Antioxidant properties of a tilapia (Oreochromis niloticus) diet with the inclusion of avocado by-product

Propiedades antioxidantes de una dieta para tilapia (Oreochromis niloticus) con inclusión de subproducto de aguacate

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ABSTRACT

Since synthetic antioxidants have been questioned for their potential carcinogenic and teratogenic effects, interest has grown in aquaculture for alternatives of natural antioxidants used in diets. This study analyzed antioxidant properties of a diet for tilapia with the inclusion of an avocado by-product (AP) paste. Avocado seed, skin and pulp, despite their proven antioxidant capacity, represent a great amount of waste in Mexico. We included four diets in this study: base diet 0 (BD), 10 (D10), 20 (D20) and 30 % (D30) of AP inclusion, and a commercial diet (CD) for comparison. 2,2'-azino-bis-(3-ethylbenzothiazoline-6-sulfonic acid (ABTS•+), iron ion reduction (Fe III) and concentration of total phenolic compounds (TPC). An increasing pattern was observed both in antioxidant activity and TPC as the level of AP inclusion increased. In some determinations, CD showed the highest levels, which were attributed to the synthetic antioxidants added and undeclared in these formulations. Therefore, AP was considered a good option for its inclusion in aquaculture diets because of its antioxidant properties and the added value granted with its use.

Keywords: avocado paste, by-product, antioxidant properties, diet, tilapia.

INTRODUCTION

Tilapia Oreochromis niloticus is a species that belongs to the Cichlidae family, native to Africa and valued in aquaculture industry because it shows tolerance to diseases and other biotic and abiotic factors, placing it as a species in high demand (Meyer et al. 2006). It is an omnivorous organism, and for this reason, its diet includes proteins, vegetable oils, aquatic plants, animal by-products, fermented yeast; in some cases, it includes agricultural-industrial by-products in commercial feed (González-Salas et al., 2014).

Currently, balanced feed is the input of the greatest economic impact in aquaculture since it represents from 40 to 60 % of the production costs (De Silva and Hasan, 2007). It mainly contains synthetic antioxidants, such as butylhydroxyxanisole (BHA) and butylhydroxytoluene (BHT), used as additives in food and susceptible products to lipid oxidation. Nonetheless, their cytotoxic and carcinogenic effects have been proven (Laguerre et al., 2007), which is why the food industry has shown a growing interest in substituting synthetic...
antioxidants for natural ones, both in diets for human and animal consumption (Thorat et al., 2013).

Furthermore, tilapia aquaculture, as for other species, can show pathological events due to management stress or adverse environmental conditions, such as high ammonium concentration (> 1 ppm), low temperature (< 20 ºC) and dissolved oxygen levels (< 2 mg/L), etc. This situation, in turn, may also trigger mortalities that affect productivity and profitability. Consequently, diet with antioxidant properties and natural bioactive compounds could be a key factor in organism health when facing events, such as oxidative stress (Tacon, 1989). Nowadays, vegetal sources with high content of proteins, lipids, and bioactive compounds (mainly antioxidants) can be utilized to generate functional diets that promote health, create stress resistance, decrease diseases within culture systems and minimize production costs (Escobar-Briones et al., 2006).

Avocado (Persea americana Mill.) is a subtropical fruit, native to southern Mexico, which can now be found in America, Australia, South Africa and Spain (FAO, 2005). In 2017, Mexico generated 2.29 million t of avocado, placing it as the main producer worldwide. Avocado Hass is the most popular variety in the international market for its by-products and industrial goals (SIAP, 2018). More than 10 % of avocado production could be used to feed different types of animal species or as material for diet formulation (Aviáes-Ríos et al., 2009). Avocado has aroused a great interest because of its high-quality nutritional content and bioactive compounds, such as essential fatty acids, high antioxidant power in its tissues, high carotenoid concentration and phenolic compounds, among others. Therefore, notable technological applications have been considered for avocado, as well as a natural additive and healthy food promoter (Rodríguez-Carpena et al., 2011). Based on the previous information, this study analyzed the inclusion of avocado by-product in functional food with antioxidant properties designed for tilapia that could promote its health and adequate growth.

### MATERIALS AND METHODS

#### Experimental diets

The avocado by-product was produced from fruit, which came from regional packing companies in Xalisco, Nayarit, Mexico, unsuitable for commercialization, for either inadequate size, alteration, spots, abrasion or lesion in the epithelium. Subsequently, the seed was removed manually to grind the pulp and skin in a hammer blade forage grinder until a homogeneous paste was obtained; then, it was vacuum packed and stored at 7 ºC until use. Four experimental diets were formulated for tilapia adults, according to the base formulation proposed by Bureau and Cho (1994), all with 35 % protein and 10 % lipids (Table 1), producing pellets in a meat grinder (Torrey Model 22) and drying for 12 h at 45 ºC. The inclusion of avocado by-product was adjusted to levels of 10, 20 and 30 % in wet basis (D10, D20 and D30). The base diet (BD) was made without the inclusion of avocado paste, while the commercial diet (CD: 35 % protein and 10 % lipids) was acquired from a local establishment (PRONUA, SA de CV, México).

#### Proximal analysis

Before formulating the experimental diets, we performed a proximal analysis of the raw materials, which consisted on determining humidity, lipids, protein, and ashes, also performed to the experimental diets, with the addition of fiber determination (A.O.A.C., 2012).

#### Ethanolic extraction

To assess antioxidant properties, three samples were taken at random of the formulated diets in the laboratory, commercial diet, and avocado paste (previously lyophilized) with a weight of 7.5 g; subsequently, samples were dissolved in 150 mL of ethanol. The extraction was performed with a Soxhlet equipment for three hours of reflux; the extracts were filtered with Whatman # 4 filters and stored at 4 ºC until subsequent analyses based on the method described by Maisuthisakul et al. (2009). In the case of avocado paste, at

<table>
<thead>
<tr>
<th>Component (g/100g)</th>
<th>Experimental diets</th>
</tr>
</thead>
<tbody>
<tr>
<td>BD</td>
<td>D10</td>
</tr>
<tr>
<td>Wheat meal</td>
<td>40.40</td>
</tr>
<tr>
<td>Soy meal</td>
<td>19.10</td>
</tr>
<tr>
<td>Fish meal</td>
<td>21.70</td>
</tr>
<tr>
<td>Gluten</td>
<td>5.00</td>
</tr>
<tr>
<td>Fish oil</td>
<td>11.30</td>
</tr>
<tr>
<td>Unflavored gelatin</td>
<td>1.50</td>
</tr>
<tr>
<td>Avocado paste</td>
<td>0.00</td>
</tr>
<tr>
<td>Vitamin Premix</td>
<td>0.50</td>
</tr>
<tr>
<td>Mineral Premix</td>
<td>0.50</td>
</tr>
</tbody>
</table>

* Vitamin Pre-mix: Vitamin A, 536 KUI/kg; Vitamin D\_3, 133.5 KUI; Vitamin E, 6,666.5 mg; Vitamin K\_1, 666.862 mg; Vitamin B\_1, 1,000.04 mg; Vitamin B\_2, 1,333.32 mg; Vitamin B\_6, 1,33 mg; vitamin C, 10,000.2 mg; Nicin, 5,334.195 mg; Panthothenic acid, 2,666.7 mg; Folic acid, 332.8 mg; Biotin, 33.3 mg; Choline chloride, 40,020.6 mg. DSM Nutritional Products México S. A. de C.V.

** Table 1. Formulation of the base diet (BD) and diets with avocado by-product inclusion at 10, 20 and 30 % (D10, D20 and D30).**

**Tabla 1. Formulación de dieta base (BD) y dietas con inclusión de subproducto de aguacate al 10, 20 y 30 % (D10, D20 y D30).**
the end of the analyses, results were converted to wet basis as it was used during diet formulation.

**Antioxidant activity**

**DPPH scavenging activity**

The determination was performed according to the methodology reported by Morales and Jiménez-Pérez (2001), using the stable free radical DPPH* as an assay method. Results are expressed in μmol equivalent to Trolox/g (μmol ET/g), making a standard Trolox (6-Hydroxy-2, 5, 7, 8-tetramethylchroman-2-carboxylic acid) curve from 0 to 500 μmol/L.

**ABTS**+- scavenging capacity

The scavenging ability of the stable cation ABTS**+- was determined according to the method reported by Re et al. (1999) and Kuskoski et al. (2004). The assay radical was determined with the ABTS**+- reactive at a concentration of 7 mM with potassium persulfate at 2.45 mM, incubated at an approximate temperature of 7 °C and in darkness for 16 h. Once the ABTS**+- radical was formed, it was diluted with ethanol until an absorbance of 0.700 (± 0.1) at 754 nm was obtained. The reference antioxidant was Trolox, assayed at a 0-1.5 mM concentration range, expressing the results in mmol Trolox equivalent/g of sample (mmol TE/g).

**Ferric Reducing Antioxidant Power (FRAP)**

Reduction ability of Fe (III) ion to Fe (II) ion was performed following the method of Hinneburg et al. (2006). Determination was performed at an absorbance of 700 nm, and ferric reduction (III) was determined as Trolox equivalent (μmol TE/g).

**Determination of total phenolic compounds (TPC)**

The phenolic compounds were analyzed using the Folin-Ciocalteu reaction and according to that reported by Stintzing et al. (2005). Compound concentration was obtained from a standard curve of gallic acid (0 to 400 mg/L), from which the results were calculated as mg gallic acid equivalent/gram of sample (mg GAE/g).

**Statistical analysis**

For data analysis, we used a completely randomized design, subjecting results to a one-way analysis of variance (ANOVA), taking as a main factor the five diet types (including the commercial diet). A Tukey’s mean comparison test was performed, considering a significance level of 5 % (p < 0.05). All analyses were performed with the software IBM SPSS Statistics 2.0.

**RESULTS AND DISCUSSION**

**Experimental diets**

Table 2 shows the proximal composition of the ingredients used for the experimental diet formulations, highlighting that the protein ingredients were unflavored gelatin, gluten and fish meal with 84.04 ± 0.48, 65 ± 0.17 and 55.92 ± 0.11 % of protein, respectively. Likewise, the ingredients with greater lipid content were fish oil (97.57 ± 0.37 %), followed by avocado paste (AP) with a percentage of 13.81 ± 0.20, which was closer to that reported by Hernández-López (2016) for AP in which pulp, skin and seed were included (14.16 %).

In the proximal analyses of the experimental diets (Table 3), no significant differences (p > 0.05) were observed for the percentage of the main components (protein and lipids). In the proximal analysis of the experimental diets (Table 3), there were significant differences (p < 0.05) for the percentage of the main components. However, these components keep with the established requirements for the study organisms (protein = 35 % and lipids = 10 %). In the case of fiber, it is considered indigestible because tilapia does not have the enzymes required for its digestion. Nonetheless, it is well-known that its contribution in diets is essential to modulate food transit through the gastrointestinal tract and facilitate nutrient absorption. Furthermore, cellulase activity is present in microorganisms that colonizes O. mossambica intestine (Saha et al., 2006). According to that indicated by the Secretaría de Agricultura, Ganadería, Desarrollo Rural, Pesca y Alimentación (SAGARPA, 2009) in Mexico, in general, crude fiber levels in food for tilapia should not be greater than 5 %. It is important to point out that the formulated diets did not go beyond the maximum composition level of crude fiber with the exception of CD.

On the other hand, CD was the one with highest

![Table 2](image)

**Table 2. Proximal composition of the components used in experimental diet formulation in wet basis (%).**

<table>
<thead>
<tr>
<th>Component</th>
<th>Humidity</th>
<th>Proteins</th>
<th>Lipids</th>
<th>Ashes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Wheat meal</td>
<td>12.71 ± 0.19</td>
<td>13.02 ± 0.28</td>
<td>1.80 ± 0.03</td>
<td>0.60 ± 0.02</td>
</tr>
<tr>
<td>Soy meal</td>
<td>10.55 ± 0.04</td>
<td>43.05 ± 0.28</td>
<td>0.55 ± 0.07</td>
<td>6.19 ± 0.17</td>
</tr>
<tr>
<td>Fish meal</td>
<td>12.03 ± 0.25</td>
<td>55.92 ± 0.11</td>
<td>2.10 ± 0.08</td>
<td>18.94 ± 0.07</td>
</tr>
<tr>
<td>Unflavored gelatin</td>
<td>11.04 ± 0.29</td>
<td>84.04 ± 0.48</td>
<td>0.08 ± 0.01</td>
<td>1.89 ± 0.04</td>
</tr>
<tr>
<td>Gluten</td>
<td>8.53 ± 0.18</td>
<td>65.00 ± 0.17</td>
<td>1.44 ± 0.20</td>
<td>0.80 ± 0.05</td>
</tr>
<tr>
<td>Fish oil</td>
<td>0.92 ± 0.48</td>
<td>ND</td>
<td>97.57 ± 0.37</td>
<td>ND</td>
</tr>
<tr>
<td>Avocado paste</td>
<td>70.93 ± 0.22</td>
<td>1.37 ± 0.01</td>
<td>13.81 ± 0.20</td>
<td>4.25 ± 0.32</td>
</tr>
</tbody>
</table>

Values represent the mean and standard deviation (n = 3).

ND: Not determined.
percentage of ashes, which depends on fish meal used and the amount of minerals added in the preparation of diet, and whose percentage is unknown due to formula confidentiality. Fish meal could have from 10-20 % ashes, but when it is higher than 23 %, protein quality and digestibility is decreased (Galleguillos and Romero, 1994). Therefore, fish meal did not represent a risk in the diets made in this study since ash percentage was 18 % and its inclusion decreased as the level of AP inclusion increased.

Antioxidant activity

DPPH•

In the DPPH• assay results, AP anti-radical activity was 5.14 ± 0.04 μmol TE/g, which is three times greater than the previously reported for avocado Hass pulp (Wang et al., 2012; Moreno et al., 2014). The previous results could be due to the presence of carotenoids, tocopherols, tocotrienols and other phenolic compounds that were present in greater proportion in skin (Rodríguez-Carpena et al., 2011; Wang et al., 2012; Daiuto et al., 2014). Likewise, the formulated diets showed significant differences (p < 0.05) among them (Figure 1) since they increased in the antiradical activity in proportion to AP inclusion. It is worth to point out that no differences (p > 0.05) were found between D30 and CD where the antiradical activity of the latter could be attributed to the addition of synthetic antioxidants (frequently undeclared) or to the concentration of synthetic analogs, for example, vitamin E (Lundebuye et al., 2010).

Previously, several experimental diets have been formulated with antioxidant properties and, some of them assessed antiradical ability (against DPPH•). Ahmed et al. (2015) prepared a diet for tilapia with the inclusion of edible mushroom waste; Carbonera et al. (2014) designed an antioxidant diet for tilapia with residual extract of acerola; Lim and Lee (2011) assessed diets for tilapia including soy meal and fermented cotton seeds with Aspergillus oryzae. Similarly, Kim et al. (2013) supplemented a diet for Paralichthys olivaceus with spirulina and quercetin; and Kim et al. (2009) utilized Korean “Meju” and fermented soy meal also with the same microorganism but in a diet for parrot fish Oplegnathus fasciatus. In general, all these studies attributed the antioxidant ability against DPPH• radical to the content of phenolic compounds in each one of the non-conventional ingredients implemented. As previously mentioned, in the case of this study, the greatest antiradical activity in comparison with the reference diet (BD) could be attributed to different bioactive phenolic compounds present in the by-product used.

ABTS++

In the ABTS++ assay, AP showed a scavenger ability of this cation of 78.13 ± 0.41 mmol TE/kg, which was greater than the reported in avocado Hass pulp by Pellegrini et al. (2003) and Daiuto et al. (2014) with 2.22 and 15.22 mmol TE/kg, respectively. The previous results can be attributed once more to incorporating skin to the paste, since an antiradical activity had been reported related to the lipophilic and hydrophilic compound concentration, mainly carotenoids, tocopherols procyanidins and other phenolic compounds found in this tissue in high concentrations (Rodríguez-Carpena et al., 2011; Wang et al., 2012).

The behavior of the experimental diets (Figure 2) showed that D30 had a maximum increase in scavenger ability against ABTS++ with a value of 8.23 ± 0.22 mmol ET/kg. For this reason, we can consider the inclusion of PA as the increase promoter with respect to the possible bioactive compounds present acting as antioxidants. It is worth...

**Figure 1.** DPPH• free radical scavenger ability of experimental diets. Bars show the average and standard deviation of three replicates. BD: base diet (0 %); D10, D20 and D30: Diets formulated with 10, 20 and 30 % of avocado paste inclusion; CD: Commercial diet. Different letters in bars indicate significant differences (p < 0.05).

**Figure 2.** ABTS++ free radical scavenger ability of experimental diets. Bars show the average and standard deviation of three replicates. BD: base diet (0 %); D10, D20 and D30: Diets formulated with 10, 20 and 30 % of avocado paste inclusion; CD: Commercial diet. Different letters in bars indicate significant differences (p < 0.05).
highly suggesting that the CD antioxidant activity in this assay was greater than all diets formulated and produced in the laboratory, which could be attributed to the synthetic antioxidants normally added and frequently underestimated.

At the present time, few studies are available where antioxidant activity in aquaculture diets has been assessed by this radical, since the majority have focused on the analysis of the antiradical DPPH• activity, in some cases together with total phenolic compound determination (Lim and Lee, 2011; Kim et al., 2013; Ahmed et al., 2015). Montoya et al. (2014) implemented diets that included tomato and lycopene extract in functional diets designed for goldfish Carassius auratus and platy Xiphophorus maculatus. The diets with the greatest antiradical ABTS+• ability were those supplemented with 0.6% (w/w) with 19.4 mmol TE/kg, with greater value than that shown by D30 (8.23 ± 0.22 mmol TE/kg). About this, it is important to mention that in the present work we used an avocado by-product, considered waste in current agricultural food industry, instead of a purified commercial product as the lycopene used in the reference study, which is also more expensive.

**Ferric Reducing Antioxidant Power**

From the FRAP assay antioxidant activity results, AP showed an ability to reduce Fe (III) ion at 20.23 ± 0.24 μmol TE/g of sample. Such capacity can be attributed to the presence of phenolic compounds and ascorbic acid in the fruit (Wang et al., 2012). According to previous studies, approximate values of 15-22 μmol ET/g have been reported in avocado Hass pulp (Gorinstein et al., 2011; Moreno et al., 2014), which can be considered close to those found in this study in AP.

The diets formulated in this study were significantly (p < 0.05) affected (Figure 3), showing an increase in their ability to reduce Fe (III) ion according to the inclusion formulation level, in which D30 recorded the greatest reduction ability in comparison with CD (12.77 ± 0.15 mg TE/g and 13.26 ± 0.42 mg TE/g respectively), without being statistically different (p > 0.05). In the case of D30 and the rest of avocado by-product formulated diets, the antioxidant activity in this assay can be attributed to the bioactive compounds in this fruit. In the case of BD, to antioxidant compounds present in some base components, such as soy and wheat meal or the pre-mixture of vitamins and minerals (Devi et al., 2009; Lv et al., 2012). With respect to CD, its high ability to reduce Fe (III) ion could be attributed in a great measure to the content of synthetic antioxidants added during its formulation.

Similar to the ABTS +• assay, few studies are available in aquaculture where the antioxidant ability in diets has been assessed through this method. This procedure is important since a clear idea of the type of compounds present can be obtained by performing different tests related to antioxidant properties, as well as the possible mechanisms of action. In general, DPPH assay is considered for hydrophobic compounds, FRAP mainly for hydrophilics, and ABTS +• for both types function as amphipathic assay (Opitz et al., 2014).

**Phenolic compounds**

The concentration of phenolic compounds in AP was 20.91 ± 0.28 mg GAE/g of sample. This concentration was 15-20 times greater than the reported by Wang et al. (2010), Moreno et al. (2004) and Daiuto et al. (2004), who found 4.9, 5.82 and 3.3 mg GAE/g, respectively, in avocado Hass pulp. This could be attributed to skin inclusion of this fruit in AP since a greater concentration of phenolic compounds have been reported in this tissue than in pulp (Daiuto et al., 2004; Wang et al., 2010; Rodriguez-Carpena et al., 2011).

On the other hand, diets showed significant differences with respect to phenolic compound content (p < 0.05), observing a clear tendency to increase its values as inclusion of AP increased (Figure 4). Regarding this, D30 was the diet with the greatest concentration of phenolic compounds, including BD and CD. As to BD, despite AP was not included during formulation, its phenolic compound concentration was considerable, and it could be attributed to the content in the components used during formulation, such as soy and wheat meal or vitamin pre-mixture (Devi et al., 2009; Lv et al., 2012). Although CD showed a lower concentration than other diets, the presence of these compounds could be attributed to BHT, BHA, BHQ that are synthetic phenolic-type substances normally included in commercial diets (Lundebye et al., 2010).

Few studies are available analyzing aquaculture diets about total phenolic compounds. Obasa et al. (2013) reported 5.6 mg GAE/g in a diet for tilapia including seed meal from fermented mango; Oniszczuk et al. (2019) designed a diet for carp utilizing eastern purple coneflower (Echinacea purpurea (L.) Moench.) as part of the components, finding a maximum value of 4.2 mg GAE/g in the diet with the greatest inclusion.
of the plant (60 g/kg). Lim and Lee (2011) experimented with a diet designed for tilapia including soy meal and fermented cotton seeds, finding values around 3.8 mg GAE/g. All these values were lower than that found for D30 (best functional diet with 12.58 ± 0.11 mg GAE/g) in this study, which could be attributed mainly to incorporating avocado by-product as component, that showed a great concentration of these bioactive compounds.

CONCLUSIONS

As previously mentioned, aquaculture diet components need to have a concentration of bioactive compounds capable of promoting antioxidant activity. They are important in feed, both for an adequate conservation and to avoid deterioration due to oxidative processes, as well as in species that consume it to mitigate the oxidative stress caused by exploitation conditions. In this context, the diet including 30 % of avocado by-product showed the greatest antioxidant properties among the diets that included AP, so it can represent a greater advantage against oxidation by reactive species. Likewise, its formulation showed the greatest percentage of prosopoploitation conditions. In this context, the diet including 30 % of avocado paste should be considered an interesting and novel by-product to be included as an alternative nutrient source in aquaculture diets, such as lipids and bioactive compounds with antioxidant properties.

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