Specific Energy: A New Approach to Ultrasound-assisted Extraction of Natural Colorants

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Abstract  The demand for natural colorants has boosted the search for innovative technologies to obtain them. Ultrasound-assisted extraction (UAE) is one of the new green techniques that has been studied due to its process advantages that include high yields, short extraction times and nonutilization of elevated temperatures. However, a nonstandardization of the UAE variables complicates comparisons and hinders progress in the studies of this topic. In this review, the focus is the verification and discussion of which UAE process conditions authors have used to obtain natural colorants. Thus, it is possible to confirm that some authors used ultrasonic systems that are not appropriate for performing a good extraction, that some used a great amount of solvent and a long extraction time, and that researchers did not express the main variables (nominal power, extraction time and sample mass) as a function of the specific energy applied to processing. Therefore, it is possible to conclude that some studies using UAE were not conducted to obtain the best results, and the expression of the variables as a function of specific energy can generate a standardization, which facilitates comparison among the results obtained by the scientific community.

Keywords  Dyes, Sonication, Emerging technology

1. Introduction

Colors are an important characteristic of food products, and they awaken people's different expectations [1]. According to Lee, Lee, Lee and Song [2], people perceive a food via their visual perception system, and through that, predict its taste before making a decision about whether or not to purchase it. Because of this, colorants are an important food ingredient, and the global food colorant market is growing—according to Markets and Markets, a growth of approximately $0.4 bn is estimated through 2020 [3].

However, the majority of industries utilize synthetic dyes, which have been associated with health problems and cause, for instance, allergies and intolerances, especially in children [4]. Because of this, synthetic food colorants have been progressively replaced by those extracted from natural matrices [5, 6]. These are mostly carotenoids, anthocyanins, betacyanins and chlorophylls obtained from fruits and others vegetables [7].

Obtaining natural colorants, generally includes, a solid-liquid extraction, which is a separation process that involves mass transference and employs a solvent [8]. The solvent utilized for extraction depends on the vegetable matrix, the chemical properties of the pigment and the technic to be employed [9]. In many separation processes performed by industry, large quantities of volatile and flammable organic solvents are used, thus, affecting the environmental and economic performance of the overall extraction [10]. Therefore, currently there is a search for nontoxic solvents that generate less waste and reduce costs [11, 12].

In view of new tendencies, green technologies have also been studied for the extraction of different compounds [13]. For the extraction of colorants, techniques include ultrasound [14], microwave [15], pulse electric field [16], pressurized liquids [17] and supercritical fluid processing [18]. Among them, sonochemistry (the principle of ultrasound) has been mentioned as a green chemistry, evidencing various advantages, such as energy savings due to the short time of operation; major yields due to the selectivity; and a reduction in the generation of waste, by the possible use of solvents, including water [19].

The variables in ultrasound-assisted extractions (UAEs) include the system utilized, the matrix and compound to be extracted, the solvent, the proportions of solvent and feed (the amount of solid sample) and the specific energy applied to processing. Specific energy is energy per unit mass expressed as a function of the nominal power and process time. As will be seen in this review, these variables have recently been studied for the obtaining of natural food colorants. However, a lack of standardization in the
expression of these conditions is observed, for example, the authors did not clearly indicate what specify energy was applied for the extraction.

Thus, in this review, a verification and discussion are performed with respect to the variables used in UAE to obtain natural food colorants. For this, articles published between 2016 and 2018 are reviewed.

2. Ultrasound Technology

The utilization of ultrasonic techniques has been increasing over time, and they can be utilized for the analysis (low-intensity) or modification of foods (high-intensity) [20]. High-intensity ultrasound is characterized by the use of a frequency of 20 - 24 kHz and high levels of power (10 - 1000 W/cm²), which physically rupture the materials [20, 21].

The principle of this technique involves acoustic cavitation that is promoted by the system. The waves of acoustic energy promote cycles of compression and rarefaction of the molecules in the solution. Through pressure changes occurs the formation and collapse of microbubbles in the medium that result in microjetting. The microjetting generates effects such as surface peeling, erosion and particle breakdown [22, 23], promoting different applications such as the extraction of different compounds [24], microbial and enzymatic inactivation [25], emulsion formation [26] and physical modifications [27]. Among the applications, UAEs have recently been increasingly studied for the acquisition of natural food colorants.

3. Variables in Colorant Extraction by UAE

In the achievement of extracts by UAE, the relevant variables include the type of system utilized, the matrix and compound to be extract, the solvent employed, the relation of solvent/feed (S/F), the temperature and the specific energy applied [28, 29].

The results obtained by some recent studies on the UAE of natural colorants are shown in Table 1, and the variables and the best values as determined by the authors are presented.

3.1. Ultrasonic System

The UAE of natural colorants has been performed using a system with a probe (Figure 1-A) or a bath (Figure 1-B) [23, 30].

The probe system (Figure 1-A) contains a power generator, a transducer, an amplifier and a probe. The power generator produces high-frequency electrical energy of 20 kHz, which is converted to a mechanical energy by the transducer. The mechanical energy is amplified, and further, the acoustic energy is dissipated by the probe in the form of waves [23]. In the bath system (Figure 1-B), consisting of a power generator, a transducer and a bath, the power generator normally produces an energy of 40 kHz [30]. The transducer or transducers dissipate the acoustic energy into bath in the form of waves. In this system, the samples do not receive the waves directly, as is the case in the probe system.

It is possible to observe in Table 1 that the current research into colorant extractions has been performed using either of the two systems, probe or bath; however, the ultrasonic bath was not the more adequate system for this. The frequencies normally employed in this system are more than 24 kHz, such as 37 kHz [31, 32], while the physical effects dominate at lower frequencies [33]. In a study to extract anthocyanins, Cai, Qu, Lan, Zhao, Ma, Wan, Jing and Li [34], using a bath
ultrasonic system, reported that the UAE is a long-time extraction method and resulted in a low anthocyanin yield. Nevertheless, they used the bath system with an operation frequency of 37 kHz, which was not the more adequate system for this purpose.

Apart from the probe and bath systems, some extraction procedures have been performed with a combination of UAE and other techniques. For example, Wizi, Wang, Hou, Tao, Ma and Yang [35] studied the ultrasound-microwave-assisted extraction of natural colorants from sorghum husk. They obtained a yield of 3.6 times that produced via the conventional shaking method. However, the authors used equipment with a frequency of 25 kHz; thus, had a system that generates a lower frequency been used, it is possible the results would have been better.

A combined treatment using a probe system and a cell grinder was realized by Jiang, Yang and Shi [28] to obtain anthocyanins from blueberry. In this work, the authors observed that the cell grinder destroyed the cell walls, which would release water-soluble anthocyanins into any concentration of ethanol.

Comparing the two systems (the probe combined with the cell grinder and the probe), the authors noted some advantages in utilizing the combined technique.

For example, they were able to use water as the only solvent, as well as acquire a higher yield (2.12 to 2.89 mg/g) and utilize a shorter extraction time (120 to 40 min). Most likely, even though the use of combined techniques to obtain the extract promotes a better yield, it is still necessary to explore only the high-intensity ultrasound approach.

| Ultrasonic System | Matrix/Compound | Solvent | S/F (w/w) Estimated | Temperature (°C) | Process Time (min) | Power (W) | Extraction Yield | Ref. |
|-------------------|----------------|---------|---------------------|------------------|-------------------|----------|-----------------|-----|
| Probe             | Rhizomes of Curcuma/ Curcumin | Ethanol | 25 | 35 | 60 | 250 | 9.18 mg/g | [36] |
| Probe             | Pomegranate wastes/ Carotenoids | Soy oil | 10 | 51.5 | 30 | 130 | 0.67 mg/100 g | [37] |
| Probe             | Fig peel/ Anthocyanin | Ethanol | 5 | 30-35 | 21 | 310 | 3.82 mg/g | [38] |
| Probe             | Gomphrena globosa L. /Betacyanins | Water | 73 | uncontrolled temperature | 22 | 500 | 46.9 mg/g | [39] |
| Probe             | Hibiscus sabdariffa calyces/ Anthocyanin | Ethanol/ water | 30 | 30-35 | 45 | 500 | 51.7 mg/g | [40] |
| Probe             | Mulberry (Morus nigra) pulp/Anthocyanins | Methanol/ Water | 8 | 48 | 10 | 200 | 149.9 μg/g | [41] |
| Probe             | Undaria pinnatifida/ Carotenoids and Chlorophylls | Water | 30 | 50 | 30 | 300 | 34 and 0.5 mg/mL | [42] |
| Probe             | Red prickly pear peels and pulps/ betanin and isobetanin | Water | 10 | uncontrolled temperature | 10 | 400 | 57.47, 89.29 and 28.25 mg/100 g | [43] |
| Probe/ Probe and Cell grinder | Blueberry /Anthocyanins | Ethanol and Acidified Water/ Acidified Water | 20 | 25 | 120 | 1800 | 2.12 and 2.89 mg/g | [28] |
| Bath              | Purple sweet potatoes/ Anthocyanins | Ethanol/ water | 10 | 60 | 60 | 200 | 214.92 mg/100 g | [34] |
| Bath              | Wine lees/ Anthocyanins | Choline–chloride-with Malic Acid/Water | 10 | 35 | 30.6 | 341.5 | 6.55 mg/g | [31] |
| Bath              | Residues of Rubus fruticosus, Vaccinium myrtillus and Eugenia brasiliensis/ Anthocyanins | Ethanol/ Water | 20 | 80 | 90 | 580 | 2.38, 2.33 and 0.87 mg/g | [32] |
3.2. Matrix and Compound

Vegetables are a typical resource in research for various applications, due to the vast diversity of molecules [44]. However, some care must be taken to avoid variations due to the matrix. For example, depending on its cultivar, the fig acquires different colors as a function of the anthocyanin concentration [45].

Furthermore, the compounds to be extracted showed different tolerances and peculiarities. For example, anthocyanins are sensitive to temperature, pH, light, oxygen and metals, which must be considered during the separation processes to avoid loss [46]. Additionally, curcumin, according Heger, van Golen, Broekgaarden and Michel [47], degrades in the presence of sunlight and visible light.

3.3. Solvent and Feed

The physical properties of the mixture (solvent and feed) strongly influence the effectiveness of the cavitational breakdown, which needs to be proper for acoustic energy transference. For example, the solvent characteristics affect the cavitational phenomenon: the steam pressure governs the intensity of the bubble collapse, and the surface tension and viscosity govern the transient threshold of cavitation [33].

With respect to the solvent, it should be well-matched with the compound [48]. For example, the anthocyanins are normally stable under acidic conditions, and because of this, employing acidified ethanol as the solvent is an option [28]. Carotenoids possess nonpolar characteristics; thus, a good option, according to Goula, Ververi, Adamopoulou and Kaderides [37], is the utilization of vegetable oils.

Machado, Pereira, Barbero and Martínez [32] demonstrated the importance in the choice of solvent to be employed. Obtaining anthocyanins from residues of Rubus fruticosus, Vaccinium myrtillus and Eugenia brasiliensis, the authors observed different results using different solvents (water or ethanol). They observed that using water as solvent, higher extracts yields were obtained, but in relation to the antioxidant activity, they verified the opposite. The colorants in ethanol showed a higher antioxidant activity (determined using in vitro methods) than did the ones obtained with water. This phenomenon, according the authors, occurs because more compounds are soluble in water, but not the target compounds.

In addition, the solvent utilized must be GRAS (generally regarded as safe) [49] and minimize the environmental impact. In Table 1, it is possible to observe that most of the research has been performed using green solvents such as water and ethanol. Nevertheless, some, including Shirasath and Sable (23), still studied solvents such as methanol and acetone to perform the ultrasound-assisted extractions. However, the majority of authors used GRAS solvents to do the extractions, which demonstrates a tendency in the utilization of green solvents over toxic organic solvents [50].

The amount of solvent employed is also very important to obtain an efficient extraction, but an excessive quantity must be avoided to minimize the environmental impact. Table 1 shows that a large amount of the solvent was still used. Considering that in some studies S/F relations of 5 and 8 were utilized, a relation of 73 is extremely high and must be reduced to avoid the generation of a great amount of effluent.

Backes, Pereira, Barros, Prieto, Genena, Barreiro and Ferreira [38] observed that the relation of S/F is extremely important to obtain a pure extract of anthocyanin pigments from Ficus carica. Out of the values of S/F studied, the of 5 obtained the best results. This result indicated the possibility of obtaining good extraction results using a low relation of S/F.

3.4. Temperature

In addition to the cavitational and mechanical effects, the thermal effects also have a significant influence on the UAE [51]. Thus, the temperature is another important variable in the extraction.

In previous studies, high-intensity ultrasound was verified as an economically feasible technology for the extraction of thermostable compounds, but with long extractions time, it is important to pay attention to this factor. [43]. During the extraction process, there is a fast rise in the temperature of the reaction system [40]. The temperature can increase considerably, and the process can be characterized as thermal.

The increase in temperature can be favorable for the extraction of some dyes, but not for others. For example, Zhu, Wu, Di, Li, Barba, Koubaa, Roohinejad, Xiong and He [42] observed that with an increase of 10°C (40 to 50°C), the carotenoid yield increased by 8%; however, with a major increase, the yield was reduced. This result was attributed to the degradation of thermostable carotenoids. On the other hand, the chlorophylls' recovery showed a positive increase with the temperature (40 to 60°C).

It is possible to observe in Table 1 that most of the processes recently studied were performed at controlled temperatures. The evaluated temperatures varied from 25 to 80°C; however, in some studies, the temperature was not controlled. It is necessary to emphasize that temperature control is particularly important, especially when working with thermostable compounds. For example, in the extraction process realized by Roriz, Barros, Prieto, Barreiro, Morales and Ferreira [39], the temperature not was controlled, but they extracted betacyanins, which are thermostable compounds [52].

3.5. Specific Energy

The energy densities applied by ultrasound in food processes have been standardized by some researchers according to Equation 1 [26, 53, 54]. It is possible to observe in Equation 1 that the energy applied in the processes depends on the nominal power, the extraction time and the sample volume. However, the volume is a function of the pressure and temperature, and because of that, another way to express the energy is as a function of mass, which does not depend on other variables. This relation is expressed in
specific energy (Equation 2), according Rajha, Boussetta, Louka, Maroun and Vorobiev [55].

\[
ED = \left( \frac{1}{mL} \right) = \frac{\text{Nominal power} (W) \times \text{Extraction time} (s)}{\text{Sample volume} (mL)} \tag{1}
\]

\[
E = \left( \frac{1}{g} \right) = \frac{\text{Nominal power} (W) \times \text{Extraction time} (s)}{\text{Sample mass} (g)} \tag{2}
\]

3.5.1. Nominal Power

The nominal power is the power provided by the ultrasound device itself; however, this is not exactly the same value that is converted into the cavitation phenomenon [56]. This occurs due to energy loss in the equipment by dissipation during the subsequent conversions of mechanical energy into cavitation. According to Mamvura, Iyuke and Paterson [57], an energy conversion from electrical to cavitation of 9% was achieved. Shirsath, Sable, Gaikwad, Sonawane, Saini and Gogate [36] also verified through a calorimetric method that the energy efficiency of the process was approximately 5.6%. Because of that, most authors have used the highest nominal power of the equipment to do the extractions [37].

A high nominal power causes great shear forces in plant materials that results from the critical pressure and temperature obtained from the oscillation and collapse of cavitation bubbles within the solvent [40]. Thus, high values of the nominal power normally result in high extraction yields.

In the majority of the studies shown in Table 1, higher nominal powers were selected as the best condition for the extractions. According to Zhu, Wu, Di, Li, Barba, Koubaa, Roohinejad, Xiong and He [42], better results for obtaining pigments are achieved with more intense ultrasonic treatments, mainly due to the cavitation effect of ultrasound.

3.5.2. Extraction Time

The extraction time is directly associated with the nominal power supplied by the ultrasound and the samples mass (solvent and feed), as given by Equation 2. Therefore, the effect of time on the extraction is one of the most important factors; if the samples are exposed to shorter or longer times than suitable, the compounds could be degraded or not be completely extracted [39].

Thus, the time to be employed depends on the other variables. For example, Espada-Bellido, Ferreiro-González, Carrera, Palma, Barroso and Barbero [41] observed that to recover anthocyanin from mulberry, the maximum time was 10 min. According to them, this time was sufficient for a quantitative extraction and to avoid anthocyanin degradation. However, the time can probably be reduced when using a higher nominal power (greater than 200 W).

It is possible to verify in Table 1 which extractions used a long time (up to 120 min). This time can be reduced substantially. From Equation 1, it is possible that a high nominal power and a small sample mass can result in a short extraction time.

3.5.3. Sample Weight or Volume

Another criticism lies with the relationship of the sample employed in the ultrasound-assisted extractions. The majority of the studies expressed values in terms of the volume and this term is not the more suitable, considering that it can change with the temperature (Boyle-Mariotte law).

Beyond that, it is possible observe that a great amount of solvent was used in relation to a little amount of feed (Table 1). This implies that little energy was applied to the sample, decreasing the friction between the particles. The friction causes cellular rupture and the release of the compounds of interest, thus, assisting the extraction.

3.5.4. The Variable Combinations

It is important to observe the combinations of the main variables through the specific energy (E) value (Equation 2), which permits an easy understanding of the UAE process employed. The E value relates the variables of time, nominal power and sample mass and clearly represents the energy involved in the process. Furthermore, typically, a bigger E value results in the best extraction yield.

It is possible to observe some combinations in Table 1. Goula, Ververi, Adamopoulou and Kaderides [37] utilized a longer time (30 min) but a smaller nominal power (130 W) for extraction; while Koubaa, Barba, Grimi, Mhemdi, Koubaa, Boussetta and Vorobiev [43], using the same S/F (10), used a shorter time (10 min) and a larger nominal power (400 W) to obtain the dye extract.

However, the authors did not express the E value utilized and doing so would have favored a comparison of the extractions realized. In the papers reviewed, it is possible to verify that only Koubaa, Barba, Grimi, Mhemdi, Koubaa, Boussetta and Vorobiev [43] approached the process by regarding the specific energy utilized in the UAE; however, the authors did not connect the specific energy to the extraction time, thus, expressing the specific energy input in kJ/kg.

4. Quality of Dyes Obtained by UAE

Extracts from vegetables contain bioactive compounds that are valuable to the nutraceutical fields, and because of this, the extraction process is a crucial step that needs to ensure that the active ingredients are not lost or destroyed during its operation [58]. Among the methods recently revised by Nâthia-Neves and Meireles [59] to obtain natural colorants, the UAE is prominent because of the high purity of the final product. Thus, it is worth noting that in addition to obtaining high yields, it is also interesting to obtain pure extracts that facilitate the material’s application.

The ultrasound efficiency was observed by Koubaa, Barba, Grimi, Mhemdi, Koubaa, Boussetta and Vorobiev...
[43], who verified a cell denaturation after ultrasound treatment via scanning electron microscopy. According to the authors, this result can provide a better recovery of the intracellular compounds with less impurities [43]. The best results were obtained by [Backes, Pereira, Barros, Prieto, Genena, Barreiro and Ferreira [38]] who related that the UAE technique led to an extract with a greater purity of cyanidin 3-rutinoside in comparison with that of extractions assisted by heat and microwave.

Machado, Pereira, Barbero and Martínez [32] observed that among the emergent methods, UAE was the least aggressive in recovering total and individual anthocyanins, followed by the others studied, UAE + pressurized liquid (PLE) and PLE, using hydroethanolic mixtures as the solvent.

Thus, it was possible verify that ultrasound-assisted extractions are a good choice to obtain adequate results in terms of yield and quality of colorant extracts.

5. Conclusions

In this review, an evaluation of the variables that have been used in the ultrasound-assisted extractions is made, and some important aspects are observed:

✓ It is important to know the stability of the compound to be extracted to avoid loss due to the extraction operation.
✓ The solvent to be employed should be compatible with the compound of interest, be a GRAS, and be in a smaller amount than the majority of the studies have been using.
✓ The variation in temperature must be controlled during the process of extraction.
✓ The extraction time can be reduced by using a high nominal power.
✓ Larger nominal power values generate higher extraction yields.
✓ The best manner is to express the amount of solvent and feed in units of mass.
✓ The colorants obtained by UAE showed a high quality.

At the end of this review, is possible to suggest that the variables of nominal power, extraction time and sample mass are expressed in terms of the specific energy to standardize the form of expressing the energy applied to the sample. Thus, a comparison among the results of ultrasound-assisted extraction of colorants would be facilitated.

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