

CHANGES IN PROTEINS AND FATS OF OLD HENS MEAT AS  
INFLUENCED BY PRE-FREEZING TREATMENTS

BY

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INTRODUCTION

Deterioration in food products of animal source during storage at low temperatures occurs by the effect of bacterial activity and tissue enzymes which cause the undesirable changes in protein and fat leading to spoilage and loss of natural flavour, taste, colour and texture (Khan and Van Den Berg, 1965; Khlebnikov, 1973; Laurie, 1974 and George and Mountney, 1976).

Concerning the tough meat, for example the meat obtained from old hens of egg laying flocks (Srimpton, 1960), many treatments may be used to increase its tenderness, specially that frozen storage increase the toughness of such meat (Lawson *et al.*, 1958; Khan, 1964; Khan and Van Den Berg, 1967 and El-Samany and Shehata, 1977). Other treatments were applied to reduce the oxidation of lipids during frozen storage (Tomson, 1964; Arnold, 1970; Mitovcva and Oskaloza, 1971 and 1972 and hao *et al.*, 1970).

This work was conducted to study the effect of some pre-freezing treatments used for increasing tenderness and decreasing lipids and protein changes during frozen storage of meat obtained from old hens.

MATERIALS AND METHODS

Harvard chicken hens 2 - 3 years old (stewing) of same weight were obtained from General Poultry Company. These birds were slaughtered, bled for 5 minutes, scalded for 2 minutes at 55°C, plucked, eviscerated, rinsed with water, strained, then jointed to breast, thigh and wing base. Pre-freezing treatments included the following: a) Untreated samples (control); b) Breast and wing base breaded cuts with spiced fine crust, black pepper and salt;

c) Soaked samples for 4 hours in cold (4°C) 0.5% pyrophosphate solution or 1.0% NaCl solution; d) Glazed samples, by immersion three times (5 seconds each) in cold about (4°C) 0.2% ascorbic acid + 0.2% citric acid solution or in cold 0.1% propolis antioxidant + 0.1% citric acid solution, between successive immersions, the meat was held for 1 hour at 10°C; e) Unsoaked or soaked samples for 4 hours in cold (4°C) 0.5% pyrophosphate solution and cold stored for 3 days at 4°C before freezing.

Frozen storage was carried out in polyethylene bags at -10°C. Samples were taken for analysis after 0, 2, 4 and 6 months of storage.

Total volatile nitrogen (TVN) and trimethylamine nitrogen (TMAN) were determined using the methods described by Winton and Winton (1958). Amino nitrogen (AN) was determined by the formal titration method of Kolencov (1952). Total bacterial count (TC) was carried out according to Frasier and Foster (1950). Thiobarbituric acid value (TBA) was determined following the method of Pearson (1970). Fatty acid composition was carried out for selected samples of thigh after 6 months of frozen storage according to the method given by the A.C.A.C. (1970). Methyl esters were prepared by the method of Ganglitz and Lehman (1963) and Pye-Unicorn gas liquid chromatography apparatus was used.

RESULTS AND DISCUSSION

A- Protein degradation.

From tables (1, 2 and 3) it could be noticed that fresh as well as frozen stored samples showed highest TVN and TC for breast followed by leg and wing base; TMAN and AN were highest for the leg, followed by breast, being lowest for the wing base. This may indicate the effect of original properties of raw materials.

Total bacterial count (TC) decreased after 2 months storage then increased after 4 and 6 months. Sokolov (1965) reported that at -18°C, TC decreased progressively, while at -10°C, the decrease was noticed only at early periods of storage, followed by progressive increase in counts. Although after 2 months of storage TC decreased, survived bacteria were able to raise TVN, TMAN and AN which increased progressively during frozen storage with no reduction.

<sup>1</sup> Commercial name, produced from bee hives wax.

After 6 months of storage, highest TC, TVN, TMAN and AN were noticed for post-rigor freezing treatment (samples frozen after cold storage and the control sample) while were lowest for pre-rigor frozen pyrophosphate, citric + ascorbic and propolis samples.

Other samples (post-rigor pyrophosphate, NaCl and fine crust sample) showed intermediate values. In case of AN, fine crust coated sample had the lowest values, possibly because of probable reaction of amino nitrogen groups with some fine crust constituents, presumably the reactive groups in carbohydrates. The reason, however, may be simple dilution of meat with fine crust.

Post-rigor freezing accelerated the break-down of proteins, nevertheless the pre-treatment with pyrophosphate before cold storage followed by freezing reduced this change, possibly because of the antibacterial effect of pyrophosphate as mentioned by Alexeyev *et al.* (1958). Citric + ascorbic acid treatment reduced the break-down of protein, possibly because of low pH value and its influence on micro-organisms. With regard to propolis effect, El-Dashlouty (1978) found that such resin had antibacterial effect during cold storage of irradiated camel meat.

B- Lipide changes:

1) Thiobarbituric acid value (TBA).

From table (4) it could be observed that before and after frozen storage, TBA was highest for leg, intermediate for wing base, while was lowest for the breast.

Post-rigor freezing samples showed highest lipids oxidation during frozen storage, because of initial higher malonaldehyde content by the effect of cold storage before freezing. Control samples followed the post-rigor frozen meat. Due to the antioxidation effect of pyrophosphate (Alexeyev *et al.*, 1958), pyrophosphate post-rigor freezing samples had lower TBA values. Nevertheless, propolis samples had highest antioxidation effect in case of pre-rigor frozen meat, followed by pyrophosphate and citric + ascorbic acid treatments. Antioxidation effects of ascorbic and citric acids were reported by Sokolov (1965) and Arnold (1970). NaCl and fine crust treatments showed intermediate TBA values; being between pyrophosphate post-rigor freezing and citric + ascorbic acid treatments.

The fatty acid composition of treatments before, as mentioned by TBA value, proved to have antioxidation effect was studied during frozen storage of the leg muscle for 6 months. Results are given in table (5).

Fresh lipids of hens showed higher contents of oleic, palmitic and linoleic acids; 48.06, 22.69 and 20.36 mol.% respectively. The fatty acid composition in general was similar to that given by Koznetsova *et al.* (1971) working on chicken lipids. These authors found that total saturated and total unsaturated fatty acids of chicken lipids were 77.80 and 22.20 mol.% which was confirmed by the present data (Table, 5); being 77.99 and 22.01 mol.% respectively. This indicated that hen's lipids may be less stable for oxidation due to higher unsaturation of fatty acids. For beef lipids, the reverse was found, where unsaturated fatty acids (41.41 mol.%) were lower than total saturated fatty acids (58.59 mol.%) as given by El-Dashlouty (1975). Nevertheless, the biological value of chicken lipids are considered to be higher than for beef, due to higher contents of essential fatty acids which are unsaturated (linoleic, lauric and arachidonic). Moreover, assimilation of chicken fat was higher than the beef fat (Sokolov, 1965).

During frozen storage apparent increase of total saturated fatty acids was found. This increase was mainly due to the increase in C<sub>16</sub> and C<sub>18</sub> mainly. The contents of C<sub>12</sub> and C<sub>14</sub> also increased while C<sub>17</sub> mostly decreased.

After 6 months of storage, the total unsaturated fatty acids decreased due to lipids oxidation which was mainly due to reduction of C<sub>18:1</sub> and C<sub>18:2</sub>. Tetraenoic acid (C<sub>18:4</sub>) also decreased, while changes of C<sub>16:1</sub>, C<sub>17:1</sub> and C<sub>18:3</sub> were less pronounced showing decrease in some treatments and increase in others.

The rate of total increase in saturated fatty acids and the rate of decrease of total unsaturated fatty acids were considered as indicators for lipids oxidation, propolis treated sample showed lowest rate of rancidity (17.29, 6.72%), followed by that of pyrophosphate (20.26, 11.97%) and citric + ascorbic acids (29.12, 11.32%), while control sample showed the highest rate of rancidity (57.88, 22.19%). Such results were in accordance with data obtained for the TBA value (Tables, 4 and 5).

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Table (1) Total volatile acids (T.V.A. mg/100 gm) and histamine nitrogen (H.N. mg/100 gm) of different parts of bone during frozen storage as affected by pre-freezing treatments.

Treatments	After 2 months			After 4 months			After 6 months			
	Wing base	Breast	Leg	Wing base	Breast	Leg	Wing base	Breast	Leg	
Control	T.V.A. mg/100 g	62.53	110.04	102.03	83.25	150.12	165.42	107.55	187.13	130.50
	H.N. mg/100 g	24.61	44.54	51.07	44.75	52.13	65.35	65.39	76.25	81.36
Pyrophosphate 0.5%	T.V.A. mg/100 gm	43.39	62.42	60.57	64.56	128.63	102.05	80.68	149.75	109.95
	H.N. mg/100 gm	18.53	37.43	39.55	38.55	45.85	52.15	52.25	64.25	77.75
NaCl 1.0%	T.V.A. mg/100 gm	55.18	95.63	74.98	70.54	139.59	99.50	90.35	156.57	130.43
	H.N. mg/100 gm	21.63	44.13	46.57	41.23	48.32	50.13	57.23	65.85	80.13
Propolis 0.1% + citric acid 0.1%	T.V.A. mg/100 gm	40.13	71.35	62.25	58.42	118.89	70.75	70.10	125.83	92.13
	H.N. mg/100 gm	20.11	33.01	44.35	40.11	47.51	55.25	54.83	60.35	79.13
Citric acid 0.2% + ascorbic acid 0.2%	T.V.A. mg/100 gm	45.40	62.19	71.55	67.58	134.85	92.93	91.75	152.83	112.89
	H.N. mg/100 gm	19.87	38.78	42.82	39.55	46.78	54.20	53.35	67.53	78.83
Pine crust	T.V.A. mg/100 gm	50.16	90.61	-	65.54	134.50	-	85.33	152.95	-
	H.N. mg/100 gm	20.55	39.63	-	40.50	47.25	-	54.05	64.83	-
Cold storage + freezing	T.V.A. mg/100 gm	93.19	166.15	126.65	123.83	183.85	143.50	143.78	216.15	169.13
	H.N. mg/100 gm	59.65	66.24	83.65	81.50	92.53	96.53	95.35	113.39	121.95
Pyrophosphate 0.5% + cold storage + freezing	T.V.A. mg/100 gm	75.50	120.30	109.75	105.35	158.38	139.57	125.45	178.32	159.65
	H.N. mg/100 gm	20.42	40.83	45.55	31.54	52.82	54.18	43.54	64.46	75.23

Zero time

	Wing base	Breast	Leg
Before freezing			
Fresh	TVH 3.77	12.86	6.73
	THAN 2.81	2.92	3.79
3 days of cold storage	TVH 73.18	146.11	106.64
	THAN 39.66	48.25	64.02
0.5% Pyrophosphate + 3 days of cold storage	TVH 55.55	108.60	89.78
	THAN 23.40	26.24	35.61

Table (2) Antimicrobial (A.M.) of different parts of bone during frozen-storage as affected by different pre-freezing treatments (mg/100 gm).

Treatments	After 2 months			After 4 months			After 6 months		
	Wing base	Breast	Leg	Wing base	Breast	Leg	Wing base	Breast	Leg
Control	665.93	730.13	750.87	1101.90	1342.80	1495.10	1102.13	1420.30	1730.19
Pyrophosphate 0.5%	970.13	565.61	700.11	837.50	871.30	1030.50	917.30	1020.10	1170.60
NaCl 1.0%	594.12	687.53	774.25	856.40	893.61	1052.32	939.63	1025.30	1190.90
Propolis 0.1% + citric acid 0.1%	1300.15	675.52	710.25	847.30	885.54	1041.32	928.45	1010.35	1185.92
Citric acid 0.2% + ascorbic acid 0.2%	165.15	570.55	704.15	842.10	876.45	1035.38	922.85	1000.25	1183.75
Pine crust	423.20	532.50	-	680.11	733.36	-	760.96	813.87	-
Cold storage + freezing	785.23	845.85	875.35	1303.43	1544.13	1677.35	1395.15	1635.50	1820.25
Pyrophosphate 0.5% + cold storage + freezing	683.52	788.35	814.28	895.48	953.65	1095.85	975.36	1101.45	1485.25

Zero time

Before freezing	Wing base	Breast	Leg
Fresh	254.10	317.60	390.20
3 days of cold storage	684.00	739.70	742.90
0.5% pyrophosphate + 3 days of cold storage	591.40	689.40	754.11

Table (3) Total bacterial counts of different parts of bone during frozen storage as affected by different pre-freezing treatments (count  $\times 10^6/g$ ).

Treatments	After 2 months			After 4 months			After 6 months		
	Wing base	Breast	Leg	Wing base	Breast	Leg	Wing base	Breast	Leg
Control	0.6	15.0	1.0	1.0	15.0	1.4	2.5	22.0	3.3
0.5% pyrophosphate	0.4	10.0	0.65	0.5	10.5	1.1	1.6	14.0	2.5
NaCl 1.0%	0.5	12.0	0.8	0.8	14.0	1.1	1.9	15.0	2.9
Propolis 0.1% + citric acid 0.1%	0.48	7.0	0.75	0.7	9.0	0.95	1.7	11.0	2.2
Citric acid 0.2% + ascorbic acid 0.2%	0.35	5.5	0.6	0.45	12.0	0.72	1.4	10.5	1.8
Pine crust	0.8	15.0	-	0.9	15.0	-	2.3	22.0	-
Cold storage + freezing	0.9	1.1	1.9	1.1	19.0	2.5	2.9	23.0	3.9
Pyrophosphate 0.5% + cold storage + freezing	0.7	11.0	1.7	0.65	13.5	1.9	1.8	14.5	3.9

Zero time

Before freezing	Wing base	Breast	Leg
Fresh	2.0	60.0	4.0
3 days of cold storage	4.0	15.5	9.7
0.5% pyrophosphate + 3 days of cold storage	3.4	16.0	4.5

Table (4) Thiobarbituric acid value (T.B.A.) of different parts of bone during frozen-storage as affected by pre-freezing treatments (mg/100 gm).

Treatments	After 2 months			After 4 months			After 6 months		
	Wing base	Breast	Leg	Wing base	Breast	Leg	Wing base	Breast	Leg
Control	0.290	0.268	0.252	0.322	0.300	0.490	0.440	0.350	0.680
Pyrophosphate 0.5%	0.218	0.150	0.210	0.224	0.150	0.260	0.235	0.225	0.310
NaCl 1.0%	0.265	0.200	0.280	0.228	0.210	0.500	0.480	0.380	0.660
Propolis 0.1% + citric acid 0.1%	0.250	0.138	0.250	0.210	0.146	0.255	0.200	0.154	0.240
Citric acid 0.2% + ascorbic acid 0.2%	0.220	0.152	0.233	0.220	0.155	0.257	0.235	0.227	0.320
Pine crust	0.210	0.171	-	0.275	0.250	-	0.300	0.260	-
Cold storage + freezing	0.325	0.333	0.445	0.400	0.363	0.515	0.475	0.492	0.750
Pyrophosphate 0.5% + cold storage + freezing	0.212	0.182	0.210	0.229	0.218	0.265	0.450	0.350	0.600

Zero time

Before freezing	Wing base	Breast	Leg
Fresh	0.175	0.135	0.232
3 days of cold storage	0.262	0.233	0.425
0.5% pyrophosphate + 3 days of cold storage	0.218	0.163	0.290

Table (5) Fatty acid composition of fresh and frozen stored lipids for 6 months (mg/100 g).

Treatments	Fatty acids	Fatty acid composition (mg/100 g)			
		Fresh	Control	Pyrophosphate 0.5% + citric acid 0.1%	Citric acid 0.2% + ascorbic acid 0.2%
Fatty acids	C <sub>12</sub> (lauric)	-	0.24	0.19	-
	C <sub>14</sub> (myristic)	0.45	0.52	0.58	0.55
	C <sub>16:1</sub> (tetradecenoic)	0.20	Trace	0.14	0.13
	C <sub>16</sub> (palmitic)	22.69	21.35	24.47	27.93
	C <sub>18:1</sub> (palmitoleic)	3.40	2.40	4.32	3.45
	C <sub>17</sub> (heptadecenoic)	0.36	0.25	0.31	0.42
	C <sub>17:1</sub> (heptadecenoic)	Trace	Trace	0.11	0.09
	C <sub>18</sub> (stearic)	4.48	6.43	5.08	7.23
	C <sub>18:1</sub> (oleic)	48.06	43.08	44.82	43.45
	C <sub>18:2</sub> (linoleic)	20.36	9.53	17.78	16.11
	C <sub>19:1</sub> (nonoleic)	-	Trace	-	0.57
	Total saturated	mg/100	27.99	41.15	32.63
Fatty acid % increase		57.88	17.29	29.12	28.26
Total unsaturated	mg/100	72.01	55.81	67.37	63.86
Fatty acid % decrease		22.49	4.72	11.34	10.97