

EFFECT OF DIFFERENT STERILIZATION MODES UPON THE QUALITY CHARACTERISTICS OF LOW-FAT MEAT PÂTÉS

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Abstract: The present work studies the effect of different sterilization modes upon the quality characteristics of meat pâté with a nutritional profile improved by fat reduction and enrichment with inulin and lentils. The changes in the texture and color parameters, the emulsion stability and the oxidative changes in the lipid fraction of low fat pâtés obtained under five sterilization modes, i.e. 111°C for 24 min, 121°C for 24 min, 111°C for 70 min, 121°C for 70 min, and 116°C for 47 min were evaluated. The lowest emulsion stability was registered in the sample with the most intensive sterilization conditions and the highest temperature applied, viz. sample 4 (98.94%). It was found that there was a relationship between hardness and sterilization mode, as well as between the temperatures applied and the formation of primary and secondary lipid oxidation products. The increase in the hardness values was more pronounced in the samples obtained under the milder sterilization conditions, while a decrease in the hardness index was observed under the most severe sterilization conditions used for sample 4.

Keywords: inulin, lentil flour, texture, emulsion stability, thermal effect

INTRODUCTION

In food preservation through heat sterilization, the set temperature is not achieved at once and simultaneously in the entire volume of the canned food (Teixeira et al., 1969; Flambert et al., 1977). The degree of heat treatment applied is directly proportional to the antimicrobial effect (Nitsch & Vuković, 2002; 2003; 2006; FAO, 2010), but the quality characteristics of the end products are affected by the temperature applied, the treatment duration and the fat content. Heat treatment that is too intensive may result in protein denaturation, in undesirable changes in the product appearance, texture and taste, and in reduction of its water holding capacity (Penaramos and Xiong, 2002).

Meat pâtés are made from poultry, beef or pork, liver and fat tissue. Apart from being a source of saturated fatty acids and cholesterol, these components are the reason for the higher calorific value of the products in which they are included (Lorenzo et. al., 2014; Ozvural & Vural, 2008). The fat reduction in meat products is often accompanied by technological and commercial problems in their production. A main reason is the fact that fats determine to a considerable extent the texture and taste of the end product (García et al., 2006;

Choi et al., 2013). Therefore, the choice of components to be used for fat substitution that could compensate for the changes in the sensory parameters of low-fat products is of great importance. Prebiotic inulin is a functional additive that has a future in this respect (Roberfroid, 2010; Shoaib, 2016).

The aim of this study was to evaluate the sterilization mode effect on the texture and color parameters, emulsion stability and oxidative changes in the lipid fraction of meat pâtés having a lower fat content.

MATERIALS AND METHODS

Pâté making: The meat pâté production was based on the use of a modified recipe including the following main ingredients: deboned turkey meat: 30%, chicken liver: 10%, egg mélange: 18%, inulin: 12.5%, soft fat: 11.25%, cornstarch: 2%, lentil flour: 1.25%, cooking salt: 1.5%, sodium nitrite: 0.005%, polyphosphates: 0.2%, black pepper: 0.3%, nutmeg: 0.05%, coriander: 0.15%, and drinking water: 15%. The Orafiti®HPX inulin used was provided by ARTEMIS OOD, Sofia, representatives of Beneo-Orafiti Ltd., Belgium. The inulin was added during cutting in the form of a gel obtained through hydration in a 1:4 (w/v) ratio as described by Latoch *et al.* (2016). The filling mass prepared was heated to 70°C and filled manually into cans having a size of H=26.5 mm and D=99 mm. The pâté prepared in this way was sterilized in a laboratory autoclave at four different modes: 111°C for 24 min, 121°C for 24 min, 111°C for 70 min, 121 °C for 70 min, and 116°C for 47 min.

Texture Profile Analysis (TPA): The texture profile analysis (TPA) of the ready product was made using a TA-XT.Plus texture analyzer (Stable Micro Systems, Surrey, GB). The cylinder used in this study was 51.75 mm in height and 30.37 mm in diameter, at sample height of 40 mm (quantity: 30 g). The samples were compressed twice, with a 5-second interval between the two compression cycles, at a rate of 2 mm.s⁻¹. The Hardness [N], Adhesiveness [Nmm], Cohesiveness and Friability [N] values were obtained as described by (Bourne, 1978).

Color metrics: The color parameters were determined spectrophotometrically using a Minolta Chroma meter (model CR 410, Osaka, Japan) according to the CIELab system.

Emulsion stability: The method described by Ockerman (1985) and Zorba et al. (1993) was used for the evaluation of the meat emulsion stability.

Determination of oxidative changes: The peroxide quantity was determined using the method described by Hornero-Méndez et al. (2006), and the thiobarbituric acid reactive substances (TBARs) were determined according to the method described by Sørensen &

Jørgensen (1996). The lipid extraction from the samples for the analysis of the peroxide quantity was performed according to the method of Bligh & Dyer (1959).

Statistical analysis: The statistical processing of the data obtained was made using the Statgraphics 16 software. The experiments were conducted with threefold repetition, and the data in the tables and graphs are arithmetic means of the indicators measured. Statistically significant differences were found at probability less than 0.05.

RESULTS AND DISCUSSION

Fig. 1 shows the thermograms during sterilization of the pâté samples, and the lethal effect (L_0) of the different samples is shown in Table 1.

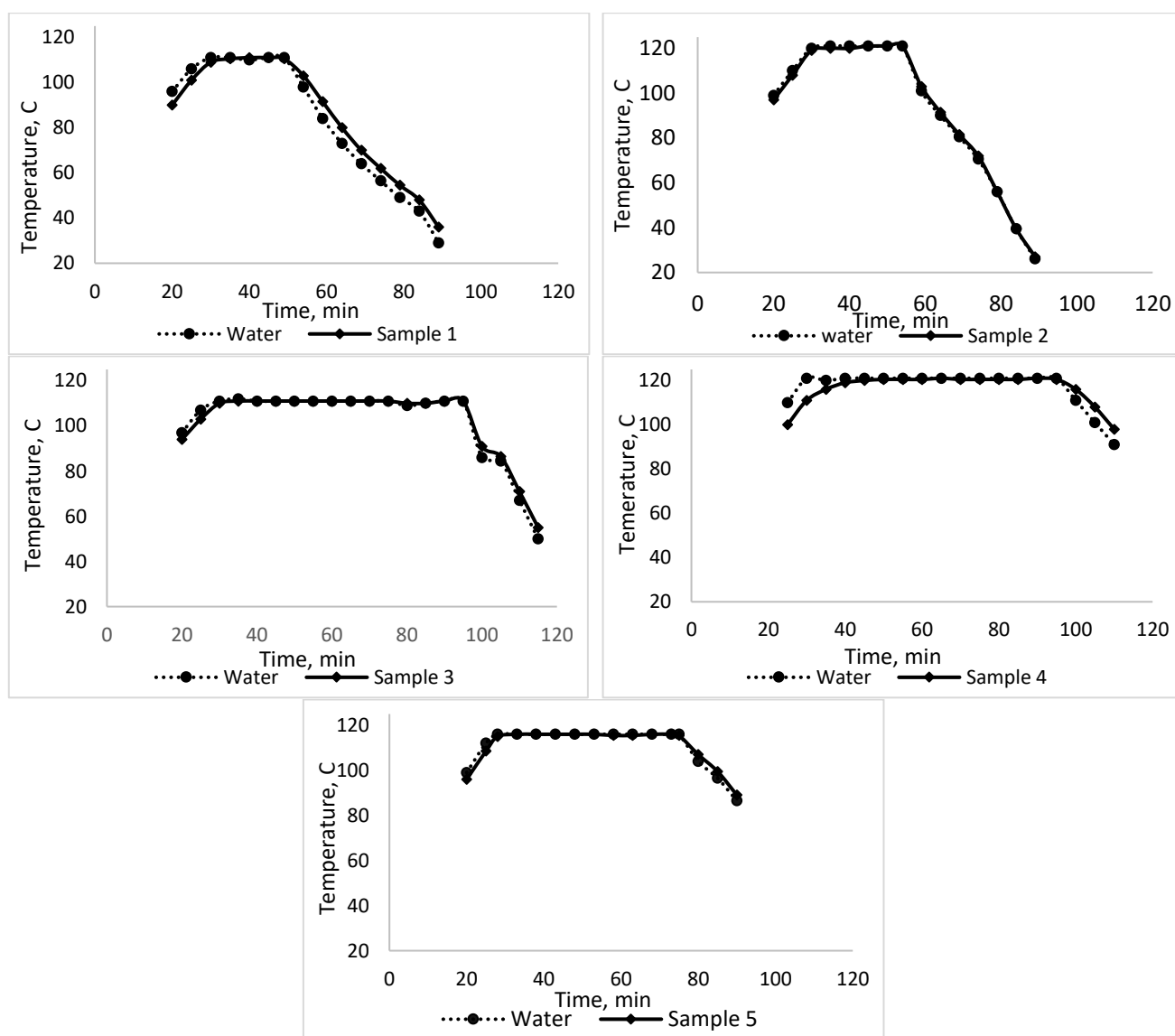


Figure 1. Heating curves of the pâté samples under different sterilization conditions

Table 1. Actual lethality of the meat pâtés sterilized under different conditions

Sample	1	2	3	4	5
L_{121}^{10}	2.57 min	24.79 min	6.52 min	55.51 min	15.74 min

Texture analysis

Texture is one of the most important qualitative parameters of food, and suitable instrumental methods of analysis have been developed for its assessment (Ivanov, G et al., 2018). The experimental data on hardness demonstrated that there were significant differences between individual samples ($P < 0.05$) (Table 2). The results indicated that the decrease in hardness was directly proportional to the increase in the actual sterilization effect (Table 1), the higher temperature and longer sterilization time leading to a product of lower hardness. These results were also in good correlation with the emulsion stability data obtained.

Table 2. Effect of sterilization conditions on the texture parameters, emulsion stability and lipolytic changes of the pâté samples

Sample Parameter	1	2	3	4	5
Hardness, N	62.02±3.78d	33.84±1.41bc	30.3±3.06b	23.91±1.96a	34.26±3.47c
Adhesiveness, Nmm	48.57±4.46b	27.53±7.8a	26.61±7.03a	19.82±3.36a	24.54±5.05a
Cohesiveness	0.25±0.03a	0.25±0.03a	0.27±0.04a	0.22±0.03a	0.25±0.06a
Friability, N	15.19±1.47c	8.56±0.71b	8.1±1.62b	5.36±0.95a	8.38±1.32b
Peroxide content, meqO ₂ .kg ⁻¹ fat	0.18±0.01a	0.22±0.01ab	0.4±0.00c	0.43±0.05c	0.23±0.02b
(TBARs), mg.kg ⁻¹ pate	1.27±0.02a	1.32±0.03b	1.33±0.02b	1.54±0.02d	1.48±0.02c
Emulsion stability, %	99.79±0.02d	99.97±0.02e	99.66±0.02c	98.94±0.02a	99.51±0.02b

a-d – the values within the same column bearing a common letter designation did not differ statistically ($P > 0.05$)

Lipolytic changes: The data on the formation of primary lipid oxidation products showed a considerable peroxide increase in the samples with the longest thermal processing, whereas sample 1 exhibited the lowest peroxide generation. A similar trend was observed in the secondary products of oxidation reactions in the lipid fraction of the samples expressed by the TBARs indicator. The lowest malondialdehyde quantities were accumulated in the sample subjected to the mildest sterilization conditions, i.e. 111°C for 24 min. Contrariwise, the largest MDA quantities were formed and accumulated in sample 4, where the sterilization process was longer and the temperature was higher (121°C). Similar results were obtained by other authors as well (Al-Abdullah & Al-Majali, 2011). These data are in conformity with Pither's report (2003) that more intensive heat treatment had a negative effect on the biological value of meat products.

Emulsion stability: According to Kim et al. (2001), the formation of a stable gel, hence a stable meat emulsion, is affected by several factors, temperature and heat treatment time being the most important ones. The lowest emulsion stability was found in the sample

with the strongest actual sterilization effect, whereas the highest values were reported for samples 2 and 1, where the retention time at the set temperature was the shortest (Table 2). The heat treatment degree together with the changes in the protein fraction probably altered the structure of the hydrocarbon chains in the lentils and inulin added, thus affecting their ability to stabilize the meat emulsion.

Color metrics: According to Jiménez-Colmenero et al. (2010), meat products having a lower fat content demonstrate higher brightness values, which may be due to the greater dispersion of light. On the other hand, Cáceres et al. (2004) reported that the inulin added to meat products imparted glossy features similarly to fats. The present study established that the increase in the sterilization effect resulted in a decrease in color brightness (Table 3). According to Jaeger et al. (2010), in foods subjected to heat treatment the darkening usually results from the Maillard reaction occurring between reducing sugars and amino acids. A further reason may be the different emulsifying and related water retention capacity of the samples under different sterilization conditions connected to the optical properties of food.

Table 3. Effect of the sterilization conditions on the color parameters and color stability of the pâté samples

Sample	(L*)	(a*)	(b*)	(C)	(h)	(ΔEab)
1	57.10+0.60c	4.99+0.29b	11.61+0.35a	12.64+0.42a	66.75+0.75b	1.36+0.006d
2	56.49+0.24b	4.31+0.1a	12.75+0.15b	13.46+0.13bc	71.33+0.54d	0.83+0.006b
3	56.23+0.24b	4.99+0.14b	11.9+0.2a	12.91+0.15ab	67.24+0.86bc	0.69+0.006a
4	52.50+0.43a	5.78+0.06c	12.64+0.94b	13.9+0.88c	65.36+1.36a	1.36+0.006d
5	56.31+0.36b	5.02+0.1b	12.61+0.19b	13.57+0.2c	68.3+0.36c	1.29+0.0065c

a-d – the values within the same column bearing a common letter designation did not differ statistically ($P > 0.05$)

Changes resulting from the sterilization effect achieved were also observed in the red and yellow color component values for individual samples. The red component value (a*) followed the trend of rising along with the increase in the sterilization time and temperature, which was somewhat contradictory to the reported oxidative changes in lipids and myoglobin pigments (Fuentes et al., 2010, Ganhão et al., 2010). This could be attributed to the different time and rate of the reactions involved in the formation of nitrosomyoglobin and its subsequent denaturation to nitroso-hemo-chromogen under the different sterilization conditions indicated by the different heating curves of the filling mass for the pâté samples (fig.1). The results obtained for b* demonstrated that the yellow component was less affected by the sterilization time than by the temperature applied.

The data on the color hue (h) showed statistically significant differences ($P < 0.05$) between individual samples, and this parameter was mainly affected by the sterilization time. The ΔE values measured at the beginning and after two hours of exposure of the opened can

samples to light and oxygen were indicative of the color stability over time. The values obtained varied from 0.69 to 1.36, the most stable sample being the one sterilized at 111 °C for 70 minutes ($L_{121}^{10} = 6,52$ min), and the most unstable ones being samples 4 (111 °C for 24 minutes) and 1 (121 °C for 70 minutes), which exhibited the highest and the lowest values of L_{121}^{10} .

Conclusion

The analysis of the results showed that the more intensive sterilization modes led to better texture parameters compared to the milder sterilization conditions. With regard to the heat treatment effect on the primary and secondary lipid peroxidation products, the longer treatment at higher temperatures was found to result in the accumulation of larger quantities of oxidated products. Shorter treatment at lower temperatures had the weakest effect on the color parameters and emulsion stability. Nevertheless, it is worth mentioning that treatment that is too short is undesirable from the point of view of the red coloring of the pâté and the preservation of the samples after opening.

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