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DAMASIO and C. SARRA Studio per l'Alimentazione degli Animali in Produzione Zootecnica - C.N.R. - Torino, Italy.

turne of age on total lipids and phospholipids in turkey meat.

factors have influence on the amount and quality of fat in meat. Among these factors the plan of nutrition factors in connection with the modern intensive raising techniques) and the sex of the animals have the importance, even though the role of the age of the animals must not be extensive investigations have put in evidence that as animals must not be underestimated. mutest investigations have put in evidence that, as animals grow and accumulate fat, the propor-fact, extensive investigations have put in evidence that, as animals grow and accumulate fat, the propor-fact, extensive in the muscles increases at the expense of phosphalicide. of triglycerides in the muscles increases at the expense of phospholipids; simultaneously there is a vaat the expense of phospholipids; simultaneously there is a va-

nation is concerned. fatty acids C16 and C18 are mainly involved in the process, which is of the utmost importance as unsaturately acids play a significant role both in limit ratty acids play a significant role both in lipid metabolism and in determining the perishability of meat. determinant of the characteristics of the as a determinant of the characteristics of the various classes of muscle lipids.

marials and Methods trust and leg samples from turkeys carcasses were used for the analysis. The large-type turkeys (White Giant) red on a standard diet and killed at 18 and 23 weeks of age.

to the method of Bligh and Dyer (1959) which of an homogenisation with chloroform-methanol-water followed by filtration and separation of the chloreform phase which contains the lipids. Part of the lipid extract was mixed - according to the method of Bunstein et al. (1967) - with activated silicic acid and a mixture of chloroform-hexane-diethylether (2:1:1) and then centrifuged in order to separate the triglycerides and free fatty acids (in the supernatant) from the phospholipids which remain bound to the silicic acid. Phospholipids are then transmethylated directly m the silicic acid by means of sodium methoxide and finally extracted with hexane. The concentrated hexane phase is analysed by GLC. The supernatant containing both triglycerides and free fatty acids (FFA) was treaand with a purified anion exchange resin Dowex 1-X8 so as to separate the two components: the resin retains the FFA. The methylation of FFA was accomplished directly on the resin by means of a mixture methanol-HCl 5-10%. The methylesters were analysed by GLC. The supernatant containing the triglyceride fraction was conified with KOH; the fatty acids were then released by means of HCl and finally extracted with hexane. This extract was treated with the anion exchange resin so as to separate the cholesterol, which was not determined, from the fatty acids which are absorbed on the resin. The fatty acids were then methylated as described for the FFA and determined, after extraction with hexane, by means of a chromatograph C.Erba 4200 equipped with a hydrogen flame ionization detector. The percentage of phospholipids was determined on the assumption that Phosphorus represents 4% of total phospholipids (Hornstein et al., 1961). Phosphorus is then determined coording the method described by Morrison (1964) based on the destruction of organic matter by wet oxidation, liberating phosphoric acid, formation of a phosphomolybdic acid complex which is quantitatively reduced to a

malysis carried out on the turkey's breast and leg muscles put in evidence some modifications in the composition of the various lipid classes as regards to the age of the animals.

In particular:

beteropoly blue color and measurement of the color by spectrophotometry.

for what triglyceride are concerned (see table 1) the fatty acids of the breast don't substantially change age, except C16:1 which increases and C17:1 which decreases (P < 0.05), while in the leg it is possible observe a greater variation, with a decrease of C14:0, C16:0, C17:0, C18:0 (P≤0.05), C20:3, C20:5 $(P \leqslant 0.01)$ and an increase of C18:1 (P \leqslant 0.01), C18:2, C18:3 (P \leqslant 0.05).

for the FFA (see table 2) there are not great differences between breast muscles from 18 weeks old turand 23 weeks old ones: there is only an increase of C18:2 (P < 0.05) and a decrease of C20:3 and C20:5 0.05), while also in this case in leg muscles there are more meaningful variations: C16:1, C18:1 and C18:2 decrease ($P \le 0.05$, $P \le 0.01$ and $P \le 0.001$ respectively) and C18:0 increases ($P \le 0.01$).

finally for what phospholipids are concerned (see table 3) in breast muscles there is a decrease of C15:0, (P \leq 0.001) and C22:6 (P \leq 0.01) and an increase of C18:0 (P \leq 0.01), whereas in leg muscles there is decrease of C17:0 (P \leqslant 0.05), C18:2 (P \leqslant 0.01) and an increase of C18:1 and C20:4 (P \leqslant 0.01).

proportion of phospholipids on the overall amount of lipids is different in leg and in breast muscles in mation to the age of animals. In fact in breast such proportion is almost the same in 18 weeks old turkeys the age of animals. In fact in preast such proportion = $\frac{1}{23}$ weeks old, while in leg muscles the phospholipids significantly decrease (P \leq 0.001).

e results are closely related to those obtained by other Authors in investigations carried out on diffetent kinds of meat.

Tab. 1 - Effect of age on fatty acid composition of triglycerides of turkey's meat. (°)

BREAST

FATTY ACIDS	18	weeks	23	weeks		F	18	weeks	23	weeks	F
14:0	2.70	(0.65)	2.52	(0.83)		n.s.	2.71	(0.83)	1.81	(0.22)	ж
15:0	0.61	(0.34)	0.33	(0.16)	×	n.s.	0.26	(0.03)	0.35	(0.18)	n.s.
16:0	24.81	(1.68)	26.28	(2.01)		n.s.	27.74	(3.44)	24.21	(0.61)	*
16:1	3.67	(0.70)	5.34	(1.19)		ж	3,94	(0.45)	4.62	(1.55)	n.s.
17:0	=		-				0.31	(0.08)	0.20	(0.06)	*
17:1	0.57	(0.21)	0.34	(0.11)		ж	-		30		
18:0	9.55	(1.67)	7.87	(1.29)		n.s.	10.82	(1.82)	8.12	(1.34)	ж
18:1	25.26	(6.64)	25.90	(4.58)		n.s.	22.45	(3.84)	29.35	(2.17)	ж ж
18:2	19.01	(5.25)	21.25	(3.14)		n.s.	20.56	(3.32)	24.97	(1.70)	*
18:3	₩.		S#				0.94	(0.37)	1.49	(0.17)	*
20:3	5.22	(4.18)	4.21	(2.33)		n.s.	2.71	(1.10)	0.92	(0.47)	* *
20:4	1.99	(1.05)	1.72	(0.80)		n.s.	3.09	(1.39)	2,35	(1.11)	n.s.
20:5	2.96	(2.44)	2.45	(1.38)		n.s.	2.68	(1.20)	0.80	(0.32)	* *

^(°) Mean and standard deviation in brackets (percent).

Tab. 2 - Effect of age on free fatty acid composition of turkey's meat. (°)

	BREAST						L E G					
FATTY						9						
ACIDS	18	weeks	23	weeks		F	18	weeks	23	weeks	F	
						2						
14:0	2.93	(1.08)	2.84	(1.14)		n.s.	2.06	(0.61)	2.18	(0.59)	n.s.	
15:0	3.73	(1.52)	2.31	(1.39)		n.s.	2.65	(1.26)	3.85	(1.87)	n.s.	
16:0	20.39	(2.25)	21.70	(2.81)		n.s.	18.94	(1.66)	18.53	(3.30)	n.s.	
16:1	2.36	(0.36)	2.65	(0.89)		n.s.	2.89	(0.31)	2.11	(0.60)	ж	
17:1	-		-				0.77	(0.23)	1.68	(1.18)	n.s.	
18:0	13.09	(1.19)	12.23	(3.21)		n.s.	9.09	(1.25)	13.56	(2.37)	* *	
18:1	13.96	(2.08)	17.08	(2.82)		n.s.	17.86	(0.95)	14.11	(2.39)	* *	
18:2	17.97	(3.57)	24.74	(5.46)		ж	28.58	(1.81)	22.97	(2.36)	米准基	
18:3	-		-77				0.80	(0.24)	0.91	(0.53)	n.s.	
20:3	7.88	(4.24)	3,52	(1.45)		ж	2.11	(0.61)	4.14	(2.78)	n.s.	
20:4	11.04	(1.23)	9.43	(5.06)		n.s.	10.39	(1.50)	11.87	(3.53)	n.s.	
20:5	3.06	(1.15)	1.39	(0,55)		ж	2,30	(1.09)	2,30	(1.21)	n.s.	

^(°) Mean and standard deviation in brackets.(percent).

Tub. 3 - Effect

FATTY ACIDS 14:0 15:0 16:0 16:1 18:0 20:4 22:6

(*) Mean and st * significant P≤0.001.

BIBLIOGRAPHY

Acosta S.O., M

Sci. 45, 169. Bligh E.G. and Physiol. 37, 9 Hornstein I., 26, 581. Mornstein I., techniques. An Katz M.A., Dug Tissues. J. Fo Link B.A., Bray during growth. Link B.A., Bra lipids during Marion J.E. and attion. J. Food Morrison W.R., Auterials. Ana Ohtake Y., Hosh ceride Composit Osborn W.E., Mc Estradiol-17-Be Salmon R.E. and Fat on the Fatt Poultry Sci. 52

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^{*} significant differences $P \le 0.05$; * * significant differences $P \le 0.01$.

^{*} significant differences $P \leqslant 0.05$; * * significant differences $P \leqslant 0.01$; * * * significant differences P≤0.001.

Effect of age on fatty acid composition of phospholipids of turkey's meat. (°)

	В	REA	. S T				L E G				
(ATTY	18	weeks	23	weeks	F	18	weeks	23	weeks	F	
ACIDS											
	0.28	(0.04)	0.32	(0.09)	n.s.	0.21	(0.05)	0.20	(0.02)	n.s.	
14:0	7.56	(0.41)	4.08	(0.67)	* * *	4.44	(0.53)	4.15	(0.63)	n.s.	
15:0	19.57	(0.47)	20.44	(2.15)	n.s.	15.10	(1.37)	14.66	(1.00)	n.s.	
16:0	0.48	(0.24)	0.73	(0.40)	n.s.	0.58	(0.06)	0.68	(0.24)	n.s.	
16:1	0.19	(0.06)	0.23	(0.10)	n.s.	0.27	(0.04)	0.21	(0.04)	ж	
17:0	1.84	(0.41)	0.84	(0.32)	* * *	0.95	(0.22)	0.86	(0.17)	n.s.	
18:0	16.52	(1.26)	19.71	(1,91)	ж ж	21.84	(0.77)	21.30	(1.52)	n.s.	
18:1	16.02	(1.53)	19.67	(3.79)	n.s.	11.56	(0.67)	14.28	(1.59)	ж ж	
18:2	18.56	(2.00)	18.90	(2.54)	n.s.	24.88	(1.06)	21.70	(1.85)	ж ж	
18:3						0,22	(0.12)	0.32	(0.06)	n.s.	
20:4	10.90	(0.51)	8.94	(2.24)	n.s.	11.99	(0.50)	13.76	(0.86)	ж ж	
22:6	2.95	(0.57)	1.79	(0.23)	ж ж	2.12	(0.63)	2.34	(0.30)	n.s.	

Mean and standard deviation in brackets. (percent).

BIELIOGRAPHY

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kosta S.O., Marion W.W. and Forsythe R.H., 1966. Total Lipids and Phospholipids in Turkey Tissues. Poultry

Bligh E.G. and Dyer W.J., 1959. A rapid method of total lipid extraction and purification. Can. J. Biochem. Physiol. 37, 912.

Momstein I., Crowe P.F. and Heimberg M.J., 1961. Fatty acid composition of meat tissue lipids. J. Food Sci. 26, 581.

Monstein I., Crowe P.F. and Ruck J.B., 1967. Separation of muscle lipids into classes by non-chromatographic techniques. Anal. Chem. 39, 352.

Matz M.A., Dugan L.R.Jr. and Dawson L.E., 1966. Fatty Acids in Neutral Lipids and Phospholipids from Chicken Masues. J. Food Sci. 31, 717.

Link B.A., Bray R.W., Cassens R.G. and Kauffman R.G., 1970 a. Lipid deposition in bovine skeletal muscle during growth. J. Anim. Sci. 30, 6.

thk B.A., Bray R.W., Cassens R.G. and Kauffman R.G., 1970 b. Fatty acid composition of bovine skeletal muscle lipids during growth. J. Anim. Sci. 30, 726.

Arion J.E. and Woodroof J.G., 1965. Lipid Fractions of Chicken Broiler Tissues and Their Fatty Acid Compoaltion, J. Food Sci. 30, 38.

trison W.R., 1964. A Fast Simple and Reliable Method for the Microdetermination of Phosphorus in Biological

aterials. Anal. Biochem. 7, 218. Y., Hoshino Y., Aoki T., Ohgane T., Ohnuki M. and Fukumari Y., 1975. Changes in Fatty Acid and Trigly-

Compositions of Porcine Tissue Lipids Associated with Growth of Pigs. Jap. J. Zootech. Sci.46(8), 460. compositions of Porcine Tissue Lipids Associated with Grown of Sex, Age and W.E., Moreng R.E. and Hartung T.E., 1969. Turkey Lipid Characteristics: Influence of Sex, Age and tradiol-17-Beta-Monopalmitate. Poultry Sci. 48, 274.

R.E. and O'Neil J.B., 1972. The Effect of the Level and Source and of a Change of Source of Dietary the Fatty Acid Composition of the Depot Fat and the Thigh and Breast Meat of Turkeys as Related to Age. bultry Sci. 52, 302.

R.C., Suess G.G. and Brungardt V.H., 1968. Fatty acids of certain bovine tissue and their association The growth, carcass and palatability traits. J. Anim. Sci. 27, 632.

significant differences P≤0.05; * * significant differences P≤0.01; * * * significant differences P\$0.001.