Study on the Profile of Fatty Acids of Broiler Chicken Raised and Slaughtered in Industrial System

MARIUS MIHAI CIOBANU¹, PAUL CORNELIU BOISTEANU¹*, DANIEL SIMEANU¹*, ALINA NARCISA POSTOLACHE², ROXANA LAZAR¹, CATALIN RAZVAN VINTU³

¹ University of Agricultural Sciences and Veterinary Medicine of Iasi, 3 Mihail Sadoveanu Alley, 700490 Iasi, Romania

² Cattle Breeding Research Station from Dancu, Iasi - 9 Ungheni Road, 707252 Iasi, Romania

³ University of Agricultural Sciences and Veterinary Medicine of Bucharest, 59 Marasti Avenue, 011464 Bucharest, Romania

Fatty acid profile and the related nutritional indices of the breast, thigh and drumstick muscles were studied at three farms, suppliers of ROSS 308 line of broilers, slaughtered at the age of 42 days. The proximate chemical composition of the commercial slaughter cuts revealed contents between 16.26-22.78% for proteins and 1.80-7.45% for total lipids, the breast having the highest protein and ash content and lowest values for fat and moisture. The obtained values were mainly affected by region (P < 0.001). Meat fatty acid profile was affected (P < 0.001) by commercial slaughter regions (CSR) and by the interactions between CSR and supplier farms (Farm A, B, and C) at different levels. The obvious findings highlighted that Farm B supplied broilers with a delivered higher content of beneficial fatty acids (LA, LNA, AA, EPA, and DHA) in breasts and drumstick, while for thigh, Farm C had the best results. The content of total saturated fatty acids (SFAs), monounsaturated fatty acids (MUFAs) and polyunsaturated fatty acids (MUFAs) had the highest level in the thigh (P < 0.001).

Keywords: broiler meat, fatty acids, lipids quality

an ever evolving and industrialized world, consumer health [1] and quality of ending meat [2, 3] and meat products [4, 5] are the two highlighted issues debated from technological point of view in all stages of food chain, from farm to table, food production being in a continuous transition [6]. With vertically integrated raising, slaughtering and processing technologies, the poultry meat sector is the most widespread from the processing and consumer point of view [7, 8], while the bird's welfare in the industrial exploitation is a commune active management practice for ensuring a positive image among the final consumers [9]. Broiler production and processing at the mass level has already been accomplished worldwide, which is why the all-time focus is relying on enhancing the quality of meat by adjusting the different meat characteristics through diet [10-18] and by keeping them at a consistent rate for the final customer at all times, especially from nutritional point of view [19].

This tendency in industrial practices methods seeks to build an equilibrium [20], since it is acknowledged that the management of poultry meat production, in hardcore, primarily reflects the consumption characteristics, such as juiciness, tenderness or meat flavor [21, 22].

The current research is a study case based on the need to evaluate the efficiency of a constant nutritional management performance of 3 supplier farms within the RTC Holding Company Ltd., which supplies Ross 308 broilers for industrial slaughter, the main issue involved being the evaluation of the quality of meat lipids in relationship with supplier and commercial slaughter regions for maintaining a constant quality of fresh meat delivered in retail market and for further processing sector.

Experimental part

Material and methods

Animals, slaughtering and meat samples

The Bioethical Committee of the Ion Ionescu de la Brad University of Applied Life Sciences and Environment, Iasi

Romania endorsed the current experiment in cooperation with Room Trading Company Ltd. The assumption of this research has passed from the ongoing need to deliver broiler chicken meat of steady quality in all aspects, including dietary. 150 chicken broilers (ROSS 308) were considered for meat quality evaluation from 3 supplier farms within the company group (50 birds/every experimental group (L1 = Farm A, L2 = Farm B, L3 = Farm C) stochastically divided in a completely randomized design at the time of slaughter, 42 days old, Table 1 describing both, the entire population of birds and the selected biological material. Pre-harvest handling, transportation (between 0.25-2.75 hours), and slaughtering procedures (stunning, decapitation and bleeding) were in accordance with the implemented good animal welfare practices approved by the E.U. rules [9] after a fasting period of 12 hours (overnight) and a resting time before slaughter of 30-90 minutes. All the chickens were weighed before being slaughtered and eviscerated, the resulted carcasses were cooled and maintained at 4°C for 24 h *postmortem* and after, the breasts, thighs and drumsticks were separated, and frosted at -18°C for use in chemical analysis (brute chemical composition and fatty acid profile).

All the farms from the experimental design applied the same industrial technology of raising: population with 1 day chicken broilers on the same day, biological material purchased from the same commercial hatchery; accommodation in a deep litter (chopped straw with wood shavings in a ratio of 60:40) in climate controlled facilities with a photoperiod of 23 hours of light/day 'till 7 days old and 18 hours light/day starting on the 8th day of life, the environment temperature ranged between 20–25°C and a stock density of 33 kg/m² in accordance with welfare E.U. Directive [9].

All birds were fed a starter, grower and finisher diet, with ad libitum access to water (using a nipple water system) and corn-wheat-soy diet (Table 2), formulated to meet the nutrient requirements for finishing. To improve feed

^{*} email: paulb@uaiasi.ro, dsimeanu@uaiasi.ro

Items	L1 (Farm A)	L1 (Farm B)	L1 (Farm C)
Broilers population, (N)	98560	105600	112640
Broilers at slaughterhouse reception, (N)	98353	105435	112552
Mortality during transport, (%)	0.21±0.01	0.16±0.01	0.08±0.01
Experimental groups			
Live weight at slaughter, (g)	2539.87±50.22	2695.01±40.10	2613.64±3.25
Hot carcass weight, (g)	1804.59±37.78	1969.02±29.58	2049.22±28.74
Hot carcass dressing percentage, (%)	71.05±0.49	73.21±1.10	78.40±1.09
Cold carcass dressing percentage, (%)	70.07±0.69	72.19±0.97	77.09±1.00
Fracture incidence, (%)	2.00	4.00	6.00
Breast with bleeding spots, (%)	0.66	3.33	2.67
Wings with bleeding spots, (%)	1.33	4.00	10.66

 Table 1

 DESCRIPTION OF THE BIOLOGICAL

 MATERIAL

Data: mean±SEM; N=birds number raised on each supplier farm and loaded for transport;

conversion, carcass yield and breast meat yield was added a supplement of Lysine. The feed supplier performed values of descriptive parameters for diet proximate chemical composition.

 Table 2

 DIET COMPOSITION AND ANALYZED PROXIMATE CHEMICAL

 COMPOSITION

Items (%)	Starter	Grower	Finisher
Com	35.92	36.81	38.12
Wheat	26	26	26
Soybean meal	25	21.3	17.6
Sunflower meal	5.8	8.76	10.1
Sunflower oil	3.9	4.2	5.2
Monocalcium carbonate	0.9	0.5	0.55
Dicalcium phosphate	0.8	0.8	0.8
Vitamin-mineral premix	1	1	1.25
Salt	0.3	0.25	0
Lysine	0.28	0.28	0.28
Dl-Methionine	0.1	0.1	0.1
Analyzed chemical composition	,(%)		
Moisture	10.92	10.73	11.16
Metabolizable Energy, (MJ/kg)	2932	2932	2932
Crude protein	23.13	18.7	20.04
Fat	3.96	5.63	6.44
Crude fiber	2.77	3.39	2.92
Ash	6.07	4.41	4.47
NaCl	0.61	0.61	-

Proximate and fatty acid analysis

Prior to the beginning of the assessment, samples maintained for 2 months in freezing conditions were thawed overnight, deboned, minced and homogenized. Meat nutrient content was determined according to AOAC (2000), by measuring dry matter, ash, fat (using ether extraction Soxthlet method) and protein (using Kjeldahl method) [23].

The concentration of individual fatty acids was determined in two extracts from all samples by gas-liquid chromatography (GLC). Nonadecanoic acid ($C_{19:0}$; 3–5 mg) was added to the sample (1.5–2.5 g) as internal standard (IS). The extraction of lipids from meat samples was performed with a mixture of chloroform and methanol (2:1 v/v), as described by Folch [24]. Next, lipid extracts were converted to fatty acid methyl esters (FAME) through a consecutive trans-esterification with methylene chloride [25-27].

Separation and quantification of the fatty acid methyl esters (FAME) was performed using CarloErba 5300 mega series gas chromatograph (GS) equipped with a flame ionization detector (FID) suited for a fused-silica Omegawax 320 capillary column type SP-2380 (60 x 0.25 mm internal diameter x 0.20 μ m film thickness, Supelco Inc., Bellafonte, PA). The chromatographic operating conditions were as follows: initial column oven temperature 160°C (programmed to increase at a rate speed of 1°C/min. and from 180°C to 260°C at a rate speed

of 5°C/min.) and then maintaining it at 260°C for 5 minutes, the total running time being 45 min. The carrier gas was helium at a flow rate of 1.2 mL/min. and the splitting ratio, 1:20. The peaks were identified by comparison with the retention times of the standard fatty acids methyl esters used -C19:0 (Supelco, 37 components FAME mix).

Indices and sums calculations

The following Eqs. were used to calculate saturated fatty acids (SFAs), monounsaturated fatty acids (MUFAs) and polyunsaturated fatty acids (PUFAs):

$$\begin{split} SFA &= C8:0 + C10:0 + C12:0 + C14:0 + C15:0 + C16:0 \\ &+ C17:0 + C18:0 + C20:0; \end{split}$$

MUFA = C14:1 + C16:1 + C17:1 + C18:1 n-9 + C18:1 n-7 + C20:1 n-9;

 $\begin{array}{l} PUFA = C18:2n{-}6cis + C18:3n{-}6 + C18:2n{-}6cis + \\ C18:3n{-}3 + C20:2 + C20:3 n{-}3 + C20:4 n{-}6 + C20:5 n{-}3 + \\ C22:5 n{-}6 + C22:5 n{-}3 + C22:6 n{-}3; \end{array}$

The algorithm of the lipids indices was based on the fatty acid composition of the intramuscular lipids extracted here, being calculated Index of Atherogenicity (IA), Index of Thrombogenicity (IT) [28, 29] and Hypocholesterolemic/ Hypercholesterolemic ratio [30] by using the next Eqs.:

IA =	(4 x C14:0 +	C16:0)/[MUFA	$+\Sigma(n-6) +$	$\Sigma(n-3)];$
$\mathbf{I}\mathbf{T} =$	(C14:0 + C16)	(0 + C18.0)/[0]	5 x MUFA +	- 0.5 n (n-

6) + 3 x (n-3) + (n-3)/(n-6)];

 $h/H = (\Sigma MUFA + \Sigma PUFA)/(C14:0 + C16:0);$

Statistical analysis

Data are presented as mean \pm SEM. All statistical analysis was performed using the software package SPSS v.20 (SPSS Inc., Chicago, IL). Fatty acid profile and meat chemical composition data were analyzed using a general linear model (GLM) with CSR (commercial slaughter region: breast, thigh, drumstick), supplier (L1 = Farm A, L2 = Farm B, L3 = Farm C) and their interaction as fixed effects. Carcass weight and muscle fat content was included as corrected covariates because of variations in fatness rate. Principal component analysis (PCA) was used to explore and understand the variability of bird's meat composition by studying the correlation among the various fatty acids indices and summarizing them in meaningful components (PCs).

Results and discussions

Processing and its impacts on poultry's dietary meat have become more worrying in recent years. Overall, literature describes the low impact of primary and further processing on the dietary significance of chicken meat, with the exception of wet chilling, where exposure can immediately influence water-soluble nutrients, but without significant impact on proteins or lipids [20].

	S			V.	v	v	v								F
	IAL CARCASS	20)	Ц3	77.290.46	16.560.49	2.9031	$1.01_{0.02}$								
	H COMMERCI	mstick (n = '	L2	76.310.40	16.260.45	$4.10_{0.49}$	0.970.04								TVEN AGENT
	ELATION WIT	Dru	L1	76.00 _{0.40}	16.930.60	3.690.41	1.020.05								
~	AE DIET, IN RI ER FARMS		L3	72.590.49	$16.92_{0.42}$	7.450.52	$1.00_{0.03}$		L3=Farm C);					ıble 4	
Table	AT, FED THE SAN AND SUPPLIE	high (n = 50)	L2	73.170.55	16.520.46	6.880.22	0.970.04		A, L2=Farm B; I					I	
	R CHICKEN MF	F	ГI	73.670.60	16.710.55	5.940.27	0.970.04		r's (L1=Farm /						TTT ITT ITT ITT
	A) OF BROILE		L3	73.50033	22.630.20	$1.80_{0.17}$	$1.11_{0.05}$		olier of broile						M TEDMC OF
	tion (mean _{se}	reast $(n = 50)$	L2	74.150.39	21.160.22	2.030.19	$1.16_{0.07}$]; NS=P>0.0 <u>'</u>	gion; S=Supj						
	al composit	B	ΓI	72.290.29	22.780.44	1.860_{07}	$1.12_{0.04}$	M [meansrm]	al slaughter re						
	CHEMIC	Traits		Moisture (%)	Protein (%)	Fat (%)	Ash (%)	Data: mean ± SE	CSR =commercia						
Resul chemica shown ir betweer lipids. Th (P<0.00 the high and mo influenc observed	ts from the l compose n Table 3, r n 16.26-22 ne comme 1) overall est proteir isture. W ed the m 1 that the i	e an itior evea .78% ercia che i an /hile nusc nter	aly als, fo als, fo als emi d a emi d a c emi c emi c emi c emi c emi c emi c emi c emi c emi c emi c e emi c fo fo fo fo fo fo fo fo fo fo fo fo fo	zeo br for pr lau cal sh ne pr ior	d m oile th pro gh co m ote s b	lea ens e n tei ter om nte ana ein oet	t sa co naj ns re po ent age cc we	am mi or an gio siti an em ont en	iple ner con id 1 ion, id lo ien ent sup	s on rcial npor .80- ad a the owes t of (P- oplie	the slau nen 7.45 a ma bro st va the < 0.0	e pro 1gh 1s, c 5% f ajor east alue e su 05), nd 1	oxin ter c cont for t imp ha es fo uppl , it resu	nate cuts ent tota pac ving or fa lier: wa:	e , si i t git s sd

CSR. X

ń

SR

Main effects

AUGHTER REGIONS

CALCULATED LIPIDS INDICES IN TERMS OF THE HEALTH NEEDS FOR CONSUMERS OF BROILER CHICKEN MEAT, FED THE SAME DIET, IN RELATION WITH COMMERCIAL CARCASS SLAUGHTER REGIONS AND SUPPLIER FARMS Table 4

		κS.	Ξ	8	01	5	₽	10	90									
	ects	CSR	0.0	0.0	×.0.∖	0.0	0.0)'0 ×	0.0									
	lain eff	Ś	NS.	0.001	0.002	0.032	0.002	< 0.001	0.016*									
	Μ	CSR	< 0.001	< 0.001	< 0.001	< 0.001	< 0.001	< 0.001 ×	< 0.001									
	()	L3	886.2361.85	191.8710.51	4.690.32	$1.13_{0.01}$	0.330.007	0.570.02	3.160.03									
	mstick (n = 5)	L^2	1228.91100.67	147.3511.79	8.350.22	$1.13_{0.03}$	0.340.006	0.69001	3.160.04		; L3 = Farm C)							
	Dru	ГI	1100.2584.27	140.08129	7.95 _{0.19}	$1.13_{0.01}$	0.340.006	0.680.01	3.160.05		L2 = Farm B;							
		L3	2280.88151.22	201.8614.57	11.37 ₀ .22	$1.17_{0.01}$	$0.31_{0.02}$	0.660.007	3.480.05		(L1 = Farm A)							
	Fhigh (n = 50)	L^2	2105.6771.70	188.455.76	$11.18_{0.2}$	$1.17_{0.02}$	0.310.01	0.660.009	3.47 _{0.05}		laughterhouse							
	F	ГI	1820.0339.13	161.497.36	11.270.19	$1.17_{0.02}$	$0.31_{0.02}$	0.660.008	3.470.05		broilers for sl	ed Fatty Acid						
	6	L3	604.1117.82	68.661.33	8.840.36	$1.17_{0.05}$	0.350.01	0.730.02	3.15±₀.11	> 0.05;	= Supplier of	olyunsaturate						
	Breast (n = 5	L2	605.0414.22	87.98227	6.900.21	$1.03_{0.02}$	0.390.001	0.740.01	2.780.06	sem]; NS. = F	ter region; S	ds, PUFA = I						
		I	576.8310.17	74.38120	7.770.17	1.050.03	0.390.001	0.770.02	2.760.07	SEM [mean	ercial slaugh	ed Fatty Aci						
	Traits		n-6	n-3	n-6 / n-3	PUFA / SFA	IA	II	H/H	Data: mean ± (CSR = comme	SFA = Saturate						
carcas and de	s cor eper	nn eff	ner fec	cia t, o	l s n f	lau at	gh (P∢	ter <0.	сц .01	its) a	ha nd	ad m	bot oist	th a ture	an e (1	ext P <	tens :0.0	sive 5).
Our indicat	resu ed wi	lts de	are rar	e c ige	on	npa val	ira	ble s fo	e to orp) () orot	th tei	er s ns, l	stuc lipio	dy ds a	fin	din I m	gs t iner	hat als,
betwee fatty ac	en 18 cid co	.4-2 mp	23.4 005	1%, itio	1.3 n a	3-6 inc	.0% l th	b, r eir	esp he	ece alt	ctiv h-1	ely ela	0.8 tec	5-1. 1 lip	2% oid	o [3 ind	1]. icat	i'he ors

of breast, thigh and drumstick meat of birds fed on a cereal-

based diet centered on corn, wheat, soy meal, sunflower meal, sunflower oil and supplemented with lysine, methionine and

The muscle fat content of these three commercial slaughter

cuts was quantitatively mainly represented by 8 fatty acids, as

follows: C18:2n-6c (LA), C18:1n-9 an it's isomer C18:1n-7,

mineral complexes is provided in Tables 4 and 5.

REV.CHIM.(Bucharest) ♦ 70 ♦ No. 11 ♦ 2019

Table 5

FATTY ACID COMPOSITION (MEAN_{SEN}) (mg/100 g) OF BROILER CHICKEN MEAT, FED THE SAME DIET, IN RELATION WITH COMMERCIAL CARCASS SLAUGHTER REGIONS AND SUPPLIER FARMS

Patro and	F	Sreast $(n = 7)$	(†)	L	Thigh $(n = 74)$	_	a	rumstick $(n = 74)$		Ma	in effects	
r any actus	ГЛ	Γ_2	L3	ΓI	L_2	L3	ΓI	L_2	L3	CSR	ŝ	CSR x S.
C8:0	$1.34_{0.09}$	$1.82_{0.13}$	$1.30_{0.12}$	3.480.16	3.75030	4.290.51	2.410.19	2.36029	2.470.17	< 0.001	NS.	NS.
C10:0	3.180.12	2.940.31	3.570.20	4.830.34	3.500.17	6.330.47	$5.13_{0.26}$	5.740.53	4.180.19	< 0.001	NS.	< 0.001
C12:0	$1.46_{0.13}$	2.29 _{0.15}	$1.31_{0.10}$	$1.96_{0.22}$	$2.21_{0.22}$	2.450.26	0.880.17	0.92015	0.700.14	< 0.001	0.021	0.010
C14:0	9.830.39	10.910.55	9.810.45	26.76132	30.981.11	33.58227	17.33137	19.751.82	14.161.08	< 0.001	NS.	0.004
C14:1	1.050.03	1.83015	$1.23_{0.12}$	3.880.18	4.42 _{0.18}	$4.84_{0.18}$	3.420.40	3.800.44	2.70027	< 0.001	0.038	0.029
C15:0	$1.75_{0.16}$	1.94021	1.340.16	3.670.32	4.19023	5.120.74	2.16 0.15	2.42 0.31	1.70 _{0.15}	< 0.001	NS.	0.020
C16:0	438.501286	477.441256	380.3011.95	1192.7640.19	1388.015354	1503.70110.11	792.92 _{67.14}	880.26 _{69.03}	635.68 45.27	< 0.001	NS.	0.002
C16:1	44.081.83	50.461.39	42.190.89	171.81_{688}	$199.22_{7.63}$	217.63 _{16.08}	145.4414.95	159.79 _{mm}	115.00 8.98	< 0.001	NS.	0.004
C17:0	$4.61_{0.14}$	6.120.50	3.91019	8.90 _{0.73}	10.170.65	$11.09_{0.83}$	5.750.74	6.37 _{0.81}	4.430.30	< 0.001	0.039	0.028
C17:1	3.51015	3.19 _{8.19}	3.35020	$19.86_{0.86}$	22.951.14	25.45200	8.80 _{0.99}	9.720.98	6.87 _{0.79}	< 0.001	NS.	0.003
C18:0	166.579.40	$173.42_{9.11}$	$180.02_{5.27}$	443.8519.36	517.4020.66	557.4440.68	289.62 _{30.98}	319.7228.14	227.58 _{15.50}	< 0.001	NS.	0.005
C18:1 n-9	485.6323.17	509.31 _{15.88}	492.389.35	1967.5896m	2277.808576	2463.23155.62	1069.1011574	$1190.92_{123.78}$	842.40 _{60.84}	< 0.001	NS.	0.003
C18:1 n-7	36.830.86	38.841.00	38.520.95	77.753.66	90.63428	98.12 _{7.86}	68.35 ₆₈₁	75.41 _{62.15}	53.89 ₁₄₄	< 0.001	NS.	0.006
C18:2 n-6c	524.3510.81	547.8314.06	558.20 _{18.26}	1745.0598.18	2014.5668.14	2181.06140.08	995.50 _{74.08}	1112.65ap.sp	803.85 _{57.26}	< 0.001	NS.	< 0.001
C20:0	2.50014	3.65023	$2.62_{0.15}$	9.520.46	10.97034	$11.96_{0.73}$	6.700.91	$7.41_{0.09}$	5.200.52	< 0.001	NS.	0.025
C18:3 n-6	$1.61_{0.21}$	2.460.40	$1.47_{0.20}$	$3.31_{0.31}$	3.76034	4.190.40	3.391,44	$3.27_{1.13}$	$1.98_{0.47}$	0.09	NS.	NS.
C18:3 n-3	38.74 _{0.81}	42.730.98	33.31 _{0.81}	$117.66_{5.01}$	137.05522	147.9310.63	68.38 _{7.17}	75.90_{782}	53.89 _{am}	< 0.001	NS.	0.004
C20:1 n-9	3.82027	$4.41_{0.26}$	3.73023	17.951.45	20.641.04	$22.30_{1.78}$	7.900.81	8.810.94	6.22038	< 0.001	NS.	0.033
C20:2	7.730.91	8.060.04	$7.40_{0.86}$.pu	nd.	nd.	28.2415.10	$27.01_{12.02}$	19.25 ₈₄₆	< 0.001	NS.	NS.
C20:3 n-3	8.89027	9.570.35	7.740.14	.pu	nd.	nd.	17.311.84	18.971.56	13.641.00	< 0.001	0.010	NS.
C20:4 n-6	47.500.89	51.241.00	42.561.16	$67.12_{7.21}$	82.13995	89.81 _{11.94}	$96.10_{8.69}$	107.1510.04	76.30 _{4.75}	< 0.001	NS.	NS.
C20:5 n-3	12.650.35	$16.22_{0.71}$	$13.22_{0.73}$	$16.34_{0.68}$	19.080.98	$20.76_{1.70}$	23.69_{233}	25.871.94	18.741.46	< 0.001	0.028	0.017
C22:5 n-6	3.370.82	3.51022	$1.88_{0.19}$	4.550.22	5.22034	5.81 _{0.50}	5.250.59	5.84 _{0.70}	4.110.29	< 0.001	0.041	0.015
C22:5 n-3	8.360.13	10.84 _{0.57}	8.28033	$13.60_{1.72}$	15.32 _{1.15}	15.881.18	$16.74_{1.52}$	18.461,47	13.330.98	< 0.001	0.032	NS.
C22:6 n-3	5.74 _{0.87}	8.620.24	$6.11_{0.26}$	$13.89_{0.69}$	17.01146	17.281.17	12.301.10	$13.48_{0.86}$	9.81 _{0.70}	< 0.001	0.002	0.024
SFA	629.7410.07	680.53 ₁₆₁₄	584.18 _{15.76}	1695.4670.89	1971.1875.48	2135.9615432	1123.12404.07	1245.49 _{101.67}	895.58 _{61.15}	< 0.001	NS	0.002
MUFA	574.9224.00	651.21 _{17.45}	538.23 _{9.81}	2258.84107.18	2615.659539	2831.58180.91	1303.02137.40	1448.45144.66	$1027.10_{72.21}$	< 0.001	NS	0.003
PUFA	658.94 _{10.51}	701.0814.55	680.1717.00	1981.52 9615	$2294.13_{76.72}$	2482.73165.18	1266.92105.70	1408.60114.35	1014.89 _{60.14}	< 0.001	NS	< 0.001
Data: mean	± SEM [mea	anszw]; NS	= P > 0.05; C	SR = commer	cial slaughter	region;						
S = Supplier	r of broilers	for slaughter	house (L1 =)	Farm A, L2 =]	Farm B; L3 =	Farm C);						
SFA = Satur	rated Fatty A	cids. MUFA	I = Monouns:	aturated Fatty /	Acids. PUFA	= Polyunsatura	ted Fatty Acids;					

C16:0, C18:0, C18:3n-3 (LNA), C16:1, C20:4n-6 (AA) and C20:5n-3 (EPA). The literature reports the same decreasing order of concentration for the main fatty acids of broiler meat, important for human nutrition (LA, LNA, AA, EPA) as a result of the dietary inclusion of oil sources rich in n-3 polyunsaturated fatty acids. [32]. More than that, because these FA are recommended for human nutrition due to their ability to minimize the probability of lifestyle-related diseases occurrence [33-35], the recent researches centered on n-3 PUFA targeted broiler meat enrichment through new (ultrasound-assisted nano-emulsion preparation [36] and consecrated strategies, like direct feed supplementation [37-46]. These approaches are used to minimize ù-6: ù-3 ratio in human diets [47, 48].

As we mentioned, in agreement with the values of total fat content, the predominant lipid fractions in broiler meat is displayed in Figure 1, monounsaturated fatty acids (MUFA) ranging between 28.97-38.04% of the total IMF. C18:1 n-9 is the most important in the MUFA group, with an average overall value of 85.18% of total MUFA.

PUFA fraction is the most important group, with average values between 31.85-39.61% of total FA, C18:2n-6c and C18:3n-3 being found to be the dominant ones, responsible for 83.77% and 5.72% of total PUFA. The total SFA content of breast, thigh and drumstick meat did not differ between suppliers (P>0.05), although thigh and drumstick had the higher content, which is unfavorable to human health [49]. The overall average values for this fraction were between



28.60–33.85% of total FA. Here, C16:0 was the most abundant in terms of quantity (70.17% of total SFA), followed by C18:0, (26.22% of total SFA).

In our study there is an evident statistically significant effect (P<0.001) of CSR on each fatty acid content, with less extensive rate for C18:3n-6 (P<0.05). The obvious findings highlighted that Farm B supplied broilers with a delivered higher content of beneficial fatty acids (LA, LNA, AA, EPA, DHA) in breasts and drumstick, while for thigh Farm C had the best results. For C20:5n-3 (EPA), it was noted that the quantity of the drumstick lipids was comparatively nearly two times greater than that of the IMF breast muscles (P<0.001).

Dietary incorporation of sunflower meal and oil lowered the C18:2n-6c (LA) content in breast, but not in thigh and drumstick meat. The same pattern is retained and for some undesirable saturated fatty acids (USFA), such as C10:0, C12:0, C14:0, C16:0, C17:0 and C20:0.

C18:1 was the major MUFA, while PUFA was mainly defined by C18:2n-6c. No statistical differences (P>0.05) were found in the whole monounsaturated fatty acids (MUFA) and total polyunsaturated fatty acids (PUFA) contents of breast, thigh and drumstick meat between suppliers, only CSR and interaction CSR x S had a significant, but specific effect on almost all FA.

Meat fatty acids profiles of chicken meat (SFA, MUFA, PUFA, n-3, n-6), and the ratios among them, showed a significant effect on meat of CSR, source or their interaction, that can be attributed to the implementation of the dietary strategy and feed consumption (Table 4).

Descriptive data of lipids health related parameters have confirmed, through PUFA n-6/n-3 index values, that the breast muscles have the lowest ratios (P<0.001), in a context where the literature indicate an optimum value ratio of 4:1 [35].

Fig. 1 Composition of fatty acids groups (%) in broiler meat, in accordance with their commercial slaughter cuts (breast, thigh, drumstick) and supplier farms (L1, L2, L3)

Fig. 2 Principal component analysis for some fatty acids' groups and indexes in different commercial slaughter regions

Lipid nutritional quality indices for adipose tissue ranged from 0.31 to 0.39 for IA, respectively 0.57 and 0.77 for IT. According to the relative contents of the particular groups of fatty acids, the thigh lipids showed the lowest atherogenic (IA; P < 0.001) and thrombogenic (IT; P < 0.001) indexes in comparison with the analyzed fats of drumstick and breast muscles (Table 4). This image is clearly represented throughout the principal component analysis (PCA), where CSR samples were displayed in the multi-dimensional space of the newly calculated variables. The first two PCs calculated from these descriptors account for 93.71% of the total data variability, as shown in Figure 2.

IT and IA indices are strongly correlated with breast muscle lipids composition, being adjacent and related with F1 plane in a negative way. The lipids of drumstick have loading values affiliated stronger to F2 plane. The thigh meat samples are positively correlate with F1 plane, having lipids strongly related and positively described by all health-related indices. As anticipated, studied CSR and their FA profiles reflected the FA composition of bird's diet fat (especially sunflower oil), this type of supplementation being company nutritional strategy, based on literature studies [50, 51].

Conclusions

The proximate chemical composition of the commercial slaughter cuts revealed breast superiority, the obtained values, especially for proteins (16.26-22.78%) and lipids (1.80-7.45%) being mainly affected by region (P<0.001). Meat fatty acid profile (P<0.001) was affected by commercial slaughter regions (CSR) and interactions between CSR and supplier farms (Farm A, B, and C) at different levels, with quantitative values comparable to

those mentioned in the literature, with emphasis on dietary manipulation. Total content of SFAs, MUFAs and MUFAs had the highest level in the thigh (P < 0.001). Farm B supplied broilers with a delivered higher content of beneficial fatty acids (LA, LNA, AA, EPA, and DHA) in breasts and drumstick, while for thigh Farm C had the best results.

Although the results of the current study demonstrate that the fatty acid profile in edible tissues (breast, thigh and drumstick) depends of the feed composition, it definitely can be influenced and by the performance management of the supplier farms in all aspects, such as: nutritional management, training of the people or degree of the good raising practices (GRP) implementation. This assessment played an active role in the future execution of the company brand growth plan.

References

1.PEREIRA, P.M., VICENTE, A.F., Meat Sci., 93, no. 3, 2013, p. 586.

2.WEBB, E.C., O'NEILL, H.A., Meat Sci., 80, 2008, p. 28.

3.SCOLLAN, N.D., PRICE, E.M., MORGAN, S.A., HUWS, S.A., SHINGFIELD, K.J., Proc. Nutr. Soc., **76**, 2017, p. 603.

4.DE SMET, S., RAES, K., DEMEYER, D., Anim. Res., 53, 2004, p. 81.

5.ZHANG, W., XIAO, S., SAMARAWEERA, H., LEE, E.J., AHN, D.U., Meat Sci., **86**, 2010, p. 15.

6.SILBERGELD, E. K., Global Transitions, 1, 2019, p. 83.

7.BOGOSAVLJEVI-BOSKOVI, S., PAVLOVSKI, Z., PETROVI, M.D.,

DOSKOVI, V., RAKONJAC, S., Afr. J. Biotechnol., 9, no. 54, p. 9177.

8.WOOD, J.D., ENSER, M., FISHER, A.V., NUTE, G.R., SHEARD, P.R., RICHARDSON, R.I., Meat Sci., **78**, 2008, p. 343.

9.* * *, E.U. REPORT "Report to the European Parliament and the Council on the application of Directive 2007/43/EC and its influence on the welfare of chickens kept for meat production, as well as the development of welfare indicators".

10.EL-KATCHA, M.I., EL-KHOLY, M.E., SOLTAN, M.A., EL-GAYAR, A.H., J. Poultry Sci., 2, 2014, p. 71.

11.BARCLAY, W., ABRIL, R., ABRIL, P., WEAVER, C., ASHFORD, A., World Rev. Nutr. Diet, **83**, 1998, p. 61.

12.FRITSCHE, K.L., CASSITY, N.A., HUANG, S., Poultry Sci., 70, 1991, p. 1213.

13.AHMAD, H., TIAN, J., WANG, J., KHAN, M.A., WANG, Y., ZHANG, L., J. Agric. Food Chem., **60**, 2012; p. 7111.

14.ALAGAWANY, M., ELNESR, S.S., FARAG, M.R., ABD EL-HACK, M.E., KHAFAGA A.F., TAHA, A.E., TIWARI, R., Animals, **9**, 2019, p. 573.

15.LEE, S.S., WHENHAM, N., BEDFORD, M.R., Animal Nutrition 5, 2019, p. 11.

16.SAEED, M., YATAO, X., HASSAN, F. U., ARAIN, M. A., ABD EL-HACK, M. E., NORELDIN, A. E., SUN, C., Int. J. Mol. Sci., **19**, 2018, p. 462.

17.YAN, L., KIM, I. H., J. Appl. Anim. Res., 41, 2013, p. 392.

18.PASCARIU, S.M., POP, I.M., SIMEANU, D., PAVEL, G., SOLCAN, C., Braz. J. Poultry Sci., 19, no. 2, p. 191.

19.MOTTET, A., TEMPIO, G., Worlds Poultry Sci. J., **73**, no. 2, 2017, p. 245. 20.MIR, N.A., RAFIQ, A., KUMAR, F., SINGH, V., SHUKLA, V., J. Food Sci. Technol., **54**, no. 10, 2017, p. 3001.

21.AHIWE, E.U., OMEDE, A.A., ABDALLH M.B., IJI, P.A., Animal Husbandry and Nutrition, Chapter 6, 2018, p. 115.

22.SUKHIJA, P. S., PALMQUIST, D. L., ýJ. Agric. Food Chem., **36**, no. 6, 1988, p. 1202 – 1206.

23.***, AOAC (2000). Official methods of analysis of AOAC. International 17th edition; Gaithersburg, MD, USA Association of Analytical Communities. 24.FOLCH, J., LEES, M., SLOANE STANLEY, G. H., J. Biol. Chem., **226**, 1957, p. 497.

25.FRUNZA, G., SIMEANU, D., POP, I. M., BOISTEANU, P. C., STEFAN, M., Rev. Chim. (Bucharest), **70**, no. 2, 2019, p. 174.

26.PAPUC, C., CRIVINEANU, M., NICORESCU, V., PAPUC, C., PREDESCU, C., Rev. Chim. (Bucharest), **63**, no. 12, 2012, p. 1198.

27.LUP, F., POP, I. M., SIMEANU, D., VICAS, S., SIMEANU, C., MIERLITA, D., Rev. Chim. (Bucharest), **69**, no. 1, 2018, p. 222.

28.ULBRICHT, T. L., SOUTHGATE, D. A. T., The Lancet, **338**, 1991, p. 991. 29.POPOVA, T., IGNATOVA, M., PETKOV, E., STANISIC, N., Arch. Anim. Breed., **59**, 2016, p. 319.

30.FERNANDEZ, M. L., WEST, K. L., J. Nutr., 135, 2005, p. 2075.

31.CULIOLI, J., BERRI, C., MOUROT, J. Sci. Aliment., **23**, no. 1, 2003, p. 13. 32.KALAKUNTLA, A.S., NAGIREDDY, A.N.K., PANDA, B.A.K, NARASIMHA JATOTH, A., RAGHUNANDAN THIRUNAHARI, A., VANGOOR, R.R., Anim. Nutr., **3**, 2017, p. 386.

33.SIMOPOULOS, A.P., Biomed. Pharmacother., 56, 2002, p. 378.

34.SIMOPOULOS, A.P., Exp. Biol. Med., 233, no. 6, 2008, p. 674.

35.SIMOPOULOS, A.P., Nutr., 8, 2016, p. 128.

36.ABBASIA, F., SAMADIA, F., JAFARIB S.M., RAMEZANPOURC, S., SHAMS-SHARGHD, M., J. Funct. Foods, **57**, 2019, p. 373.

37.LOPEZ-FERRER, S., BAUCELLS, M.D., BARROETA, A.C., GRASHORN, M.A., Poultry Sci., **80**, no. 6, 2001, p. 741.

38.LOPEZ-FERRER, S., BAUCELLS, M.D., BARROETA, A.C., GALOBART, J., GRASHORN, M.A., Poultry Sci., **80**, no. 6, 2001, p. 753.

39.CHEONG, A. M., TAN, C. P., NYAM, K. L., Ind. Crop. Prod., 87, 2016, p. 1.

40.BETTI, M., PEREZ, T.I., ZUIDHOF, M.J., RENEMA, R.A., Poultry Sci., 88, 2009, p. 1740.

41.BETTI, M., SCHNEIDER, B.L., WISMER, W.V., CARNEY, V.L., ZUIDHOF, M.J., RENEMA, R.A., Poultry Sci., **88**, 2009, p. 1085.

42.JIA, W., ROGIEWICZ, A., BRUCE, H.L., SLOMINSKI, B.A., Can. J. Anim. Sci., **90**, 2010, p. 203.

43.ELKIN, R.G., YING, Y., HARVATINE, K.J., J. Agr. Food Chem., **63**, 2015, p. 2789.

44.APPERSON, K.D., CHERIAN, G., Poultry. Sci., 96, 2016, p. 1228.

45.KONIECZKA, P., CZAUDERNA, M., SMULIKOWSKA, S., Anim. Feed Sci. Tech., **223**, 2017, p. 42.

46.SHAHIDI, F., AMBIGAIPALAN, P., Ann. Rev. Food. Sci. Technol., 9, 2018, p. 345.

47.GIVENS, D.I., GIBBS, R.A., Lipids, 40, 2005, p. 121.

48.GIVENS, D.I., GIBBS, R.A., P. Nutr. Soc., 67, 2008, p. 273.

49.KANAKRI, K., CARRAGHER, J., HUGHES, R., MUHLHAUSLER, B., GIBSON, R., Br. Poultry Sci., **58**, no. 3, 2017, p. 283.

50.SCAIFE, J.R., MOYO, J., GALBRAITH, H., MICHIE, W., CAMPBELL, V., Br. Poultry Sci., **35**, 1994, p. 107.

51.HRDINKA, C., ZOLLITSCH, W., KNAUS, W., LETTNER, F., Poultry Sci., 75, 1996, p. 208.

Manuscript received: 12.11.2019