

Total phenolic content, antioxidant activity, shelf-life and reconstitutability of okra seeds powder: influence of milling and sieving processes

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Total phenolic content, antioxidant activity, shelf-life, and reconstitutability of okra seeds
 powder: influence of milling and sieving processes

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13 Abstract

This study aimed at relating phytochemical properties (total phenolic content and antioxidant activity), reconstitutability (water absorption capacity and water solubility index), and preservation ability (water activity) of okra seed powders to physicochemical properties (particle size distribution and moisture content). Okra seed powders were produced at three different milling frequencies, then each obtained sample was sieved with 180, 315, and 500 µm sieves. An increased milling frequency resulted in lower median particle size, water activity, and moisture content. The maximal water activity and moisture content were 0.583 ± 0.010 and 12.07 ± 0.05 % (w/w), respectively, showing the good preservation ability of okra seed powders. Total phenolic content and antioxidant activity were raised at higher milling frequency. Reconstitutability was significantly enhanced at higher milling frequency and/or for smaller median particle size. Thus, the successive milling and sieving process was successful in improving physicochemical and functional properties of okra seed powders, especially for smaller particles.

Keywords: okra seed powder; particle size; polyphenols; antioxidant activity; preservation;
 powder reconstitution.

⁵ 29 **1. Introduction**

Okra fruit (*Abelmoschus esculentus* (L.) Moench) is widely consumed around the world and it
 is cultivated in tropical and subtropical regions (Torkpo *et al.*, 2009; *Jarret et al.*, 2011). Okra

is renowned for its nutritional potential and health benefits: chemical composition analyses of okra revealed its richness in proteins, fat, minerals (Mn, Ca, Mg, Fe, Cu), and vitamins (B2, B3, B6, B9, C) (Moyin-Jesu et al., 2007; Xia et al., 2015). Mature okra seed is a good source of oil rich in unsaturated fatty acids such linolenic and linoleic acids, which are both essential for humans (Ndangui et al., 2010; Kumar et al., 2013). It has been reported in the literature that okra pods have high contents in polysaccharides and phenolic compounds including flavonoids (Arapitsas, 2008). Phenolic compounds are known to possess strong antioxidant and anti-stress effects (Ahmed et al., 2016; Sabitha et al., 2011). Although the high phytochemical-activity content of okra pods has recently been proved (Hu et al., 2014; Xia et al., 2015, 2018), little is known about the distribution of bioactive compounds within granulometric classes of okra seeds powdered by milling.

Nowadays, milling is commonly used for the production of new food ingredients (Karam *et al.*, 2016; Becker et al., 2017; Deli et al., 2019). Powdered plant parts are used as ingredients by many industries in the food, pharmaceutical, and cosmetics fields (Leuenberger et al., 2005). Moreover, food ingredients and pharmaceutical excipients are often manufactured in powder form (Siliveru et al., 2017; Deli et al., 2019). Advantages of the powder technology resides in the convenience offered by powdered products, especially regarding higher shelf-life and easier transportability (Forny et al., 2011). Successive drying, milling, and sieving processes have shown a great potential in the production of food powders with enhanced functional properties (Brewer et al., 2014; Becker et al., 2017; Deli et al., 2019; Waiss et al., 2020). It leads to the production of microparticles from plant material (whole plant, aerial parts, stems, roots, seeds, and/or fruits). It is worth noting that drying, milling, and sieving processes have a great impact on physicochemical and functional characteristics of produced powders (Landillon et al., 2008), such as particle size and shape distributions, surface roughness, density, cohesion, compressibility, water solubility, wettability, surface reactivity, and hygroscopicity (Becker et al., 2017; Deli et al., 2019). A previous study performed with the same okra powders as in the current study showed a strong correlation between median particle size and proximate composition: smaller granulometric classes contained more proteins and fat, but less water, carbohydrates, and minerals (Waiss et al., 2020). Also, the influence of milling frequency and median particle size of okra seeds powder on flow properties was highlighted (Waiss *et al.*, 2020): powders with large median particle sizes were not cohesive and had better flowability. Functional properties of powders (flowability, preservation ability, reconstitutability, etc.) are very important for the pharmaceutical and food industries (Räsänen et al., 2003), as they

determine the conditions by which they should be manufactured, handled, stored, and reconstituted with a view to protect bioactive molecules (Deli et al., 2019a; Meghwal et al., 2014; Petit et al., 2017). Powder physicochemical characteristics are often strongly influenced by particle size and physical structure (Meghwal and Goswami, 2014; Deli et al., 2019). Thus, it is of great importance to determine the physical and hydration characteristics of food powders in order to predict their behavior during storage and reconstitution and use them efficiently for an efficient use as functional food ingredients (Benković et al., 2017). The current study was intended to increase knowledge about the influence of particle size on phytochemical properties, reconstitutability, and preservation ability of okra seeds powder. Okra seeds were ground at three milling frequencies (6 000, 12 000, and 18 000 rpm); then, each powder sample was fractionated in four granulometric sizes by sieving: $\leq 180 \mu m$, 180 - 315 μm , 315 - 500 μ m, and \geq 500 μ m. Last, physicochemical properties (particle size distribution and moisture content), ability to interact with water (water activity, water absorption capacity, and water solubility index), and phytochemical properties (total phenolic content and antioxidant activity) of powders were evaluated.

80 2. Material and methods

2.1. Material: okra seeds

Okra seeds were purchased from farmers of Impfondo locality, in the north of the Republic of
the Congo (1° 32'28" N; 18°07'10" E). Then, the okra seeds were removed from the pods and
sun-dried in the open air at 30 °C. Dry seeds were stored at 10 °C until use for powder
production.

2.2. Drying and milling of okra seeds

87 500 g dried okra seeds were milled with an ultra-centrifugal mill ZM 200 (Retsch, France) using
88 a 10-cm sieve drilled with 1-mm trapezoidal holes at three milling frequencies (6 000, 12 000,
89 and 18 000 rpm). Then, obtained powder was collected in sealable polyethylene bags and stored
90 at 10 °C until sieving.

4849 91 2.3. Sieving of okra seed powders

Each okra seed powder issued from the milling step was separated in two aliquots. The first aliquot was immediately put in sealed polyethylene bags and stored at 10 °C until analysis; these powders were not sieved, thus they will be named unsieved powders hereafter. The other aliquot was sieved with an Analyzer 3 Spartan sieve shaker (Fritsch, Idar-Oberstein, Germany) with three superimposed sieves of 180, 315, and 500 µm mesh sizes, at 0.5 mm vibration amplitude during 24 min. Hence, \leq 180 µm, 180 - 315 µm, 315 - 500 µm, and \geq 500 µm

98 granulometric classes were obtained by sieving. These granulometric classes were finally put
99 in sealed polyethylene bags and stored at 10 °C until analyses.

2.4. Physical properties

101 2.4.1. Particle size distribution

Powder granulometric characteristics define the granular state of a powder. In the pharmaceutical industry, particle size analysis is of utmost importance as it partly conditions the bioactivity of medicines (Carmona *et al.*, 2003). The particle size distribution of unsieved powders and granulometric classes of okra seed powders was determined by laser diffraction using dry dispersion (Mastersizer 3000, Malvern Instruments France, Orsay, France). Dispersion conditions for all powders except for the $\leq 180 \,\mu\text{m}$ powder samples were as follows: 100 % air pressure at 4 bar, 3 mm hopper gap, and 40 % feed rate. For the \leq 180 µm powder samples, the following dispersion conditions were employed: 100 % air pressure at 4 bar, 4 mm hopper gap, and 70 % feed rate. The chosen size estimator was the equivalent diameter in volume. Classical granulometric parameters were determined: d10, d50, and d90, which correspond to the first decile, the median, and the ninth decile of the particle size distribution, and the span, which quantifies the width of the particle size distribution (equation 1):

2.4.2. Moisture content and water activity

Moisture content was measured by weight loss after drying 3 g powder sample at 103 °C for
5 h ('AFNOR, Agence Française de Normalisation, 1976).

(1)

 $\text{Span} = \frac{\text{d90} - \text{d10}}{\text{d50}}$

42 118 Water activity of okra powders was evaluated with a water activity-meter (HygroPalm23-AW-43 119 A, Rotronic, France): 15 g powder was put in a WP-40 sample holder and the quick analysis 46 120 mode was employed at 22 ± 3 °C, leading to water activity stabilization in about 5 min.

- 48 121 2.5. Phytochemical properties
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- ⁵⁰
 122 2.5.1. Preparation of hydromethanolic extracts

Hydromethanolic extracts of okra seed powders were prepared by maceration at room temperature (22 °C) in order to preserve the bioactivity of extracted molecules by minimizing their thermal alteration (Ćujić et al., 2015; Becker et al., 2016). It is worth noting that this method permitted to obtain a solution of extractable phenolic compounds, but it was not able to extract chemically-bound phenolic compounds. 2 g powder was put in 25 mL methanol-distilled

water mixture (70/30 % (v/v)) during 24 h under stirring at 300 rpm. Then, the resulting extract was centrifuged (Thermo Scientific, Heraeus Megafuge 8R Centrifuge) at 3 460 × g for 20 min at 20 °C and the obtained supernatant was filtrated using Whatman filter paper n° 1 (Sigma Aldrich, France). The filter paper was washed with 20 mL methanol-distilled water mixture (70/30 % (v/v)) and the filtrate was also centrifuged at 3 460 × g for 20 min at 20 °C. Last, the resulting supernatant was collected and stored at 4 °C until analysis.

⁴ 134 **2.5.2. Total phenolic content**

In test tubes were added 30 µL okra powder hydromethanolic extract, 2.67 mL distilled water, 500 µL Folin-Ciocalteu reagent (1 N), and 300 µL sodium carbonate (Na₂CO₃, 20 % (w/v)). The mixture was vortexed and dark-incubated during 30 min at room temperature (22 °C). Finally, absorbance of obtained solutions was measured at 750 nm with a Cary 50 Scan UV/visible spectrophotometer (Agilent, Santa Clara California, USA). Standard solutions of gallic acid (from 0.01 to 0.03 g/L by 0.005 mg/mL steps) were used for calibration ($R^2 = 0.99$) and total phenolic content was expressed in terms of milligrams of gallic acid equivalents per gram of dry matter (mg GAE/g DM).

31 143 2.5.3. Antioxidant activity evaluated by DPPH scavenging

The 1,1-diphenyl-2-picrylhydrazyl (DPPH) free radical scavenging method offers a first approach for evaluating the antioxidant potential of a compound, an extract, or other biological sources (Kitagaki et al., 1999). The DPPH' scavenging method was performed according to the method of Masuda et al. (1999). 500 µL of 0.1 mM DPPH solution was mixed with 500 µL okra seed powder hydromethanolic extract. This mixture was vortexed, then dark-incubated at 30 °C for 30 min, and finally its absorbance was measured at 517 nm against a blank solution composed of 500 µL DPPH solution mixed with 350 µL methanol and 150 µL distilled water that was left 30 min at rest. The antioxidant activity was evaluated as the percentage of DPPH radical inhibition and calculated from the absorbance decrease using (equation 2) (Buijnsters et *al.*, 2001):

% Inhibition of DPPH radical =
$$\left[\frac{ABS - abs}{ABS}\right] \times 100$$
 (2)

with ABS (-) representing the absorbance of the blank solution and abs (-) the absorbance of
the solution obtained from okra seeds powder extract after the DPPH assay.

⁵⁹ 157 **2.5.4.** Antioxidant activity evaluated by ABTS scavenging

ABTS⁺ radical was produced by a reaction between an aqueous solution of ABTS at 7 mM and a potassium persulfate K₂S₂O₈ solution at 4.9 mM. This mixture was stirred for 16 h in the dark at room temperature (22 °C), leading to a blue green-colored solution containing the ABTS⁺ cation radical (Cai et al., 2015). Afterwards, the ABTS^{+•} solution was then diluted with ethanol until reaching between 0.65 and 0.70 absorbance at 734 nm. For analysis, 100 µL powder sample hydromethanolic extract was mixed with 900 µL ABTS⁺ radical solution and the mixture was vortexed. After 10 min incubation at room temperature (22 °C), the absorbance of the mixture was measured at 734 nm against a blank solution (mixture of 100 µL phosphate buffer at pH 7.4 and 900 µL ABTS⁺ radical solution). The results were expressed in terms of inhibition percentage according to (equation 3):

% Inhibition of ABTS radical =
$$\left[\frac{A_0 - A_1}{A_0}\right] \times 100$$
 (3)

with A_0 (-) representing the absorbance of the blank solution and A_1 (-) the absorbance of the solution obtained from okra seeds powder extract after the DPPH ABTS⁺⁺ assay.

2.6. Hydration properties

Water absorption capacity (WAC) and water solubility index (WSI) were evaluated according to Deli *et al.* (2019). Water absorption capacity (WAC) was determined as follows: first, 1 g okra seeds powder and 10 mL distilled water were mixed in a centrifuge tube, then the mixture was stirred at 300 rpm (Variomag Poly) for 30 min at 20 °C and centrifuged (Thermo Scientific, Heraeus Megafuge 8R Centrifuge) at 1 500 × g for 30 min at 20 °C. The supernatant was carefully removed. The pellet was weighed, dried in an oven at 103 ± 2 °C for 6 h, cooled in a desiccator, and weighed again. WAC was evaluated as follows (equation 4):

WAC =
$$\frac{\text{mass of wet sediments}}{\text{mass of dried sediments}}$$
 (4)

49 180 Where masses of wet and dry sediments were expressed in grams.

The following protocol was used to determine the water solubility index of powder samples: 2.5 g (M₁) okra powder and 20 mL distilled water were mixed in a centrifuge tube, then the mixture was stirred at 300 rpm (Variomag Poly) for 30 min at 20 °C. After centrifugation (Thermo Scientific, Heraeus Megafuge 8R Centrifuge) at 1 500 × g for 20 min at 20 °C, the supernatant was carefully removed and the pellet was dried in an oven at 103 ± 2 °C for 6 h, cooled in a desiccator, and weighed (M₂). WSI was calculated according to equation 5:

WSI (%) = $\left(\frac{M_1 - M_2}{M_1}\right) \times 100$ (5)

188 2.7. Statistical analysis

One-factor analysis of variance, the calculation of Pearson's correlation coefficients, and principal components analysis (PCA) were performed by using Microsoft Excel 2016 and significant differences were assessed with Tukey's HSD at a 5 % confidence level with the DSAASTAT add-on (Andrea ONOFRI, Department of Agricultural, Food and Environmental Sciences, University of Perugia, Italy). The determination of Pearson's correlation coefficients and PCA were performed to examine correlations between characterized properties of okra seed powders: median particle size, span, total phenolic content, results of DPPH and ABTS assays, water activity, moisture content, WAC, and WSI. Results were expressed as means \pm standard deviations.

26 198 3. Results and discussion 27

28 29 199 **3.1. Particle size distribution**

The analysis of the particle size distribution was performed in order to check if the successive milling and sieving processes were efficient in micronizing okra seeds and separating unsieved powders into sufficiently different granulometric classes. Ttable 1 shows that the median particle size decreased when the milling frequency was increased: for instance, for unsieved powders, median particle size was lowered value, from 723 µm at 6 000 rpm to 478 µm at 12 000 rpm and 229 µm at 18 000 rpm. The median particle size of granulometric classes was found between the mesh sizes of their respective upper and lower sieves as expected when sieving is completed, confirming the efficiency of the sieving process to separate powders in granulometric classes differing in median particle size.

Span of unsieved powders obtained at 6 000, 12 000, and 18 000 rpm were equal to 2.11 ± 0.02 , 2.03 ± 0.01 , and 2.96 ± 0.16 , respectively, showing that the width of the particle size distribution was larger at higher milling frequency. This could be explained by the higher proportion of fine particles when using a higher milling frequency, leading to increased heterogeneity of unsieved powders.

⁵⁶ 214

Table 1

59 215 **3.2. Total phenolic content**

Figure 1 presents the total phenolic content of unsieved samples and granulometric classes of okra seed powders obtained at the three different milling frequencies. Higher total phenolic contents were observed for samples composed of smaller particles. Moreover, a greater milling frequency induced an increase in total phenolic content: for example, for unsieved powders, the total phenolic content equaled 2.34, 3.44, and 7.40 mg GAE/g DM at 6 000 rpm, 12 000, and 18 000 rpm milling frequencies, respectively. This can be explained by the fact that a more intense milling (here at higher milling frequency, but it could also result from a longer milling duration) enhances the size reduction of plant material, which further facilitates the accessibility of solvents for the extraction of bioactive compounds owing to the higher specific surface area of smaller particles (Rosa et al., 2013). This observation was confirmed by the moderate negative correlation between total phenolic content and median particle size d50 evidenced by a Pearson's correlation coefficient of r = -0.71 (p < 0.05). Indeed, the bioactive compounds are often associated with proteic and fatty fractions in plant extracts (Becker et al., 2016; Deli, Baudelaire, et al., 2019). As indicated in the introduction, a previous study focused on the influence of granulometric characteristics of these powders on their proximate composition revealed that higher protein and fat contents were found in the smaller particle size fractions of okra powder seeds (Waiss et al., 2020). Thus, the discriminating power of the sieving process in matter of total phenolic content and proximate composition was enhanced at higher milling frequency.

Figure 1

3.3. Antioxidant activity

The DPPH[•] and ABTS^{•+} assays are spectrophotometric techniques based on quenching of stable colored radicals which are widely used for the assessment of antioxidant activities of complex biological mixtures such as plant or food extracts.

3.3.1 DPPH' radical scavenging activity assay

Results of antioxidant activity of okra seed powders obtained by DPPH test are presented in Figure 2. All extracts were able to scavenge DPPH[•] radicals. The percentage of DPPH inhibition was higher for $\leq 180 \ \mu m$ and $\geq 500 \ \mu m$ granulometric classes whatever the milling frequency. The $\leq 180 \,\mu\text{m}$ granulometric classes were richer in lipids and proteins and the $\geq 500 \,\mu\text{m}$ granulometric classes contained more carbohydrates, as shown in a previous study performed with the same powders as investigated in the present study (Waiss et al., 2020). It has been showed that bioactive molecules such as polyphenols are often associated with lipids, proteins,

and polysaccharides in plant tissues (Zaiter et al., 2016). This observation partially meets the results obtained in the present study, as at 6 000 and 12 000 rpm milling frequencies, DPPH. scavenging activity decreased with particle size for the ≤ 180 , 180 - 315, and 315 - 500 μ m granulometric classes; however, at a given milling frequency, the DPPH[•] scavenging activity of the \geq 500 µm granulometric classes were greater than for the other granulometric classes and unsieved powders, indicating that bioactive compounds may be mainly associated with carbohydrates, likely fibers, in okra seeds. The increasing trend of DPPH' scavenging activity with particle size at 18 000 rpm confirms this interpretation, as a more intense milling may have permitted to break fibers and decrease their particle size, increasing the DPPH' scavenging activity of the granulometric classes of higher median particle size. Besides, micronization of plant material generally leads to an increase in the antioxidant activity of plant extracts (Rosa et al., 2013), and this was confirmed by the present study where milling at 18 000 rpm induced a marked increase in the DPPH[•] inhibition percentage. Milling is known to alter the physical structure of the plant matrix, increasing its specific surface area and leading to the exposition of antioxidant compounds of the plant material (Hemery et al., 2010). Last, the fact that the DPPH scavenging activity was systematically higher for granulometric classes than for unsieved powders (Figure 2) shows that hydromethanolic extraction was improved after sieving, maybe by decreasing powder cohesion and then facilitating its dispersion and the exposure of bioactive molecules to the solvent (Salameh et al., 2016).

Figure 2

3.3.2 ABTS cation radical scavenging activity assay

The results of ABTS⁺⁺ radical scavenging activity of granulometric classes and unsieved samples of okra seed powders obtained at 6 000, 12 000, and 18 000 rpm milling frequencies are reported in Figure 3. High antioxidant activity of all okra seed powders was observed during the ABTS assay. No great difference of ABTS⁺⁺ inhibition was denoted between samples, in particular, no significant difference could be attributed to milling frequency, but a trend of increased antioxidant activity for powder samples having a smaller median particle size was apparent. This was confirmed by the Pearson's correlation coefficient between median particle size d50 and ABTS⁺⁺ inhibition percentage of r = -0.64 (p < 0.05), which denoted a moderate negative correlation between these parameters. Besides, the antioxidant activity obtained by the ABTS cation radical scavenging was significantly higher than for DPPH radical scavenging for all granulometric classes and unsieved powders of okra seeds (cf. figures 2 and 3). Indeed,

previous studies showed that the ABTS test leads to higher inhibition percentage than the DPPH test for fruits, vegetables, and beverages (Floegel et al., 2011), owing to the better reactivity of high-pigmented and hydrophilic antioxidants with ABTS⁺⁺. Obtained results are consistent with the fact that antioxidant activity of plant products is generally attributed to radical scavenging activity of phenolic compounds (Rahman et al., 2007; Becker et al., 2016; Deli, et al., 2019). Indeed, the phenolic compounds such as flavonoids, polyphenols, and tannins are the main bioactive molecules that contribute to antioxidant activity of plant extracts (Rosa et al., 2013; Brewer et al., 2014; Becker et al., 2017).

Figure 3

20 289 **3.4. Moisture content and water activity**

Moisture content is an important parameter that is indirectly related to powdered food ingredient stability during storage. Low levels of moisture content (up to 12 % (w/w)) are known to limit microbial growth and extend food shelf-life (Aryee et al., 2006; Kaur et al., 2013). Results displayed in Ttable 2 show that the moisture content of granulometric classes and unsieved samples of okra seed powders ranged from 8.61 ± 0.08 to 12.07 ± 0.05 (w/w), presuming a good stability during storage. The moisture content was generally decreased at smaller median particle size and/or at higher milling frequency (cf. Table 2), as evidenced by the moderate positive correlation between moisture content and median particle size (Pearson's correlation coefficient of r = 0.73, p < 0.05). Moisture content is not sufficient to characterize food stability during storage, as food structure also contributes to water availability for microorganisms and alteration reactions: the concept of water activity has been developed to account for these combined influences of moisture content and food structure (Tapia, et al., 2007; Bhandari et al., 2013). Water activity ranged between 0.493 ± 0.010 and 0.583 ± 0.010 for investigated okra seed powders (cf. Ttable 2), which was below the threshold of 0.650 permitting the growth of microorganisms like xerophilic mold and bacteria (Tapia et al., 2007). Water activity was generally decreased at lower median particle size, similarly to water content. This was expected, as a more intense milling leads to higher stress, more friction, and thus higher heating of the plant material, leading to smaller particles with a lower moisture content (Becker et al., 2017)

56 309 3.5. Water interactions

The water absorption capacity (WAC) and water solubility index (WSI) results of all
granulometric classes and unsieved samples of okra seed powders are presented in Table 2.

WSI significantly differed according to the milling frequency. When milling frequency was increased, the median particle size of powders was decreased, enhancing water solubility index. This was confirmed by the highly negative correlation between WSI and median particle size (Pearson's correlation coefficient of r = -0.89, p < 0.05), indicating that interaction with water was improved for smaller particles (cf. Ttable 2). This can be explained by the fact that milling induces particle size reduction, leading to the improvement of material functionalities such as particle surface specific area and surface energy, enhancing the interactions of plant material with water (Ye et al., 2016). In addition, this may result from the exposure of hydrophilic groups (coming from cellulose and hemicellulose) at the surface of particles obtained by plant micronization (Zhao et al., 2009; Ting et al., 2014). Zhao et al. (2009) also reported increased dispersibility and solubility for ginger powders produced with a more intense milling, owing to improved interactions with water evidenced by a higher WSI (Zhao et al., 2009). Water absorption capacity represents the ability of a product to absorb water. Table 2 shows that all powders had a high WAC whatever the granulometric class and the milling frequency. This can be explained by the high proportion of mucilage of okra seeds that highly contribute to water absorption (Aboubakar et al., 2008). No clear influence of median particle size on WAC was denoted, except for a small trend of higher WAC for intermediate granulometric classes. Thus, performed milling was sufficient to break cell walls and expose hydrophilic groups of all okra seed components. The results of the current study are consistent with the work of (Deli, Petit, et al., 2019), who reported that WAC and WSI were enhanced for smaller particles of Boscia senegalensis seeds, Dichostachys glomerata fruits, and Hibiscus sabdariffa calyxes. Hence, it can be hypothesized that okra seed powders obtained at higher milling frequency should interact more easily with water and be more soluble, which can facilitate their intestinal absorption during digestion.

Table 2

337 3.6. Principal component analysis of physicochemical, water interaction, and

338 phytochemical properties

PCA is used to highlight similarities or oppositions between variables and identify the variables that are most correlated between them. All granulometric classes and unsieved samples of okra seed powders obtained at the three milling frequencies were compared using PCA on the basis of their physicochemical (moisture content, median particle size), water interaction (water activity, water solubility index, and water absorption capacity), and phytochemical (total phenolic content, and antioxidant activity deduced from DPPH and ABTS tests) properties. The

result of this PCA is presented in Figure 4. Variables are presented in red color and principal components are represented on F1 and F2 axes which respectively explained 43.4 % and 20.7 % variability among powder properties, allowing the PCA to explain a total variability of 64.1 % which was sufficient to outline the major links between physicochemical and phytochemical properties of okra seed powders. F1 axis was positively correlated with ABTS⁺⁺ inhibition, WSI, and total phenolic content, and negatively correlated with median particle size and moisture content. F2 axis was positively correlated with span and WAC, and negatively correlated with DPPH inhibition and water activity. It is worth noting that median particle size was negatively correlated to total phenolic content, WSI, and ABTS⁺⁺ inhibition percentage, and positively to moisture content, confirming the previous discussion of experimental results.

The representation of okra powder samples on the F1 \times F2 biplot revealed a separation of powder samples according to median particle size (Figure 4). Smaller granulometric classes $(\leq 180 \ \mu m \text{ at } 6\ 000,\ 12\ 000,\ and\ 18\ 000\ rpm\ milling\ frequency\ and\ 180\ -\ 315\ \mu m\ at\ 18\ 000\ rpm$ milling frequency) were located at the bottom right of the F1 \times F2 biplot and characterized by their high total phenolic content and WSI, and obviously by their low median particle size. Higher granulometric classes (\geq 500 µm at 6 000, 12 000, and 18 000 rpm milling frequency and 315 - 500 µm at 18 000 rpm milling frequency) were found at the left of the F1 × F2 biplot and characterized by high moisture content and median particle size, and low total phenolic content and WSI. The other granulometric classes of intermediate median particle size and unsieved powders were located in the top middle of $F1 \times F2$ biplot and they were positively correlated with span and WAC, and negatively to water activity and DPPH[•] inhibition.

Figure 4

4. Conclusion

 The combination of milling and sieving processes was successful in generating granulometric classes with significantly different physicochemical, water interaction, and phytochemical properties. The results evidenced a considerable influence of median particle size of okra seeds powder on its total phenolic content, antioxidant activity deduced from ABTS and DPPH tests, moisture content, water activity, and water solubility index. Powders with smaller median particle sizes (i.e. lower size granulometric classes and/or powders obtained at higher milling frequency) had lower moisture content, water activity, and water absorption capacity, and higher total phenolic content, antioxidant activity, and water solubility index. This can result

from the higher level of disintegration of plant matrix structure induced by a more intense milling (i.e. undergone by smaller particles and/or at higher milling frequency) which makes its compounds more accessible to solvents, thus improving phenolic compounds extraction and powder solubility, and decreases the number of pores of the solid matrix, thus lowering water absorption capacity. For all samples, moisture content and water activity were sufficiently low to avoid food spoilage. Hence, okra seed powders were biochemically and microbiologically stable, ensuring a good shelf-life during storage. Water interaction parameters showed high levels of water absorption and solubility, which seems to indicate that the reconstitutability of okra powder could be improved by the successive milling and sieving process. Finally, all obtained results suggest that okra seed powders can be used in food manufacturing as functional ingredients enhancing antioxidant activity and water interactions.

3 388 **5. Acknowledgments**

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6. Conflict of interest

396 The authors declare that they have no conflict of interest.

7. Ethical approval

No experiment was performed on humans or animals: no ethics approval was not required forthis research.

48 400 **8. Data availability statement**

4950 401 Research data are not shared.

⁵² 402 **9. References**

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Figure 4: Representation of all granulometric classes (≤ 180 ; 180 - 315; 315 - 500; $\geq 500 \mu m$) and unsieved samples of okra seed powders obtained at 6 000 rpm (green), 12 000 rpm (black), and 18 000 rpm (blue) milling frequencies and their physicochemical and functional characteristics (red) on the principal components F1 and F2 deduced from PCA.

Milling speed (rpm)	Granulometric classes	d10 (µm)	d50 (µm)	d90 (µm)	Span (-)
· • ·	≤ 180 μm	23.1 ± 0.2^{ab}	85.3 ± 0.6^{a}	181.3 ± 0.5^{a}	1.87 ± 0.01^{f}
	180 - 315 μm	152.0 ± 1.0^{de}	$257.0 \pm 1.0^{\circ}$	406.0 ± 1.0^{b}	0.97 ± 0.01^{d}
	315 - 500 μm	326.6 ± 6.0^{g}	498.3 ± 2.5^{e}	$745.3 \pm 2.5^{\circ}$	0.87 ± 0.03^{bcd}
6 000	≥ 500 μm	589.3 ± 18.5^{i}	1100.0 ± 43.5^{h}	2173.3 ± 73.7^{g}	1.44 ± 0.02^{e}
	Unsieved powder	141.0 ± 7.8^{d}	$723.3 \pm 29.9^{\rm f}$	$1670.0 \pm 65.5^{\rm f}$	2.11 ± 0.02^{h}
	≤ 180 μm	21.9 ± 0.1^{ab}	75.6 ± 2.8^{a}	167.3 ± 6.3^{a}	1.95 ± 0.06^{fg}
	180 - 315 μm	165.6 ± 1.5^{e}	$264.6 \pm 1.5^{\circ}$	409.3 ± 9.8^{b}	0.92 ± 0.03^{cd}
	315 - 500 μm	312.0 ± 1.0^{g}	463.3 ± 1.1^{de}	$671.6 \pm 4.9^{\circ}$	$0.78 \pm 0.01^{\rm bc}$
12 000	≥ 500 μm	542.6 ± 5.6^{h}	$801.3 \pm 25.0^{\text{g}}$	$1243.3 \pm 75.0^{\text{e}}$	0.90 ± 0.05^{bcd}
	Unsieved powder	$52.9 \pm 0.5^{\circ}$	457.6 ± 5.1^{de}	983.0 ± 7.8^{d}	2.03 ± 0.01^{gh}
	≤ 180 μm	17.8 ± 0.2^{a}	52.1 ± 0.6^{a}	149.3 ± 1.5^{a}	0.77 ± 0.01^{b}
	180 - 315 μm	37.4 ± 0.1^{bc}	181.0 ± 0.0^{b}	332.6 ± 2.0^{b}	0.48 ± 0.01^{a}
18 000	315 - 500 μm	290.0 ± 4.3^{f}	442.6 ± 12.8^{d}	$644.6 \pm 25.9^{\circ}$	0.80 ± 0.02^{bc}
	≥ 500 μm	527.6 ± 4.7^{h}	781.0 ± 14.1^{g}	1186.6 ± 37.8^{e}	0.84 ± 0.02^{bcd}
	Unsieved powder	25.7 ± 0.3^{ab}	228.6 ± 7.5^{bc}	702.67± 55.7°	2.96 ± 0.16^{i}
Tukey's HSD	test.				

Table 1: Granulometric parameters of granulometric classes ($\leq 180 \,\mu$ m, 180 - 315 μ m, 315 - 500 μ m, $\geq 500 \,\mu$ m) and unsieved samples of okra seed powders obtained at 6 000, 12 000, and 18 000 rpm milling frequencies (Waiss *et al.*, 2020).

 Table 2: Moisture content, water activity, water absorption capacity, and water solubility index of granulometric classes ($\leq 180 \ \mu m$, 180 - 315 μm , 315 - 500 μm , $\geq 500 \ \mu m$) and unsieved samples of okra seed powders obtained at 6 000, 12 000, and 18 000 rpm milling frequencies.

Milling frequency	Powder samples	Moisture content	Water activity (-)	Water absorption	Water solubilit
(rpm)		(g/g) (Waiss et al.,		capacity, WAC (g/g)	index, WSI (g/10
		2020)			g)
	\leq 180 μ m	9.59 ± 0.12^{cd}	$0.555 \pm 0.001^{\text{ef}}$	4.16 ± 0.46^{abc}	$32.71\pm0.53^{\rm f}$
	180 - 315 μm	9.01 ± 0.67^{ab}	$0.556 \pm 0.002^{\text{ef}}$	6.46 ± 0.20^{e}	$24.06\pm0.10^{\text{cd}}$
6 000	315 - 500 μm	9.21 ± 0.11^{bc}	$0.540\pm0.001^{\text{cd}}$	5.58 ± 0.25^{cde}	22.74 ± 0.74^{bc}
	\geq 500 μ m	12.07 ± 0.05^{i}	0.524 ± 0.003^{b}	4.76 ± 0.24^{abcd}	15.98 ± 0.23^{a}
	Unsieved powder	$11.10 \pm 0.07^{\text{fg}}$	$0.583 \pm 0.007^{\rm h}$	5.07 ± 0.75^{cde}	22.75 ± 0.37^{bc}
	\leq 180 μ m	9.15 ± 0.10^{abc}	0.565 ± 0.008^{efg}	4.09 ± 0.28^{ab}	$37.75\pm0.46^{\text{g}}$
	180 - 315 μm	8.61 ± 0.08^{a}	0.524 ± 0.002^{b}	5.77 ± 0.41^{de}	$32.53 \pm 1.06^{\rm f}$
12 000	315 - 500 μm	9.22 ± 0.05^{bc}	0.503 ± 0.004^{a}	5.12 ± 0.30^{bcde}	25.05 ± 0.57^{d}
	\geq 500 μ m	11.79 ± 0.06^{hi}	0.531 ± 0.004^{bc}	4.98 ± 0.23^{bcde}	17.64 ± 0.16^{a}
	Unsieved powder	11.27 ± 0.08^{fgh}	0.550 ± 0.001^{de}	5.19 ± 0.52^{bcd}	22.00 ± 0.78^{b}
	\geq 500 μ m	9.56 ± 0.08^{bc}	0.572 ± 0.002^{gh}	3.52 ± 0.12^{a}	$41.12\pm1.16^{\rm h}$
10.000	315 - 500 μm	10.15 ± 0.02^{de}	0.564 ± 0.005^{fgh}	4.06 ± 0.87^{abc}	36.77 ± 0.66^g
18 000	180 - 315 μm	11.32 ± 0.06^{gh}	0.560 ± 0.002^{efg}	5.60 ± 0.50^{cde}	23.50 ± 0.07^{bc}
	≤180 μm	13.20 ± 0.17^j	0.550 ± 0.010^{ef}	4.67 ± 0.12^{abc}	17.00 ± 0.14^{a}
	Unsieved powder	$10.69 \pm 0.09^{\text{ef}}$	0.493 ± 0.009^{a}	5.55 ± 0.55 bcde	30.58 ± 0.80^{e}

620 Means \pm standard deviations followed by the same superscripted letters in the same column were not significantly different at p < 0.05 according to Tukey's 621 HSD test.

 For peer Review



400x400mm (96 x 96 DPI)

Institute of Food Science and Technology

Conflict of interest

The authors declare that they have no conflict of interest.

for per period

Data availability statement

Research data are not shared.

for per period



Figure1: Total phenolic content of granulometric classes ($\leq 180 \ \mu m$; 180 - 315 μm ; 315 – 500 μm ; $\geq 500 \ \mu m$) and unsieved okra seed powders. DM: dry matter, GAE: gallic acid equivalents. Bars topped by different letters were statistically different at p < 0.05 according to Tukey's HSD test.

Figure 2



Figure 2: Inhibition of DPPH• radical by hydromethanolic extracts of granulometric classes (\leq 180 µm; 180 - 315 µm; 315 - 500 µm; \geq 500 µm) and unsieved okra seed. Bars topped by different letters were statistically different at p < 0.05 according to Tukey's HSD.



Figure 3: Inhibition of ABTS++ by hydromethanolic extracts of granulometric classes ($\leq 180 \mu m$; 180 - 315 μm ; 315 - 500 μm ; $\geq 500 \mu m$) and unsieved powders of okra seed. Bars topped by different letters were statistically different at p < 0.05 according to Tukey's.



Figure 4: Representation of all granulometric classes (≤ 180 ; 180 - 315; 315 - 500; $\geq 500 \ \mu m$) and unsieved samples of okra seed powders on the principal components F1 and F2 deduced from PCA.

Milling speed	Granulometric classes	d10 (µm)	d50 (µm)	d90 (µm)	Span (-)
(rpm)					
	≤ 180 μm	23.1 ± 0.2^{ab}	85.3 ± 0.6^{a}	181.3 ± 0.5^{a}	$1.87\pm0.01^{\rm f}$
	180 - 315 μm	152.0 ± 1.0^{de}	$257.0 \pm 1.0^{\circ}$	406.0 ± 1.0^{b}	0.97 ± 0.01^{d}
	315 - 500 μm	326.6 ± 6.0^{g}	498.3 ± 2.5^{e}	$745.3 \pm 2.5^{\circ}$	0.87 ± 0.03^{bcd}
6 000	\geq 500 μ m	589.3 ± 18.5^{i}	1100.0 ± 43.5^{h}	2173.3 ± 73.7^{g}	1.44 ± 0.02^{e}
	Unsieved powder	141.0 ± 7.8^{d}	$723.3 \pm 29.9^{\rm f}$	$1670.0 \pm 65.5^{\rm f}$	2.11 ± 0.02^{h}
	≤ 180 μm	21.9 ± 0.1^{ab}	75.6 ± 2.8^{a}	167.3 ± 6.3^{a}	$1.95\pm0.06^{\rm fg}$
	180 - 315 μm	165.6 ± 1.5^{e}	$264.6 \pm 1.5^{\circ}$	409.3 ± 9.8^{b}	0.92 ± 0.03^{cd}
	315 - 500 μm	312.0 ± 1.0^{g}	463.3 ± 1.1^{de}	$671.6 \pm 4.9^{\circ}$	0.78 ± 0.01^{bc}
12 000	\geq 500 µm	542.6 ± 5.6^{h}	$801.3 \pm 25.0^{\text{g}}$	$1243.3 \pm 75.0^{\text{e}}$	0.90 ± 0.05^{bcd}
	Unsieved powder	$52.9 \pm 0.5^{\circ}$	457.6 ± 5.1^{de}	983.0 ± 7.8^{d}	$2.03\pm0.01^{\text{gh}}$
	≤ 180 μm	17.8 ± 0.2^{a}	52.1 ± 0.6^{a}	149.3 ± 1.5^{a}	0.77 ± 0.01^{b}
	180 - 315 μm	37.4 ± 0.1^{bc}	181.0 ± 0.0^{b}	332.6 ± 2.0^{b}	0.48 ± 0.01^{a}
18 000	315 - 500 μm	290.0 ± 4.3^{f}	442.6 ± 12.8^{d}	$644.6 \pm 25.9^{\circ}$	0.80 ± 0.02^{bc}
	\geq 500 µm	527.6 ± 4.7^{h}	781.0 ± 14.1^{g}	$1186.6 \pm 37.8^{\circ}$	0.84 ± 0.02^{bcd}
	Unsieved powder	25.7 ± 0.3^{ab}	228.6 ± 7.5^{bc}	702.67± 55.7°	2.96 ± 0.16^{i}

Table 1: Granulometric parameters of granulometric classes ($\leq 180 \ \mu m$, 180 - 315 μm , 315 - 500 μm , $\geq 500 \ \mu m$) and unsieved samples of okra seed powders obtained at 6 000, 12 000, and 18 000 rpm milling speed (Waiss et al., 2020).

For each milling speed separately, means \pm standard deviations followed by the same superscripted letters in the same column were not significantly different at p < 0.05 according to Tukey's HSD test.

Table 2: Moisture content, aw (-): water activity, WAC: Water Absorption Capacity(g/) and WSI: Water Solubility Index (g/100g) of \geq 500 µm; 500–315µm, 315–180 µm,180 µm \geq and for unsieved seeds. Means \pm standard deviations followed by the same superscripted letters in the same row were not significantly different at P < 0.05 according to Tukey's HSD test.

Milling speed (r.p.m.)	granulometric classes	Moisture content (g/g)	aw (-)	WAC (g/g)	WSI (g/100 g)
		(Waiss et al., 2020)			
	≥ 500 μm	$12.07 \pm 0.05^{\circ}$	$0,524 \pm 0,00^{a}$	$4,76 \pm 0,24^{ab}$	15,98 ± 0,23 ^b
	315 - 500 μm	9.21 ± 0.11^{a}	$0,540 \pm 0,00^{b}$	$5,58 \pm 0,25^{b}$	$22,74 \pm 0,74^{b}$
6 000	180 - 315 μm	9.01 ± 0.67^{a}	0,556 ± 0,00 °	$6,46 \pm 0,20^{\circ}$	$24,06 \pm 0,10^{b}$
	\leq 180 μ m	9.59 ± 0.12^{a}	$0,555 \pm 0,00^{\circ}$	$4,16 \pm 0,46^{a}$	32,71 ± 0,53°
	Unsieved powder	11.10 ± 0.07^{b}	$0,583 \pm 0,01^{d}$	$5,07 \pm 0,75^{b}$	$22,75 \pm 0,37^{b}$
	\geq 500 μ m	11.79 ± 0.06^{d}	$0,531 \pm 0,00^{b}$	$4,98 \pm 0,23^{b}$	$17,64 \pm 0,16^{a}$
	315 - 500 μm	9.22 ± 0.05^{b}	$0,503 \pm 0,00^{a}$	$5,12 \pm 0,30^{b}$	$25,05 \pm 0,57^{b}$
12 000	180 - 315 μm	8.61 ± 0.08^{a}	$0,524 \pm 0,01^{b}$	$5,77 \pm 0,41^{b}$	32,53 ± 1,06°
	\leq 180 μ m	9.15 ± 0.10^{b}	$0,565 \pm 0,00^{d}$	$4,09 \pm 0,28^{a}$	$37,75 \pm 0,46^{d}$
	Unsieved powder	$11.27 \pm 0.08^{\circ}$	$0,550 \pm 0,00^{\circ}$	$5,19 \pm 0,52^{b}$	$22.00\pm0,78^{ab}$
	\geq 500 μ m	13.20 ± 0.17^{e}	$0,550 \pm 0,01^{b}$	$4,67 \pm 0,12^{ab}$	$17,00 \pm 0,14a$
10.000	315 - 500 μm	11.32 ± 0.06^{d}	$0,560 \pm 0,00^{\circ}$	$5,60 \pm 0,50^{b}$	$23,50 \pm 0,07^{b}$
18 000	180 - 315 μm	10.15 ± 0.02^{b}	0,564± 0,00 ^{cd}	$4,06 \pm 0,87^{ab}$	$36,77 \pm 0,66^{d}$
	\leq 180 μ m	9.56 ± 0.08^{a}	$0,572 \pm 0,00^{d}$	$3,52 \pm 0,12^{a}$	$41,12 \pm 1,16^{e}$
	Unsieved powder	$10.69 \pm 0.09^{\circ}$	$0,493 \pm 0,01^{a}$	$5,55 \pm 0,55^{b}$	$30,58 \pm 0,80^{\circ}$

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Dear Editor-in-chief of International Journal of Food science and Technology,

Please find the following manuscript 'Total phenolic content, antioxidant activity, shelf-life, and reconstitutability of okra seeds powder: influence of milling and sieving processes.', by 'Idriss MIGANEH WAISS, André KIMBONGUILA, FATOUMA MOHAMED ABDOUL-LATIF, Laurette Brigelia NKELETELA, Joël SCHER, Jérémy PETIT which we wish to submit for publication as an original article in 'International Journal of Food science and Technology'.

The objective of this work is to determine the impact of the particle size of okra seed powders on physicochemical, water interaction, and phytochemical properties. To our knowledge, this is the first time that a study focuses on characterization of okra seed powders with granulometric method.

The results of this study permitted to identify a granulometric class achieving a compromise between particle size of powders and bioactive compound content, powders reconstitution with interaction of water.

We confirmed that the manuscript or its contents in some other form have not been published previously by any of the authors and/or is not under consideration for publication in another journal at the time of submission.

Kind regards,

Jérémy PETIT

Becker, L. et al. (2017) 'How do grinding and sieving impact on physicochemical properties, polyphenol content, and antioxidant activity of Hieracium pilosella L. powders?', Journal of Functional Foods, 35, pp. 666–672. doi: 10.1016/j.jff.2017.06.043.

This reference was cited in the introduction as matters the impact of grinding and sieving on physicochemical and phytochemical properties of resulting powders. It was useful to discuss results of chemical composition and antioxidant activity.

Deli, et al. (2019) 'Effect of sieved fractionation on the physical, flow and hydration properties of Boscia senegalensis Lam., Dichostachys glomerata Forssk. and Hibiscus sabdariffa L. powders', *Food Science and Biotechnology*. Springer Singapore, 22, pp. 1–15. doi: 10.1007/s10068-019-00597-6.

This reference was cited because it reports information on the influence of sieving and grinding on the physical properties (particle size, shape, and specific surface area), aptitude for preservation (water activity), and reconstitutability (solubility, water absorption capacity) of plant powders. Similar grinding and sieving conditions and methods for the determination of proximal composition were employed.

Arapitsas, P. (2008) 'Identification and quantification of polyphenolic compounds from okra seeds and skins', Food Chemistry, 110(4), pp. 1041–1045. doi: 10.1016/j.foodchem.2008.03.014.

This reference, cited in introduction, reports information regarding contents of phenolic compounds in Ours seed.

Tapia MS, et al. (2007) 'Effects of water activity (aw) on microbial stability: as a hurdle in food preservation', in Fundamentals and Applications. Woodhead Publishing Limited, pp. 599–630. doi: 10.1533/9781845690243.3.599.

This article was useful in the discussion section. It was cited for interpreting okra seed powders stability during storage from water activity values.

The authors wish to thank the reviewers for their helpful comments. Detailed point-by-point response can be found hereafter, with reviewers' comments in black (line numbers associated with reviewers' comments correspond to submitted manuscript) and answers in blue (line numbers related to manuscript revision correspond to revised manuscript). In revised manuscript, additions were performed in blue and deletions in red crossed words.

Editor comments

Please pay particular attention to:

• The Graphical Abstract should be designed to be read in conjunction with the text abstract, it should be approximately square and contain a high impact Figure, Graph or Photograph that summarises the key findings of your research.

The graphical abstract was corrected.

• UK spelling should be used.

This was made in whole the manuscript.

• Tables & Figures must be in separate files, at the end of the manuscript.

This was made in the manuscript.

• References must follow the Harvard system of referencing and should be shown at the end of the paper, listed in alphabetical order.

This was made in the manuscript.

• Annotated References: authors are asked to annotate approximately 4 references indicating why the article was cited.

This was included in the submission file.

• Author statements on Conflict of Interest; Ethics and Data Availability.

This was included in the submission file.

• Figures and Tables should be uploaded as separate files. Tables should be in editable format and Figures. Scans (TIFF only) should have a resolution of at least 300 dpi (halftone) or 600 to 1200 dpi (line drawings) in relation to the reproduction size

This was included in the submission file.

 \emptyset Printing charges: Printing up to 7 journals pages is free to authors. Articles longer than this will incur an excess printing charge of £100 GBP per page. Unless your Institution is prepared to pay excess charges you should aim for approximately 4,500 words and 6 display items. Extra items may be assigned to Supplementary Files.

Reviewer #1

This is an interesting work that contributes with important results considering the influence of the particle size of milling in relation to the antioxidant activity and extractable phenolic levels. This is then related to the accessibility of these important compounds as well as of proteins and polysaccharides into the gut. This work complements a previous one of these authors where a complete determination of the chemical composition of these fractions with determined particle size ranges was performed.

This manuscript is acceptable for publication after correction according to the following revision.

Please, have the manuscript grammar and writing revised.

Amendments can be also found in the pdf file of the manuscript.

When a Figure number and a Table number is mentioned along the text of the manuscript, they also should begin written in Capital letter:

Examples: "as shown in Figure 2....." "as summarized in Table 2,"

This should be corrected along the manuscript.

This was corrected in the whole manuscript.

ABSTRACT.

The Abstract was in general well written.

Please, better write: "The influence of the particle size derived by milling of okra seeds at three different speed velocities (6,000; 12,000 and 18,000 rpm) on functionality (total phenolic content and antioxidant activity), reconstitutability (water absorption capacity, water solubility index), and preservation (water activity) of powders was studied. Okra seed powders were sieved with 180, 315, and 500 μ m sieves. An increased"

The abstract was clear, complete and grammatically sound, thus the authors preferred to keep it unchanged. In particular, 6 000, 12 000, and 18 000 rpm are values of milling frequencies, not milling velocities. Also, reconstitutability and preservation ability are powder functionalities.

Please, could you express the water activity value here and along the manuscript with three figures after the decimal point? Water activity meters are supposed to measure the water activity with three figures after the decimal point. It is important this complete value.

This was corrected in the whole manuscript.

"preservation" (not preservability)

"Preservability" was replaced by "preservation ability" in L21.

KEYWORDS.

Please, replace "okra seeds" by "okra seed powder".

This was corrected in L27.

INTRODUCTION.

It was well written indeed.

L38. Please, correct to: "of oil rich in unsaturated fatty acids including linolenic and linoleic acids, which are both essential for humans"

Concerned sentence was corrected (cf. L35).

L40. Please, correct to: "contents in polysaccharides and phenolic compounds including flavonoids (Arapitsas, 2008)."

Concerned sentence was corrected (cf. L37-38).

L41. "Phenolic compounds are known to"

Concerned sentence was corrected (cf. L38).

L42. Please, correct to: "Although the high phytochemical content of

What is phytochemical "activity"?"

Concerned sentence was corrected (cf. L39-40).

All compounds in a plant are "phytochemicals", many of which are not precisely healthy (e.g. curare, valerian, alkaloids, etc., etc.).

L64. Please, correct to: "okra seed powder" ("seed" in singular because here it functions as an adjective).

Concerned sentence was corrected (cf. L61).

L73. Please, change to: "predict their powder behavior during storage and reconstitution for an efficient use as functional"

Concerned sentence was corrected (cf. L70-71).

L76. "as well as on preservation"

Concerned sentence was corrected (cf. L73).

L92. "The powder obtained was then collected...."

As concerned sentence appeared clear and grammatically sound, it was kept unchanged.

L104-106. These two sentences should be passed to the Results and Discussion section.

Sentence in L102-104 was used to introduce the interest of analysing particle size distribution of powders. So, authors judged preferable to keep it at the beginning of the method subsection related to granulometry.

L106. Please, define clearly here in the text the meaning of 'unsieved' powders. It is really not understandable.

This was added in L92-94 when defining how unsieved powders were prepared.

L113-115. Please, re-write to: "d10, d50, and d90, which signifies the point in the size distribution, up to and including which, 90% or 50% or 10% of the total volume of material in the sample is 'contained'. For example, if the d90 is 315 micrometres, this means that 90% of the sample has a size of 315 micrometres or smaller. The span of a volume-based size distribution is defined as indicated by equation (1), which quantifies the...."

The authors preferred to redefine granulometric parameters from a statistical point of view for accuracy purposes.

L120. Please, correct to: "5 h (AFNOR, Agence Française de Normalisation, 1976)."

Concerned sentence was corrected (cf. L117)

L121. The water activity meter reported gave the water activity value with three decimal figures after the decimal point. Please, report the water activity values in the manuscript with three decimal figures after the decimal point, including in the Abstract.

Indeed, an additional significant digit was added to water activity values in the abstract and in the results and discussion sections.

L125. With this method only the extractable phenolics are being obtained, but not the chemically bound phenolics. Please, indicate this fact under 'Results and Discussion' section.

This was added in L125-127.

L136. "In test tubes, 30 μL of okra powder"

This was kept unchanged, as it was grammatically sound (cf. the numerous occurrences of "X mL distilled water" for instance : https://www.linguee.com/english-french/search?source=auto&query=ml+distilled+water).

L136. "2.67 mL of distilled"

Idem.

 L137. "500 μL of Folin-Ciocalteu"

Idem.

L137. "and 300 μL of sodium carbonate (Na2CO3, 20 % w/v) were added."

Idem.

L141. "for calibration curve (R2 = 0.99)"

The initial sentence was grammatically sound; thus, it was kept unchanged. Also, the "2" should be in exponent to mean "power 2", it was also kept unchanged.

L145-148. "The 1,1-diphenyl-2-picrylhydrazyl (DPPH) free radical scavenging method was used according to Matsuda et al. (1999). Five hundred microlitres of 0.1 mM DPPH"

Concerned sentence was corrected (cf. L144-146).

L148. "with 500 μL of okra seed powder"

As previously indicated, this was kept unchanged, as it was grammatically sound.

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L151. "of DPPH solution"
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Idem.

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L151. "and 150 \mu\text{L} of distilled water"
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Idem.

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L155. "with ABS"
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The authors did not understand this comment. "(-)" means dimensionless and unit should be given in brackets for parameters used in presented equations.

L171. "after the ABTS+• assay" (no "DPPH")

This was corrected in L170.

L173-174. "(WAC) was determined by addition of 10 mL of distilled water to 1 g of okra seed powder contained into a centrifuge tube. The mixture was stirred at 300 rpm"

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Concerned sentences are grammatically sound, thus authors preferred to keep it unchanged.

L192. "at a 5 % of confidence level"

"X % confidence level" can be employed: cf. https://www.linguee.fr/anglaisfrancais/traduction/confidence+level.html

L193. "Please, report the version, trade mark and country of origin of DSAASTAT add-on software of Excel."

It was added in L192-193.

L201. Please, also clarified here what is the meaning of "unsieved"

Unsieved is now defined in L91-94.

L202-203. Please, clarify the meaning of d50, at least between parentheses to remember it.

"d50" was replaced by "median particle size" in L202-203.

L205-206. Please, indicate here the mesh size of the 'upper' and the mesh size of the 'lower' sieves used.

"their respective" was added in L206 to indicate that upper and lower mesh sizes depend on the considered granulometric class.

L206. It was written: "confirming the efficiency of the sieving process." Why? Please, these explanations and conclusions are not understandable. Please, explain first these facts, based on the literature too.

The efficiency of the sieving process concerns its ability to separate powders into granulometric classes differing in median particle size (this was added L207-208). The fact that median particle size is found between upper and lower mesh sizes indicates that sieving was completed, i.e. particles were separated in granulometric classes according to their size comprised between the upper and lower mesh sizes ("sieves as expected when sieving is completed" was added in L203-204 to improve clarity).

L208. "in Table 1."

Concerned sentence was corrected (cf. L202).

L208. Nothing has been here reported and explained about the span (equation 1) results obtained and shown in Table 1. The values of span obtained are high, low, (????).

A discussion about span results was added in L209 -213.

L234. "The DPPH^{.+} and ABTS^{.+} assays"

The order was inverted in L237.

L239. "Figure 2."

 Figure was capitalised in L242.

L239. "All extracts were able to scavenge DPPH• radicals and this capacity increased with the speed of milling from 6,000 to 18,000 rpm, that is, with the decrease in particle sizes."

Concerned sentence was kept unchanged because it was grammatically sound, and the influence of particle size was discussed in L251,254.

L248-249. As observed in Figure 2, this is not true when comparing >=500 microns granulometry and <=180 microns for 6,000 and 12,000 rpm.

"at a given milling frequency" was added in L251 to remind that this observation concerns granulometric classes and unsieved powder obtained at the same milling frequency.

L251. "The general increase in DPPH• scavenging"

Concerned sentence was corrected (cf. L 254).

L260. "unsieved powders (Fig. 2) shows"

"(Figure 2)" was added in concerned sentence (cf. L264).

L269. "in Figure 3."

F of figure was capitalised in L271.

L290-291. It was written: "Moisture content is an important parameter that determines powdered food ingredient stability during storage." This is not true!!! It is a wrong physicochemical concept. The 'water activity' is which determines the stability and NOT the water content or moisture content. Please, correct it.

Indeed, moisture content is indirectly related to food stability during storage, as moisture content and food structure both impact food water activity, and the latter parameter is the one that is determinant for food stability during storage. Thus, it is not false to write that "moisture content is an important parameter that determines powdered food ingredient stability during storage" but it can be misleading. Concerned sentence was reformulated (cf. L290, 293), and additions were made in L300-302.

Please, begin this part (3.4) with the information of the water activity results obtained. Afterwards, report the moisture contents. The authors preferred to keep the original order: moisture content results first, and water activity results after. Indeed, this permitted to remind that water activity is the good parameter to evaluate water availability for microorganisms and alteration reactions.

L291-292. It was written: "are known to limit microbial growth and extend food shelf-life" Again, this is the same mistake. The 'water activity' is which determines the growth of bacteria, yeasts and molds. Please, see the stability map of Labuza (1977). Be careful, this is a physicochemical concept. Labuza (1977) - Figure 10: https://doi.org/10.1111/j.1745-4549.1977.tb00321.x Indeed, as indicated in a previous answer, water activity is the good parameter to evaluate water availability for microorganisms and alteration reactions. However, it is true that "low levels of moisture content are known to limit microbial growth and extend shelf-life". The threshold of 12 % (w/w) is often reported in the literature as a global limit for all types of food. Thus, it was decided to keep concerned sentence unchanged, as additions were made in L290 and 299-302 to better define the link between water moisture, water activity, and food preservability (Tapia, *et al.*, 2007; Bhandari *et al.*, 2013).

L293-295. Please, re-analyzed all results with the water activity of each granulometry.

The influence of median particle size (directly linked to granulometry) on water activity was already discussed in L290-293, and the influence of particle size was similar for all milling frequencies. Thus, no addition was made in the previous discussion of the influence of median particle size on water activity.

The moisture content is a function of the water activity of equilibration and the sorption capacity of the solid material, where the latter depends on the chemical composition and porosity.

Indeed, this was briefly added with the concept of "food structure" in L308-311.

L293. "in Table 2"

Concerned sentence was corrected (cf. L293).

L309. "in Table 2."

Concerned sentence was corrected (cf. L296).

L310-312. WAC and WSI results are not reported in Table 2. Please, complete the Table. It is then not possible to revise this part.

The result of WAC and WSI were added in table 2. The comments were added in L323-335.

L312. "high" (not 'highly', an adverb)

Concerned sentence was corrected (cf. L314).

L314. "smaller particles (Table 2)."

Table was capitalized (cf. L316).

L339. "in Figure 4."

Figure 4 was deleted. The comments were added in L345-349.

L340-341. The total explained varianza obtained is certainly lower than that recommended by the PCA method (80% to 90%).

It was added that this was sufficient to outline the main links between parameters in L348-349.

CONCLUSIONS.

L371-372. It was written: "Powders with smaller median particle sizes had lower moisture content and water activity," I'm thinking about: If powders with smaller median particle sizes had lower moisture content and also lower water activity, hence the conclusion is not that 'as the particle size was reduced bellow XXX microns, the aDsorption capacity of the solid matrix of the powder decreased.' I believe that the true conclusion is that the porous were in part collapsed by the milling process. Please, treat to indicate possible explanations for the results observed.

This was described in more details in L374-382.

L374. "food spoilage by microbial"

Concerned sentence was corrected (cf. L382).

L375. Why biochemically???

Biochemically was indicated here to refer to enzymatic reactions.

TABLES.

Footnotes: Please, delete "according to Tukey's HSD test" since this was already explained under 'Statistical analyses' section.

This information must be indicated for each table where samples were sorted by Tukey's HSD test, as it explains the meaning of superscripted letters.

Table 2. It was written in the legend: "and water solubility index of granulometric classes" Indeed, the water solubility index was not reported in the Table. Please, add a column with these results. Please, also comment and discuss them in the text.

 WAC and WSI were added in table 2 and discussed in L323-330 and 333-335.

Referee # 2

Comments to the Author

I think this is an interesting and well-written study.

A few questions:

- what is the definition of superfine milling in terms of particle size?

Superfine milling permits to reach particle sizes up to tens of micrometers. This term has been removed from the manuscript.

- why didn't you repeat with more than one okra sample? How general are the results?

Okra seeds were imported from the Republic of Congo by the first author of the article and only a limited amount of material was allowed to cross borders. It appeared important to work with a single lot of okra seeds in order to limit the impact of proximal composition variability among lots. Thus, the authors chose to work with a single lot and to produce a single replicate of each sample, but to replicate analyses three times to ensure result accuracy.

- How did you control and measure moisture content during milling and sieving? It must have a very big effect on results.

Equipments employed for milling and sieving were not equipped to control moisture content of product. This was achieved after milling and sieving steps by a thermogravimetric method (cf. L122-123). Milling and sieving steps were performed the same day for all powders in a close room where air thermohygrometry was controlled (20 °C, 40 - 50 % relative humidity). This permitted to limit the influence of surrounding air relative humidity on moisture content of produced powders.

- It is very much preferable, if you report total phenolics, to also measure and report the chemical composition of the phenolics.

The laboratory is not equipped with HPLC able to separate and quantify phenolic compounds, thus it was chosen to determine the total phenolic content only.

Please format every reference accurately and consistently -

- Scientific names in italics, and words like 'in vitro'. Genus name – upper case initial letter, species name, lower case initial letter.

This was corrected in the whole manuscript.

- Upper and lower case initial letters as required for article title, and for journal name.

This was corrected in the whole manuscript.

Pay attention to accurate format for book or chapter citation

This was corrected in the whole manuscript.

Referee #3

Comments to the Author

The scientific data is relevant and should be shared with others. Nonetheless, the manuscript has several problems.

1. All the affiliation of authors should be in English.

Affiliations are standardised by the universities of authors and should be kept in their native language.

2. In "3.4. Moisture content and water activity", \geq 500 µm granulometric classes at 6 000, 12 000, and 18 000 rpm milling speed have higher moisture content, but why did the powders have lower water activity in table 2?

Indeed, > 500 μ m granulometric classes have the highest moisture contents but intermediate values of water activity. This may be due to the fact that components of > 500 μ m granulometric classes interact more with water and decrease its bioavailability. This is consistent with the hypothesis that these higher size granulometric classes may contain more fibres, which are known to strongly retain water.

3.In Line 308-309, "The water absorption capacity (WAC) and water solubility index (WSI) results of all granulometric classes and unsieved samples of okra seed powders are presented in table 2.". However, no WAC and WSI in table 2. The data was not found in the manuscript.

WAC and WSI were added in table 2 and discussed in L323-335.

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