Bakteriologische Zusammenhänge in rohem und verarbeitetem Fleisch und ähnlichen Produkten

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Lusammenfassung

Ergebnis einer Vergleichsstudie durch Ermittlung der Bakteriengesamtzahl und der Anwesenheit von Esch. coli und Salmonellen spp. in Fleisch und adnlichen Produkten.

Einhundertfunfzehn Muster wurden von einer ortlichen Fabrik gesammelt, im Zeitraum 1976/1977. Die Informationen Wurden analysiert um herauszufinden ob zwischen der Gesamtzahl der Bakterien und den Krankheitstragern irgendein Zusammenhang besteht.

In der Hauptzahl der Falle war keine Wechselbeziehung zwischen der Gesamtzahl und Esch. coli festzustellen. Bedeutende negative Wechselbeziehungen existierten jedoch fur Steakburger (P < 0.05), Frankfurter Wurstchen (P < 0.01), und Fruhstucksfleisch (P < 0.05) was darauf hindeutet dass bei steigender Gesamtzahl die Zahl der Each. Esch. coli abnahm.

Sieben Prozent der Proben waren Salmonella-positiv. Die durchschnittszahlung <u>Esch. coli</u> von der letzteren (log₁₀ 4.51 g⁻¹) war bedentend hoher (P < 0.01) als die duchschittszahlung <u>Esch. coli</u> bei den Salmonella-negativ proben (log₁₀ 2.89 g⁻¹). Dass lasst darauf shliessen dass die Proben mit hoher <u>Esch. coli</u> zahlungen sind mit ^{grosserer} währschinlichkeit Salmonella-positiv als die proben with einer niedrigen anzahl von <u>Esch. coli</u>.

Bacteriological relationships in raw and processed meats and associated products

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Abstract

A study was made to collate information on the total plate count and the presence of Esch. coli and Salmonella spp. in meats and associated products.

One hundred fifteen samples were collected from a local factory during 1976/1977. The data were analysed to entry the samples and pathogens. t_0° ^{NUn}dred fifteen samples were collected from a local factory outling and pathogens. establish if any relationship existed between total bacterial numbers and pathogens.

No significant correlation existed between total numbers and Esch. coli in the majority of cases. However, significant negative correlations existed for steak-burgers (P < 0.05), frankfurters (P < 0.01) and luncheon coll (P < 0.05) indicating that as total numbers increased, numbers of Esch. coli decreased.

Seven percent of the samples were Salmonella-positive. The mean Esch. coli count of the latter $(\log_{10} 4.51/g)$ was significantly higher (P < 0.01) than the mean Esch. coli count of the Salmonella-negative samples $(\log_{10} 2.01)$ that the mean Esch. coli counts are more likely to be Salmonella-positive. $(\log_{10} 2.89/g)$. This implies that samples that samples of Esch. coli. This implies that samples with high Esch. coli counts are more likely to be Salmonella-positive

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Rapports bacteriologiques dans les viande crues, viandes de conserve et produits associes

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Resume

Une étude a été menée, visant à comparer des informations sur le denombrement total de colonies et la presence d'<u>Esch. coli</u> et de <u>Salmonellae spp</u>. dans les viandes et produits associés.

115 échantillons ont été prélevés d'une usine locale durant les années 1976 et 1977. Les données ont été analysées afin d'établir l'existence de rapports entre le dénombrement bactérier total et des pathogenes. corrélation significative existait entre les dénombrements totaux et d'<u>Esch. coli</u> dans la majorité des ces. Cependant, une corrélation négative significative existait pour les steakburgers (P < 0.05), saucisees de Francfort (P < 0.01), saucises 'luncheon-rolls' (P < 0.01) indiquant une diminution des nombres d'<u>Esch. coli</u> inversement proportionelle a l'accroissement des dénombrements totaux.

Sept pour cent des échantillons étaient salmonelle positifs. Le moyen compte d'Esch. <u>coli</u> de la dernier $(\log_{10} 4.51 \text{ g}^{-1})$ était significativement plus haut (P < 0.01) que le moyen compte d'<u>Esch.</u> <u>coli</u> des salmonelle negatif échantillons (log₁₀ 2.89 g⁻¹). D'ici, les échantillons avec hauts comptes d'<u>Esch.</u> <u>coli</u> auront plus de possibilite d'être salmonella-positif que les échantillons avec les bas comptes d'<u>Esch.</u> <u>coli</u>.

Бактериалогические соотношения в сырон и переработанног чиссе и сопутствующих продуктах.

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Peziore.

Исспедавание сделано для сравнения сведении о сужгарног числе бактерий и о присутствии <u>Esch. coli</u> и <u>salmonella spp.</u> в чиле и сопутствующих продуктах

В техение 1976/1977 года из честного завода собрани 115 образещков. Данные быти акализированы стобы установить мобые соотношения существующие нежду общит числог баштерий и патогенании.

D'éalbunnembe cryraeb de douro grarumentenoù hoppensignu vienegy gywraputour ruarori danmeputi a <u>Esch. whi</u> Bré-manu zharumentenas omputyamentenas hoppensignes cyneembebare que pytrenoù nomrembe (steak-bungers) (P<0.05), warou (frankfurters) (P<0.01) a vischor pyrema (huncheon roll) (p<0.05) yhaz bibas rmo mak kak vorigee ruaro yberwanoce, ruaro <u>Esch. whi</u> grentenurvoce. Cerus nywyennob odpazranob tomu Salmonelle noroskumenstate (reguee ruaro <u>Esch. whi</u> noareguex (hog₁₀ 4,51 g⁻¹) towo zhorumentates boune (P<0.01) cheopiero ruara <u>Esch. whi</u> gue Salmonelle - omputyamensiere otpazrano bere zamo odpazana c becouw ruaror <u>Esch. whi</u> gue Salmonelle - omputyamensiere otpazrano bere zamo samo odpazana c becouw ruaror <u>Esch. whi</u>

Bacteriological relationships in raw and processed meats and associated products

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Introduction

Among the most popular "indicator" tests used to determine the bacteriological quality of a food are the total ^{Agrobic} plate count, [®]coliform[®] count and <u>Esch. coli</u> count (Miskimin et al 1976). However, a clear distinction ^{has} been made between bacteriological quality of food and safety of food by Read and Baer (1974) who stated that standards for quality and safety are quite separate as are the tests for them. Thus the presence of coliforms or large numbers of bacteria in meat have a much different meaning than they would have, for example, in water And milk (Goepfert, 1976) where a single test (Esch. coli) assesses both "cleanliness" or quality and safety in Water (Kitchell, 1967). The presence of Esch. coli in meat does not indicate there are pathogens present. A search of the literature by Goepfert, (1976) failed to reveal any published data of a correlation between Esch. Coli and salmonella organisms in raw beef and Hagberg et al (1973) noted no such relationship in turkey Processing plants. According to Roberts (1976) there is no constant ratio of Esch. coli or coliforms or Enterobacteriaceae to salmonella and salmonella organisms may be found on clean or dirty carcass meat.

However, Tompkin and Kueper (1973) observed that the rate of 'salmonella' isolations increased in rendered ^{Augusta} Tompkin and Kueper (1973) observed that the rate of "salmonella" isolations increased in removed animal by-products from 12% to 70% as the aerobic plate count increased from <10 /g to 10 /g. Furthermore, when the aerobic count of barbacued chicken was $<10^{6}$ /g entercoccci, among other "indicator" organisms, were abaent, but when the plate count increased to 10^{4} , 10^{5} and 10^{6} /g, pathogens were observed in 5%, 9% and 39% of chickens respectively, Seligman and Frank-Blum, 1974.

This study was therefore undertaken to collate information on the total aerobic plate count and the presence of Pathogens (<u>Esch. coli</u> I and <u>Salmonella spp</u>.) in raw and cooked meats and associated products and to establish if any relationship existed between them.

Materials and Methods

Samples.

Samples. Seventy-one samples (group 1) were collected from a local factory between May - September 1976. They were transported to the laboratory in plastic pouches within 1 hr and held at +4°C before examination. Of the 71, 20. ^{2ens}ported to the laboratory in plastic pouches within 1 hr and neid at 74 c before examination of 20 had received some form of heat treatment. To increase the number of samples for statistical analysis, a ^{further} 44 (group 2) were collected between February - May 1977 in the same manner. None of these had received ^a heat treatment. They were frozen (-18°C) and were between 16 and 22 days old when examined, Table 1.

(b) Total aerobic count

Ten gram aliquots of each sample (group 1) were weighed and transferred to 100 ml Ringers diluent +0.1% of added Peptone in a plastic pouch. The sample was homogenised for 60 sec. in a Colworth Stomacher 400 (A.J. Seward, Bury St. Edmunds, Suffolk, England). Serial decimal dilutions were made and plated in duplicate using a 1 ml "Oxformation 'Oxford' $v_{xf}^{\text{Ord}^{\circ}}$ sampler pipette with a disposal tip by the method of Bousfield, Smith and Trueman (1973). Pre-dried Plates of Oxoid Plate Count Agar (PCA) were divided into quarters and inoculated with replicate 0.025 ml amounts (w_{y}/v_{y}) of added salt (NaCl) was used in the count medium.

(c) <u>Escherichia</u> coli

This was estimated by membrane filtration. One millilitre of the homogenate described above was serially diluted and filtered by the method described by Dempster et al (1973). Confirmation of type I Esch. coli was obtained by growing typical lactose-fermenting colonies in peptone water and MacConkey broth (18h/44°C) since indel. Indole production in the former and acid+gas in the latter are indicative of Esch. coli faecal type I (Howe and Linton 1976).

(d) Salmonella

¹⁰ Salmonella Fifty grams of sample were weighed and aseptically combined with 100 ml Ringer's diluent + 0.1% peptone in a plastic pouch and homogenised as described above. One hundred millilitres of double strength Mannitol Selenite broth (Oxoid Cm 39a) were added, thoroughly mixed and the total volume (200 ml) of single strength broth transferred to a sterile screw capped jar. Incubation was for 48h at 41°C. Another 50g sample was similarly homogenised in Ringer's diluent and 100 ml double strength tetrathionate broth (CM 29) added and transferred to a similar jar for incubation at 37°C for 48h. A loopful of culture from each bottle was streaked after incubation on to Brilliant Green Agar plates (Oxoid CM 263) and the plates incubated at 37°C for 24h. Suspect colonies were picked into lysine decarboxylase medium (Moeller 1955) and streaked on to freshly poured PCA plates (Oxoid) for biochemical and serological recognition of salmonelles by the procedure of Georgala and plates were picked into lysine decarboxylase medium (Moerier 1955) and streamed on to receive of Georgala and Books (Oxoid) for biochemical and serological recognition of salmonellae by the procedure of Georgala and Boothroyd (1969).

Group 2 samples were thawed for 24h at room temperature (C.18°C) and examined for total numbers, Esch. coli and ^{salmone}lla in the same fashion as Group 1 samples.

(e) Statistical analysis

The Correlations between total count and Esch, coli count were calculated for each set of data and for all the

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data. The mean Esch. coli count was calculated for samples with salmonella present and for samples with salmonella absent.

Results and Discussion

Group I samples

Of 71 products, 24 (34%) contained less than 20 Esch. coli/g and 13 (18%) less than 1×10^6 /g (total aerobic count). Only 4 (5.6%) samples were salmonella-positive, these were chicken/ham mix (2), pork sausage (1) and pork streak (1) Table 2.

Six out of 8 samples of luncheon roll (75%) and 9/20 (45%) pork sausage samples had less than 20 Esch. <u>coll/9</u>, whereas 4/6 (66%) chicken/ham; 4/4 (100%) pork skin, 7/20 (35%) pork sausage and 4/5 (80%) samples of rew beer contained more than 1,000 <u>Esch</u>. <u>coli</u>/9, despite the fact that some of these products had been heat-treated. Possibly this high incidence was a result of improper handling post-heating, inadequate storage or in the case of unheated samples, to a carry-over contamination effect from the slaughter hall.

Only one sample (pork streak) exhibited organoleptic spoilage at examination, i.e., it had an off-odour and the meat was discoloured. Characteristically, the total aerobic count was 47×10^{-1} /g. Perhaps surprisingly it contained only 10 Esch. coll/g and it was salmonella-negative despite the generally held opinion that a high total viable count in a food is indicative of a greater risk of pathogens being present (Miskimin et al. 1976). One other "pork streak" was salmonella-positive (Salm. panama) and had an Each. coli count of 90/g and a total count of 182 x 10⁻⁴/g. The remaining four samples of pork streak were salmonella-negative and had total count of 182 x 10⁻⁴/g. The remaining four samples of pork streak were salmonella-negative and had total count of 182 x 10⁻⁴/g. The remaining four samples of pork streak were salmonella-negative and had total count of 182 x 10⁻⁴/g. The remaining four samples of pork streak were salmonella-negative and had total count of 182 x 10⁻⁴/g. The remaining four samples of pork streak were salmonella-negative and had total count of 182 x 10⁻⁴/g. The remaining four samples of pork streak were salmonella-negative and had total count of 182 x 10⁻⁴/g. The remaining four samples of pork streak were salmonella-negative and had total count of 182 x 10⁻⁴/g. The remaining four samples of pork streak were salmonella-negative and had total count of 182 x 10⁻⁴/g. The remaining four samples of pork streak were salmonella-negative and had total count of 182 x 10⁻⁴/g. The remaining four samples of pork streak were salmonella-negative and had total counts and the stated ... "there is no constant ratio of Esch. coli or coliforms or Enterobacteriaceae to count indicates safety has been shown to be not always true. Furthermore, the multiplicity of factors involved make generalisations on the value of total counts virtually impossible according to Silliker (1962). In feetimany workers have stated that total numbers of organisms are unrelated to the presence or absence of pa

In the majority of cases, no correlation existed between total viable count and <u>Esch. coli</u> count (Table 3), only two cases was there a negative correlation (steakburgers P < 0.05 and frankfurters P < 0.01) indicating that as the total count increased, so the <u>Esch. coli</u> count decreased. The correlation was non-significant (R = 0.13) when the data were pooled. Table 1. Type and number of products examined and heat treatments applied

Braduat		
Product	n	Heat Treatment
Chickes /Hen six		
	0	
Pork sausage	20	
Steakburger	7	
Raw beef	5	
Pork lean	4	
Pork skin	4	Cooked at 185 [°] F until pliable, then passed through colloid mill and filled into shallow trays for storage at 4 [°] C
Plasma ²	3	and the second of the second
Luncheon roll	8	$2\frac{1}{2}$ hr/165°F
Pork streak ³	6	
Frankfurter	5	$2\frac{1}{2}$ hr/165° + 30 min/165°F
White pudding	3	45 min/196 ⁰ F
Total	71	
Group 2 (Held 16 - 22	2_days/-18 ⁰ C	before testing) 1 = Shoulder meat
Chicken/Ham mix	5	2 = Equal amounts of lard + ear scalpo + 40 kg dehvdrated blood.
Chicken/Ham mix	17	- 3 = Cut from pork carcase 2 days post-mortem
Chicken/Ham mix	11	a day it on port carcade 2 days por
Luncheon roll	11	-
Total	44	

Group 1 (Held 1 hr/4°C before testing)

press.	-		-
L			6
		-	
	-	-	-

Table 2.	Total aerobic count	and specific	pathogens	in pork,	beef	and	associated	products	(Group	l samples)	
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Sample n		Total count g					coli 1 g ⁻¹		Salmonella			
		< 10 ⁶	>10 ⁶ <10 ⁷	>10 ⁷	>10 ⁸	< 20	20-10 ²	>10 ²	>10 ³			_
Chicken/Ham mix	6	-	2	3	1	2	-	-	4	Salm.	newport	(2)
ork sausage	20	3	2	6	9	9	-	4	7	Salm.	newport	: (1)
Steakburger	7	-	-	4	3	3	-	2	2			
Raw beef	5	1	-	1	3	1	-	-	4			
Pork lean	4	1	1	1	1	-	2	1	1			
Pork skin	4	-	2	2	-	-	-	-	4			
Plasma	3	1	1	1	-	-	-	2	1			
uncheon roll	8	3	1	3	1	6	-	-	2			
^p ork streak	6	2	1	-	3*	1**	2	2	1	Salm.	panama	(1)
Frankfurter	5	1	-	1	3	2	1	1	1			
White pudding	3	1	1	-	1	-	-	1	2			
Total	71	13	11	22	25	24	5	13	29	1.2.3	4	
% Total	100	18.3	15.5	31.0	35.2	33.8	7.0	18.3	40.8		5.6	

One sample had 47 x 10¹⁰ g⁻¹ total count **

10 Esch. coli g

Table 3. Correlation between Esch. coli count and total aerobic count $(\log_{10} g^{-1})$ of 71 samples (Group 1)

Sample	n	Correlation coefficient	Significance level
Chicken/Ham mix Pork sausage Steakburger Raw beef Pork lean Pork skin plasma Luncheon roll Pork atreak Frankfurter white pudding All data	6 20 7 5 4 4 3 8 6 5 3 71	-0.66 0.43 -0.78 0.73 0.82 0.65 0.23 0.55 -0.28 -0.96 0.76 0.13	N.S. N.S. 5% N.S. N.S. N.S. N.S. N.S. 1% N.S. N.S.

Group 2 sample

The sample of the sample of the sample of the sample of the second seco Pooled, Table 5.

05

For both groups 1 and 2, the mean Esch. coli count $(\log_{10} 4.51/g)$ for the eight salmonella-positive samples was $\frac{\log_{10} \log_{10} \log_{10} 2.89/g}{\log_{10} 2.89/g}$ for the salmonella-negative $\frac{\log_{10} 2.89/g}{\log_{10} 3.0/g}$ are more likely $\frac{\log_{10} 2.09}{\log_{10} 3.0/g}$ are more likely $\log_{10} 3.0/g$

1 9 Table 4 Total aerobic count and specific pathogens in processed meats (Group 2 samples)

Sample	n	<10 ⁶	Total Count ≻10 [°] <10 [′]	s g ⁻¹ >10 ⁷	>10 ⁸	<20	Each. coli 20 - 10 ²	1 g ⁻¹ >10 ²	>10 ³	Salmonella
Chicken/Ham mix	5	-	-	1	4	-	-	-	5	Salm. typhi murium (1)
Chicken/Ham mix	17	-	-	6	11	-	-	-	17	Salm. derby
Chicken/Ham mix	11	-	-	6	5	-	-	-	11	Salm. typh: murium (1)
Luncheon roll	11	-	-	5	6	-	-	4	7	-
Total	44	-	-	18	26	-	-	4	40	4
% Total	100	-	-	41.0	59,0			9.1	90,9	9.0

Table 5. Correlation between Esch. coli count and total aerobic count $(\log_{10} g^{-1})$ of 44 samples (Group 2)

n	Correlation coefficient	Significance level
5	-0,08	N.S.
17	0,17	N.S.
11	0,37	N.S.
11	-0,61	5%
44	0.05	N.S.
	n 5 17 11 11 44	n Correlation coefficient 5 -0.08 17 0.17 11 0.37 11 -0.61 44 0.05

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