Rancidity as Lipolytic Index in Poultry Meat Cuts
Faten, S. Hassanin; Hassan M.A.; Nabila, I. Elsheikh; Tereza, H. Amin

1 Food Control Department, Faculty of Veterinary Medicine, Benha University
2 Animal Health Research Institute, Tanta Gharbia

ABSTRACT

Ninety random samples of fresh poultry meat cuts represented by chicken thigh & breast, duck thigh & breast and turkey thigh & breast (15 of each) were collected from different slaughtered shops Tanta city, Gharbia government. The collected samples were analyzed for determination of thiobarbiturc acid (TBA), perioxide value (P.V.) free fatty acids (FFAs) and fractionation of fatty acids to determine their lipolytic indices for their keeping quality as well as their composition of fatty acids. The obtained results showed that mean values for duck thigh, chicken thigh and turkey thigh samples were 0.52 ± 0.03, 0.33 ± 0.02 and 0.15 ± 0.01 (mg/kg) for TBA & 0.85 ± 0.09, 0.57 ± 0.07 and 0.33 ± 0.04 (mgO₂/kg) for PV, respectively. Plus, 0.69 ± 0.07, 0.45 ± 0.06 and 0.28 ± 0.02 (mg %) for FFAs, respectively. Concerning breast of both duck, chicken and turkey samples, the mean values were 0.40 ± 0.02, 0.19 ± 0.01 and 0.12 ± 0.01 (mg/kg) for TBA, where, 0.64 ± 0.07, 0.41 ± 0.05 and 0.27 ± 0.02 (mgO₂/kg) for PV. As well as, 0.52 ± 0.05, 0.36 ± 0.03 and 0.20 ± 0.03 (mg %) for FFAs respectively. Finally, samples subjected to fractionation of fatty acids the ratio between total unsaturated fatty acids and total saturated fatty acids were 0.96, 1.23 and 1.66 for thigh of duck, chicken and turkey, respectively. While breast of duck, chicken and turkey ratios were 1.15, 1.45 and 2.23 respectively. Turkey breast meat had the highest keeping quality and nutritive value compared with the other samples.

Keywords: TBA, PV, FFA, Poultry

with chicken is associated with significant decrease in a polypoprotein B and total cholesterol levels in microalbumin uric type 2 diabetic patients this effect is probably related to the higher PUFA (polyunsaturated fatty acids) content of chicken meat in comparison to beef (Gross et al., 2002).

The microbial activities are considered the major factor of foods alteration during storage and manipulation, when these activities are under efficient control, the foods alteration will be of chemical nature, lipid oxidation is a principal chemical changes of foods, which depends on the level of oxygen and metals. Lipid oxidation products are responsible for the development of rancidity by the production of low molecular weight compounds that cause undesirable flavor (Frankel et al., 1987).

Cholesterol is also oxidized in similar reaction mechanisms to those observed of fatty acids. Many of the cholesterol oxidation products have adverse effect such as cytotoxicity and modifications of enzyme activity (Bosinger et al., 1993) atherosclerosis (Kumar and Singal, 1991) carcinogenicity and mutagenicity (Ansari and Smith, 1979).

Oxidative deterioration results in losing the quality of poultry meat due to development of rancid odor and taste. Moreover, the rancid flavor can develop rapidly during refrigerated storage of such frozen poultry meats, which are more susceptible to rancidity because of their high contents of unsaturated fatty acid (Ang, 1988).

The thiobarbituric acid (TBA) method is the most widely used test for measuring the extent of lipid peroxidation in red meat and poultry, due to its speed and simplicity (Raharjo and Sofos, 1993).

Therefore, the aim of this present study was to evaluate the lipolytic indices for keeping quality and fat composition for nutrition value of poultry meat.

2. Materials and methods

2.1. Collection of samples:

Ninety random samples of fresh poultry meats represented by chicken thigh & breast, duck thigh & breast and turkey thigh & breast (15 of each) were collected from different slaughtered poultry shops in Tanta city, Gharbia government. All collected samples were kept in a separated sterile plastic bag and preserved in an ice box then transferred as quickly as possible to the laboratory with a minimum limit of delay and then subjected to following examination.

2.2. Determination of Thiobarbituric Acid Number (TBA) Pikul et al. (1989):

2.3. Determination of Peroxide value (Asakawa and Matsushita, 1978):

2.4. Determination of Free Fatty Acids (FFA):

2.4.1. Lipid Extraction Using the Folch Method (Folch et al., 1957):

2.4.2 Titration procedure (Brake and Fennema, 1999):

2.5. Fractionation of fatty acids:

2.5.1. Extraction of fat from chicken meat:

One hundred grams of the sample were placed in a 500 ml closed stopper flask then, 300 ml of n-hexane were added, and the flask was shacked for 30 min. using horizontal shaker and left for 24 hours at room temperature. The homogenated mixture was filtered and the residue was re-extracted as mentioned above. The combined filtrates were evaporated under reduced pressure, according to AOAC, (2000).

2.5.2. Identification and determination of fatty acids:

Fatty acids were determined in meat by Gas Chromatography technique (GC) according to Aura et al. (1995).

2.5.3. Isolation and extraction of fatty acids:

The fats under study were saponified with ethanolic potassium hydroxide (40%, w/v) for
24 hours at room temperature according to the method of AOCS (1993).

The aqueous layer (containing potassium salt of fatty acids and free from unsaponifiable matter) was acidified with HCL (0.5N), and then it was extracted three times with petroleum ether. The petroleum ether extract was washed several times with distilled water, and dried over anhydrous sodium sulphate.

2.5.4. Methylation of fatty acids (Vogel, 1975):

2.5.5. Separation of fatty acid methyl esters:
The fatty acids methyl esters were analyzed by Hewlett Packard gas chromatography (5890 series) equipped with flame ionization detector. The chromatograph was fitted with FFAP (2.5m × 0.30µm film thickness and 0.32mm diameter).

Capillary column coated with polyethylene glycol. The column oven temperature was programmed from 50°C to 240°C (7°C /min.) and finally kept at 240°C for 30 minutes. Injector and detector temperature were 250 and 260°C, respectively. Gases flow rates were 33, 30 and 330 ml/min. for N2, H2 and air, respectively.

The flow rate inside column was 2ml/min. Under these conditions, all peaks from C8 to C22 homologous series well defined. Peak identification was performed by comparison of the relative retention time (RTT) for each peak with those of standard chromatograms. The peak was measured by triangulation and the relative proportions of the individual compound were therefore obtained by determination of the partial areas in relation to the total area.

3. RESULTS

From achieved results in table (1) it is showed that mean values of TBA “mg/kg” in examined thigh samples were 0.52±0.03 for duck samples, 0.33±0.02 for chicken samples and 0.15±0.01 for turkey samples. Concerning breast sample, the mean value of TBA were 0.40±0.02, 0.019±0.01 and 0.12±0.01 for duck, chicken and turkey, respectively.

The data recorded in table (2) declared that perioxide mean values were 0.85±0.09, 0.57±0.08 and 0.33±0.04 for duck, chicken and turkey thigh samples, respectively. While breast sample mean values were 0.64±0.07, 0.11±0.05 and 0.27±0.02 for duck, chicken and turkey, respectively.

It is evident from results obtained in table (3) that free fatty acids mean values were 0.69±0.07, 0.45±0.04 and 0.28±0.02. For duck, chicken and turkey thigh samples, respectively. On the other hand breast samples mean values were 0.52±0.05, 0.36±0.03 and 0.20±0.03 for duck, chicken and turkey, respectively.

It is obvious from results obtained in table (4) that total saturated fatty acids (TSFAs) were 2481 & 2253; Total mono unsaturated fatty acids (TMUFAs) were 1660 & 1741. Plus, total poly unsaturated fatty acids (TPUFAs) were 745 & 869 in thigh and breast of duck samples respectively. Ratio between total unsaturated FAs and total saturated FAs was 0.969 for thigh samples and 1.15 for breast samples.

Concerning results in table (4), total saturated fatty acids (TSFAs) were 2164 & 1959, total mono table unsaturated fatty acids (TMUFAs) were 1793 & 1872 and total poly unsaturated fatty acids (TPUFAs) were 884,967. For thigh and breast of chicken samples, respectively. Ratio between total unsaturated fatty acids to total saturated fatty acids was 1.23 for thigh samples and 1.45 for breast samples.

Also, in table (4) the results achieved in thigh and breast of turkey samples of total saturated fatty acids, total mono unsaturated fatty acid and total poly unsaturated fatty acid were 1754 & 1408, 1885 & 2012 and 1028 & 1136, respectively. Ratio between total
unsaturated fatty acids to total saturated fatty acids was 1.66 for thigh samples and 2.23 for breast samples.

Table (1): Statistical analytical results of Thiobarbituric acid value (TBA) "mg/Kg" in the examined samples of thigh and breast of various poultry meats (n=15).

<table>
<thead>
<tr>
<th>Cuts</th>
<th>Species</th>
<th>Min</th>
<th>Max</th>
<th>Mean ± S.E</th>
<th>Min</th>
<th>Max</th>
<th>Mean ± S.E</th>
</tr>
</thead>
<tbody>
<tr>
<td>Thigh</td>
<td>Duck</td>
<td>0.36</td>
<td>0.64</td>
<td>0.52 ± 0.03</td>
<td>0.29</td>
<td>0.51</td>
<td>0.40 ± 0.02</td>
</tr>
<tr>
<td></td>
<td>Chicken</td>
<td>0.18</td>
<td>0.39</td>
<td>0.33 ± 0.02</td>
<td>0.11</td>
<td>0.25</td>
<td>0.19 ± 0.01</td>
</tr>
<tr>
<td></td>
<td>Turkey</td>
<td>0.07</td>
<td>0.21</td>
<td>0.15 ± 0.01</td>
<td>0.06</td>
<td>0.19</td>
<td>0.12 ± 0.01</td>
</tr>
</tbody>
</table>

S.E = standard error of mean  
High significant differences (P<0.01)

Table (2): Statistical analytical results of Peroxide value (PV) "meqO2/kg" in the examined samples of thigh and breast of various poultry meats (n=15).

<table>
<thead>
<tr>
<th>Cuts</th>
<th>Species</th>
<th>Min</th>
<th>Max</th>
<th>Mean ± S.E</th>
<th>Min</th>
<th>Max</th>
<th>Mean ± S.E</th>
</tr>
</thead>
<tbody>
<tr>
<td>Thigh</td>
<td>Duck</td>
<td>0.68</td>
<td>1.03</td>
<td>0.85 ± 0.09</td>
<td>0.47</td>
<td>0.83</td>
<td>0.64 ± 0.07</td>
</tr>
<tr>
<td></td>
<td>Chicken</td>
<td>0.35</td>
<td>0.76</td>
<td>0.57 ± 0.08</td>
<td>0.23</td>
<td>0.58</td>
<td>0.41 ± 0.05</td>
</tr>
<tr>
<td></td>
<td>Turkey</td>
<td>0.21</td>
<td>0.53</td>
<td>0.33 ± 0.04</td>
<td>0.16</td>
<td>0.44</td>
<td>0.27 ± 0.02</td>
</tr>
</tbody>
</table>

S.E = standard error of mean  
High significant differences (P<0.01)

Table (3): Statistical analytical results of free fatty acids (mg %) in the examined samples of thigh and breast of various poultry meats (n=15).

<table>
<thead>
<tr>
<th>Cuts</th>
<th>Species</th>
<th>Min</th>
<th>Max</th>
<th>Mean ± S.E</th>
<th>Min</th>
<th>Max</th>
<th>Mean ± S.E</th>
</tr>
</thead>
<tbody>
<tr>
<td>Thigh</td>
<td>Duck</td>
<td>0.49</td>
<td>0.87</td>
<td>0.69 ± 0.07</td>
<td>0.45</td>
<td>0.71</td>
<td>0.52 ± 0.05</td>
</tr>
<tr>
<td></td>
<td>Chicken</td>
<td>0.30</td>
<td>0.64</td>
<td>0.45 ± 0.06</td>
<td>0.21</td>
<td>0.48</td>
<td>0.36 ± 0.03</td>
</tr>
<tr>
<td></td>
<td>Turkey</td>
<td>0.19</td>
<td>0.36</td>
<td>0.28 ± 0.02</td>
<td>0.15</td>
<td>0.32</td>
<td>0.20 ± 0.03</td>
</tr>
</tbody>
</table>

S.E = standard error of mean  
High significant differences (P<0.01)
Table (4): Fractionation of fatty acid composition (mg/100 g) in the examined samples of duck, chicken and turkey meat cuts.

<table>
<thead>
<tr>
<th>Fatty acids</th>
<th>Duck meat</th>
<th>Chicken</th>
<th>Turkey</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Thigh</td>
<td>Breast</td>
<td>Thigh</td>
</tr>
<tr>
<td>Lauric acid (C12:0)</td>
<td>59</td>
<td>42</td>
<td>41</td>
</tr>
<tr>
<td>Myristic (C14:0)</td>
<td>106</td>
<td>95</td>
<td>95</td>
</tr>
<tr>
<td>Palmitic (C16:0)</td>
<td>1582</td>
<td>1430</td>
<td>1378</td>
</tr>
<tr>
<td>Stearic (C18:0)</td>
<td>734</td>
<td>686</td>
<td>650</td>
</tr>
<tr>
<td>Total Saturated F.As</td>
<td>2481</td>
<td>2253</td>
<td>2164</td>
</tr>
<tr>
<td>Palmitoleic (C16:1)</td>
<td>290</td>
<td>347</td>
<td>356</td>
</tr>
<tr>
<td>Oleic (C18:1)</td>
<td>1370</td>
<td>1394</td>
<td>1437</td>
</tr>
<tr>
<td>Total Mono- Unsaturated F.As</td>
<td>1660</td>
<td>1741</td>
<td>1793</td>
</tr>
<tr>
<td>Linoleic (C18:2)</td>
<td>516</td>
<td>558</td>
<td>597</td>
</tr>
<tr>
<td>Linolenic (C18:3)</td>
<td>74</td>
<td>83</td>
<td>89</td>
</tr>
<tr>
<td>Eicosadienoic acid (C20:2)</td>
<td>10</td>
<td>17</td>
<td>14</td>
</tr>
<tr>
<td>Dihomo-γ-linolenic (C20:3)</td>
<td>16</td>
<td>24</td>
<td>20</td>
</tr>
<tr>
<td>Arachidonic (C20:4)</td>
<td>92</td>
<td>129</td>
<td>135</td>
</tr>
<tr>
<td>Eicosapentaenoic “EPA” (C20:5)</td>
<td>11</td>
<td>15</td>
<td>12</td>
</tr>
<tr>
<td>Docosapentaenoic “DPA” (C22:5)</td>
<td>14</td>
<td>26</td>
<td>24</td>
</tr>
<tr>
<td>Docosahexaenoic “DHA” (C22:6)</td>
<td>12</td>
<td>17</td>
<td>19</td>
</tr>
<tr>
<td>Total Poly- Unsaturated F.As</td>
<td>745</td>
<td>869</td>
<td>884</td>
</tr>
<tr>
<td>Total Unsaturated F.As/ Total</td>
<td>0.96</td>
<td>1.15</td>
<td>1.23</td>
</tr>
<tr>
<td>Saturated F.As</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

4. DISCUSSION

From achieved results in table (1) it is showed that duck meat samples had the highest TBA mean values followed by chicken then turkey which indicate that oxidative rancidity is more in duck followed by chicken then turkey.

High significant differences were associated with the examined samples of poultry species (P<0.01) and poultry cuts.

Comparing obtained data of chicken thigh and breast samples, it showed that they were in accordance with results of both of Afifi-Jehan, (2000), lower than Hassanin-Fatin and Hassan, (2003) and higher than Fathy-Eman, (2013).

TBA test has widely used for measuring oxidative rancidity as it is very sensitive for...
evaluating products of unsaturated fatty acids (Melton, 1983).

The data recorded in table (2) declared that peroxide value recorded higher in duck than chicken and turkey indicate faster development of oxidative rancidity.

The obtained results disagreed with Saad et al., (2013) who recorded 0.09±0.01, 0.16±0.01 and 0.03±0.01 (mg/kg) for TBA and 0.12±0.01, 0.30±0.01 and 0.08±0.01 for PV for chicken breast, duck breast and pigeon, respectively.

High significant differences (P<0.01) were obtained as results of poultry species and cuts. Peroxides values used to determine the quality of fat and it is used as indicator for lipid oxidation (Masoud et al., 2008).

It is evident from results obtained in table (3) that free fatty acids values increased in duck compared with chicken and turkey samples indicating hydrolytic rancidity.

The differences associated with free fatty acid value in examined sample of poultry meat cuts were highly significant (P<0.01) as results species.

Free fatty acids test used as indicator for hydrolytic rancidity (Melton, 1983).

Hydrolytic rancidity is mainly due to the presence of poly unsaturated fatty acids (Brenner, 2002).

All breast samples have higher keeping quality than thigh samples.

It is evident from results in table (4) that palmitic acid (c16:0) as saturated fatty acid (SFAS) was the highest in duck followed by chicken then turkey meat samples.

On the other hand, turkey has the highest content of oleic acid (C18:1) and linoleic acid (C18:2) as poly unsaturated fatty acids (PUFAS) then chicken and finally duck samples.

The results obtained disagreed with those recorded by (Saad et al., 2013) who recorded the ratio between total unsaturated fatty acid and total saturated fatty acid 0.78, 0.89 and 0.74 for chicken breast, duck breast and whole pigeon, respectively.

The presence of unsaturated fatty acids (UFAS) in food particles increases its nutritive value and contrast, presence of high content of saturated fatty acids (SFAS) lower (decrease) its nutritive value and increase shelf life of the product (Pearson, 1984).

Finally according to the results of examined samples, it is obvious that duck meat has lower nutritive value besides high susceptibility to rancidity (especially thigh samples).

Turkey meat has the highest nutritive value due to high content of PUSFAs and the highest keeping quality as compared with chicken and duck meat samples.

5. REFERENCES


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