USE OF MIXTURE DESIGN TO IMPROVE A TROPICAL MIXED FRUIT NECTAR

PAULO HENRIQUE MACHADO DE SOUSA*
AFONSO MOTA RAMOS**
EDY SOUZA DE BRITO***
GERALDO ARRAES MAIA****
HENRIETTE MONTEIRO CORDEIRO DE AZEREDO*****
GIOVANA MATIAS DO PRADO******
MARIA LEÔNIA DA COSTA GONZAGA*******

This study aimed to develop beverages of mixed nectars of tropical fruits and to determine the best accepted formulation. The puree blend contents were: cashew apple (12.25-21.00 g 100 g⁻¹); acerola (1.75-10.50 g 100 g⁻¹); and mango (12.25-21.00 g 100 g⁻¹), varying according to a mixture design. The evaluated responses were based on physicochemical, chemical and sensorial determinations. The models for antioxidant activity, overall acceptance, ascorbic acid content, phenolic content and viscosity were statistically significant (P ≤ 0.05). The formulation with 12.25 g of cashew apple puree 100 g⁻¹; 21.00 g of mango puree 100 g⁻¹ and 1.75 g of acerola puree 100 g⁻¹ was the best accepted by the tasters. Mango and cashew apple purees must be present in a greater proportion in the mixture, since they have high flavour acceptance. The antioxidant activity of mixed nectars was most increased by acerola, followed by cashew apple. A high correlation was observed between antioxidant activity and acerola puree content and there was also a high correlation between antioxidant activity and both ascorbic acid and total phenolic contents.

KEY-WORDS: TROPICAL FRUITS; SENSORY EVALUATION; ANTIOXIDANT ACTIVITY.
1 INTRODUCTION

There is a growing market for beverages made from mixed fruits, especially tropical fruits. Such products may be carbonated or not, with a varying content of fruit puree. Mixed fruit formulations have many advantages, such as the possibility of combining different flavours, as well as nutritional and functional components, such as ascorbic acid, phenolic compounds and carotenoids.

Tropical fruits are widely accepted by consumers, and they are important sources of antioxidant compounds (REDDY, SRERAMULU & RAGHUNATH, 2010). Acerola fruit (Malpighia emarginata D.C.) has very high ascorbic acid levels and is also rich in anthocyanins and carotenoids: antioxidant pigments whose combination is responsible for its red colour (LIMA et al., 2005; ROSSO and MERCADANTE, 2005). Mangoes contain considerable levels of phenolic compounds and carotenoids (LUXIMON-RAMMA, BAHORUN & CROZIER, 2005). Cashew apple is a good source of ascorbic acid, carotenoids and phenolic compounds (KUBO et al., 2006). Cashew apple is mainly used as juice, but high astringency tends to impair its acceptance. The formulation of mixed fruits and nectars may be an efficient way of reducing the negative impact caused by cashew apple astringency. Mangoes have also been used as components of mixed nectars because of the high viscosity of their juice and their exotic and much appreciated flavour. In addition, mangoes are an important source of β-carotene, minerals and fibre (MOSTAFA, ABD-EL-HADY & ASKAR, 1997). Some authors have mentioned the possibility of using acerola puree to increase the ascorbic acid contents of several juices and nectars (SOUZA et al., 2010; JAIN and KHURDIYA, 2004; MATSUURA et al., 2004; AKINWALE, 2000).

The emphasis on the antioxidant capacity of foods is supported by indications that oxidative stress (which occurs when the rate of formation of free radicals exceeds that of their inactivation) is an etiologic factor for several chronic diseases. In physiological conditions, the aggressor compounds are controlled by an integrated and harmonic action of enzymes that depends on antioxidant nutrients provided mainly by vegetable foods, such as vitamin C and other bioactive compounds. The antioxidant action of polyphenols, the most important antioxidant group, is independent of the enzymatic system (ARAYA, CLAVIJO & HERRERA, 2006). The ingestion of foods rich in antioxidant compounds, such as vitamins C and E, carotenoids and phenolics, prevents the development of certain diseases and there is evidences which corroborates the hypothesis that dietetic antioxidants enhance the antioxidant defence system, decreasing oxidative damage (LE CORE et al., 2004).

The experimental design is the most important stage of experiments, since it makes an experiment capable of providing the exact desired information. Several designs can be used, such as complete or fractioned factorial designs, central composite designs and mixture designs. The mixture design is an important methodology for the development and optimisation of food formulations. The quality of a food product normally depends on the proportions of the single ingredients present in the formulations. Unlike classic factorial design methods, this method takes into consideration the interaction between the mixture components, which are also considered during planning and analysis of the results. The proportions of the several components of a mixture are not independent variables because the sum of the components is always 100 % (DINGSTAD, WESTAD & NAES, 2004; BJERKE, NAES & ELLEKJAER, 2000). This technique, when combined with sensory analysis, is a relevant tool to design new food products and such a combination has been used in the optimisation of fruit juice and fruit juice blends (SOUZA et al., 2007; VIEIRA and SILVA, 2004).

This study aimed to develop formulations of mixed tropical fruit nectars enriched with ascorbic acid present in acerola puree, using a mixture design to determine the proportions of cashew apple, mango and acerola purees for the best accepted formulation, and to evaluate the antioxidant activity and its correlation with the bioactive compounds of these nectars.
2 MATERIAL AND METHODS

2.1 RAW MATERIAL

Cashew apple, mango and acerola puree, freshly extracted and pasteurised by a local fruit juice industry (Ceará, Brazil) were used. The formulated blended juices were diluted in potable water and the total soluble solids were standardised with sucrose.

2.2 FRUIT PUREE CHARACTERISATION

The following determinations were performed on the purees out accordance with Instituto Adolfo Lutz (2008): pH, soluble solids (SS), titratable acidity (TA), SS/TA ratio and ascorbic acid contents.

2.3 NECTAR FORMULATION AND EXPERIMENTAL DESIGN

The formulations were composed of 35 % puree blend and sucrose and potable water up to 11 °Brix. The proportions of cashew apple, acerola and mango puree in each treatment were defined according to a simplex mixture design, with 10 treatments (MYERS and MONTGOMERY, 2002) as shown in Table 1. Since the sum of the proportions of a mixture has to be 1.0 (100 g 100 g⁻¹), the values of the fruit puree contents (whose sum was 35 g of puree 100 g⁻¹) were normalised to make the sum equal to 1.00.

The puree proportion ranges were established by preliminary tests. Acerola puree was added in the lowest proportions, due to its limited sensory acceptance (MATSUURA et al., 2004).

The fruit purees were added with water and sugar, and homogenised. Three replications were performed for each formulation. The nectar from each treatment was heated (90 °C, 60 s), hot-filled in 200 mL glass containers, closed with a plastic screw cap and cooled in running water. Samples from all the treatments were submitted to sensory evaluations, as well as physicochemical and chemical determinations.

2.4 SENSORY EVALUATION

Sensory acceptance tests were carried out according to Stone and Sidel (1993) in order to evaluate the acceptance of each formulation by potential consumers. Fifty-five non-trained tasters were enlisted. The tests were applied in individual booths equipped with daylight fluorescent lamps. The nectars were served monadically, under controlled conditions. All the tasters evaluated samples from all the treatments. The samples (30 mL) were served at normal consumption temperature (10 ± 1 °C), in glasses codified with random three digit numbers. The order of presentation was balanced according to the design proposed by Macfie et al. (1989). Global impression was evaluated by using a nine-category structured hedonic scale (PERYAM and PILGRIM, 1957). For data analysis, numerical values were associated to each category, from 1 (“disliked very much”) to 9 (“liked very much”).

2.5 PHYSICOCHEMICAL DETERMINATIONS

The chemical and physicochemical determinations were conducted at least in duplicates for each replication of the 10 treatments. The pH was determined by using a pH meter (Hanna Instruments, model HI 9321), periodically calibrated with buffered solutions (pH 4.0 and 7.0) (IAL, 2008). Soluble solids (SS) contents were determined by refractometry, by using a manual Atago refractometer model N-50E, at room temperature; the results were expressed in °Brix (IAL, 2008). Titratable acidity (TA) was determined by titrimetry by using a NaOH solution (0.1mol L⁻¹), according
to the technique described by the AOAC (1995), using phenolphthalein (1 g 100 mL⁻¹) as pH indicator; the results were expressed as g of citric acid 100 mL⁻¹ of sample. The suspended insoluble solids content determination was conducted according to the International Federation of Fruit Juice Producers (IFFJP, 1991), using an Excelsa II, model 206MP centrifuge. A 50 mL sample was centrifuged at 370 g for 10 min and the suspended insoluble solids content was measured directly in the tube, the result being expressed as mL 100 mL⁻¹ of suspended insoluble solids. Viscosity measurements were performed using a digital Brookfield rheometer (model DV-III+) coupled to a water circulation thermostatic bath (FANEM, model 111) at 25.0 °C ± 0.1 °C, and equipped with a CP52 spindle, using 0.5 mL of sample, by applying a 0.5 rpm cycle and a shear rate of 1s⁻¹ (FERRARI and ROCHA-FILHO, 2011).

2.6 CHEMICAL DETERMINATIONS

Total phenolics were determined according to the Folin-Ciocalteu method (ZIELINSKI and KOZLOWSKA, 2000). In a dark room, the samples (5 mL) were dissolved in 40 mL of distilled water (6:4 v:v) and placed in a water bath for 2 h at 85 °C for the elimination of ascorbic acid, according to Georgé et al. (2005). After cooling, the samples were placed in a 100 mL volumetric flask and the volume was completed with distilled water. The extracts were filtered under reduced pressure through filter paper (Whatman No. 1). An aliquot of 5 mL of extracts was added to 15 mL distilled water, 5 mL of Folin-Denis reagent, and 10 mL of a saturated solution of sodium carbonate, completing the volume to 100 mL with distilled water. After standing for 30 min at room temperature, the absorbance was measured at 760 nm using a UV-vis spectrophotometer (Micronal, Model B582, São Paulo, Brazil). All determinations were made in triplicate and values were calculated from the calibration curves obtained with five concentrations of gallic acid. Linearity was obtained between 0 and 5 mg mL⁻¹, corresponding to absorbance values between 0.0 and 0.5. The total phenolics were expressed as milligram of gallic acid equivalents (GAE) 100 g⁻¹ of fresh weight. The ascorbic acid (AA) content was determined by the 2,6-dichlorophenol-indophenol titration method, described in Strohecker and Henning (1967). The samples (5 mL) were diluted with 40 mL of 4 g 100 mL⁻¹ oxalic acid aqueous solution. The solution was titrated by adding the 2,6-dichlorophenol-indophenol solution until a distinct rose-pink colour persisted. Several precautions were taken in order to perform all the operations under reduced light and at 4 °C. The L-ascorbic acid was used to prepare a standard solution (0.5 mg mL⁻¹) and the ascorbic acid concentration was calculated by comparison with the standard and expressed as mg 100 mL⁻¹ sample.

2.7 ANTIOXIDANT ACTIVITY

For the antioxidant activity determination, the nectars were homogenised in a blender and centrifuged at 15000 rpm for 15 min. The supernatant was collected, filtered and its antioxidant activity was determined through the 2,2’-azinobis(3-ethylbenzothiazoline-6-sulfonic acid) (ABTS) method of Re et al. (1999) modified by Almeida et al. (2011). The ABTS radical cation (ABTS⁺) was generated by the reaction of 5 mL of aqueous ABTS solution (7 mM) and 88 µL of 140 mM (2.45 mM final concentration) of potassium persulfate solution. The mixture was held in the dark at 29 °C for 14 h before use, and then it was diluted with ethanol to obtain an absorbance of 0.7±0.02 units at 734 nm using a UV-vis spectrophotometer (Micronal, Model B582, São Paulo, Brazil). Fruit extracts (30 µL) or reference substance (Trolox) were allowed to react with 3 mL of the resulting blue-green ABTS radical solution in dark conditions. The decrease of absorbance at 734 nm was measured at the endpoint of 6 min. The standard curve was linear between 0-15 µM of Trolox 1 mL⁻¹ (final concentration). The results were thus expressed as Trolox Equivalent Antioxidant Capacity (TEAC). The activity of extracts was estimated at a minimum of three different concentrations. All tests were performed in triplicate.
2.8 STATISTICAL ANALYSIS

The mathematical description of the mixture modelling was performed by using the Statistica software, version 5.0 (STATSOFT, 1995). The models obtained for the experimental responses were evaluated in terms of their significance \((P \leq 0.05)\) and determination coefficients \((R^2)\). Correlations between antioxidant capacity and phenolics, carotenoids and ascorbic acid contents were determined using Pearson's Correlation Coefficient Test (ALMEIDA et al., 2011).

3 RESULTS AND DISCUSSION

The results of the physicochemical evaluation of the fruit puree used as raw materials are shown in Table 1. The pH, SS, TA and SS/TA ratio values from the fruit purees were consistent with those observed by other authors for acerola (VENDRAMINI and TRUGO, 2005), mango (AKINWALE, 2000; MOSTAFA, ABD-EL-HADY & ASKAR, 1997) and cashew apple (AKINWALE, 2000).

<table>
<thead>
<tr>
<th>DETERMINATIONS</th>
<th>Purees</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Cashew apple</td>
</tr>
<tr>
<td>pH</td>
<td>3.92</td>
</tr>
<tr>
<td>Soluble solids content (SS) (°Brix)</td>
<td>16.0</td>
</tr>
<tr>
<td>Titratable acidity (TA) (g of citric acid 100 mL(^{-1}))</td>
<td>0.8</td>
</tr>
<tr>
<td>SS/TA ratio</td>
<td>12.3</td>
</tr>
<tr>
<td>Ascorbic acid (mg of ascorbic acid 100 mL(^{-1}))</td>
<td>296.4</td>
</tr>
</tbody>
</table>

The acerola puree contained the highest ascorbic acid content \((688.1 \text{ mg 100 mL}^{-1})\), followed by the cashew apple \((294.4 \text{ mg 100 mL}^{-1})\) and mango \((49.0 \text{ mg 100 mL}^{-1})\) purees. The cashew apple puree presented higher ascorbic acid levels than those reported by Assunção and Mercadante (2005). The ascorbic acid content for mango was within the range reported previously by Akinwale (2000). On the other hand, the ascorbic acid levels of the acerola puree were much lower than those reported by Assis, Lima and Oliveira (2001).

The models of antioxidant activity, overall acceptance, ascorbic acid, phenolics and viscosity were statistically significant \((P \leq 0.05)\). The total acidity, pH, suspended insoluble solids and carotenoids were not significant \((P > 0.05)\). The contour graphics shown in Figure 1 represent the surface response generated by the coefficients of equations in a diagram of triangular coordinates. The experimental responses of each mixture design treatment are shown in Table 2. In some cases, the linear model was the only one with a significant F value \((p \leq 0.05)\); in others, the quadratic or special cubic models were found to be more adequate to represent attribute variations.

Analysis of Figure 1 indicates that overall acceptance was greater in mixtures with higher proportions of mango puree, followed by cashew apple puree, while the acerola puree received the lowest acceptance grades. However, the least accepted formulation still presented a good acceptance level. Only formulation 3, which had the highest acerola proportion, presented a hedonic value below 7 (“liked moderately”). The other formulations presented average hedonic values ranging from 7.0 to 7.7 (Table 2). Although the high ascorbic acid content in acerola puree is an advantage in terms of antioxidant properties, its presence in higher concentrations tended to impair the acceptance of
the nectar. This observation is in accordance with results reported by Matsuura et al. (2004). The higher acceptance of blends with higher mango puree proportions were in agreement with results obtained by Mostafa, Abd-El-Hady and Askar (1997), who observed that the mixture with mango puree enhanced the acceptance of a papaya juice.

### TABLE 2 - PROPORTIONS OF THE FRUIT PUREE COMPONENTS FOR EACH TREATMENT AND MEANS OF THE SENSORY AND PHYSICOCHEMICAL DETERMINATIONS OF THE MIXED NECTARS OBTAINED FROM THE MIXTURE DESIGN

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Independent variables</th>
<th>Dependent variables</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Cashew apple</td>
<td>Mango</td>
</tr>
<tr>
<td>1</td>
<td>21.00</td>
<td>12.25</td>
</tr>
<tr>
<td>2</td>
<td>12.25</td>
<td>21.00</td>
</tr>
<tr>
<td>3</td>
<td>12.25</td>
<td>12.25</td>
</tr>
<tr>
<td>4</td>
<td>16.80</td>
<td>16.80</td>
</tr>
<tr>
<td>5</td>
<td>16.80</td>
<td>12.25</td>
</tr>
<tr>
<td>6</td>
<td>12.25</td>
<td>16.80</td>
</tr>
<tr>
<td>7</td>
<td>15.05</td>
<td>15.05</td>
</tr>
<tr>
<td>8</td>
<td>18.20</td>
<td>13.65</td>
</tr>
<tr>
<td>9</td>
<td>13.65</td>
<td>18.20</td>
</tr>
<tr>
<td>10</td>
<td>13.65</td>
<td>13.65</td>
</tr>
</tbody>
</table>

TA = titrable acidity (g of citric acid 100 mL⁻¹); SIS = suspended insoluble solid (%); η = viscosity (mPa s); TC = total carotenoids (mg 100 mL⁻¹); TP = total phenolics (mg of gallic acid 100 mL⁻¹); Ascorbic acid (mg of ascorbic acid 100 mL⁻¹); Antioxidant activity (μM trolox mL⁻¹).

Acidity and pH varied little among the formulations, which was attributed to the similar pH and acidity values of the purees. However, a small acidity increase and pH reduction was observed and this was attributed to the acerola puree. Such tendencies became very small due to the low proportion of acerola puree used in the formulations. The acidity and pH values were in accordance with results reported by other authors (MATSUURA et al., 2004; AKINWALE, 2000). The suspended insoluble solid contents presented little variation among the formulations, ranging from 39.3 mL 100 mL⁻¹ to 45.7 mL 100 mL⁻¹ (Table 2), with the cashew apple puree contributing the most, followed by mango puree. The acerola puree contributed most to increases in the viscosity of the mixtures, followed by the mango puree (Figure 1).

The acerola puree increased the content of total phenolics, despite the small amount of puree used in the formulations. The total phenolic contents varied from 53.25 to 111.20 mg of EAG 100 mL⁻¹ of nectar and were higher when more acerola puree was used (Table 1; Figure 1).
Acerola puree was the component that most contributed to enhance the ascorbic acid content, followed by cashew apple puree. The ascorbic acid content varied considerably (66.5-202.5 mg of ascorbic acid 100 mL$^{-1}$) between the formulations (Table 1; Figure 1). Even so, a daily portion of 200 mL (volume of the package used) of the formulation presenting the lowest ascorbic acid would provide 295 % of the recommended daily intake (RDI) of this vitamin, i.e., 45 mg day$^{-1}$ (FAO/WHO, 2001). Matsuura and Rolim (2000) enhanced ascorbic acid levels of pineapple juices by mixing them with acerola juice, with little or no impairment of the sensory properties of the product when using low proportions (2.5-5 g 100 g$^{-1}$) of acerola juice. Other authors utilised cashew apple juice to enhance ascorbic acid levels of mixed fruits (AKINWALE, 2000; INYANG and ABAH, 1997). It is evident from the ascorbic acid contents of the formulations that...
acerola and cashew apple purees can be used to fortify the nutritional quality of fruit blends that are low in ascorbic acid.

The antioxidant activity varied considerably (5.1-14.7 μM trolox mL⁻¹) between the formulations and was better in the mixtures with higher proportions of acerola puree, followed by cashew apple and mango (Table 1; Figure 1). A great correlation was observed between the antioxidant activity and proportion of acerola puree; the nectars with greater amounts of ascorbic acid were responsible for the increase in the antioxidant activity.

The Pearson correlation was performed regarding antioxidant activity and the other variables studied. The antioxidant activity (TEAC) showed a significant correlation with total phenolic (r = 0.934; P ≤ 0.05) and ascorbic acid (r = 0.938; P ≤ 0.05) content, at a 5 % level of probability, presenting a highly positive correlation; while for the carotenoid content the correlation was not significant (r = 0.127; P > 0.05). Similar results were observed by Kuskoski et al. (2005), who detected a positive correlation between phenolic compounds and the antioxidant activity in fruit purees, as well as by Gardner et al. (2000), who observed that antioxidant activity was well correlated between total phenolic and ascorbic acid contents, and not with carotenoids, in several fruit juices. However, Hassimoto, Genovese and Lajolo (2005) did not find a correlation between vitamin C content and the antioxidant activity in fruits.

4 CONCLUSION

Mango and cashew apple juices must be present in a greater proportion in the mixture, since they have high flavour acceptance. On the other hand, the presence of acerola puree, even in a smaller proportion, contributes to increase ascorbic acid content in the mixture. The formulation with 21.00 g of acerola puree 100 g⁻¹, 12.25 g of cashew apple puree 100 g⁻¹ and 1.75 g of acerola puree 100 g⁻¹ received the best acceptance.

Higher proportions of acerola in the mixed fruit nectar contributed to increase the antioxidant activity, probably due to the high amounts of ascorbic acid and total phenolics in these formulations. The formulation with 12.25 g of cashew apple puree 100 g⁻¹, 12.25 g of mango puree 100 g⁻¹, and 10.50 g of acerola puree 100 g⁻¹ has the highest antioxidant activity.


31 ROSSO, V.V.; MERCADANTE, A.Z. Carotenoid composition of two Brazilian genotypes of acerola (*Malpighia punicifolia* L.) from two harvests. *Food Research International*, v.38, n.8-9, p.1073-1077, 2005.


ACKNOWLEDGMENTS

The authors would like to thank the CNPq for granting a doctorate scholarship, research scholarship and for funding this project through the Universal Edital CNPq number 019/2004.