSC, 20 to 120°C, Perkin-Elmer instrument) was used to
138 measure the thermal characteristics of the isolated starch. Both the acha starches and wheat
139 starch (3.5 mg, dry weight) were separately loaded into a 40 µl capacity aluminium pan
140 (Mettler ME-27331) and distilled water added using a Hamilton micro syringe to achieve a
141 70% starch–water suspension. Samples were sealed and allowed to stand for 5 h at room
142 temperature before it was heated in the DSC. The DSC analyzer was calibrated using
143 indium, and an empty aluminium pan was used as a reference (Sandhu and Narpinder, 2007).
144 Sample pans were heated at a rate of 10°C/min from 20°C to 100°C. Thermal transition of
145 starch samples was defined as T_o (onset temperature), T_p (peak of gelatinization), T_c
146 (conclusion temperature) and the enthalpy of gelatinization (ΔH_{gel}). Enthalpies were
147 calculated on a starch dry weight basis automatically. The gelatinization temperature range
148 (R) and peak height index (PHI) were calculated as described in equations 1 and 2.

$$149 \quad R = 2(T_p - T_o) \quad 1$$

$$150 \quad PHI = \Delta H / (T_p - T_o) \quad 2$$

151 After conducting thermal analysis, the samples were stored at 4°C for 7 days, for
152 retrogradation studies. The aluminium sample pans containing the starches were reheated at
153 the rate of 10°C/min from 25°C to 100°C to measure retrogradation. The enthalpies of
154 retrogradation (ΔH_{gel}) T_o , onset temperature; T_p , peak temperature; HI, peak height; H,
155 height; ΔH , enthalpy of gelatinization; T_c , end (conclusion) temperature, ΔT_r gelatinization
156 temperature range ($T_c - T_o$), were calculated automatically by the Pyris R9.1 software.

157

158 **Functional properties of the Starches**

159 **Textural properties of starch gels**

160 Texture profile analysis was conducted as described by Sandhu and Narpinder (2007) on an

161 Instron apparatus (model 2519, UK, 3300 series, 2000 N capacity) to evaluate the textural
162 properties of acha and wheat starch gels. A suspension of 45 g of starch in 455 ml of cold
163 water was prepared. The suspension was then boiled on a hot plate (Model Z341 Ohaus Instr.
164 US) with stirring until a gel was formed. The hot starch gel formed was then poured into
165 small aluminium dishes (10 cm in diameter) and stored at 4°C. The gels were evaluated for
166 their textural properties by means of texture profile analysis. These were calculated
167 automatically with the Blue-Hill 2 software.

168

169 **Turbidity of starch suspensions**

170 The turbidity of the acha starch suspension was measured as described by Perera and Hoover
171 (1999). A 1% aqueous suspension of acha starch as well as the reference wheat starch was
172 heated separately, in a water bath at 90°C for 1 h with constant stirring. The samples were
173 then stored for 5 days at 4°C and turbidity was determined after 5 days by measuring the
174 absorbance at 640 nm (UV-VIS Split Beam 8 Auto Cell Model UVS-2800 spectrophotometer
175 Shanghai Jingke Scientific Instruments Co. Ltd. China) against a water blank.

176

177 **Water binding capacity**

178 Water binding capacity (WBC) of both the acha starch as well as the reference starch was
179 measured as described by Medcalf and Gilles (1965). A suspension of 5 g starch in 75 ml
180 distilled water was agitated for 1 h and centrifuged at 3000 rpm for 10 min. The free water
181 was drained for 10 min and the drained wet starch was then weighed. The water binding
182 capacity was then expressed as grams per hundred grams.

183

184 **Pasting properties**

185 The method of Holm *et al.* (1988) was used to determine the pasting properties of acha starch
186 as well as the reference sample using a rapid visco analyzer (RVA 4 from Newport Scientific,
187 Warriewood, Australia). The viscosity profiles of WTAS, BLAS and WHTS were recorded
188 using starch suspensions of 9% (w/w, dry basis) and 500 g total sample weight. The starch
189 suspensions were prepared and then transferred to a rapid visco analyzer (RVA4 from
190 Newport Scientific, Warriewood, Australia). The capacity of the RVA was 50 – 50 000 cP at
191 80 rpm. A programmed heating and cooling cycle were used where the samples were held at
192 25°C for 1 min and heated to 95°C at 1.5°C/min, a holding at 95°C for 30 min before cooling
193 from 95° to 50°C, 1.5°C/min and holding at 50°C for 2 min. Parameters recorded were peak

194 viscosity, trough viscosity, final viscosity, breakdown viscosity and setback viscosity.
195 Analysis was conducted in triplicate.

196

197 **Data analysis**

198 Analysis of variance (ANOVA) was used to establish significant differences among
199 treatments. Duncan's multiple range tests was used to separate means where significant
200 difference existed (IBM SPSS 2010).

201

202 **Results and Discussion**

203 **Physicochemical properties**

204 **Proximate composition of the starches**

205 Table 1 details the proximate and amylose composition of the starches. The moisture content
206 of WHTS, BLAS and WTAS was 11.0, 11.9 and 10.7% respectively. There was no
207 significant difference between the WTAS and WHTS. The moisture content recorded for
208 acha starches was slightly lower than that reported for two acha cultivars (Fonio Hothia and
209 F. Koulli), 15 and 13% respectively. The moisture content for WTAS (10.7%) was in the
210 same range as that reported for corn starch 10.1% (Gonzales-Reyes et al 2003).

211 The protein content of WTAS (1.7%) and BLAS (1.2%) differed significantly (Table
212 1) from that of WHTS (0.5%). There was also significant difference in the protein content
213 between WTAS (1.7%) and BLAS (1.2%). Protein content of acha starch reported in
214 literature ranged between 0.3 and 0.7% for Hothia and Koulli, respectively (Carcea and
215 Acquistucci 1997). Classical studies at the University of Illinois demonstrated that the
216 variability of various chemical compounds such as the protein content in maize is of genetic,
217 environment and cultural origin and that chemical composition can thus be changed through
218 appropriate manipulation (FAO 2012). The protein content of starch is one of the critical
219 characteristics when starch is used in the manufacture of glucose syrup. The lower the
220 protein contents the better the chances of avoiding Maillard reaction (Gonzales-Reyes *et al.*,
221 2003). Although the protein content of acha starch (1.2% for BLAS and 1.7% for WTAS)
222 being significantly higher than that of wheat starch, the protein content is within the
223 acceptable range in starches that would possibly avoid Maillard reaction.

224 There was no significant difference ($p > 0.05$) between WHTS and BLAS in terms of
225 dietary fibre, 2.2% and 1.5% respectively (Table 1). There was however a significant
226 difference between BLAS (1.5%) and WTAS (0.45%). The difference in dietary fibre could
227 be attributed to genetic difference between acha cultivars. Dietary fibre recorded for WTAS

228 (0.45%) is higher than that reported in literature for lentil starches (0.21%), (Gonzales and
229 Perez 2002).

230 The fat content of WTAS and BLAS (Table 1) differed significantly from that of
231 WHTS. This difference could be due to their genetic differences. The fat content of BLAS
232 (2.9%), WTAS (2.0%) and WHTS (1.8%) were higher than that reported for corn and Okenia
233 starch, 0.17% and 0.1% respectively (Gonzales-Reyes et al 2003). Higher fat content could
234 affect gelatinization due to the formation of amylose-lipid complexes (Gonzales-Reyes et al
235 2003). The carbohydrate content of WTAS, BLAS and WHTS was 86.0, 86.0 and 88.0 g/100
236 g respectively. The carbohydrate content for WHTS and the two acha varieties did not differ
237 significantly ($p > 0.05$).

238 There are no significant differences in amylose content between BLAS (42.1%),
239 WTAS (61.4%) and WHTS (45.8%) [Table 1]. Lower amylose content was reported in
240 literature for two other varieties of acha [Hothia (22.6%) and Koulli (26.1%)] (Carcea and
241 Acquistucci 1997). In this instance the amylose content was measured in the presence of
242 lipids which may complex with amylose and reduce its iodine binding capacity (Morrison,
243 1988). Taylor et al (1997) however reported amylose content of high pasting peak viscosity
244 sorghum starch (35.7 to 36.9%) and 27.1 to 47.3% for low peak viscosity sorghum starch.
245 Environmental effects may exert more influence on amylose content than genetic differences
246 (Taylor *et al.*, 1997). A negative correlation between glycemic index, resistance starch and
247 amylose content of starch (Chung *et al.*, 2008a) was noted. Higher amylose content may
248 contribute to a lower hydrolysis index (HI) and estimated glycemic index (eGI).

249

250 **Functional properties of the starches**

251 **Viscosity**

252 Pasting properties of acha starch is indicated in Table 2. There were significant differences
253 ($p < 0.05$) in peak viscosity and final viscosity between the two acha gels. Peak viscosity is
254 the ability of starch to swell before their physical breakdown, an indication of the water
255 binding capacity of starch (Ikegwu et al 2010). The peak viscosity was 3506 cP for WHTS,
256 3994 cP for BLAS and 4936 cP for WTAS. There was a significant difference ($p < 0.05$) in
257 the peak viscosity between WTAS and BLAS. Acha starch had a higher peak viscosity than
258 that reported in literature for corn starch (Sandhu and Singh 2007). The peak viscosity of
259 various corn starches ranged between 804 and 1552 cP. The higher peak viscosity of acha
260 starch is indicative of their higher water binding capacity and thus higher degree of starch
261 swelling, an indication that acha starch may be good for products requiring high gel strength

262 and elasticity. Trough viscosity is the measurement of the holding strength of the starch paste
263 before it breaks down and viscosity decreases. This depends on the temperature and degree
264 of mixing or shear stress. The trough viscosity for two acha starch varieties was similar,
265 3125 cP and 3137 cP for BLAS and WTAS. There was however a significant difference
266 between the two acha starch varieties and wheat starch trough viscosity, 1960, 3125, 3137 cP
267 for WHTS, BLAS and WTAS, respectively. This significant difference ($p < 0.05$) in trough
268 viscosity implies a difference in the paste holding strength of wheat starch and the two acha
269 starch varieties. The holding strength of BLAS and WTAS were similar while the holding
270 strength of WHTS paste is much lower than the two acha starch paste varieties. This
271 signifies that acha starch gels may be the better option for products requiring high holding
272 strength without breaking down. The trough viscosity are much higher than the values
273 recorded for African Tall (662 cP) and Ageti (652 cP) corn starch (Sandhu and Singh 2007).

274 The break down viscosity for WHTS, BLAS and WTAS were 369, 869, 2976 cP
275 respectively. The break down viscosity for WTAS was significantly higher ($p < 0.05$) than
276 those of WHTS and BLAS. Breakdown viscosity is the measure of disintegration of cooked
277 starch. It is the difference between the peak viscosity and the trough viscosity. The higher
278 the breakdown viscosity; the lower the ability of the sample is to withstand heating and shear
279 stress during cooking (Adebowale et al. 2004). This indicates that WTAS being higher in
280 break down viscosity will have a lower ability to withstand heating and shear stress during
281 cooking.

282 The final viscosity for BLAS was 5129 cP, WTAS 5470 cP and WHTS 4436 cP. The
283 final viscosity, which is the measure of the ability of starch to form a viscous paste for
284 WTAS, was significantly higher ($p < 0.05$) than those of BLAS and WHTS. This difference
285 could be due to the difference detected in the peak time and pasting temperatures of all three
286 starches, WTAS (6.7 min, 85.9°C), BLAS (5.7 min, 80.2°C) and WHTS (4.2 min, 78.0°C).
287 The higher pasting temperatures for acha starches indicated the higher resistance of their
288 starches to swelling. The increase in viscosity with temperature may be contributed by the
289 removal of water from the exuded amylose as the starch granules swell (Sandhu & Singh,
290 2007).

291 The setback viscosity is the measure of the degree of syneresis of starch upon cooling
292 of the cooked starch paste, for WTAS, BLAS and WHTS were 2476 cP, 2332 cP and 2004 cP
293 respectively. There was no significant difference between the acha starches in setback
294 viscosity. However, the WHTS was significantly lower ($p < 0.05$) in setback viscosity
295 compared to the acha starches. The high setback values for acha starches make them

296 unsuitable for use where low syneresis rate is required, such as in frozen or refrigerated
297 foods. The setback values are also indicative of the retrogradation tendency of starch gels.
298 The higher the setback viscosity the lower the retrogradation during cooling of the products
299 made from the flour (Ikegwu et al 2010). This implies a significant difference in
300 retrogradation tendency between wheat starch and the two acha starch varieties. Acha
301 starches have lower retrogradation tendencies during cooling compared with wheat starch,
302 suggesting that the degree of re-association of the wheat starch molecules was higher than
303 that of acha starch molecules, upon cooling. Sandhu and Singh (2007) reported that the
304 pasting properties of starch depend upon various factors such as the rigidity of starch
305 granules, which in turn affects the degree of swelling of the starch granules.

306

307 **Turbidity and water binding capacity (WBC) of the starches**

308 The turbidity for WHTS, BLAS and WTAS are reported in Table 3. The values were 0.2097
309 NTU, 0.0003 NTU and 0.0193 NTU, respectively for WHTS, BLAS and WTAS. There was
310 significant difference ($p < 0.05$) in the turbidity of the starches, with WHTS being the highest
311 followed by WTAS and lastly BLAS. The turbidity of starch in foods is important when used
312 as thickener in sauces, as a carrier of flavours in beverages or as a suspending agent in liquid
313 foods. The higher the NTU value the lower the clarity of the suspension. This signifies that
314 both acha starch varieties are clearer than wheat starch, and would be the better option to use
315 in products where starch clarity is required. Sandhu and Singh (2007) reported that turbidity
316 development of starches is affected by inter-related factors such as starch granule swelling,
317 leached amylose and amylopectin, amylose and amylopectin chain length.

318 There were significant differences in the WBC (Table 3) between WHTS, BLAS and
319 WTAS (0.83, 1.33 and 1.36 g/100g, respectively), whereas there were no significant ($p >$
320 0.05) differences between BLAS and WTAS. This difference in WBC can be due to
321 structural differences in starch and proteins of the starches (Celik et al 2005). Different
322 proportions of crystalline and amorphous regions within the granules may be the result of the
323 variations in WBC. Thus weakly bonded amorphous starch granules will imbibe less water
324 (Carcea and Acquistucci, 1997). It appears that acha starch granules possess stronger bonded
325 amorphous granules compared to wheat starch. The WBC of the starch affects important
326 physical attributes such as the viscosity of sauces and batters and the texture of baked
327 products. Sandhu and Singh (2007) reported WBC for starches from different corn varieties
328 in the range of 0.82 to 0.97 g/100 g. The water binding capacity for acha starch is higher than
329 the values reported by Sandhu and Singh (2007). Acha is known to swell far more than other

330 cereals (Carcea and Acquistucci, 1997). Acha starch can thus be included as part of the
331 ingredients in the manufacturing of sauces, batters, dough's and baked products. It can be
332 used as thickener, bulking agent and most importantly due to its high WBC as water retention
333 agent.

334

335 **Textural properties of the starches**

336 The textural characteristics of the acha and wheat starches are indicated in Table 4. The
337 hardness for WTAS, BLAS and WHTS were 0.9367 N, 0.9467 N and 0.9500 N, respectively.
338 No significant difference in gel hardness was observed between BLAS gel and WTAS gel or
339 between the WHTS and the two acha starch gels. These low hardness values indicate that
340 both acha and wheat starches produced soft gels. Acha starch would thus be suitable for
341 manufacturing of foodstuff where soft gels are required. Sandhu and Singh (2007 reported
342 that the gel firmness is mainly caused by retrogradation which is associated with
343 crystallization of amylopectin, leading to harder gels. The resilience value is the first force
344 before the maximum force (F) which is the capability of the gel to return to an original shape
345 or position after having been compressed. The resilience for BLAS and WTAS and WHTS
346 gels were 1.1994, 1.2399 and 1.2402, respectively. There was no significant difference ($p >$
347 0.05) in resilience between the gels. Springiness for BLAS gel was 0.7090 mm, WTAS gel
348 was 0.7090 mm and WHTS was 0.3746 mm. There was no significant difference ($p >$ 0.05)
349 in the springiness of both acha starch gels and WHTS gels. Springiness refers to the degree
350 of gel rubberiness in the mouth (Lau et al 2000). A high degree of springiness is depicted by
351 gel structure that breaks into large pieces during the initial compression; whereas a low
352 degree of springiness is represented by a gel that breaks into smaller pieces during the initial
353 compression test (Huang et al 2007). The starch gels broke into large pieces during
354 compression. Hence, both acha and wheat gels are highly springy. Highly springy gels
355 would not break down easily during mastication (Lau *et al.*, 2000). Acha starch gels were
356 springy and resilient. This indicates that both acha starch varieties can be used for hot and
357 cold desserts as well as for soft jelly like sweets and confectionery toppings.

358

359 **Thermal properties of acha starch**

360 **Gelatinization of acha starch**

361 The gelatinization temperatures (T_o , T_p , and T_c) and enthalpy (ΔH) of wheat and acha
362 starches are shown in Table 5. There was no significant difference ($p >$ 0.05) in peak and
363 onset temperature between the acha and wheat starches. WHTS had the highest onset (T_o)

364 and peak (T_p) temperatures 34.8°C and 74.6°C, although not significant. WTAS and BLAS
365 were 25.1, 26.8°C and 68.2, 70.8°C for T_o and T_p respectively. Acha starches and the wheat
366 starch differed significantly ($p < 0.05$) in conclusion T_c and T_p temperatures. For WHTS it
367 was 96.0°C, BLAS was 87.8°C and for WTAS was 93.7°C. The gelatinization temperatures
368 of waxy maize starch were 59.9, 69.1 and 78.1°C for T_o , T_p and, T_c respectively (Miao et al
369 2009). The onset gelatinization temperature of acha starch is much lower than that of waxy
370 maize starch. This means that less energy is needed to induce gelatinization of acha starches.
371 The peak gelatinization temperature of acha starch is the same as that reported in literature
372 (Miao et al 2009). Tester and Morrison (1990) reported that due to the structural differences
373 in amylopectin, starches with the low gelatinization temperature has less crystallinity than the
374 high gelatinization temperature starches, this was attributed to structural differences in their
375 amylopectins. Hence, acha starches may have lower degree of crystallinity compared to
376 wheat starch. Ezekiel et al (2007) reported that low transition temperatures of between 50-
377 67°C was due to the lower crystallinity, meaning that the higher the transition temperatures
378 the higher the degree of crystallinity. This would lead to structural stability causing the
379 starch granules to have a higher degree of resistance to gelatinization. Acha starches have
380 low transition temperature which implies a low crystallinity and high amorphous regions
381 (Miao et al 2009). The gelatinization temperature range ($T_c - T_o$) for WTAS and BLAS was
382 66.9 and 62.8°C respectively, whereas that of WHTS was 58.3°C. The similarity of the
383 gelatinization temperature range amongst the starches could be attributed to possible
384 similarity in protein content and starch structure. The melting enthalpy (ΔH) of gelatinization
385 of BLAS was 188.41 J/kg and WTAS was 691.6 J/kg whereas that of WHTS was 213.21
386 J/kg. The melting enthalpy values for WTAS and BLAS differed significantly ($p < 0.05$).
387 The melting enthalpy for both acha starch varieties as well as WHTS was higher than that
388 reported for waxy maize starch (Chung et al 2008b). The higher melting enthalpy could be
389 due to the difference in the alignments of the hydrogen bonds in the starch molecules. This is
390 caused by the difference in the bonding forces between the double helices that form the
391 amylopectin crystallites (Sandhu and Singh 2007). Sandhu and Singh (2007) reported
392 melting enthalpy values between 11200 and 12700 J/kg for various corn starches. This is
393 much higher than the melting enthalpy of acha starch.

394

395 **Retrogradation properties of acha starch**

396 The retrogradation of acha starch compared to that of wheat starch is summarised in Table 6.

397 Retrogradation is the hydrogen bonding between starch chains that occurs after cooling of
398 gelatinized starch paste (Hoover and Zhou 2003). The transition temperatures of
399 retrogradation for both acha starches and WHTS are much lower than the transition
400 temperatures of gelatinization. The onset temperature for WHTS, WHTS and BLAS were
401 19.97, 17.58 and 19.05°C respectively. WHTS had the highest onset temperature followed
402 by BLAS and WTAS. However, the differences were not significant ($p > 0.05$). Sandhu and
403 Singh (2007) reported onset temperatures for retrogradation in the range between 41.50 and
404 43.10°C for two corn starch varieties. These values are much higher than that recorded for
405 white acha and black acha starches. According to Sandhu and Singh (2007) this could be due
406 to a less organized manner of recrystallization of the amylopectin branched chains of the gels.
407 The lower onset temperature for the acha starches could be due to weaker molecular bonding
408 of the starch chains.

409 Values for peak temperature of retrogradation were 64.13, 67.05 and 64.84°C for
410 WHTS, WTAS and BLAS, respectively. There was no significant difference ($p > 0.05$)
411 between the starches in peak temperatures. Literature values for different corn varieties
412 varied between 52.40 and 54.50°C for African Tall corn starch and Partap corn starch
413 respectively. These values were much lower than that recorded for white acha and black acha
414 starch. Higher peak temperatures of acha starches could be indicative of the need for higher
415 thermal energy for retrogradation.

416 Retrogradation enthalpy is an indication of the disentanglement and melting of the
417 double helices formed during cooling (Adebowale et al 2004). The retrogradation enthalpy
418 (ΔH) for WTAS was 37.61 J/kg and for BLAS was 118.00 J/kg. There was a significant
419 difference between BLAS and WTAS, with BLAS being higher than that of WTAS. No
420 significant difference was observed for retrogradation enthalpy between WHTS and WTAS.
421 The differences in the retrogradation enthalpy could be due to the difference in the degree of
422 starch crystallinity of the retrograded starches (Sasaki et al 2000).

423 The conclusion melting temperature values for WHTS, WTAS and BLAS were
424 94.73°C, 94.82°C and 95.95°C respectively. There was no significant difference in
425 conclusion temperature of retrogradation between the acha starches. Whenever a material
426 undergoes a change in physical state for example melting, or transforms from one crystalline
427 form to another, or when it reacts chemically, heat is either absorbed (endothermic) or
428 liberated (exothermic) (Karim et al 2000). Wheat and both acha starch cultivars presented an
429 endothermic transition (retrogradation). Retrogradation has been identified as the
430 dissociation of the amylopectin crystallites (Gonzales et al 2002). The retrogradation

431 phenomenon is a process of re-crystallization of the starch polymers (Gonzales-Reyes *et al.*,
432 2003). In the case of retrograded starch, ΔH (J/kg) provides a quantitative measure of the
433 energy transformation that occurs during the melting of recrystallized amylopectin as well as
434 precise measurement of the transition temperatures (onset, T_o ; peak, T_p and conclusion (end),
435 T_c) (Karim et al 2000). There is a significant difference in the ΔH and peak height values
436 between white and black acha starches. There were however no significant difference
437 amongst the retrogradation temperature profile (T_o , T_p and T_c) for both white and black acha
438 starch.

439

440 **Conclusion**

441 Acha starches differ in moisture, dietary fibre and carbohydrate content and are similar to
442 wheat starch in amylose content. Water absorption capacity of acha starches were higher
443 than that of wheat starch and hence acha starch can be included as part of the ingredients in
444 the manufacturing of sauces, batters, and dough's and baked products. It can be used as
445 thickener, bulking agent and most importantly due to its high WBC as water retention agent.
446 Acha starch gels were springy and resilient indicating its potential use in hot and cold
447 desserts as well as for soft jelly like sweets and confectionery toppings. Acha starch cultivars
448 and wheat starch are similar in thermal properties. Acha starch has similar retrogradation
449 temperature profiles as that of wheat starch. Wheat and both acha starch cultivars presented
450 an endothermic transition (retrogradation).

451

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457

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535 Table 1. Proximate and amylose composition of acha starch^{1,2}

Starch	Proximate (%)				Amylose (%)
	Moisture	Fat	Carbohydrate	Protein	
Wheat	11.0 ± 0.00 ^a	0.5 ± 0.00 ^a	88.0 ± 0.00 ^a	0.47 ± 0.47 ^a	45.8 ± 16.56 ^a
Black acha	11.9 ± 0.25 ^b	0.8 ± 0.00 ^a	86.0 ± 0.00 ^a	1.23 ± 1.23 ^b	42.1 ± 16.35 ^a
White acha	10.7 ± 0.26 ^a	1.3 ± 0.00 ^a	86.0 ± 0.00 ^a	1.73 ± 1.73 ^c	61.3 ± 15.46 ^a

536 ¹Values are mean ± standard deviation.537 ²Any two means with different superscript in each column differ significantly (p < 0.05).

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Table 2. Pasting properties of Acha starch^{1,2}

Sample	Peak1 (cp)	Trough1 (cp)	Breakdown Viscosity (cp)	Final Viscosity (cp)	Setback Viscosity (cp)	Peak Time (min)	Pasting Temp. (°C)
Wheat	3506 ± 16 ^a	1960 ± 41 ^a	369 ± 21 ^a	4436 ± 82 ^a	2004 ± 62 ^a	4.18 ± 0.04 ^a	78.03 ± 0.46 ^a
Black acha	3994 ± 36 ^b	3125 ± 15 ^b	869 ± 34 ^b	5129 ± 60 ^b	2332 ± 54 ^b	5.78 ± 0.10 ^b	80.15 ± 0.44 ^b
White acha	4936 ± 58 ^c	3137 ± 6 ^b	2976 ± 95 ^c	5470 ± 50 ^c	2476 ± 190 ^b	6.69 ± 0.03 ^c	85.88 ± 0.49 ^c

¹Values are mean ± standard deviation.

²Any two means with different superscript in each column differ significantly ($p < 0.05$)

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551 Table 3 Turbidity and water binding capacity of acha starch^{1,2}

Starch	Turbidity(NTU) ³	Water binding capacity (g/100 g)
Black acha	0.0003 ± 0.001 ^a	1.333 ± 0.06 ^b
White acha	0.0193 ± 0.002 ^b	1.367 ± 0.06 ^b
Wheat	0.2097 ± 0.009 ^c	0.833 ± 0.23 ^a

552 ¹Values are mean ± standard deviation.553 ²Any two means with different superscript in each column differ significantly (p < 0.05).554 ³NTU = Nephelometric turbidity units

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560 Table 4. Textural characteristics of acha and wheat starch gels^{1,2}

Cereal starch	Hardness(N)	Resilience(ratio)	Springiness (mm)
White acha	0.936 ± 0.041 ^a	1.199 ± 0.090 ^a	0.709 ± 0.112 ^a
Black acha	0.946 ± 0.015 ^a	1.239 ± 0.087 ^a	0.709 ± 0.132 ^a
Wheat	0.950 ± 0.010 ^a	1.240 ± 0.033 ^a	0.374 ± 0.273 ^a

561 ¹Values are mean ± standard deviation.562 ²Any two means with different superscript in each column differ significantly (p < 0.05).

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Table 5 Gelatinization of acha starch ^{1,2}					
Starch	To (°C)	Tp (°C)	Tc (°C)	ΔH_{gel} J/kg	$\Delta T_r = (T_c - T_o)$
Wheat	34.7 ± 25.20 ^a	74.56 ± 0.97 ^a	96.03 ± 0.49 ^b	213.21 ± 169 ^a	58.33 ± 24.71 ^a
White acha	25.05 ± 12.12 ^a	68.20 ± 1.64 ^a	93.67 ± 0.57 ^b	188.42 ± 250 ^b	66.92 ± 11.55 ^{ab}
Black acha	26.75 ± 7.56 ^a	70.54 ± 6.07 ^a	87.85 ± 2.15 ^a	691.04 ± 106 ^c	62.8 ± 5.41 ^a

570 ¹Values are mean ± standard deviation

571 ²Any two means with different superscript in each column differ significantly ($p < 0.05$). To,
572 onset temperature; Tp, peak temperature; H1, peak height; ΔH , enthalpy of gelatinization; Tc,
573 end (conclusion) temperature, ΔT_r gelatinization temperature range ($T_c - T_o$)

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Table 6		Retrogradation of cereal starch ^{1, 2}			
Starch	To (°C)	Tp (°C)	Tc (°C)	Δ Hretro J/kg	ΔTr = (Tc- To)
Wheat	17.97 ± 0.02 ^a	64.13 ± 5.35 ^a	94.73 ± 1.78 ^a	56.87 ± 32.62 ^{ab}	76.76 ± 1.76 ^a
White acha	17.58 ± 0.48 ^a	67.06 ± 4.58 ^a	94.83 ± 1.48 ^a	37.61 ± 12.88 ^a	77.25 ± 1.00 ^a
Black acha	19.05 ± 1.30 ^a	64.84 ± 0.53 ^a	95.95 ± 0.52 ^a	118.00 ± 40.58 ^b	76.90 ± 0.78 ^a

583 ¹Values are mean ± standard deviation.

584 ²Any two means with different superscript in each column differ significantly (p < 0.05). To,
585 onset temperature; Tp, peak temperature; H1, peak height; H, ΔH, enthalpy of gelatinization;
586 ΔHf, Tc, end (conclusion) temperature

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