Title	STUDIES ON CHICK SALMONELLOSIS : II. SALMONELLA SENFTENBERG INFECTION IN CHICKS
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STUDIES ON CHICK SALMONELLOSIS

II. SALMONELLA SENFTENBERG INFECTION IN CHICKS

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The writers^{2,6,13)} have already reported that in Japan not only S. pullorum but also S. senftenberg, S. thompson, S. bareilly and S. new brunswick respectively play a part in chick salmonellosis. Moreover S. montevideo, S. paratyphi B, S. typhi murium, S. cholerae suis, S. potsdam, S. enteritidis, S. give and S. newington in poultry have been detected by IWAMORI and SHIMAKURA, OCHI et al.^{10,11)} and MIURA et al.

However, as S. senftenberg among these salmonella organisms, except S. pullorum, seems to infect chicks the most frequently, some experiments to ascertain the mode of infection and the influence of S. senftenberg on baby chicks were undertaken.

Some parts of these studies were reported at the Symposium⁴⁾ of "Enteric Organisms of Domestic Animals" of the 35th Meeting (1953) and at the 39th Meeting (1955)⁵⁾ of the Japanese Society of Veterinary Science.

On the Mode of S. SENFTENBERG Infection in Newly Hatched Chicks

As reported previously, though both newly hatched chicks and dead-in-shell-chicks came from the same incubator with a difference of only $3\sim5$ days, the rate of S. senftenberg infection in newly hatched chicks which died was always superior to that in dead-in-shell-chicks^{2,4}). In systematic bacteriological investigation on baby chicks which died of a pullorum-like disease, chicks under $5\sim10$ days old showed the highest rate of infection with S. senftenberg. Though some positive reactors to the rapid agglutination test for S. senftenberg infection were discovered in field work, the detection of S. senftenberg in these reactors' bodies resulted negative with one exception of a 3 months old chick^{3,4}). S. senftenberg has not yet been found in any egg laid by reactors positive to the agglutination test⁴).

When the above noted results were taken into consideration, it was presumed that the mode of S. senftenberg infection to eggs and baby chicks is different from that of S. pullorum infection, and that baby chicks may have the most frequent opportunities

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TABLE 1. Results in T and Ts

							-1.			t CO W	10 111 1		, 10
												DATE	OF
POINT OF OBSER	VATIO	N					T 3	Hatche	ry				
			14/I	24	2/ II	5	13	17	19	23	2/III	7	13
Frequency of De Salmonella Organ			Origin	ated	from	A I	ncu.	В		C			D
Newly Hatched						P.3		P.3		0			s.
Killed Immediate Hatching*	ely art	er	•	•	•	10	•	10	•	10	•	•	10
Minutes for Expo Agar Plate in I	osure o	f SS or	80	80	80	80	80	80	80	80	70	50	30
M N C. C.		A	0	0	0	0	•	•	0	•	•	0	•
Mean No. of Sanella Organisma Detected in Hate	s	В	0	•	0	0	0	0	0	0	0	0	0
Compartment of Incubator		С	0	•	•	0	0	•	0	0	0	0	0
		D	0	•	•	•	•	•	•	0	0	0	0
			Origin	ated	from	ΑI	neu.	В		C		A~D)
Salmonella (S	Surface					0	•	P.2	•	0	•	P.3	
Organisms	Embry	70				30		30		30		136	
~	Yolk-Sa	ıc	•	•	•	0	•	0	•	0	•	P.2	•
Chicks*	Surface & Yolk		•	•	•	P.1	•	P.2	•	0	•	P.4	•

^{*} The denominator: the number of samples examined, the numerator: the number Notes: $S \cdots S$. senftenberg. $P \cdots S$. pullorum. $B \cdots S$. bareilly.

to be infected with S. senftenberg in an incubator. Therefore, some experiments to prove these presumptions were planned as follows. That is, in T, Ts and F hatcheries in Sapporo, before the beginning of hatching the insides of all 12 incubators were thoroughly cleaned up, disinfected with formaldehyde gas and then were confirmed to be free from salmonella organisms. With the beginning of hatching the conditions of contamination with S. senftenberg in hatching compartments of incubators, dead-in-shell-chicks and newly hatched chicks were investigated bacteriologically at frequent intervals.

For detection of S. senftenberg in incubator hatching compartments 4~9 Endo or SS agar plates were used and cultured for 24~48 hours at 37°C after being exposed in hatching cabinets for 80~10 minutes. All surfaces of egg-shells, some amount of egg-white on embryos and some amount of egg-yolk of embryos in regard to dead-in-shell-chicks, and, pieces of respiratory, digestive and other organs of baby chicks were respectively examined

Hatch	prine	n	7953

НАТС	HING IN	1											
							,	Гs На	tcher	у			
19	25	31	1/V	24/11	5/III	10	17	26	2/IV	8	11	17	21
	A~D												
•	S. 16 20	eri neu ritug diffusel i film di fil	•	•	•	•	5	P. 2	•	•	S.9 10	S.1 10	$\begin{pmatrix} S.3 \\ P.3 \\ \hline 6 \end{pmatrix}$
15	10	10	10	80	80	80	80	80	80	60	30	30	10
•	S.23	S.23	(S.18 P.0.5	0	0	0	0	0	•	S.0.3	S · 0.3	S. 0.3 P. 0.3	S.7
•	(S.74 P.0.2	S.92	S.19										
0	$(^{S.53}_{P.0.3}$	S.62	S.30										
S.1	S.58	S.34	S. 4										
A ~ D	$A \sim D$	A ~ D											
S.4	S.5	B. 2					0		0		0	P.2	S.1
116	106	78	•	•	•	•	64	•	41	.•	42	60	62
$\begin{pmatrix} S \cdot 1 \\ P \cdot 2 \end{pmatrix}$	S.5	S.2	•	•	•	•	0	•	0	•	0	P.1	P.3
$\left(egin{matrix} $	$\backslash P. 1$	${\scriptsize \begin{pmatrix} S.6\\P.1\\B.1\end{pmatrix}}$	•	•	•	•	0	•	P.3	•	0	P.3	P.4

of Salmonella organisms positive cases.

by direct and enrichment culture methods.

Results obtained are tabulated as tables 1, 2 and 3.

In 1953, materials originating from T hatchery equipped with 4 incubators and Ts hatchery with 1 incubator were inspected. As shown in table 1, in T hatchery S. senftenberg was suddenly found first in D incubator on the 24th day (on 19, March) after the beginning of hatching (incubator A which began to hatch at first had already worked for 63 days up to this time), and after 6 days S. senftenberg could be found not only in D but also in A, B and C incubators. Afterwards S. senftenberg was found out in all incubators in the following inspections.

On the other hand, in regard to dead-in-shell-chicks, on that day when S. senftenberg was first detected in D incubator, S. senftenberg had already been found out from dead-in-shell-chicks originating not only from D but also from each of the other incubators.

TABLE 2. Results in T Hatchery in 1954

	_									DATE	OF HAT	CHING					
POINT OF OBSE	RVATION		19/I	30	6/ II	11 ·	13	17	23	1/III	6	12	18	24	30	24/V	29
Frequency of			Origin	ated	from	B.C I	ncu.	B.C		A~D		A~D		A~D			A~D
of Salmonella (from Newly Ha	atched					P.2		P.1		S.2 P.2		S.5 B.4		S.7 B.3			S.5 T.2
Chicks Killed In after Hatching		ıy	•	•	•	17	•	12	•	15	• 	10	·	7			5
Minutes for Ex SS Agar Plate i	posure on Incubat	of tor	60	60	60	40	30	30	30	20	15	15	15	15	15	15	15
	{	A	•	0	0	0	0	0	. 0	0	$\begin{pmatrix} \mathbf{S} \cdot 1 \\ \mathbf{P} \cdot 0.3 \\ \mathbf{B} \cdot 0.5 \end{pmatrix}$	(S.38 B.2.5	(S.6 B.2.5	$\begin{pmatrix} \mathbf{S.12} \\ \mathbf{B.} & 1 \end{pmatrix}$	(S.10.5 B. 2	(S. 2.5 B. 0.5	(S. 31 B. 0.5
Mean No. of nella Organism	ns	В	0	0	0	0	0	0	0	0	$\begin{pmatrix} S.3 \\ P.0.3 \\ B.1.3 \end{pmatrix}$	(S.8 B.1	$\begin{pmatrix} S.6 \\ P.0.5 \\ B.0.5 \end{pmatrix}$	(S. 8 B. 1	$\binom{\text{S.11.5}}{\text{P. 3.5}}_{\text{B.11.5}}$	(S.74 B.45	(S.37. B. 1.
Detected in Ha Compartment of Incubator		C	0	0	0	0	0	0	0	S.0.3	(P.0.8	(S.9.8 B.0.5	(S. 65 B. 2.5	S.14	(S.24.5 B. 9.5	(S.31 B.1.8	(S.6 B.1.8
	{	D	•	•	0	0	0	0	0	0	(S.8 P.0.3	(S. 2.3 B. 2.8	S. 5.5 P. 0.5 B. 0.5	(S.82.5 B. 1.5	(S.12.5 B. 6	$\begin{pmatrix} \mathbf{S.8} \\ \mathbf{B.3} \end{pmatrix}$	$\begin{pmatrix} S.4 \\ B.5. \end{pmatrix}$
			Origin	ated	from	B.C	ncu.		A~D	A~D	A~D	A~D	A~D	A~D	A~D	A~D	A~D
	Egg-She	11	•	•	•	•	•	•	•	•	•	•	•	•	•	S. 22 B. 2	S. 12 B. 12
Salmonella		_														30	30
Organisms Detected from	Surface Embry		•	•	•	0/46	•	•	0/48	0/44	0/47	0/50	0/50	S. 1/50	0/47	S.1	0
Dead-in-Shell-	Yolk-Sa	c	•	•	•	0	•	•	0	0	P.2	0	$\begin{pmatrix} \mathbf{P.4} \\ \mathbf{B.1} \end{pmatrix}$	B.1	S.1	0	0
Chicks	Surface &Yolk		•	•		0	•	•	0	0	P. 1	P.1	$\binom{S.1}{P.2}$	S.2	P.1	$\begin{pmatrix} S.1 \\ P.1 \end{pmatrix}$	${ m S.2} m_{P.1} m_{B.1}$

Note: T...S. thompson.

Afterwards S. senftenberg was always found out from some materials originated from all incubators.

As concerned with newly hatched chicks, S. senftenberg had already been detected from 5 of 10 chicks hatched in D incubator only on 13, March, namely before 19, March on which day S. senftenberg was first found in D incubator and dead-in-shell-chicks originated from each incubator.

That is, in T hatchery, S. senftenberg was found out in newly hatched chicks prior to the finding in incubator or dead-in-shell-chicks. However on the other hand, in Ts hatchery the first detection of S. senftenberg was in an incubator on 44th day after beginning of hatching, and then in newly hatched chicks and later in dead-in-shell-chicks. (It is much to be regretted that the chicks hatched on 2 and 8, April, and the air of hatching compartment of incubator on 2, April were not inspected.)

Furthermore in successive inspections in T hatchery equipped with 4 incubators and F hatchery with 3 incubators in 1954, as shown in table 2, S. senftenberg in T hatchery was suddenly found out in C incubator as well as in newly hatched chicks on 1, March, namely on 42nd day after beginning of hatching. Detection of S. senftenberg in dead-in-shell-chicks was about 17 days later.

As shown in table 3, first detection of *S. senftenberg* in F hatchery was on egg-shells of dead-in-shell-chicks as well as in A incubator on 15, March, on the 45th day after starting to hatch, but in newly hatched chicks *S. senftenberg* was found out in 6 of 20 chicks 5 days later. (Unfortunately, chicks hatched on 15, March were not examined.)

On the ground of the above described results, it may be said that in the incubator S. senftenberg in the hatching compartment begins to be found if it continues to operate for about 50 days. And if S. senftenberg is found out in an incubator once, it is usually detected from that incubator successively onward from that time. And soon after S. senftenberg came to be found out in any incubator once, it will certainly be found in all incubators operating in the same room. This phenomenon is presumed result in following facts: when the leaf of the incubator in which S. senftenberg had multiplied, was opened to inspect the incubated eggs, S. senftenberg flowed out and mixed in the air of the incubator room and then invaded other incubators. In fact it was certified that the number of S. senftenberg which multiplied in an incubator decreases by opening the leaf of that incubator, but if the leaf of that incubator is closed and the incubator is continued to hatch as before, S. senftenberg again multiplies in it. On the other hand, at that time when the leaf of incubator with multiplied S. senftenberg was opened, in bacteriological examination of the air of the room containing such incubator, S. senftenberg was found out. And in a more severe case S. senftenberg comes to be found out too in the stockroom of hatching eggs separated from the incubator room.

One more important cause of transmission of S. senftenberg from one incubator to another is directly traceable to workers in hatchery. They infect their fingers, test-lamp etc. with S. senftenberg unintentionally by touching the infected cabinets, eggs, chicks etc. at first, and then transmit S. senftenberg to the cabinets, eggs, chicks etc. of other steriled incubators by retouching with their dirty fingers or instruments.

In spite of absolute non-finding of S. senftenberg in chicks hatched during the term

TARIE 3 Results in F

						T	ABLI	€ 3.	Resu	ts in F
DOWN OF COL									DA	TE OF
POINT OF OBSE	ERVATION	29/1	5/11	8	15	19	25	3/III	10	15
Frequency of I	Detection of	Orig	inated	from		A Ir	ıcu.	A.B	A.B	
Salmonella Orga Newly Hatched Killed Immediat	d Chicks					0		0	P.1	•
Hatching	cery arter	•		·	·	22		20	20	,
Minutes for Exp Agar Plate in I		80	80	80	60	60	40	30	15	15
Mean No. of Sa	1	0	0	0	0	0	0	P.3	0	S. 1.3 P. 4
Organisms Det Hatching Comp	partment F	3 .	0	0	0	0	0	0	0	0
of each Incubat	tor		•	0	0	0	0	0	0	0
		Origin	ated from_	A Inc	u. A	A.B	A.B	A.B	A.B	A.B.C
	(Egg-Shell	_		0_	0	0	0	0	0	S.2
Salmonella Organisms		•	•	10	10	66	68	65	70	80
Detected from	Surface of Embryo	•	•	0	0	0	0	0	0	0
Dead-in-Shell-	Yolk-Sac	•	•	0	0	0	0	0	0	0
Chicks	Surface & Yolk	•	•	0	0	0	0	0	0	P.2

which S. senftenberg is never detected in any incubator, if S. senftenberg once begins to be found out in an incubator, immediately before or almost at the same time S. senftenberg comes to be detected in newly hatched chicks originated from such infected incubator too. Moreover afterwards, it is successively detected in day-old chicks as well as in the incubators.

On the ground of the above mentioned observations, one of the important causes of S. senftenberg infection in newly hatched chicks is presumed to exist in an incubator. That is, it may be concluded that, in regard to the S. senftenberg infection in newly hatched chicks, many baby chicks are infected with S. senftenberg through their respiratory and digestive organs in the incubators in parallel with the grade of contamination of hatching compartments of incubators.

On the Appearance of S. SENFTENBERG in an Incubator

Though the bacteriological examinations were carried out under the presumption that hatching eggs may bring S. senftenberg into an incubator, as shown in tables 1, 2 and 3,

Hatcheru	in	1954

HATCHIN	īG								
20	26	1/ IV	6	13	19	26	2/V	7	13
A.B.C		A.B.C	A.B.C		A.B.C				
(S.6 P.6	_	(S. 15 P. 12	(S.3 P.1		S.5				
20	•	20	7	•	5	•	•	•	•
15	15	15	15	15	15	15	15	15	15
(S. 0.5 P. 0.5	P. 0.3	P. 3.3	P.8	S.3	(S.17 P. 2	$\left(egin{matrix} ext{S.3} \\ ext{P.0.3} \end{aligned} \right)$	$\left(egin{smallmatrix} ext{S. 15} \\ ext{P. 0.8} \end{matrix} ight)$	S. 0.5	S. 6.5
0	P. 0.8			S.7	(S. 18 P. 1.5	S.1	S.2	(S. 2.5 P. 1	S.18
P. 0.3	0	$\left(\begin{smallmatrix} \mathrm{S.0.3} \\ \mathrm{P.0.3} \end{smallmatrix} \right)$	$\begin{pmatrix} S. 0.8 \\ P. 0.8 \end{pmatrix}$	S.2	$\begin{pmatrix} S.2 \\ P.1 \end{pmatrix}$	S. 28.7	S.8	S. 2.2	$\left(\begin{smallmatrix} \mathrm{S} \cdot 12 \\ \mathrm{P} \cdot 0.8 \end{smallmatrix} \right)$
A.B.C	A.B.C	A.B.C	•	A.B.C	A.B.C	A.B.C	A.B.C	A.B.C	A.B.C
S.5	(S.37 P. 2	(S.7 P.7		S. 13	S. 37	S. 67	S. 60	S.8	S. 12
79	75	77	·	73	7 5	78	64	40	38
0	0	0	•	0	0	0	0	S.1	0
0	0	0	•	S.1	0	0	. 0	S.1	0
0	0	P. 1	•	0	P.4	$\begin{pmatrix} \mathrm{S.2} \\ \mathrm{P.1} \end{pmatrix}$	S.1	$\left(egin{smallmatrix} ext{S.2} ext{P.1} \end{aligned} ight.$	S.1

S. senftenberg has not yet been detected in dead-in-shell-chicks originated from the incubator in which S. senftenberg had not yet been found out, but if S. senftenberg once appears in an incubator, it also comes to be detected in dead-in-shell-chicks originated from that incubator. And in field work, when slide tests or tube tests for detection of S. senftenberg infection were used, some positive reactors are often discovered. However, S. senftenberg had not yet been detected from egg-shell, egg-white and yolk of eggs laid by such positive reactors⁴, and the writers could not yet find out S. senftenberg in any part of bodies of positive reactors except in under 3 months old chicks^{3,4}).

If one takes into consideration the above described results, he finds that there seems to be a difference between the manner of *S. senftenberg* infection and that of *S. pullorum* infection in adult fowls. Therefore there is a tendency to deny that *S. senftenberg* as well as *S. pullorum* is brought into an incubator through the egg. However, one must not forget the reports introduced by IWAMORI and SHIMAKURA and OKAZAKI et al. That is, as IWAMORI and SHIMAKURA (1954) reported that they had found *S. senftenberg* in an unabsorbed yolk-sac of a 1-year old hen and OKAZAKI et al. (1956) reported that *S. senftenberg* had been detected in ovary, oviduct and left kidney of a 1.5-year old hen, the

transmission of S. senftenberg to egg can not be denied completely. Therefore there seems to be need for further and more careful investigations to ascertain how S. senftenberg is brought into the hatching compartment of an incubator.

On the ones with flawless egg-shell among dead-in-shell-chicks infected with S. pullorum only or S. senftenberg only, the frequency of existence of salmonella organisms on surfaces of embryos and in yolk-sacs drawn into abdominal cavities of embryos was investigated. The results are shown in table 4.

TABLE 4. Frequency of Detection of Salmonella Organisms from Surfaces of Embryos and Yolk-Sacs Drawn into Abdominal Cavities of Embryos in Dead-in-Shell-Chicks

	ON SURFACES			ON SURFACE IN YOLK-SACS		
ТҮРЕ	ONLY OF EMBRYOS	IN YOLK- SACS ONLY	Nos. on Surfaces >Nos. in Yolk-Sacs	Nos. on Surfaces = Nos. in Yolk-Sacs	Nos. on Surfaces <nos. in<br="">Yolk-Sacs</nos.>	TOTAL
S. pullorum	5	17	2	24	5	53
S. senftenberg	12	12	8	20	3	55

The data show that there is a tendency for the rate of infection of surfaces of embryos with S. senftenberg to be higher than that with S. pullorum. And the possibility of penetrating egg-shell of S. senftenberg was experimentally certified by the senior author³. If the assumption that S. senftenberg can penetrate egg-shell in incubators operating with normal temperature and humidity is allowed, taking into consideration the above mentioned results, it may be said that the egg itself will more likely be attacked by S. senftenberg in an incubator than the staining of an incubator with S. senftenberg. However, on the other hand, as shown in table 4, S. senftenberg also is detected in yolk-sacs only of some dead-in-shell-chicks as well as S. pullorum, though the frequency of detection of it is inferior to that of S. pullorum. This fact may suggest that S. senftenberg has an ability to penetrate into a yolk-sac in adult hen's body as well as S. pullorum.

Anyhow, it is sure that *S. senftenberg* can grow and multiply in the hatching compartments of incubators working with normal temperature and humidity; however, it can not yet be proved how *S. senftenberg* is brought into the hatching cabinet of an incubator or the hatchery premises. It is necessary to pay attentions to rats, wandering birds, flies, feeds and visitors with regard to such bringing of *S. senftenberg* into a hatchery premises. Especially rats and over-wintered flies may be one of the important infective origins in the succeeding year. But

the possibility that S. senftenberg is brought into hatching compartments of incubators by egg should also not be neglected.

As shown in tables 1, 2 and especially 3, it was proved that *S. pullorum* also comes to appear in the hatching compartments of incubators with the repetition of hatching. Therefore the inhalation-infection of *S. pullorum* among newly hatched chicks in an incubator may also occur as well as in *S. senftenberg* infection. But the multiplication of *S. pullorum* in the hatching compartments of incubators being operated under the normal temperature and humidity is not so remarkable as in *S. senftenberg*.

Especially as shown in table 2, S. bareilly too appears in the hatching compartments of incubators.

All species of these salmonella organisms which appear in incubators are effectively destroyed by the fumigation of formaldehyde gas. As Frank and Wright reported, when use was made of 1.5 ml of formalin for every cubic foot of incubator space all salmonella organisms were completely killed within 30 minutes.

Nowadays in some special hatcheries paying their attention to hygienic methods of poultry raising, the raisers are preventing the infection with S senftenberg and others of newly hatched chicks in incubators by the periodic disinfection of the hatching compartments using formalin at intervals of $30\sim40$ days after the beginning of hatching.

On the Presence or Disappearance of S. SENFTENBERG in Chicks' Bodies after Infection

In the former chapter, it was stated that an incubator plays an important role in S. senftenberg infection in newly hatched chicks. Table 5 explains the relation of S. senftenberg infection between the incubators and the newly hatched chicks in more detail.

As shown in this table, soon after the beginning of first hatching in T hatchery, the distribution of S. senftenberg in baby chicks' bodies killed immediately after hatching as well as the appearance of S. senftenberg in incubators were investigated. S. senftenberg was not detected in chicks' bodies hatched on 11th and 17th February when no S. senftenberg was found in the hatching compartments of the incubators, but the frequency of detection of S. senftenberg in baby chicks' bodies hatched after 1st March when S senftenberg began to be found out in one incubator increased in parallel with the multiplication of S. senftenberg in the incubators. Further, it is interesting that the frequency of detection of S. senftenberg from digestive organs of newly hatched chicks was higher than that from respiratory organs. This phenomenon is not considered to have resulted from the absorbing S. senftenberg contained in yolk-sac before hatching. S. senftenberg which have invaded into the blood stream from respiratory organs may rather settle and multiply in digestive organs, or it may settle in digestive organs as effects of pulling feather, picking vent

Table 5. Appearance of Salmonella Organisms in Newly Hatched Chicks' Bodies in Relative to Presence and Multiplication of Salmonella Organisms in Hatching Compartments of Incubators in T Hatchery

																D	AT:	E C	F	HA	TC	н	NG	;												
POINT OF OBSERVATI	ION		11/	II '	'54		17/I	I			-	1/	/11	[•					12	/IIJ	[24	/II	I		
No. of Chicks Kille Immediately after Hatching	d	1	5	6	12	13 } 17	1		7	6	7 ≀ 9	10	11	12	13	14	15	1	. 2	2 :	3	4	5	6	7	8		9 10) :	1	2	3	4	5	6	7
{Trachea {Lung		•	р		:	•	p •	•	•	:	•		:		•	•	•	•	•	•	,	•	•	•	s •			•	ķ	3	s S	•	•	s •	•	s S
Heart Blood Liver Spleen Kidney Testicle Egg-Yolk Esophagus Crop Proventriculus				•	; p			•	•	•		P P	•	•	•	p	•	•	•	•		•	•	•	•	· s	•	•		3	S · · · ·	•	•	•	s . S	S s
Esophagus Crop Proventriculus Gizzard		•	•	•	•	•	•	•	•	s	:	:		•	•	•	•	•	•	•		3	•	Š s	•	s s.	b :	•	S	3	S S s	s :	s		s	S s
Small Intesting Caecum Rectum	e	•	•	:	:	•	· ·	•	•	s s	:	•	•	•	•	•	•	b b	b	•		•	• •	• b	•	:	•	•	20707	5.b 5	S.t S.t	s S	s	s S	s •	S.B S.B S.B
Mean No. of Sal- monella Organisms	$\begin{pmatrix} A \\ B \end{pmatrix}$	-		0			0						()								((B	. 38 . 2	.5						,	B	. 12	l 3		
Detected in Hatchery Compartment of	$\begin{pmatrix} 1 & 1 & 1 \\ 1 & 1 & 1 \end{pmatrix}$			0			0					5) 0. 8									(SB	. 1	.8 .5							S	. 14	Į		
Each Incubator	(D			0			0)								(В	. 2	.8							B	. 82		_	
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and pecking feces or spicules of fluff contaminated with that organism before feeding. Besides there is a possibility of thrusting S. senftenberg into bowels of newly hatched chicks by the using of non-disinfected chick sexing instrument.

Then investigations were carried out to ascertain how S. senftenberg settled in respiratory or digestive organs of a baby chick disappears as a chick grows on healthily. That is, studies were made of the distribution of S. senftenberg in bodies of $2\sim5$ apparently healthy chicks killed immediately after hatching or killed at one-day intervals or killed at regular intervals after brooding. These investigated chicks were sampled at random from 2 groups of male White Leghorns originated from F or T hatchery. All materials examined were cultivated on media directly and for enrichment. The regions of chick's body inspected and the results obtained are shown in tables 6 and 7.

It was proved that the frequency of detection of *S. senftenberg* is especially high in respiratory and digestive organs in both F and T groups. In F group, the frequency of detection of *S. senftenberg* in baby chicks less than about 9 days old is considerably high but no *S. senftenberg* came to be found out in bodies of chicks sacrificed when more than 25 days old. On the other hand, in baby chicks under 3 days old of T group, *S. senftenberg* was always detected with considerable frequency; in chicks' bodies of 5 to 30 days of age it was seldom detected; in over 40 days old chicks it came to be not found out at all.

In brief, it was ascertained through the results obtained on baby chicks of F and T groups that S. senflenberg invaded into newly hatched chicks' bodies in incubator was detectable with considerable frequency during the first about 7 days and afterwards, though the frequency of detection decreased, S. senftenberg itself was carried for 20~30 days, and it tended to lurk in the digestive and respiratory organs especially.

It was concluded that, even in the newly hatched chicks infected with S. senftenberg in incubator, so far as they were hatched in good health and maintained their health during rearing under good feeding, management and sanitation, most of the chicks usually overcame S. senftenberg and grew up soundly without being influenced by S. senftenberg.

When such chicks grew up to more than 40 days old, they generally showed no reaction to S. senftenberg in rapid whole blood agglutination test.

The fact that, even if the baby chicks in first about 7 days of age looked to be completely healthy, they always carry S. senftenberg in considerable numbers is in close agreement with the writers' previous report¹³ that, when large numbers of baby chicks are brooded by poultry raisers in field, the outbreak of a sickness caused by S. senftenberg brings the greatest loses in chicks under $5\sim10$ days of age to the raisers. That is, it is not so difficult to presume that, in the newly hatched chicks that had weak constitution or are brooded under poor management and sanitation, S. senftenberg which invaded into chick's body in incubator comes to be active and multiplies to result in death by septicemia within the first $5\sim10$ days of life.

Table 6. Distribution of Salmonella Organisms in Apparently

DA	YS AFTER HATCHING	1	(19)/I\	√'5	4)			2					3					5		
	NO. OF CHICKS EXAMINED	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20
	{ Trachea { Lung	:	:	• s	:		s	S	s •	• s	•,	•	:	:	S	:		s	• s	s S	•
ED	(Heart Blood Liver		:	•	•			:			:	:		:	•	•	:		:	s •	:
ZIZ	Spleen Kidney	•	•	٠	•	•	•	•	•	•	•	•	•	•	•	•	•	•	•	•	•
EXAMINED	Testicle Egg-Yolk	•	•	•	•	:	•	•	•		:	•	•	•	•		•	•	•	:	•
REGIONS	Esophagus Crop Proventriculus Gizzard	s ·	•	s :	s S ·	s :	s S ·	s ·	s S s	s s S S	s :	s s s	s S	; ;	$ss \cdot s$	s :	s	s :	; ;	s s	•
	Small Intestine Caecum Rectum	•	s s	:	s s	s s S	· s	8 8	:	s s	SSS	s S	s s	ś	· s	:	•	s s	s s	$\overset{\mathbf{S}}{\cdot}$	s

Notes: S \cdots Direct culture, s \cdots Enrichment culture of S. senftenberg.

... Negative result.

Table 7. Distribution of Salmonella Organisms in Apparently

DAY	S AFTER HATCHING	1	(29	9/ V	'54	1)		·	2					3					5					7
ì	NO. OF CHICKS EXAMINED	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23
	{ Trachea } Lung	s s	S	s	· s	S.	s S		s s	•	· s	s s		:			• s	s	•		• s	s		:
INED	Heart Blood Liver Spleen	:	:	s •	•	S s	:	:	s s	:	:	•	•	:	•	:	:	•	•	:	•	•	•	:
EXAMINED	Kidney Testicle Egg-Yolk	•	:	:	:	ś	:	:	:	:	•	s :	:	:	:	:	•	•	:	:	:	:	:	•
REGIONS	Esophagus Crop Proventriculus Gizzard	s •	s s	$\overset{\text{s}}{\cdot}$	s_s .	S S S S	s s •	$\overset{\text{s}}{\cdot}$	$\overset{\text{s}}{\overset{\text{s}}{\cdot}}$; ;	s s	S S . s	8 8 8	s •	s s •	:	· · ·	s :	:	:	•	•	•	•
	Small Intestine Caecum Rectum	s S	s	s s S	SSS	s s	s s	· s	s s	: s	s s s	s s	:	: s	SSS	SSS	•	s •	s :	:	:	•	•	:

Notes: S ... Direct culture, s... Enrichment culture of S. senftenberg.

···Negative result.

Healthy Chicks' Bodies Originated from F Hatchery

7								15						20		25	30	40	50	60							
21 22 23 24 25				26 27 28 29 30					31 32 33 34 35					36 37 38 39 40					43 41 42 \ 45		46	51	56 } 60	61	66 70		
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\mathbf{S}	•	S	•	•	•	•	•	•	•		S	•	•	s	•		•	•	•						•		•
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•	\mathbf{S}	•	•	•	•	•	٠	s	•	•	•	•	•	•	•	•	•	٠	\mathbf{S}	•	•	•	•	٠	•	•	•
	\mathbf{S}							\mathbf{s}									\mathbf{S}		\mathbf{S}		•						
\mathbf{S}	\mathbf{S}	\mathbf{S}	\mathbf{s}	\mathbf{S}	\mathbf{S}	\mathbf{S}	•	\mathbf{S}	\mathbf{S}		•		•	•	•	•	S	•	•	•	•	•	•	•			•
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Healthy Chicks' Bodies Originated from T Hatchery

24 25				9			11					15					20						25		30			40	50	60
	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	7	55 } 57	?
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SUMMARY

Experimental studies were carried out to clarify the mode of S. senftenberg infection in newly hatched chicks in some hatcheries situated in or near Sapporo and to clarify the presence or disappearance of S. senftenberg which invaded into baby chicks' bodies. At the same time the reasons for the appearance of S. senftenberg in hatching compartments of incubators and the influence of S. senftenberg upon baby chicks were discussed.

The conclusions obtained are summarized as follows:

- 1. In any hatchery so far as the writers have investigated, S. senftenberg came to be detected in the hatching compartments of incubators at time over $6\sim7$ weeks after the beginning of hatching (Tables 1, 2 and 3).
- 2. The newly hatched chicks originated from such incubators are usually infected in the incubators with S. senftenberg through their respiratory and digestive organs, and the rate of S. senftenberg infection among chicks tends to run parallel with the rate of multiplication of S. senftenberg in the incubator (Tables 1, 2 and 3).
- 3. These infected chicks carry the greatest numbers of S. senftenberg during first about 1 week of life, but so far as they are brooded under good feeding, management and sanitation, they remain healthy in appearance. They usually eliminate S. senftenberg from their bodies in $4\sim6$ weeks without therapeutic measures (Tables 6 and 7) and may be expected to exhibit no sequelae by S. senftenberg in future.
- 4. The regions where S. senftenberg lurks in baby chicks' bodies are the respiratory and digestive organs (Tables 6 and 7).
- 5. It is sure that S. senftenberg multiplies in hatching compartments of incubators working with normal temperature and humidity. However it is not yet proved what may bring, and how S. senftenberg is brought into the incubator for the first time.
- 6. As S. senftenberg contained in incubator is readily killed by formaldehyde gas, the hatching compartments of incubators should be cleansed and disinfected with the fumigation of formalin at intervals of 6 weeks after the beginning of hatching.

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