COMPOSITION OF LIVER PATES. LETERMINATION OF TRANS FATTY ACIDS, NITRATES AND NITRITES.

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Background

The patterns of Portuguese diet are changing. There are a lot of factors contributing to this issue: a new lifestyle, a wide range of Processed food products in the market, the increasing interest of the population with regard to nutritional labeling and fears ^{conc}erning undesirable components in food.

Pates are not a new processed food, although recently their consumption has been increasing probably due to marketing reasons Considering the wide variety and brands on the market. Despite these, a poor work has been done in our country on the evaluation ^{of the} composition of these products [1].

As a form of health protection, the reduction of the fat ingestion and specifically of trans fatty acid contents in the diet are nutritional bjectives. In an attempt to inform the population on the nutritional value of liver pâtés and on the potential intake levels of some of the undesirable components carried over by them, this work was developed.

Energy, lipids, proteins, cabohydrate contents and fatty acid composition have been determined.

the recent years nitrates and nitrites have attracted a great deal of attention because of their potential role in producing hittosamines in the human body [2] and also for regulamentary reasons [3].

the indispensability of nitrates and nitrites as antimicrobial agents against Clostridium botulinum and their abilities to give the desirable characteristics of colour and other organoleptic properties in cured meat justify that the determination of these salts was in this first approach of pâté study. For determination of these salts a very accurate, rapid and precise methodology by FIA has been used [4].

Methods

October (Drying of the sample at 100 -102° C in air oven (AOAC [5] 24.003); Proteins (Kjeldahl digestion with Selacion); October (Evaluation) was made by calculation); Catalyst (AOAC 2.058); Lipids (Soxhlet extraction (AOAC 24.005); Carbohydrates (Evaluation was made by calculation); actalyst (AOAC 2.030), Lipius (Souther Catalons), and Composition and trans fatty acids - GLC/FID Analysis was used [6]. A Chromatograph Pye Unicam fitted with a flame Composition and trans tatty acids - ODCIT ID (May 313 Mas documents) was used for detector (FID) and a 0,25mm x 50m fused silica capillary column coated with CP-Sil 88 (Chrompack) was used for the injected volume was aration of fatty acids methyl esters. The column was maintained at 185° C. 1:50 was the split ratio and the injected volume was Peak areas were processed by using a Pye Unicam CDP4 computing integrator.

he methyl esters were prepared by transesterification with BF3/MeOH. Before metilation the samples were hydrolysed with (0H/MeOH (11g/l)) and the methyl esters were extracted with n-heptan; Nitrates and nitrites - A multidetection flow injection Was used for the automatic sequential determination of nitrate and nitrite. This provided an alternative method of analysis to was used for the automatic sequential determination of inflate and investigation was used for the automatic sequential determination of inflate and in divided. An on-line copper-cadmium reductor column reduces nitrate to nitrite in part of the sample plug. Spectrofotometric An on-line copper-cadmium reductor column reduces muale to make the sequential th determination of nitrite and nitrate plus nitrate is made and a discount and nitrate present in different concentration ranges in the sample with only one detector.

brands of liver pâtés in a total of 15 samples were randomly purchased from the local market.

Results and Discussion

Table 1 summarises the results of water, proteins, fat, carbohydrates, *trans* fatty acids, nitrates, nitrites contents and caloric value of the 5 commercial brands of liver pâtés analysed in the total of 15 samples.

The European Community legislation for liver pâtés allows 100 mg/kg for residual nitrites (expressed as NaNO₂) and 250 mg/kg for residual nitrates (expressed as Na NO₃). Under optimal conditions for the nitrates' and nitrites' analyses [4] the levels are ranging from 4.9mg/kg ± 1.10 to 22.3mg/kg ± 0.16 and from 11.73mg/kg ± 0.088 to 130.1mg/kg ± 0.016 , respectively. A dispersion (p < 0.01) around the mean values was observed and for some brands it was significant. Despite the small sampling the results found show that in terms of averages the levels of concentrations of these constituents are below the allowable limits.

Concentrations of *trans* fatty acids are scarce in all samples (ranging from $0.37\% \pm 0.049$ to $1.64\% \pm 0.025$) when compared with levels determined in other fatty foods [6]. A significant dispersion in the results was observed in some brands.

The great variability pertaining to the *trans* fatty acids, nitrate and nitrite contents may indicate a poor uniformity of manufactoring within the same brand.

Conclusions

In terms of contents of nitrites, nitrates and runs fatty acids the liver pâtés are safe. Although being unbalanced foods and highly energetic is recommended that liver pâtés should be consumed with moderation.

References

- [1] Gonçalves Ferreira, F.A., Silva Graça, M.E. (1985), Tabela de composição dos Alimentos Portugueses, Instituto Nacional de Saúde Dr. Ricardo Jorge (Lisboa-Porto), pág 34.
- [2] F. Schweinsberg, Catalysis of nitrosamine in P. Bogovski and E.A. Walkers (Eds) N Nitroso Componds in the Environment, International Agency for Reserch on Cancer, Leao (1974) 80-85..
- [3] 95/2/EC Directive, relating with food additives excluding colouring and sweeteners, J.O.E.C., # L 61/1-25,18.3.95.
- [4] Ferreira, I.M.P.L.V.O.; Lima, J.L.F.C; Montenegro, M.C.B., Perez-Olmos, R., (1996) Analist, (submited).
- [5] Official Methods of Analysis (1980) 13th Ed., AOAC, Arlington, VA.
- [6] Oliveira, M. Beatriz P. P. and Ferreira, M. A. (1994), Grasas y Aceites, 45, 3, 113-118.

Table 1: Contents of water, protein, fat, carbohydrates, trans fatty acids, nitrates, nitrites and energy of 5 commercial brands of liver patés in the total of 15 samples.

Liver Pâté ¹ Brand / lot		Water (x ± sd) %	Protein (x ± sd) %	Carbohydrate (x ± sd) %	Fat (x ± sd) %	fatty acids (x ± sd) %	Nitrite mg/kg	Nitrate mg/kg	Energy KJ/100
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	1	47 ± 0.0	9 ± 0.7	3 ± 0.1	41 ± 0.0	1.03 ± 0.012	4.9 ± 0.10	33.7 ± 6.38	1745
A	2	48 ± 0.2	9 ± 0.5	3 ± 0.3	40 ± 0.0	0.87 ± 0.110	5.1 ± 0.37	53.9 ± 4.71	1707
	3	46 ± 0.0	7 ± 0.5	4 ± 0.5	42 ± 0.4	1.64 ± 0.025	5.3 ± 0.11	38.2 ± 5.37	1912
							*	- A	119707
	4	53 ± 0.1	16 ± 0.2	8 ± 0.1	23 ± 0.0	0.50 ± 0.060	13.7 ± 0.11	130.3±16.06	1268
В	5	57 ± 0.7	16 ± 0.0	8 ± 0.7	20 ± 1.4	0.37 ± 0.049	12.5 ± 0.10	114.9 ± 8.91	1155
	6	52 ± 0.1	16 ± 0.3	7 ± 0.1	25 ± 0.0	0.42 ± 0.031	12.7 ± 0.28	123.7 ± 9.65	1326
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	7	50 ± 0.0	12 ± 0.1	6 ± 0.6	33 ± 0.7	1.28 ± 0.066	9.4 ± 1.16	25.2 ± 5.93	1544
С	8	51 ± 0.0	13 ± 0.5	4 ± 0.2	32 ± 0.7	1.41 ± 0.059	10.4 ± 0.09	28.8 ± 4.46	1490
	9	48 ± 0.0	13 ± 0.2	4 ± 0.9	36 ± 0.7	0.79 ± 0.006	9.4 ± 0.37	39.3 ± 5.58	1640
		*		Charles 1977	W. (1980)				1981 20 20
	10	49 ± 0.0	10 ± 0.1	6 ± 0.1	35 ± 0.0	0.57 ± 0.017	7.5 ± 0.58	58.3 ± 0.62	1586
D	11	45 ± 0.0	11 ± 0.5	14 ± 0.2	31 ± 0.7	0.43 ± 0.092	10.3 ± 1.00	54.1 ± 1.20	1586
	12	48 ± 0.0	11 ± 0.3	7 ± 0.3	34 ± 0.0	0.47 ± 0.064	14.0 ± 1.51	55.6 ± 3.25	1582
						*	*	*	
	13	54 ± 0.0	13 ± 0.4	5 ± 0.0	28 ± 0.4	0.61 ± 0.113	11.7 ± 0.35	35.5 ± 0.96	1356
E	14	55 ± 0.0	10 ± 0.6	6 ± 0.8	29 ± 0.1	0.73 ± 0.069	11.8 ± 0.11	32.0 ± 2.24	1360
	15	53 ± 0.0	11 ± 0.4	6 ± 0.1	31 ± 0.3	1.25 ± 0.102	22.3 ± 0.16	11.7 ± 0.89	1452

¹ A, B, C, D, E - are different brands of commercial pates liver, the numbers represent different lots.

^{*} Significant differences between the results were determined by ANOVA methodology followed by Fisher's PLSD test. Differences were considered significant for p < 0.01.