<u>Original Research Article</u> Effect of Acetylation on the Physical and Functional Properties of Industrial and Laboratory Cassava (*Manihot esculenta*) Starches

ABSTRACT

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> Aim: Acetylated potato starch (APS) is commercially available and used widely in the food industries. It is imperative to study to the physical and functional properties of acetylated cassava (Manihot esculent Qultivar Tropical Manihot Series (TMS) 30572 and industrial starches for possible substitution/ replacement of expensive APS in food system. Study Design: The properties of acetylated cassava starches were compared with those of commercially available acetylated potato starch (APS) and native cassava starches. Place and Duration of Study: The experiment was performed in the Department of Food Science and Technology, Federal University of Technology, Akure Nigeria from June 2011 and January 2013. Methodology: Industrial starch and starch extracted from cassava TMS30572 were acetylated by standard procedure. The acetylated starches were analyzed for the physical as well as functional properties. Results: The yields after acetylation ranged between 96-98% and 80-93% for TMS30572 and industrial starches, respectively. Acetylated cassava starches showed improved physical and functional properties over the native cassava starch and these increased with increasing concentration of acetic anhydride in the reaction medium. At >2.50% acetylation, starch concentration of 5.5% had the same hot paste viscosity of 1500cPa.s with commercial APS at 5% concentration. Also at 2.50% acetylation the starch was stable until the third freeze-thaw cycles and exhibited better stability than commercial APS. Conclusion: Acetylation improved the yield of starch from cassava during processing. The industrial starch showed higher degree of acetylation than TMS30572 starch under the same experimental condition. Acetylated cassava starches (at >2.50-2.70% acetylation) has improved functional properties and lesser tendency towards retrogradation thus could be a potential replacement to the more expensive APS as ingredient in food system. Keywords: Keywords: cassava cultivar; food industry; freeze-thaw stability; retrogradation **1. INTRODUCTION**

20 21 Cassava (*Manihot esculenta* Chtz) is widely cultivated along the tropical belt for its starchy 22 tubers which are used as food, feed or as an industrial raw material. Starches are used in a 23 wide range of food products and its incorporation into food system is primarily governed by 24 factors such as gelation, pasting properties, apparent viscosity, solubility, swelling power and 25 clarity [1].

Only few percentages of the world crop of starch are used in their native state. Modification is usually carried out to introduce the desired properties and or remove certain inherent undesirable characteristics of the native starches. Genetic and chemical modification produces functionally tailored starch products that meet specific application in the food industries and this has led to expanded usage of starch and its products [2, 3]. Modified (acetylated) starch cannot be readily broken down by digestive enzymes [4], thereby making it a desirable functional resistant starch with great health benefits [5].

Acetylation involves esterification of the hydroxyl functional group of starch to produce starch with altered polarity, lower pasting temperature and improved paste clarity and freeze-thaw properties. The detrimental syneresis and retrogradation effects are greatly reduced in acetylated starch thus making it very useful in frozen foods [6-8].

There is limited information on the effects of using graded amount of acetic anhydride to modify cassava starch and the properties of such modified starches. The present study aims at extraction and modification of cassava starch by acetylation technique using graded levels of acetic anhydride, and investigation of the changes in the functional and physical properties of the starch due to acetylated potato starch in order to determine the suitability of the cheaply available cassava starch as alternative to APS in the food industry.

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45 2. MATERIAL AND METHODS

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47 **2.1 Source of Materials**

48 Cassava cultiva PIS 30572 was planted and harvested after 18 months on the research 49 farm of The Federal College of Agriculture, Akure, Nigeria. The industria parch was 50 obtained from Matna Food Company Limited, Akure, Ondo State, Nigeria. A 2.5% acetylated 51 potato starch was obtained from FAN MILK Plc. Ibadan, Nigeria.

52 2.2 Starch Extraction

53 Starches were extracted from cassava tubers according to the procedure described by ORDYLAS (1988).[9] The starch obtained was dried in a hot air oven (Labcon air oven 55 model) at 55±2 °C for 48 h, then pulverized and sieved using 254µm sieve.

56 2.3 Starch Acetylation

57 Starch acetylation was carried out using the method of Wurzburg [10] as modified by 58 Golachowski [7]. Two hundred grams (200g; dry weight basis) of native cassava starch or 59 TMS30572 cultivar was dispersed in distilled water the pH was adjusted to 8.0 with 3% 60 Naph Predetermined volume of acetic anhydride was added to the slurry at a rate of 1m m. The pH was finally adjusted to 5.4 with 10% HCl. The starch was centrifuged at 61 1,000 x g and the residue washed with distilled water and dried at 30°C for 24 h to determine 62 the yield. The dried acetylated starch samples were then pulverized, sieved to pass through 63 254µm sieve packaged in plastic containers and kept in cool dry place for further analyses. 64 65 In order to determine the starch (dry weight basis) was added to 65m bistilled water followed by 25mL of 0.5M NaOH with continuous 66 mixing using a magnetic stirrer for 30 min as described by Golachowski [7]. 67

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70 2.4 Determination of Bulk Density and Sedimentation Volume

The bulk density was determined using the procedure of Narayana & Narasinga [11] with slight modification. Sedimentation volume was determined as described by Raja et al. [12].

73 2.5 Determination of Water and Oil Absorption Capacity

A suspension of 1 g of prch (dry weight basis) was made in 10 ml of distilled water [or 10 mL of oil (executive chef® oil with density of 0.92 g/mL)]. The water and oil absorption capacities of the starch samples were determined as described by Sathe et al. [13].

77 2.6 Determination of Viscosity, Swelling Power and Solubility

Starch suspension of 5% (w/v) in distilled water was heated 0°C in a water bath with continuous stirring. The paste was transferred to a roterry viscometer and paste viscosity was measured from 90°C through to 30°C at 10°C reval of the cooling phase and expressed as centi Pascal secon ing the method of Amani et al. [14]. The method described by Osundahunsi & Mueller [15] was then used in the determination of solubility and swelling wer of the starches.

84 2.7 Determination of Paste Clarity and Freeze-Thaw Stability

85 The paste clarity was determined using the procedure of CRAIG et al. (1989) [16]. On dry weight basis, 1% aqueous dispersion of starch in distilled water was boiled at 100°C for 30 86 min under constant stirring. The paste was cooled the of and the percentage transmittance 87 was measured at 640 nm. The freeze-thaw stability was investigated using the method of 88 89 Singh & Kaur [17]. Aqueous suspension of 5% w/v (dry weight basis) was prepared using 90 distilled water. The suspension was heated at 95°C for 30 min in a water bath and then 91 booled with continuous stirring in order to prevent skin formation on the paste. The paste 92 was then subjected to a 5-cycle alternate freezing and thawing of 18 h and 3 h respectively. At the end of each cycle, the paste was centrifuged at 5,000 x g for 10 min and the amount 93 of exudates calculated in percentage was determined and plotted against the number of 94 95 freeze-thaw cycle.

96 **2.8 Statistical analyses**

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The data were subjected to analysis of variance (ANOVA) and the means were separated using Duncan's multiple range tests using SPSS statistical package version 17.

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106 **3. RESULTS AND DISCUSSION**

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The degree of acetylation increased as the volume of acetic anhydride increased (Fig. 1). This may partly be due to the larger surface area for diffusion and absorption of the acetyl groups on the starch molecules. High concentration of acetic anhydride resulted in high molecular collision rate leading to greater availability of acetic anhydride molecules in the vicinity of starch [18]. Similar observations have been reported for complete rasping processing of industrial starch led to higher reaction efficiency compared to the **IS30572**

114 **Outivar**, which resulted in higher degree of acetylation when the same amount of acetic 115 anhydride was added. Mechanical processing of industrial starch causes damage to the 116 granule structure thus increasing the susceptibility to enzymatic digestion than the 117 undamaged starch granules [20].



Fig. 1: Fig. 1

121 3.1 Effect of Acetylation on Yield, Sedimentation and Bulk Density

As shown in Table we yields of acetylated industrial starches (80 - 93%) were lower than TMS30572 (96-98%) starches due to the high concentration of impurities in industrial starches which were removed during the acetylation. The highest yield was observed at acetylation of 2.99% and 3.98% for TMS30572, and 3.60% for the industrial starch. The yields obtained were higher than the range of 82 - 88% reported earlier [18]. The sulphuric acid used as a catalyst has the tendency to cause extensive solubilisation of the crystalline structure during acetylation process and might have contributed to the observed higher yield.

129 The bulk density and sedimentation volume increased with increase in the degree of 130 acetylation of the starches (Table 1). Narayana & Narasinga [11] reported similar result for 131 winged beans flour. The sedimentation volume of the acetylated industrial starches was 132 significantly higher than the native starches. Hence, acetylated starches will exhibit better 133 cold water swelling than the native cassava starch.

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Table 1. Tweffect of acetylation on the yield, sedimentation and bulk density of cassava starches

| | Laboratory | / (TMS 30572) stare | ch | Industrial starch | | | |
|------------|------------|----------------------|---------------------------|-------------------|-----------|-----------------------|---------------------------|
| DoA (%) | Yield (%) | Sedmentation (mL) | Bulk Density (g/mL) | DoA (%) | Yield (%) | Sedimentation (mL) | Bulk Density (g/mL) |

| 0 | 98.70±0.04 | 6.00±0.82 ^c | 0.81±0.01 ^b | 0 | 92.00±1.21 | 4.50±0.16 ^d | 0.71±0.03 ^c | | | |
|---|------------|------------------------|------------------------|------|------------|------------------------|------------------------|--|--|--|
| 0.43 | 96.00±0.08 | 7.00±0.41 [°] | 0.81±0.01 ^b | 0.65 | 80.00±1.70 | 6.50±0.82 ^c | 0.72±0.01 [°] | | | |
| 1.4 | 96.00±0.05 | 7.50±0.94 ^b | 0.83±0.02 ^a | 1.5 | 88.00±0.80 | 7.00±0.16 ^c | 0.78±0.04 ^b | | | |
| 2.5 | 96.00±0.05 | 7.50±0.84 ^b | 0.83±0.02 ^a | 2.74 | 90.00±0.80 | 7.50±0.14 ^b | 0.82±0.02 ^a | | | |
| 2.99 | 98.00±0.08 | 7.50±0.62 ^b | 0.83±0.05 ^a | 3.6 | 93.00±1.40 | 7.50±0.08 ^b | 0.82±0.05 ^ª | | | |
| 3.98 | 98.00±0.08 | 7.90±0.82 ^a | 0.84±0.05 ^a | 4.09 | 90.00±0.90 | 7.70±0.05 ^b | 0.83±0.01 ^ª | | | |
| 4.57 | 97.00±0.05 | 8.00±0.62 ^a | 0.84±0.02 ^a | 4.62 | 90.00±0.80 | 8.10±0.05 ^a | 0.84±0.03 ^a | | | |
| APS | ND | 8.00±0.47 ^a | 0.78±0.02 ^c | APS | ND | 8.00±0.04 ^a | 0.78±0.02 ^b | | | |
| Values ar \mathbb{D} ean ±SEM (n=3). Mean values with different letters in a column are significantly different (p≤0.05).DoA: Degree of acetylation; APS: acetylated potato starch; ND: not determined. | | | | | | | | | | |
| (p=0.05).DoA. Degree of acelylation, AFS. acelylated polato starch, ND. not determined. | | | | | | | | | | |

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137 **3.2 Swelling Power, Solubility, Water and Oil Absorption Capacities**

138 The swelling powers of the starches were investigated at temperature range of $50 - 90^{\circ}$ C, representing the pasting range of most starches. Native starch exhibited an increase in 139 140 swelling power with increasing temperature as shown in Fig. 2A and 2B. Swelling power has 141 been reported to correlate with solubility in different varieties of sweet potato starches [21]. Acetylation increased the swelling power at any given temperature perhaps due to the 142 loosening of intra- and inter-molecular bonds in the starch molecules, which eventually 143 144 increased the swelling capacities of the starches. However, there was no significant 145 difference in the swelling power of the acetylated starch at temperature >60°C when the 146 degree of acetylation is above 2.50% for TMS30572 cultivar. Limited water available within 147 the mixture at higher temperature may affect the swelling power as complete gelatinization 148 and swelling do occur in excess water [20]. Solubility increased with increase in the degree 149 of acetylation. There was a sudden rise in the solubility of the starch acetylated at 1.4% and 150 1.5% for TMS 30572 and industrial starches respectively at 60°C. At temperatures >60 °C, the solubility could not be measured as shown in Fig. 2C and 2D. This might be due to total 151 percolation of water into the starch granules, resulting in limited amount of water available in 152 the mixture since acetylation lowers the gelatinization temperature. Singh et al. [22] have 153 reported that acetylation decreased the gelatinization temperature of acetylated sorghum 154 155 starch.

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Fig. 2: The effect of temperature on the swelling power and solubility of cassava starches at different degrees of tylation for TMS 30572 (A and C) and industrial starches (B and D). Values represent mean \pm SD, (n=3). Mean values with the same alphabet within a particular temperature are not significantly (p \leq 0.05) different according to Duncan multiple range test.

163 The water and oil absorption capacities (WAC and OAC) of the starches are shown in Fig. 3. 164 All the starches absorbed more oil than water when acetylated. Similar result has been 165 reported for acetylated sweet potato starch [23]. The result showed that acetylation altered 166 the starch polarity and leads to the introduction of hydrophobic group into the starch 167 molecules.



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Fig. 3: T fig.

174 3.3 Paste Viscosity and Clarity

175 The viscosity of cooked starch is important to the food industry. The peak viscosity of APS is significantly higher than the cassava starches. As the temperature drops from 90°C to 70°C 176 177 the viscosity remained fairly constant before it began to increase as the temperature falls below 70°C for both starches (Figure 4A and B). The pative starch showed higher paste 178 viscosity than the acetylated starches when cooled to C. Both acetylated TMS30572 179 180 cultivar and industrial starches showed lesser cooling viscosity suggesting reduced tendency towards retrogradation than the commercial acetylated potato starch (APS). Similar 181 182 observation had been reported for sorghum starch where acetylation led to a decrease in the 183 paste viscosity [22]. Substituent groups formed at lower temperature restricted the tendency of the starch molecules to realign after cooling, thus facilitating lower setback value and 184 185 retrogradation for acetylated starches [24].



Fig. 4: The effect of temperature and starch concentration on paste viscosity of cassava starches (A) TMS 30572, (B) industrial starch at different degrees of acetylation, (C) the fect of starch concentration on the paste viscosity of TMS 30572 starch acetylated at 2.5% and industrial starch acetylated at 2.74% compared with APS (acetylated potato starch) at 5% starch concentration. Values represent mean±SEM, (n=3). Mean values with the same alphabet across the temperature ranges are not significantly (p≤0.05) different according to Duncan multiple range test.

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In attempt to determine the starch concentration that will match-up in hot paste viscosity to the acetylated potato starch, cassava starch at different concentration were acetylated at 2.5% and the hot paste viscosity determined at 90°C. At 5% starch concentration, the hot paste viscosity was lower than that of the acetylated potato starch at 5% concentration (Fig. 4C). However as the starch concentration increased, the hot paste viscosity also increased. The light transmittance values of the pastes increased progressively with increase in the degree of acetylation for all the starches (Fig. 5). Increase in the degree of swelling and

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dispersion of the starch granules as well as reduced retrogradation tendency may also be responsible for the observed results [8, 17].



Fig. 5: 🗠

effect of acetylation on the paste clarity of laboratory (TMS 30572) starch 205 (A) and industrial starch (B) measured as percentage transmittance. Values represent 206 mean \pm SD, (n=3). Mean values with the same alphabet are not significantly (p≤0.05) 207 208 different according to Duncan multiple range test.

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210 3.4 Freeze-Thaw Stability

211 The pastes of roots and tubers starches are known to exhibit great stability under normal 212 condition but cannot survive freeze-thaw cycles [25]. This poor freeze-thaw stability of native 213 starch limits its application as a thickening agent in frozen foods [26]. Cassava starches 214 exhibited unusual stability at higher degree of acetylation. TMS 30572 cultivar at acetylation 215 level of 2.50% did not show any syneresis until the third freeze-thaw cycle when it exuded 216 13% water. At ≥3.98% degree of acetylation, there was no syneresis until the fourth freeze-217 thaw cycle with 16% exudates (Fig. 6). The waxy corn and Peruvian carrot starch gels 218 exhibited similar freeze-thaw stability [26]. The percentage exudates decreased as the 219 degree of acetylation increased. Jacobson et al. [27] have observed that in native pastes, 220 the hydroxyl groups of starch gradually re-associate through hydrogen bonding and 221 crystallize out of solution leading to syneresis when paste is allowed to cool or when 222 refrigerated. Commercial acetylated potato starch exhibited the lowest stability with greater 223 exudates due to syneresis when compared with all the acetylated cassava starches.

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Fig. 6: The effect of acetylation on the freeze-thaw stability of laboratory (TMS 30572) starch (A) and industrial starch (B). Values represent mean \pm SD, (n=3). Mean values with the same alphabet at a particular day are not significantly (p≤0.05) different according to Duncan multiple range test.

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239 4. CONCLUSION

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241 Processing methods used for cassava starch isolation have influence on reaction efficiency 242 of the final products. There was over 80% yield after acetylation, with improvement on the physical properties of native cassava starch. The functional properties such as swelling, 243 solubility, viscosity, clarity, and freeze-thaw stability were also enhanced; making it a 244 potential substitute to commercial APS in food system since acetylated starch has very low 245 246 impact on the sensory properties of food. Also, due to increase in freeze-thaw stability, 247 cassava starches with acetylation level ≥2.50% may serve as a good substitute for APS in 248 frozen food industry.

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