# STUDIES REGARDING THE CHEMICAL COMPOSITION OF SOME POULTRY EDIBLE ORGANS

Paula Viorela Druc<sup>1\*</sup>, Carmen Irimia (Gavrilescu)<sup>1</sup>, C. Spridon<sup>1</sup>, M.G. Usturoi<sup>1</sup>

<sup>1</sup>Universiy of Agricultural Sciences and Veterinary Medicine from Iaşi, Romania

#### Abstract

Even if at world level are consumed important quantities of poultry organs, research regarding their quality are extremely rare.

The aim of the study was to establish if are some correlations between slaughtering age of poultry and chemical composition of two edible organs (heart and gizzard).

So were settled up 6 experimental batches (3 for gizzards and 3 for hearts), differencing by the age of slaughtered poultries, as follows:  $P_1$  and  $I_1$  = gizzards and hearts gathered from chickens slaughtered at 35 days; P<sub>2</sub> and I<sub>2</sub> = gizzards and hearts gathered from chickens slaughtered at 40 days;  $P_3$  and  $I_3$  = gizzards and hearts gathered from chickens slaughtered at 42 days. Determinations were made on fresh products and aimed content in water, dry matter, proteins and NES; also we calculate the energetic value.

Regarding protein content of gizzards and hearts, our data shown a decreasing of the values at the same time with the increasing of slaughtering age; the differences between those 3 batches being without statistical significance (P>0.05).

Regarding fat content, both for hearts and gizzards, the best values were founded at chickens slaughtered at 42 days, higher with 0.91-2.78% in case of hearts and with 3.08-7.20% for gizzards face to the situation for the others slaughtering. Statiscally speaking, the differences were very significant (P<0.01) for gizzards ( $P_1$  vs.  $P_3$ ) as well as for hearts ( $I_1$  vs.  $I_3$ ).

Through the obtained results for hybrid Ross 308, we could say that slaughtering age didn't influence very significant the chemical composition of gizzards and hearts, with the exception of fat content, which increase with increasing of slaughtering age.

Key words: poultry, slaughtering, gizzards, hearts, chemical composition

#### INTRODUCTION

Meat and meat products form an important segment in human diet, because those ones provide essential nutritive elements, which couldn't be obtained from vegetables and derived products [3].

Consumption of poultry meat and edible organs considerably increased in the last years, in Romania. Poultry gizzards and hearts are consumed on a large scale due to their low price, low content in fat and due to a short period of time for their preparation [2].

The importance of edible by products is underlined by their nutritive value which is suitable for consumers [6].

The quality of animal edible organs depend of various factors: breed, race,

rearing system, physiological and health state of animal, sex, age, weight, manipulation and post-mortem processing [4].

Being known the fact that intensive rearing of meat poultry has as result the obtaining of meat in a short period of time and with a maximum economical efficiency, the aim of the current study was to establish if there are correlations between slaughtering age of the birds and chemical composition of two poultry edible organs (heart and gizzard).

Till now determination of chemical components of gizzard and heart gathered from poultry slaughtered at different ages, wasn't a priority in the research regarding quality and human consumers' food safety.

## MATERIALS AND METHODS

The studied material was represented by gizzards and hearts gathered from hybrid Ross 308. Research were effectuated based

<sup>\*</sup>Corresponding author: paula.druc@yahoo.com The manuscript was received: 09.10.2017 Accepted for publication: 19.12.2017

on an experimental design, organized on 6 experimental batches (3 for gizzards and 3 for hearts), differentiating by slaughtering age of poultry broiler, encoded as follows: P1 and  $I_1$  = gizzards and hearts gathered from poultry slaughtered at 35 days;  $P_2$  and  $I_2$  = gizzards and hearts from poultry slaughtered at 40 days; P<sub>3</sub> and I<sub>3</sub> = gizzards and hearts from poultry gathered at 42 days.

For each studied batch were formed 5 samples, each of them with a weight of 0.5 kg, which were minced and homogenised and from which were gathered samples for determinations.

Determinations were realized on fresh products and aimed the content in water, dry proteins, fats, ash and nitrogenous extractive substances (NES), and also was calculated the energetic value.

Determination of dry matter content was realised by drying in oven [7], and water content resulted as a difference in according with the formula: Water (%) = 100% - DM (%).

Proteins from those two poultry edible organs were determined by Kjeldahl method

Establishing of lipids content was effectuated with Soxhlet method [12].

Ash content was determined in according with SR ISO 936:1998 [10], and nonnitrogenous extractive substances calculated as difference with the formula: NES (%) = 100 - (Water% + Ash% +Proteins% + Lipids%) [8].

Energetic value of poultry gizzard and heart (kcal/100g) was calculated using the formula: (4.27 kcal \* Proteins%) + (9.02 kcal \* Lipids%) + (3.87 kcal \* NES%) [9].

The obtained data were subjected to some calculations, statistical using ANOVA algorithm included in MsExcel.

### RESULTS AND DISCUSSION

# • Chemical composition of gizzards

The gathered samples from poultry gizzard show a decrease of water content at poultry slaughtered at advanced ages. So, samples gathered from poultry slaughtered at 35 days (batch P<sub>1</sub>) recorded the highest mean

value, respectively 79.76±0.04%, while the samples from poultry slaughtered at 40 days (batch P2) had a water content of 79.64±0.03%, and the ones gathered from poultry slaughtered at 42 days (batch P<sub>3</sub>) of 79.55±0.02%. The calculated values for variation coefficient were inside interval 0.06-0.10% fact which shown a very homogenous character. Statistically speaking enlightened significant statistical differences between batches P<sub>1</sub> vs. P<sub>2</sub> (P<0.05) and distinct significant differences between batches  $P_1$  vs.  $P_3$  (P<0.01) (tab. 1).

Normally, dry matter content recorded an increasing evolution, at the same time with increasing of slaughtering age. The calculated means were of 20.24±0.04% for batch P<sub>1</sub>, 20.36±0.03% for batch P2 and 20.42±0.04% for batch P<sub>3</sub>. The studied character presented a very good homogeneity, a proof being the values of variation coefficient of 0.23-0.41%. Statistically speaking were recorded significant differences between batches P<sub>1</sub> vs. P<sub>2</sub> (P<0.05) and distinct significant differences between batches  $P_1$  vs.  $P_3$  (P<0.01) (tab. 1).

Regarding protein content in gizzards, P<sub>1</sub> recorded the highest level (17.28±0.02%), followed in a descendant order by batch P2 (17.23±0.03%) and by batch  $P_3$  (17.20 $\pm$ 0.03%). The analysed parameter was homogenous, all calculated values for variation coefficient being under the level of 10% (VC% = 0.28-0.40%). The differences between batches were without statistical significance (P>0.05) (tab. 1).

Regarding fat content, could be observed that this one suffered a low increasing, in parallel with increasing of slaughtering age. So, batch P<sub>3</sub> recorded the highest value, 1.34±0.02%, being followed by batch P<sub>2</sub> with a value of 1.30±0.01% and respectively by batch P<sub>1</sub> with a level of only 1.25±0.02%. The studied character was very homogenous inside each batch, its values being between 1.48 and 3.65%. From statistical analysis observed significant differences between batches P<sub>1</sub> vs. P<sub>2</sub> (P<0.05) and distinct significant between batches P<sub>1</sub> vs. P<sub>3</sub> (tab. 1).

Table 1 Chemical composition of gizzards provided from hybrid Ross 308

		An	alysed batc	ANOVA					
Quality parameters	P <sub>1</sub> (slaughtered at 35 days)		P <sub>2</sub> (slaughtered at 40 days)				P <sub>3</sub> (slaughtered at 42 days)		
	$\overline{X}\pm s_{\overline{X}}$	VC%	$\overline{X}\pm s_{\overline{X}}$	VC%	$\overline{X} \pm s_{\overline{X}}$	VC%	Compared batches	P value	Signification
Water (%)	79.76±0.04	0.10	79.64±0.03	0.10	79.55±0.02	0.06	P <sub>1</sub> vs. P <sub>2</sub>	0.04487	* (P<0.05)
							P <sub>2</sub> vs. P <sub>3</sub>	0.05409	ns (P>0.05)
(/*/							P <sub>1</sub> vs. P <sub>3</sub>	0.00114	** (P<0.01)
Dry matter (%)	20.24±0.04	0.41	20.36±0.03	0.38	20.42±0.04	0.23	P <sub>1</sub> vs. P <sub>2</sub>	0.04487	* (P<0.05)
							P <sub>2</sub> vs. P <sub>3</sub>	0.05409	ns (P>0.05)
							P <sub>1</sub> vs. P <sub>3</sub>	0.00114	** (P<0.01)
Proteins (%)	17.28±0.02	0.28	17.23±0.03	0.40	17.20±0.03	0.40	P <sub>1</sub> vs. P <sub>2</sub>	0.25941	ns (P>0.05)
							P <sub>2</sub> vs. P <sub>3</sub>	0.35654	ns (P>0.05)
							P <sub>1</sub> vs. P <sub>3</sub>	0.08975	ns (P>0.05)
Fats (%)	1.25±0.02	3.65	1.30±0.01	1.48	1.34±0.02	2.36	P <sub>1</sub> vs. P <sub>2</sub>	0.04028	* (P<0.05)
							P <sub>2</sub> vs. P <sub>3</sub>	0.05081	ns (P>0.05)
							P <sub>1</sub> vs. P <sub>3</sub>	0.00593	** (P<0.01)
Ash (%)	0.95±0.01	2.35	0.99±0.01	1.94	1.03±0.02	2.83	P <sub>1</sub> vs. P <sub>2</sub>	0.01292	* (P<0.05)
							P <sub>2</sub> vs. P <sub>3</sub>	0.04103	* (P<0.05)
							P <sub>1</sub> vs. P <sub>3</sub>	0.00124	** (P<0.01)
NES (%)	0.76±0.01	1.98	0.83±0.02	2.99	0.87±0.02	4.09	P <sub>1</sub> vs. P <sub>2</sub>	0.00300	** (P<0.01)
							P <sub>2</sub> vs. P <sub>3</sub>	0.07366	ns (P>0.05)
							P <sub>1</sub> vs. P <sub>3</sub>	0.00061	***(P<0.001)

ANOVA within rows, between groups for different superscripts, one by one comparison: ns = not significant (P>0.05); significant = \* (P<0.05); distinguished significant = \*\* (P<0.01); highly significant = \*\*\* (P<0.001).

The effectuated analysis for identification of ash content in gizzard lead to obtaining of some values situated inside the interval 0.95±0.01% (gizzards gathered from hybrid Ross 308, slaughtered at 35 days) and 1.03±0.02% (gizzards gathered from hybrid Ross 308, slaughtered at 42 days). Even if the studied character was very homogenous inside those three batches (VC% = 1.94-2.83%), between them were recorded differences with statistical significance (P1 vs. P<sub>2</sub>, P<0.05; P<sub>1</sub> vs. P<sub>3</sub>, P<0.01; P<sub>1</sub> vs. P<sub>3</sub> P<0.01) (tab. 1).

Gizzard content in non-nitrogenous extractive substances (NES) varied between a minimum of 0.76±0.01% (batch P<sub>1</sub>) and a maximum of 0.87±0.02% (batch P<sub>3</sub>). The values for variation coefficients were lower (VC% = 1.98-4.09%), which prove that this

character had a very good homogeneity. Testing of statistical significance show distinct significant differences between batches P<sub>1</sub> vs. P<sub>2</sub> (P<0.01) and very significant between batches P<sub>1</sub> vs. P<sub>3</sub> (P<0.001) (tab. 1).

Calculus of energetic value presented the lowest value of only 87.97 kcal/100g for gizzards gathered from poultry slaughtered at 35 days (batch P<sub>1</sub>), followed by gizzards of poultry slaughtered at 40 days (batch P<sub>2</sub>) with 88.54 kcal/100g and the ones of poultry slaughtered at 42 days (batch P<sub>3</sub>) with 82.92 kcal/100g (fig. 1).

The increasing of gizzards' energetic value with the increasing of slaughtering age could be explain by higher fat content existent in gizzards gathered from poultry with an advanced slaughtering age.

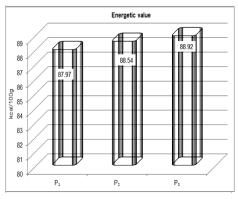


Fig. 1 Energetic value of gizzards gathered from hybrid Ross 308

## • Chemical composition of hearts

The obtained results regarding water content of poultry hearts gathered from hybrid Ross 308 slaughtered at different ages show an increasing of this parameter at the same time with increasing of slaughtering age. So, the hearts gathered from poultry slaughtered at 35 days (batch I<sub>1</sub>), had a water content of 78.12±0.06%, the ones gathered from poultry slaughtered at 40 days (batch I<sub>2</sub>) of 77.99±0.06%, and the ones from poultry slaughtered at 42 days (batch I<sub>3</sub>) of 77.94±0.04%. The studied character was very homogenous inside batches, the values of variation coefficient being inside interval 0.12-0.18%. Statistically speaking only in the case of batches I1 vs. I3 were recorded significant differences (P<0.05) (tab. 2).

Table 2 Chemical composition of hearts gathered from hybrid Ross 308

	Analysed batches (n=5)								
Quality parameters	I₁ (slaughtered at 35 days)		I <sub>2</sub> (slaughtered at 40 days)		I <sub>3</sub> (slaughtered at 42 days)		ANOVA		
	$\overline{X}\pm s_{\overline{X}}$	VC%	$\overline{X}\pm s_{\overline{X}}$	VC%	$\overline{X} \pm s_{\overline{X}}$	VC%	Compared batches	P value	Signification
Water (%)	78.12±0.06	0.16	77.99±0.06	0.18	77.94±0.04	0.12	$I_1$ vs. $I_2$	0.15489	ns (P>0.05)
							l <sub>2</sub> vs. l <sub>3</sub>	0.55430	ns (P>0.05)
							l <sub>1</sub> vs. l <sub>3</sub>	0.03564	* (P<0.05)
Dry matter (%)	21.88±0.06	0.57	22.01±0.06	0.62	22.06±0.04	0.43	l <sub>1</sub> vs. l <sub>2</sub>	0.15489	ns (P>0.05)
							$l_2$ vs. $l_3$	0.55430	ns (P>0.05)
							l <sub>1</sub> vs. l <sub>3</sub>	0.03564	* (P<0.05)
Proteins (%)	13.79±0.03	0.46	13.74±0.02	0.28	13.71±0.02	0.32	l <sub>1</sub> vs. l <sub>2</sub>	0.12979	ns (P>0.05)
							l <sub>2</sub> vs. l <sub>3</sub>	0.32766	ns (P>0.05)
							l <sub>1</sub> vs. l <sub>3</sub>	0.42073	ns (P>0.05)
Fats (%)	4.32±0.03	1.40	4.40±0.02	1.01	4.44±0.01	0.75	l <sub>1</sub> vs. l <sub>2</sub>	0.04889	* (P<0.05)
							l <sub>2</sub> vs. l <sub>3</sub>	0.18561	ns (P>0.05)
(70)							l <sub>1</sub> vs. l <sub>3</sub>	0.00623	** (P<0.01)
Ash (%)	1,03±0,01	3,07	1.08±0.02	4.69	1.10±0.03	5.85	l <sub>1</sub> vs. l <sub>2</sub>	0.07829	ns (P>0.05)
							l <sub>2</sub> vs. l <sub>3</sub>	0.71209	ns (P>0.05)
							l <sub>1</sub> vs. l <sub>3</sub>	0.06631	ns (P>0.05)
NES (%)	2.73±0.06	4.92	2.79±0.03	2.49	2.81±0.05	3.67	l <sub>1</sub> vs. l <sub>2</sub>	0.44795	ns (P>0.05)
							l <sub>2</sub> vs. l <sub>3</sub>	0.36945	ns (P>0.05)
							I <sub>1</sub> vs. I <sub>3</sub>	0.75400	ns (P>0.05)

ANOVA within rows, between groups for different superscripts, one by one comparison: ns= not significant (P>0.05); significant =\* (P<0.05); distinguished significant =\*\* (P<0.01); highly significant =\*\*\* (P<0.001).

Proportionally dry matter content had an inversely variation being higher at hearts gathered from poultry slaughtered at 42 days (batch I<sub>3</sub>) (22.06±0.04%) and lower at the ones gathered from poultry slaughtered at 35 days (batch  $I_1$ ) (21.88±0.06%). Also were recorded significant differences only at the level of batches I<sub>1</sub> vs. I<sub>3</sub> (P<0.05), in conditions of a very good homogeneity of the studied characteristic (VC% = 0.43-0.62) (tab. 2).

Protein content presented values situated in interval 13.79% (batch I<sub>1</sub>) and 13.71% (batch I<sub>3</sub>), but without statistical significance between batches. The analysed parameter presented a profound homogeneity character, all the calculated values for variation coefficient being under the level of 10% (tab. 2).

Fat content varied very significant (P<0.01) with the increasing of slaughtering age from 4.32±0.03% (batch I<sub>1</sub>) to 4.44±0.01% (batch I<sub>3</sub>), while the hearts gathered from batch I2 presented a value of 4.40±0.02%. The values for variation coefficient were inside interval 0.75-1.40%, fact which shown a very good homogeneity of the character inside each analysed batch (tab. 2).

Regarding ash content of poultry hearts, this one presented a minimum of 1.03±0.01% at batch I<sub>1</sub> and a maximum of 1.10±0.03% at batch I2. The calculated variation coefficients for this parameter described the homogeneity of batches inside interval 3.07-5.85%. Analysing statistically the obtained data weren't recorded differences with statistical signification at the level of those 3 batches (P>0.05) (tab. 2).

After calculating the content in nonnitrogenous extractive substances (NES) of poultry hearts, batch I1 recorded the lowest mean value of 2.73±0.06%, being followed by batch I<sub>2</sub> (2.79±0.03%) and by batch I<sub>3</sub>  $(2.81\pm0.05\%)$ . The values for variation coefficient (VC% = 2.49%-4.92%) show a very good homogeneity of the character inside experimental batches. The results of the statistical analysis realised between those 3 experimental batches shown differences but without statistical significance (tab. 2).

Increasing of lipid content, at the same time with increasing of slaughtering age leads also to increasing of energetic value (fig. 2).

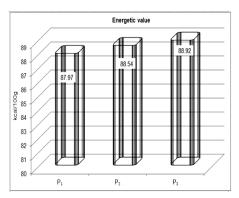


Fig. 2 Energetic value of hearts gathered from hybrid Ross 308

So, the minimum value of 108.47 kcal/100g was recorded at hearts gathered from poultry slaughtered at 35 days (batch I<sub>1</sub>), and the maximum one of 109.45 at the ones gathered from poultry slaughtered at 42 days (batch I<sub>3</sub>).

Even if the obtained results regarding chemical composition of poultry gizzard and heart are strictly relevant for the studied hybrid type, those values are close to the values obtained by other authors on other types of hybrids at different slaughtering ages (tab. 3).

Table 3 Comparison of chemical content with other published values

	Current study	Literature						
	42 days	Jokanović et al. (2014) [5]	Seong et al. (2015) [6]	Abdullah et al. (2016) [1]				
Water								
Gizzard	79.55	81.50	79.94	78.60				
Heart	77.94	73.10	77.36	74.82				
Prot	Protein							
Gizzard	17.20	13.60	17.26	17.34				
Heart	13.71	11.30	13.83	13.77				
Fat								
Gizzard	1.34	1.50	0.81	0.76				
Heart	4.44	12.50	4.53	6.97				
Ash								
Gizzard	1.03	0.90	-	0.97				
Heart	1.10	0.90	-	0.98				

Some differences could be explained by genetic variations or by nutrition of the birds.

#### CONCLUSIONS

The global analysis chemical composition characteristic to those analysed poultry edible organs (hearts and gizzards) show a high content in nutritive substances and a high energetic value.

The second important conclusion was that at the same time with the increasing of slaughtering age also increase the content in dry matter from poultry edible organs, in parallel with decreasing of water content.

#### REFERENCES

- [1] Abdullah F.A.A., Buchtova H., 2016: Comparison of qualitative and quantitative properties of the wings, necks and offal of chicken broilers from organic and conventional production systems. Veterinarni Medicina, nr. 61, p. 643-651. [2] Alvarez-Astorga M., Capita R., Alonso-Calleja C., Moreno B. and Garcia-Fernandez M. del Camino, 2002: Microbiological quality of retail chicken by-products in Spain, Meat Sci., nr. 62, p.
- [3] Byers T., Nestle M., Mctiernan A., Doyle C., Currie-Williams A., Gansler T., Thun M., 2002: American Cancer Society Guidelines on Nutrition and Physical Activity for Cancer Prevention: Reducing the Risk of Cancer with Healthy Food Choices and Physical Activity. CA Cancer J. Clin., nr. 52, p. 92–119.
- [4] Florek M., Litwinczuk Z., Skałecki P., edzierska-Matysek K.M., Grodzicki T., 2012 Chemical Composition and Inherent Properties of Offal from Calves Maintained Under Two Production Systems. Meat Sci., nr. 90, p. 402-409. [5] Jokanović M.R., Tomović V.M., Jović M.T., Škaljac S.B., Šojić B.V., Ikonić P.M., Tasić T.A., 2014: Proximate and Mineral Composition of Chicken Giblets from Vojvodina (Northern Serbia). International Scholarly and Scientific Research & Innovation nr. 8, p. 982-985.
- [6] Seong P.N., Cho S.H., Park K.M, Kang G.H, Park B.Y., Moon S.S., and Ba H.V., 2015:: Characterization of Chicken By-products by Mean of Proximate and Nutritional Compositions Korean J. Food Sci. An., vol. 35, nr. 2, p. 179-188. [7] \*\*\*AOAC, 2005: Official methods of analysis of the Association of Official Analytical Chemists, 18th Ed. Arlington, VA, USA, p. 26-25.
- [8] \*\*\*FAO, 2003a: Calculation of the energy content of foods - energy conversion factors (chapter 3). In: FAO (ed.) FAO Food Nutrition Paper, Food energy - methods of analysis and conversion factors. Food and Agriculture Organisation of the United Nations, Rome, Italy.

- [9] \*\*\*FAO, 2003b: Methods of Food Analysis (chapter 2). Food and Agriculture Organisation of the United Nations, Rome, Italy.
- [10] \*\*\*SR ISO 936:1998. Meat and meat products. Determination of total ash.
- [11] \*\*\*SR ISO 937:2007. Meat and meat products. Determination of nitrogen content.
- [12] \*\*\*SR ISO 1443:2008. Meat and meat products. Determination of total fat content.