

Report on an enquiry into rat plague in East Anglia, during the period July-October 1911 / by Drs A. Eastwood and F. Griffith ; [and] Appendix containing report upon the postmortem examination of rats at Ipswich / by Drs G.H. Macalister and R. St John Brooks.

Contributors

Eastwood, A.
Griffith, F.
Macalister, G. H.
Brooks, R. St. John.

Publication/Creation

London : H.M.S.O., 1912 ([London] : Darling & Son.)

Persistent URL

<https://wellcomecollection.org/works/unbcyrdz>

License and attribution

The copyright of this item has not been evaluated. Please refer to the original publisher/creator of this item for more information. You are free to use this item in any way that is permitted by the copyright and related rights legislation that applies to your use.
See rightsstatements.org for more information.

**wellcome
collection**

Wellcome Collection
183 Euston Road
London NW1 2BE UK
T +44 (0)20 7611 8722
E library@wellcomecollection.org
<https://wellcomecollection.org>

Confidential.
FOR OFFICIAL USE ONLY.

REPORTS
TO THE
LOCAL GOVERNMENT BOARD
ON
**PUBLIC HEALTH AND MEDICAL
SUBJECTS.**

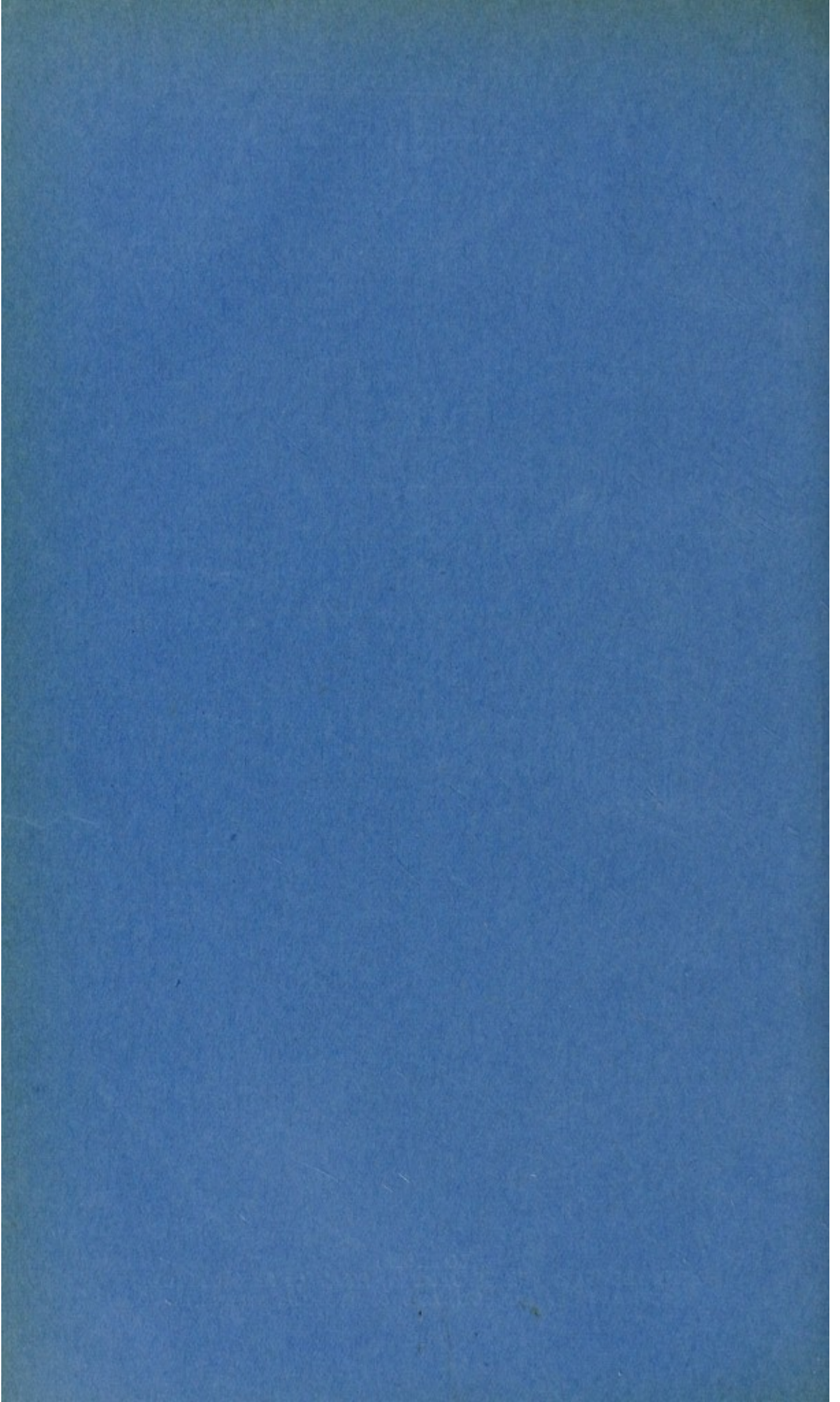
Report on an Enquiry into Rat Plague in East Anglia, during the period July—October, 1911 ; by Drs. A. Eastwood and F. Griffith.

Appendix containing Report upon the Post-mortem Examination of Rats at Ipswich ; by Drs. G. H. Macalister and R. St. John Brooks.



LONDON :
PRINTED FOR HIS MAJESTY'S STATIONERY OFFICE,
By DARLING AND SON, LTD., 34-40, BACON STREET, E.

1912.



Report to the Local Government Board on an
Enquiry into Rat Plague in East Anglia
during the Period July-October, 1911.
By Drs. A. Eastwood and F. Griffith.

CONTENTS.

	Page
Introduction	2
Organisation :—	
Police Assistance in Rat Collection	2
Arrangements by Mr. Huddart	2
The Ipswich Laboratory	3
The Board's Pathological Laboratory	3
Routine	3
Bacteriological Diagnosis in the Board's Laboratory :—	
Methods	4
Results	6
Analysis of Results in Positive Cases :—	
Preliminary Data	6
Results of Culture Work :—	
<i>Isolation of the Plague Bacillus by Culture</i>	7
<i>Characteristics of the Plague Cultures</i>	7
<i>Fermentation Tests</i>	8
Results of Animal Experiments :—	
<i>Inoculation of Rats with Plague-infected Tissues</i>	9
<i>Inoculation of Guinea-pigs with Plague-infected Tissues</i>	9
<i>Inoculation of Rats and Guinea-pigs with Plague Cultures</i>	10
Tabular Statement of Results	11
The Virulence of the Plague Strains	16
Analysis of Results in Negative Cases :—	
Preliminary Data	17
Results of Cultures and Animal Experiments	19
Tabular Statement... ..	20
Bacteria of some Special Interest :—	
<i>Bacilli belonging to the Gärtner Group</i>	24
<i>A Cocco-bacillus from a Rabbit</i>	24
<i>Colonies bearing some Resemblance to Plague</i>	25
Evidence as to Distribution and Limitation of Infection :—	
Map	26
The Area from which Rats were Collected	26
Comparison with the Area previously Investigated	26
The Supply of Rats	27
Acknowledgments	28
Summary	29
Recommendations	29

INTRODUCTION.

This enquiry was conducted in accordance with the instructions of the Board, who directed that attention should first be confined to the area where plague-infected rats had previously been found and that, if such rats were again discovered, an endeavour should then be made to determine the limits of the infection. The area shown by previous investigations to have been infected comprised the rural districts of Samford and Woodbridge, the urban districts of Woodbridge and of Felixstowe and Walton, and the borough of Ipswich. The present enquiry commenced on July 1st, and was continued until October 31st, 1911.

ORGANISATION.

Police Assistance in Rat Collection.

The Board obtained, through the Home Office, the assistance of Captain Mayne, Chief Constable of East Suffolk, throughout the enquiry. Captain Mayne issued instructions to the following effect:—

(1) The police officers throughout East Suffolk were to make confidential enquiries and then to report answers to the following questions:—(a) Are rats known to be dying in any part of the officer's district? (b) If so, in what parishes and on whose premises? (c) What is the supposed cause of death? (d) Has any sort of rat poison or virus been recently laid down? (e) Any other information?

(2) The officers in charge of every police station in that part of East Suffolk selected by the Board for the collection of rats were requested (a) to ascertain, by local enquiry, in what parts of their districts rats were likely to be found and to inform the Board's rat-catchers accordingly, (b) to receive at the police station all rats brought in by these rat-catchers, (c) to urge all residents in their districts to catch rats and bring them in to the police station, and to pay, on behalf of the Board, 2*d.* per rat, (d) to disinfect all rats brought in, (e) to see that a disc bearing a number was attached to each rat, (f) to fill in for each rat a card of identification, bearing specified particulars, and (g) to forward the rats and the cards to the Municipal Laboratory at Ipswich.

When it was found necessary to extend the area of enquiry into portions of West Suffolk and the north of Essex, similar assistance was obtained, through the Home Office, from the Chief Constables of West Suffolk, Essex, and Colchester.

Arrangements by Mr. Huddart.

Mr. Huddart, Assistant Inspector to the Board, (1) made all arrangements with the Chief Constables and their superintendents for the work of the police, (2) engaged rat-catchers and assigned them work week by week, (3) obtained and distributed all material (disinfectants, cards, boxes, rat-traps, &c.) required by the police and the rat-catchers, (4) visited all the police stations and explained to the constables what was required,

(5) made periodical tours of inspection throughout the enquiry for the purposes of supervising the rat-catchers and keeping in touch with the work of the police, and (6) took charge of the clerical work in connection with the enquiry.

The Ipswich Laboratory.

On the recommendation of Dr. Pringle, Medical Officer of Health for Ipswich, the Council of that Borough placed their Municipal Laboratory gratuitously at the disposal of the Board throughout the enquiry. All rats were received and dissected in this laboratory by Drs. G. H. Macalister and R. St. John Brooks, who were engaged by the Board from July 1st to October 31st for the performance of this work. The routine clerical work was conducted in a room adjacent to but separate from the laboratory by Mr. Leach, a temporary clerk, who worked under the direction of Mr. Huddart.

The Board's Pathological Laboratory.

In the case of any rats which on post-mortem examination did not appear to Drs. Macalister and Brooks to be free from suspicion of plague, the liver, spleen, and any other suspected material from each of these animals were sent to the Board's laboratory for diagnosis. They were there examined microscopically, culturally, and by animal inoculation. Dr. Klein acted as consultant in cases where it was thought that the diagnosis needed his confirmation.

Routine.

When a rat was caught by one of the Board's rat-catchers he attached to it with string a perforated circular metal disc bearing a stamped number. This number and the place where the rat was caught he recorded in a note-book. It was to his interest to keep these records accurately, as his weekly claims for payment were based on them. He caught his rats with the aid of ferrets, dogs, and traps, but was not allowed to use poison. In addition to the rats so obtained he added to his collection any dead rats he could find; these latter were labelled and recorded in the same way as the caught rats, and, in addition, a piece of coloured tape was tied to each as a distinctive mark.

He brought in his day's bag, in which he was not permitted to include any rats less than half-grown, to the nearest police station. There the constable disinfected them in 2 per cent. lysol and entered on a separate card for each rat the following particulars:—(1) the number on the metal disc (termed "local number"); (2) the date; (3) name of catcher; (4) parish where caught or found; (5) exact place where caught or found; (6) whether caught or found dead; (7) whether the occupier had previously laid down virus; (8) if so, when and what kind.

When rats were brought in by voluntary workers, the constable affixed to each a disc bearing a "local number," tied coloured tape to any found dead, and then dealt with them in the same way as with those brought in by the professionals.

The constable packed the rats in tin boxes, each of which was enclosed within an outer wooden box, and forwarded them,

generally by carrier, to the Ipswich laboratory. The cards bearing details of each rat were sent to the laboratory by post.

At Ipswich the cards were received in the clerk's room, the rat-boxes in the laboratory.

When the examination of each rat was completed, the circular disc bearing the local number was removed by cutting the string attaching it to the rat, and a square disc bearing a serial number was attached by wire to the circular disc. The serial numbers ran in sequence from 1 upwards, and registered the total number of rats received in the laboratory. The pairs of circular and square discs were collected in two lots, one bearing the numbers of the rats found free from plague, the other bearing the numbers of those reserved for further examination. After being sterilised, these batches of discs were sent up to the clerk's room. No articles of any sort which might have been in contact with infective material were sent up to the clerk's room without being sterilised.

On receiving the pairs of discs, the clerk identified each card by its local number and copied on to it the corresponding serial number. Each rat was then entered up, according to its serial number, as "normal" (*i.e.*, free from plague) or "reserved" (*i.e.*, specimens sent to the Board's laboratory for examination). At the end of each day's work the clerk drew up a detailed statement of all the rats examined during the day, giving the names of the parishes from which the rats were received, the numbers of rats received from each, the serial numbers of all "normal" and all "reserved" rats and of all rats found dead, and, in the case of each "reserved" rat and of every rat found dead, the exact address where it was obtained. One copy of this information was posted each evening to the Board's laboratory and another to Mr. Huddart.

In the case of each "reserved" rat Drs. Macalister and Brooks sent to the Board's laboratory the liver, spleen and any other material considered suspicious, together with a full account of the post-mortem appearances of the rat, including microscopic evidence obtained from smear preparations. This material and information, and also the clerk's daily record, were received the next morning at the Board's laboratory. The specimens were there investigated microscopically, culturally, and by animal inoculation. When a case of plague was fully established, the Board was at once notified of the fact and of the exact place whence the rat was obtained. At the same time Mr. Huddart was asked to stop further supplies from the parish whence the rat was obtained, to notify the occupier of the premises where the rat was found of the presence of rat-plague, to warn him of the danger, and to advise him to take energetic measures to destroy any rats on his premises.

BACTERIOLOGICAL DIAGNOSIS IN THE BOARD'S LABORATORY.

Methods.

The routine method of staining film preparations from tissues, fluids, or cultures was as follows: After drying in the air and then fixing for two minutes in absolute alcohol, the smears were stained, as recommended by Dr. Klein, with Czinzinski's

mixture of methylene blue and eosin. The formula for this mixture is:—

Methylene blue (conc. aqueous solution)	...	50 cc.
Eosin (soluble in alcohol)5 gm.
Alcohol (absolute)	70 cc.
Water (distilled)	130 cc.

The stain was applied for at least five minutes, and at the beginning and the end of this period was warmed over the flame until steam rose. The specimens were then washed in water, dried, and mounted in Canada balsam.

For the isolation of cultures slanted agar tubes and agar plates were employed, the medium consisting of ordinary nutrient agar made with beef broth, reaction + 10 to phenolphthalein. The surface of the agar was fairly dry. In the earlier part of the investigation some use was made of MacConkey's neutral-red, bile salt medium, with the addition of mannite or lactose. It was found, however, by comparative experiments with plague-infected tissue, that growth on a bile salt medium was less certain and slower than on agar, and that the bile salt had some inhibitory effect, fewer plague colonies being produced than on ordinary nutrient agar plates inoculated with the same dose of the same material. Again, the fact that intestinal bacteria, the growth of which was not inhibited by bile salt, were commonly present in such contaminated and decomposed tissues as had to be dealt with rendered bile salt preparations of no great advantage as selective media for isolating plague bacilli from a mixture of organisms.

For culture work the spleen was usually taken. Cultures were also made from other material which, after microscopic examination, seemed suitable. After the surface of the spleen, or other organ, had been well seared, a portion of the interior was cut out with a sterile knife and placed on the first plate or culture tube. After this material had been rubbed over the surface two or three more plates or tubes were inoculated from it in series. The temperature of incubation was 30° C.

In addition to the cultural investigation, the specimens in nearly every case were tested by inoculation upon white rats or guinea-pigs or both. Animals were also inoculated, when thought necessary, with cultures or with such material from the first experimental animals as required further investigation. The inoculation was usually made subcutaneously, but in some cases cutaneously.

In the case of the specimens sent from Ipswich for diagnosis the spleen and liver were usually selected for inoculation. After the surfaces of these organs had been thoroughly seared, portions of the interior were transferred to a small test tube, well rubbed up with a glass rod and then emulsified with normal saline. Other material, such as lymphatic glands, pleural effusion or blood, was also used when thought requisite.

No case was diagnosed as positive until a typical plague culture and typical plague infection of an inoculated animal had been obtained. With the exception of 12 cases where animal inoculation was not considered necessary, no case was

settled as negative until it was proved that the material was incapable of producing plague in an experimental animal.

Enquiry was also made into the action of the plague bacilli, after isolation in pure culture, upon various carbohydrates and allied substances and into their virulence, in measured dosage of culture, for experimental animals. Broth cultures showing typical stalactites were also obtained with several of the viruses.

Results.

Out of the 15,332 rats which were dissected at the Ipswich laboratory by Drs. Macalister and Brooks, specimens from 151 were sent to the Board's laboratory for diagnosis. They were there examined by the methods described above.

Thirty-five of the rats were found to be plague infected, the remaining 116 being proved negative as regards plague.

ANALYSIS OF RESULTS IN POSITIVE CASES.

Preliminary Data.

Out of the 35 rats proved to be plague infected, 21 had been caught and killed, the remaining 14 had been found dead.

At the time of the post-mortem examination the condition, as regards preservation, of 30 of the above rats was described by Drs. Macalister and Brooks as either "good" or "fair"; the remaining five were either "bad" or "putrid."

Accompanying the selected material submitted to the Board's laboratory for diagnosis, full details were sent of the post-mortem and microscopic appearances which Drs. Macalister and Brooks found.

These data may be classified as follows:—

Group I.—In this group, to which 28 of the positive cases belong, the appearances were either typical of plague or very strongly suggestive of it. The predominant features were marked subcutaneous congestion and general distribution in the tissues of bacilli, usually very numerous, which were morphologically identical with *B. pestis*. A good example is Rat 2,876, which was described in the following detail:—*Macroscopically*: The right inguinal gland was the size of a pea, with a necrotic centre and thick walls; the pelvic glands were enlarged; subcutaneous congestion was intense in the region of the right inguinal gland and was very well marked all over the trunk; no hæmorrhages were seen; there was some clear pleural effusion; the liver was large, firm, generally pale, mottled, and showed numerous punctate necrotic foci; the spleen was large and firm, with well-marked granulation. *Microscopically*: A few bacilli, with some involution forms, were found in the necrotic centre and in the walls of the right inguinal gland; some bipolar bacilli were seen in the right pelvic gland; no organisms were found in the pleural effusion; scanty bipolar bacilli were found in the liver and spleen; a few plague-like bacilli were found in the heart's blood.

This case proved to be, in most respects, typical of the group. But in the remaining 27, though glandular enlargement and congestion were usually found, necrosis of a lymph gland only occurred once, and plague-like bacilli were numerous in the

organs in 22 cases. Other differences which occurred in some of these 27 cases were minor and were not sufficiently marked to justify classification in a separate group.

Twelve of these 28 cases were rats which had been found dead; three of the rats were found at the post-mortem examination to be "bad" or "putrid."

Group II.—To this group seven positive cases belong. In general, the macroscopic and microscopic data cannot be regarded as affording more than slight suspicion of plague. Some bipolar bacilli were found in each case, but subcutaneous congestion was only marked in one case and was absent altogether in three. Rat 1,030 belongs to this group. As this was the first case in the present enquiry which turned out to be positive, the full description of its post-mortem and microscopic appearance is of particular interest. *Macroscopically*: There was some enlargement of the submaxillary glands; there was neither subcutaneous congestion, hæmorrhage, nor pleural effusion; the liver showed punctate, white necrotic foci; in the spleen there were remains of a white infarct. *Microscopically*: Numerous bipolar bacilli were found in the liver and heart's blood. This is the only case, among the seven, in which bipolar bacilli were recorded as numerous; in one case, in which the tissues were decomposed, "mixed organisms" including bipolar bacilli were found; in the remaining five cases bipolar bacilli were stated to be scanty.

Two of these seven cases were rats which had been found dead; two of the rats were found on post-mortem examination to be "bad" or "putrid."

Results of Culture Work.

Isolation of the Plague Bacillus by Culture.

In 29 out of the 35 positive cases *Bacillus pestis* was obtained by direct culture of the original material, and in many cases without admixture of any contaminating organisms. Cultures were obtained from the spleen in 26 of these cases; in the remaining 3 they were obtained from the liver; additional cultures were obtained in 8 cases from livers and in 2 cases from lymphatic glands.

In the remaining 6 out of the 35 cases direct cultures from the original material failed. In three instances they were overgrown; in two other cases, where bile salt media alone were used, no plague colonies appeared; in the sixth case no culture was attempted, as the original material was very putrid.

In each of the above six cases cultures of *B. pestis* were obtained from the animals inoculated with the original material.

Characteristics of the Plague Cultures.

The following were the main characteristics of the cultures, which were obtained by the methods described above (p. 4).

On examination the next day, within 24 hours after inoculation on agar, plague colonies were usually visible with the aid of a hand lens as minute transparent points; when the material inoculated contained numerous plague bacilli, a thin growth over the surface could be seen by the naked eye. The colonies,

transparent at first, soon became denser and then exhibited a delicate but characteristic ground-glass appearance; as the colonies grew older they became completely opaque. The typical plague colony, two to three days old, was greyish white, with a smooth, shining surface; the centre of the colony was slightly raised, the margin thin, grey, translucent and irregular; in this margin minute colonies sometimes developed. The adhesive nature of the culture was an important characteristic, to which special attention has been called by Dr. Klein. This peculiarity, which developed early when incubation took place at 30° C., was readily demonstrated by touching the growth with the platinum needle; on gently raising the needle, the culture adhered to it and was drawn out in the form of a long thread.

In subculture on agar slants, and also in primary culture from tissues rich in plague bacilli without admixture of other organisms, the surface became covered with a continuous greyish-white layer, exhibiting the characteristic stickiness when touched with the platinum needle. Not uncommonly discrete, dense colonies of plague bacilli grew up above the rest of the surface of a pure culture.

As the cultural results obtained on agar were unequivocal and distinctive and were corroborated in every instance by the production of typical plague in an experimental animal, no further culture work was considered necessary for the purpose of diagnosis.

Incidentally, a few experiments were made on the production of stalactites in broth covered with a thin layer of butter fat and kept at room temperature. Out of the six strains, selected at random, which were tested five produced typical stalactites, the sixth failed to do so.

Fermentation Tests.

For the purpose of a comparative study of plague bacilli with other organisms which are often differentiated from each other by means of fermentation tests, each of the 35 plague viruses was submitted to a series of these tests.

To ordinary peptone water, made with tap water and coloured with 10 per cent. litmus solution, .5 per cent. of one of the following chemically pure carbo-hydrates or other compounds was added:—glucose, fructose, galactose, maltose, lactose, saccharose, raffinose, iso-dulcitol, glycerin, mannitol, dulcitol, adonitol, inulin, and salicin. Each of these media was put up in Durham's fermentation tubes. Litmus milk and malachite green peptone water were also used as test media.

With all 35 viruses acid, but no gas, was produced in—glucose, fructose, galactose, maltose, mannitol, adonitol, and salicin. The reactions with the last two were slow, salicin taking about 6 days and adonitol about 12 days.

With all 35 viruses no change of reaction was produced in—lactose, saccharose, raffinose, dulcitol, glycerin, inulin, and litmus milk. The period of observation, in each case, was 28 days. During this period no appreciable change was noted in the malachite green medium, but after about 5 weeks' incubation many strains slightly decolourised this medium.

With 4 of the viruses acid was formed, in 20 days, with iso-dulcite; with the other 31 viruses during this period no change was produced with this compound, but in the case of two of these 31 acid was formed after a further incubation of 20 days.

For the purpose of control, all the above tests were performed also with two previously authenticated strains of plague bacilli; for one of these we are indebted to Dr. Klein, for the other to Dr. Rowland, of the Lister Institute. These two strains produced no change in iso-dulcite; in all the other tests they corresponded exactly to the 35 strains mentioned above.

Results of Animal Experiments.

Inoculation of Rats with Plague-infected Tissues.

With the exception of a few cases where the cutaneous method of inoculation was adopted, the material was inoculated subcutaneously in the right groin.

The day after inoculation the rats usually looked ill, with staring coat, and local tenderness and swelling could be felt on palpation. They rapidly became worse, dyspnœa being marked, and died in from 2 to 8 days; death usually occurred between the second and the fourth day. In every case subcutaneous inoculation produced a fatal result, a tumour being formed at the site of inoculation.

The local lesion was invariably found to be necrotic and œdematous. It was surrounded by an area of congestion which was often intense. There was general redness and vascular engorgement throughout the ventral subcutaneous tissues. These features were constant and particularly conspicuous.

The inguinal and iliac glands were usually enlarged and often congested and necrosed, the extent of the lesions being proportionate to the duration of the animal's life. The axillary and submaxillary glands were sometimes enlarged and congested.

In several cases the spleen did not appear abnormal in size, colour, or consistency. In others it was enlarged, firm, and dry on section. In one rat, which lived for 8 days, the spleen was peppered with grey foci.

In animals which died in from 2 to 3 days the liver generally showed nothing more than congestion. In later cases it showed either small, irregular patches of necrosis or minute irregular grey foci.

The lungs were frequently œdematous and congested, but no pleural effusion was found.

Plague bacilli were almost invariably numerous in the local lesion and the nearest glands, being usually irregular in size and shape and often showing involution forms. In the spleen they were often scanty, especially when the animal died early with marked local reaction. In some cases the bacilli were abundantly disseminated throughout the body and in the blood.

Inoculation of Guinea-pigs with Plague-infected Tissues.

The material was inoculated subcutaneously in the right groin, in the same manner as with the rats.

The day after inoculation it was usually found that the animals were quiet and that local tenderness and some swelling

could be detected on palpation. Day by day the animals rapidly became more ill and lost flesh, whilst the local swelling increased in size. At death there was usually a firm, prominent swelling and the right hind leg was drawn up. Death occurred between the third and the tenth day; in the majority of cases it was between the fifth and the sixth day.

At the site of inoculation there was found in the subcutaneous tissue an infiltrating tumour which, on section, was seen to be centrally necrotic and to be surrounded by an area of intense congestion and œdema. The size of the tumour varied, being usually larger in the animals which lived longer; the largest was the size of a pigeon's egg and was found in a guinea-pig which had lived for ten days after inoculation.

Subcutaneous congestion and œdema extended for a considerable distance beyond the local lesion, with engorgement of the subcutaneous capillaries and veins; but the œdema was never found to spread completely over the ventral subcutaneous tissues.

The inguinal and iliac lymphatic glands were enlarged and congested, particularly in the cortex; their interior usually showed extensive necrosis.

The spleen was usually enlarged, congested, and either mottled with pale, irregular necrotic areas or studded with discrete, soft, grey nodules ranging in size up to 1 mm. in diameter. In cases of very acute infection ending fatally in about three days the spleen remained small and did not exhibit macroscopic lesions.

The liver, as a rule, was enlarged; sometimes it showed nodules similar to those found in the spleen; in other cases it was peppered with irregular, grey foci. In cases which terminated fatally in three days no lesions were visible.

The lungs were usually congested and œdematous, sometimes with patches of consolidation. They were frequently studded with soft, greyish-white nodules with a congested periphery. The largest of these nodules were found in the less rapidly fatal cases; in one animal, which had lived for 10 days, they attained the diameter of 5-6 mm. Pleural effusion was found sometimes.

In almost every instance the tissues of the animals contained enormous numbers of plague bacilli; in smears from the spleen or the lung nodules this feature was particularly noticeable.

Inoculation of Rats and Guinea-pigs with Plague Cultures.

In rats inoculated with culture, the œdema, congestion, and necrosis of the local lesion, the general subcutaneous congestion, and the enlargement, congestion, and necrosis of the inguinal and iliac glands were all features which corresponded closely with the results obtained by the inoculations of infected tissues. As compared with tissue emulsions, pure cultures produced less macroscopic change in the spleen and liver; the spleen, though sometimes enlarged and often firm and dark on section, did not show foci of necrosis; the liver was generally normal in appearance, though occasionally showing a few grey foci. The lungs were generally œdematous and sometimes congested; in a small proportion of the cases (less than a quarter) there was a little pleural exudate.

In guinea-pigs culture inoculations produced very much the same effects as emulsions of tissue.

Tabular Statement of Results.

Details of Cultural Results and Animal Inoculations in the Diagnosis of the 35 Positive Cases.

* The Roman numerals refer to the grouping on pp. 6 and 7.

Number and Condition of Rats.		Results of Cultures from Original Material, as regards Growth of Plague Bacilli.	Results of Animal Experiments.			
Number.	*Condition.		Material inoculated.	Animals.	Duration of Life.	Post Mortem Results.
1030	II.	Discrete colonies from liver on mannite bile salt.	Liver	Rat 30	Died, 3 days.	Acute plague. Bacilli numerous in liver, spleen, and blood.
			Spleen	G.P. 507	Died, 5 days.	Acute plague. Bacilli abundant in spleen.
1264	II. (<i>putrid.</i>)	Cultures, on bile salt media, failed.	Liver	Rat 31	Died, 4 days.	Acute plague. Bacilli numerous in spleen.
			Liver	Rat 32	Died, 6 days.	Acute plague. Bacilli numerous in liver, spleen, and blood. Cultures from Rats 31 and 32 typical of plague.
1691	I. (<i>found dead.</i>)	Cultures, on bile salt media, failed.	Liver	Rat 36	Died, 3 days.	Acute plague. Bacilli numerous in spleen and liver.
			Liver	Rat 37	Died, 3 days.	Acute plague. Bacilli (not numerous) in spleen and liver.
			Spleen	G.P. 514	Died, 8 days.	Typical plague. Bacilli fairly numerous in spleen. Cultures from Rat 37 typical of plague.
1694	I. (<i>found dead.</i>)	Discrete colonies from liver on lactose bile salt.	Liver	Rat 38	Died, 3 days.	Acute plague. Bacilli fairly numerous in liver and a few in spleen.
			Liver	Rat 39	Died, 3 days.	Acute plague. Bacilli numerous in liver and spleen.
			Spleen	G.P. 515	Killed, 16 days.	Healthy. Cultures from Rat 38 typical of plague.
1769	I.	Discrete colonies from liver and spleen on mannite bile salt.	Liver	Rat 40	Died, 2 days.	Necrotic local lesion; much oedema, spreading over thoracic wall; nothing definite in organs. Two bipolar bacilli seen in spleen, none in liver.

Number and Condition of Rats.		Results of Cultures from Original Material, as regards Growth of Plague Bacilli.	Results of Animal Experiments.			
Number.	*Condition.		Material inoculated.	Animals.	Duration of Life.	Post Mortem Results.
1769	I.	Discrete colonies from liver and spleen on mannite bile salt.	Pleural fluid Spleen	Rat 41 G.P. 540	Died, 6 days. Died, 6 days.	Typical plague. A few bacilli in spleen, fairly numerous in liver. Typical plague. Bacilli numerous in spleen. Cultures from Rat 40 typical of plague.
2064	II.	Pure culture from spleen on agar slants.	Liver Liver	Rat 51 G.P. 561	Died, 4 days. Died, 4 days.	Acute plague. Bacilli rather scanty in spleen. Acute plague. Bacilli abundant in spleen.
2156	II. (found dead.)	Pure cultures from spleen and liver on agar slants.	Liver	Rat 52	Died, 5 days.	Acute plague. A few bacilli in spleen.
			Liver	Rat 53	Died, 3 days.	Acute plague. Bacilli rather scanty in spleen.
			Liver	G.P. 566	Died, 6 days.	Acute plague. Bacilli abundant in spleen.
2522	II. (putrid; found dead.)	No culture attempted. ...	Spleen and Liver Spleen and Liver	Rat 72 (cutaneous). Rat 73 (cutaneous).	Killed, 13 days. Died, 6 days.	Healthy. Acute plague. Bacilli scanty in spleen, many in axillary gland. Cultures from Rat 73 typical of plague.
2876	I. (found dead.)	Discrete colonies from inguinal bubo on mannite bile salt slants. Abundant colonies on agar slants from spleen.	Spleen Liver	Rat 94 G.P. 652	Died, 3 days. Died, 3 days.	Acute plague. Bacilli moderately numerous in spleen. Acute plague. Bacilli abundant in spleen.

3346	I. (<i>found dead.</i>)	Overgrown (liver and spleen).	Spleen and Liver Spleen and Liver	Rat 116 Rat 117	Died, 3 days. Died, 4 days.	Acute plague. Bacilli moderately numerous in spleen Soft caseous nodule at site of inoculation, without congestion; spleen mottled with greyish necrotic patches; two similar patches in liver. A few bipolar bacilli in spleen. Cultures from liver and spleen of Rat 116 typical of plague. Cultures from spleen of Rat 117 overgrown with coliform growth.
3430	I. (<i>found dead.</i>)	Pure cultures from spleen on agar slants; cultures almost pure from liver on same medium.	Spleen and Liver Spleen and Liver	Rat 118 Rat 119	Died, 2 days. Died, 3 days.	Acute plague. Bacilli moderately numerous in spleen. Acute plague. No bacilli seen in spleen, fairly numerous in iliac gland. Pure cultures from spleen and liver.
4849	I.	Numerous colonies in pure culture from spleen and almost pure from liver—on agar slants.	Spleen and Liver	Rat 143 G.P. 706	Died, 5 days. Died, 10 days.	Acute plague. Bacilli moderately numerous in spleen and axillary gland. Plague lesions well marked. Bacilli in spleen.
5381	I.	Pure culture from spleen on agar slants.	Spleen and Liver	Rat 144	Died, 8 days.	Purulent sinus at site of inoculation, slight injection of subcutaneous vessels; right inguinal gland enlarged; liver and spleen peppered with irregular grey foci. No bipolar bacilli seen in spleen; two seen in liver. Coliform colonies from spleen but no plague colonies. Acute plague. Bacilli abundant in spleen.
5382	I. (<i>found dead.</i>)	Discrete colonies from liver on one agar slant. Three other tubes overgrown.	Spleen and Liver Liver	G.P. 743 G.P. 744	Died, 6 days. Died, 4 days.	Acute plague. Bacilli abundant in spleen. Pure culture isolated from spleen.
7643	I.	Abundant growth on agar slants from spleen and liver, almost pure.	Spleen and Liver	Rat 157	Died, 2 days.	Acute plague. Bacilli moderately numerous in spleen, from which a pure culture was obtained.
8086	I. (<i>found dead.</i>)	Discrete colonies on agar slants from spleen.	Spleen and Liver	Rat 160	Died, 3 days.	Acute plague. Bacilli rare in spleen, moderately numerous in iliac gland.

Number and Condition of Rats.		Results of Cultures from Original Material, as regards Growth of Plague Bacilli.	Results of Animal Experiments.			
Number.	*Condition.		Material inoculated.	Animals.	Duration of Life.	Post Mortem Results.
10169	I.	Pure culture on agar slants from spleen.	Spleen and Liver	Rat 169	Died, 2 days.	Acute plague. Bacilli moderately numerous in spleen, abundant in iliac gland.
10340	I. (<i>found dead.</i>)	Pure culture on agar slants from spleen.	Spleen and Liver	Rat 175	Died, 2 days.	Necrosis at site of inoculation, with oedema but no congestion; slight enlargement of iliac gland; spleen firm; liver lobules outlined. Numerous bacilli in iliac gland.
10848	I.	Discrete colonies on agar slants from spleen and liver.	Spleen and Liver	Rat 176	Died, 4 days.	Acute plague. Bacilli scanty in spleen, moderately numerous in iliac gland. Pure culture obtained from spleen.
11407	II. (<i>bad.</i>)	Discrete colonies on agar slants from spleen; contaminating organisms numerous.	Spleen and Liver	Rat 180	Died, 4 days.	Oedematous necrotic tissue at seat of inoculation, without congestion; iliac gland enlarged; spleen firm; liver peppered with irregular grey foci. One doubtful bacillus seen in spleen. Cultures from liver and spleen overgrown. G.P. 876, inoculated from the liver, died in 5 days from acute plague.
11724	I. (<i>found dead.</i>)	Pure cultures on agar slants from spleen.	Spleen and Liver	Rat 184	Died, 4 days.	Acute plague. No bacilli seen in spleen, but pure culture was obtained from it.
12012	I.	Abundant, pure culture on agar slants from spleen.	Spleen and Liver	Rat 189	Died, 3 days.	Acute plague. Abundant bacilli in iliac gland.
12013	I.	Abundant, pure culture on agar slants from spleen.	Spleen and Liver	Rat 190	Died, 3 days.	Acute plague. Abundant bacilli in iliac gland.
12245	I.	Overgrown.	Spleen and Liver	Rat 192	Died, 3 days.	Very marked local congestion, necrosis, and oedema; organs apparently normal. Bacilli scanty in spleen, abundant in local lesion. Culture isolated from spleen. Rat 197, fed with organs of Rat 192, died in 5 days from plague, with submaxillary bubo and abundant bacilli in spleen.

13663	I. (<i>found dead.</i>)	Pure culture on agar slants from spleen.	Spleen and Liver	Rat 199	Died, 3 days.	Acute plague. Bacilli abundant in iliac gland and numerous in spleen.
13712	I.	Pure culture on agar slants from spleen.	Spleen and Liver	Rat 200	Died, 2 days.	Highly congested, necrotic local lesion; slight general subcutaneous congestion; iliac gland slightly enlarged; spleen small and rather firm. Bacilli abundant in iliac gland, a few in spleen.
13713	I.	Confluent pure culture on agar slants from spleen.	Spleen and Liver	Rat 201	Died, 3 days.	Acute plague. Bacilli moderately numerous in spleen and abundant in iliac gland. Culture obtained from spleen.
13817	I.	Confluent pure culture on agar slants from spleen.	Spleen	Rat 202	Died, 2 days.	Acute plague. Bacilli abundant in iliac gland and spleen.
14499	I.	On agar slants pure confluent growth from lymphatic gland, discrete colonies from spleen and liver.	Spleen and Liver Gland	Rat 209 G.P. 897	Died, 3 days. Died, 6 days.	Acute plague. Bacilli moderately numerous in spleen, abundant in iliac gland. Acute plague. Bacilli abundant in spleen.
14816	II.	Discrete colonies from spleen on agar slants.	Spleen	Rat 220	Died, 4 days.	Slightly congested necrotic local lesion; nothing else abnormal. No bacilli seen in smear from spleen. Pure culture of plague from spleen.
14854	I.	Pure cultures from liver and spleen on agar slants.	Spleen	Rat 221	Died, 3 days.	Acute plague. Bacilli moderately numerous in spleen.
14855	I.	Discrete colonies from spleen on agar slants.	Spleen	G.P. 902	Died, 5 days.	Acute plague. Bacilli abundant in spleen.
14911	I. (<i>found dead.</i>)	Pure cultures on agar slants from liver and spleen.	Spleen	G.P. 909	Died, 4 days.	Acute plague. Bacilli abundant in spleen.
15079	I.	Discrete colonies from spleen on agar slants.	Spleen and Liver	Rat 230	Died, 4 days.	Purulent nodule at site of inoculation; general subcutaneous congestion; necrotic foci in iliac gland. Two bacilli found in smear from iliac gland.
15219	I.	Discrete colonies from spleen on agar slants.	Spleen	Rat 229	Died, 2 days.	Acute plague. Bacilli moderately numerous in inguinal gland.

The Virulence of the Plague Strains.

In the earlier part of the enquiry the cultures isolated from each case of rat plague were tested for virulence and were compared in this respect with two previously authenticated strains of plague bacilli received respectively from Dr. Klein and from Dr. Rowland at the Elstree Laboratory. It was found that pure cultures of all the strains were of high, and of about equally high, virulence for rats and guinea-pigs. It was not considered necessary to continue these culture tests systematically throughout the enquiry because in no instance was a virus obtained which, when inoculated in the form of a tissue emulsion, afforded any indication of being lower in virulence than the viruses already tested by inoculation both of cultures and of tissue emulsions.

The estimation of dosage was made by weighing on a chemical balance a platinum loop charged with culture and then subtracting the weight, previously ascertained, of the platinum loop. The weighed culture was finely emulsified with the requisite amount of normal saline solution to afford a convenient volumetric measure of dosage.

Most of the inoculations were made subcutaneously, but virulence for rats was also tested by the cutaneous method, one or two loopfuls of dilute emulsion of culture being smeared over the shaved and lightly scarified skin at the root of the tail.

The results of these virulence tests are tabulated below.

Virulence of the Plague Cultures for Rats inoculated subcutaneously.

Number or Name of Virus.	Dose inoculated, in mgms.	Number of Rat inoculated.	Duration of life, in days.
1,264	.1	46	3
1,030	.1	48	5
1,694	.01	42	3
1,691	.01	44	3
1,769	.01	57	5
2,064	.01	58	4
" Klein "	.01	60	3
" Elstree "	.001	62	4
2,156	.001	131	3
3,346	.001	133	3
3,430	.001	135	4
4,849	.0001	150	[remained healthy.]
4,849	.00001	151	5

Virulence of the Plague Cultures for Guinea-pigs inoculated subcutaneously.

Number or Name of Virus.	Dose inoculated, in mgms.	Number of Guinea-pig inoculated.	Duration of life, in days.
1,030	1	559	4
1,030	.1	560	5
1,694	.1	554	5
1,691	.1	556	6
1,264	.1	557	6
1,769	.1	591	3
2,064	.1	593	5
"Klein"	.1	595	5
1,691	.01	555	5
1,264	.01	558	4
1,769	.01	592	5
2,064	.01	594	3
"Klein"	.01	596	7
"Elstree"	.01	597	4
"Elstree"	.001	598	5
2,156	.001	689	13
3,346	.001	690	8
3,430	.001	692	8
4,849	.0001	760	[killed, 18 days ; local disease only.]
4,849	.00001	761	8

Virulence of the Plague Cultures for Rats inoculated cutaneously.

Number or Name of Virus.	Number of Loopfuls of Dilute Culture Emulsion used.	Number of Rat.	Duration of life, in days.
1,694	1	43	6
1,691	1	45	[Killed, 15 days ; healthy.]
1,264	2	47	4
1,030	?	50	5
1,769	1	56	4
2,064	1	59	3
"Klein"	1	61	4
"Elstree"	1	63	4
2,156	1	132	5
3,346	1	134	3
3,430	1	136	5
5,381	2	152	4
11,407	?	174	[killed, 24 days ; healthy.]
14,816	?	231	3

ANALYSIS OF RESULTS IN NEGATIVE CASES.

Preliminary Data.

Out of the 116 cases which were under more or less definite suspicion at the post-mortem examination but proved negative

when further investigation was made at the Board's laboratory, 105 had been caught and killed; the remaining 11 had been found dead.

At the time of the post-mortem examination the condition, as regards preservation, of 93 of the above rats was described by Drs. Macalister and Brooks as either "good" or "fair"; the remaining 23 were either "bad" or "putrid."

The post-mortem and microscopic details furnished from the Ipswich laboratory may be classified as follows:—

Group I.—Five cases fall within this group and resemble the positive cases found in Group I. (page 6) in so far as their descriptions are strongly suggestive of plague. Subcutaneous congestion was present in all and was very marked in one. In four cases bipolar bacilli were numerous and generally distributed; in the fifth they were generally distributed in small numbers. With the exception of one liver, necrotic or granular changes are described in the livers and spleens of all the cases. Pleural effusion was found in every case and glandular enlargement in all but one. One rat (4,843) was in bad condition and one (10,510) was found dead.

Group II.—Sixty-one of the negative cases fall into this group. They resemble the positive cases in Group II. (page 7) in that the macroscopic and microscopic data afford slight, but no more than slight, suspicion of plague. The following are the main general facts concerning these 61 cases:—

Macroscopically:—Subcutaneous congestion was present in 39 cases but absent in 22; pleural effusion was present in 42 but absent in 19; mottling or necrotic foci were present in 30 livers but absent in 31; granulation or necrotic foci were present in 26 spleens but absent in 35; enlargement of lymphatic glands was seen in the majority of cases.

Microscopically:—In each case bipolar bacilli were found in some part of the body, though usually they were not numerous; in 54 cases the liver or spleen, or both organs, were examined, and in 53 of these cases bipolar bacilli were found in one or both of these organs.

In eight cases there were found—subcutaneous congestion, bipolar bacilli in the liver and spleen, pleural effusion, and the above-mentioned changes in the liver and spleen.

Eight of the sixty-one rats had been found dead; 13 were "bad" or "putrid" when dissected.

Group III.—This group comprises 50 cases, each presenting some element of doubt at the time of the post-mortem examination, though it was obvious that further investigation was needed before plague infection could be proved or even definitely suspected.

The following are the main features of these cases:—

Macroscopically:—Subcutaneous congestion was present in 18 but absent in 32; pleural effusion was present in 21 but absent in 29; liver changes were present in 25 but absent in 25; spleen abnormalities were present in 21 but absent in 29; enlargement of lymphatic glands was noted in all but 12.

Microscopically:—Bipolar bacilli were only found once in the liver or spleen; they were present in other regions of the body in 33 cases; in 16 cases no bipolar bacilli were found in any part of the body.

Two of the above 50 cases had been found dead; 9 were "bad" or "putrid" when dissected.

Results of Cultures and Animal Experiments.

Cultures were usually made from the spleen; under special circumstances the liver, a lymphatic gland, pleural exudate, or other material was also used.

Cultures were attempted from the original material of all except four of the cases which turned out to be negative, these four being omitted because the material was too putrid to give any prospect of success. In 20 cases the agar media inoculated remained sterile; in 16 the primary cultures were overgrown, so that it was impossible to decide whether or no they contained plague bacilli; and in 76 discrete colonies were obtained, which rendered it possible to decide, either directly or after subculture of any doubtful colonies, that no growth of plague bacilli had been obtained.

The animals inoculated with original material remained healthy in 71 out of the 116 negative cases, and after being kept alive for a sufficient period (from one to two weeks) were killed, and proved by post-mortem examination to be free from disease.

In 12 cases it was not thought necessary to inoculate animals with original material, and in these the microscopic and cultural evidence appeared conclusive. In 2 of these cases the culture media inoculated remained sterile; in 8 discrete colonies, unlike plague, were obtained; and in 1, though the agar cultures were overgrown, cultures on bile salt media yielded discrete colonies but failed to develop any resembling plague.

In the twelfth case, Rat 818, a culture from the submaxillary gland, though more luxuriant than plague, was somewhat sticky in character; it was therefore tested on a guinea-pig and a rat, but failed to produce any disease. It was also observed that the culture fermented raffinose and saccharose.

In 33 cases the original material inoculated produced, in one or more of the animals, lesions, sometimes fatal, which were distinguished from plague either directly by macroscopic and microscopic examination or after further investigation, including the raising of cultures and, where necessary, animal inoculation.

Tabular Statement.

Details of the 33 Negative Cases where Lesions were produced in Experimental Animals.

* The Roman numerals refer to the grouping on p. 18. † "B. Gärtner," throughout this table, means a bacillus belonging to the Gärtner group.

Number and Condition of Rats.		Results of Cultures from Original Material.	Animals.	Results of Animal Experiments.	
Number.	*Condition.				Results.
544	III.	Discrete colonies from spleen and blood.	G.P. 501	Died, 6 days; only slight local reaction, necrotic foci in liver, spleen apparently normal. B. coli isolated from spleen.	
711	II.	Discrete colonies from spleen and liver. B. Gärtner isolated from six.	G.P. 504 Rat 27	Died, 2 days; septic cellulitis. † B. Gärtner isolated from blood and spleen. Killed, 5 days; necrosis at site of inoculation, spleen peppered with minute, irregular, grey foci. B. Gärtner isolated.	
1444	III.	Discrete colonies from spleen, liver, and blood.	G.P. 513 Rat 33	Died, 1 day; acute infection. B. coli isolated from inguinal gland. Killed, 7 days; like Rat 27. B. Gärtner isolated from spleen.	
2353	II.	Discrete colonies from pleural fluid. A coccus isolated.	G.P. 611 Rat 66 (cutaneous.) Rat 67	Died, 3 days; necrotic local lesion and subcutaneous congestion. B. coli isolated. Killed, 6 days; healthy.	
2891	II. (putrid.)	Overgrown.	G.P. 656 G.P. 657	Killed, 6 days; healthy. Died, 1 day; septic cellulitis. Died, 3 days; septic cellulitis. Culture of colon type isolated from spleen.	
3023	II.	Discrete colonies from spleen. Adhesive culture isolated which fermented saccharose.		Rat 113 was inoculated cutaneously with culture isolated from original material. Died, 7 days; grey foci in liver. B. Gärtner isolated from liver.	

3023	II.	Discrete colonies from spleen. Ad- hesive culture isolated which fermented saccharose.	G.P. 658	Died, 1 day; intense oedema and subcutaneous congestion. Culture isolated was inoculated into 2 guinea-pigs which died in 1 and 3 days from acute disease unlike plague, and into 2 rats which remained healthy.
3030	II.	Discrete colonies from spleen. A coccus isolated.	Rat 106	Killed, 8 days: no local lesion, grey foci in liver and spleen. B. Gärtner isolated.
3051	II.	Discrete colonies from spleen. A coccus isolated.	Rat 108	Killed, 8 days; purulent nodule at site of inoculation, grey foci in liver and spleen. B. Gärtner isolated.
3052	II.	Discrete colonies from spleen and liver. A coccus isolated.	Rat 109	Killed, 8 days; grey foci in liver and spleen. B. Gärtner isolated.
3053	II.	Discrete colonies. A coccus isolated	Rat 110	Ditto ditto ditto.
3056	III. (<i>putrid.</i>)	Discrete colonies from liver. A coccus isolated.	Rat 111	Ditto ditto ditto.
3138	II.	Discrete colonies from spleen and liver.	Rat 112	Died, 9 days; no lesions beyond slight local congestion at site of inocu- lation; colon-like bacillus isolated from liver.
3495	III.	Discrete colonies from liver.	Rat 126	Died, 2 days; slight local congestion only. Some bipolar bacilli in spleen, from which colon-like culture was obtained.
3615	III.	Discrete colonies from gland at root of penis, yielding B. coli.	Rat 127	Died, 3 days; no local reaction, grey foci in spleen. B. Gärtner isolated.
3649	II.	Discrete colonies from gland at root of penis. A coccus isolated.	Rat 128	Killed, 13 days; local abscess, spleen speckled with grey foci. B. Gärtner isolated.
3762	III.	Overgrown.	G.P. 687	Died, 2 days; septic cellulitis; microscopically, no plague-like bacilli.
4187	III.	Overgrown.	G.P. 697	Killed, 13 days; normal except 2 yellow foci in spleen. Motile bacilli isolated from spleen.

Number and Condition of Rats.		Results of Cultures from Original Material.	Results of Animal Experiments.	
Number.	*Condition.		Animals.	Results.
4242	II.	Discrete colonies from liver.	G.P. 696 G.P. 695	Died, 3 days; acute cellulitis; no organisms found in spleen. Killed, 13 days; healthy.
4659	II. (<i>putrid.</i>)	Discrete colonies from spleen.	Rat 141	Died, 2 days; subcutaneous oedema, necrosis and congestion. Bipolar bacilli found in spleen and iliac gland. Saccharose fermenter, isolated from spleen, inoculated in doses of 1 mg. into Rat 149 and G.P. 759; both animals killed, 18 days,—healthy.
6427	II. (<i>putrid.</i>)	Discrete colonies from spleen.	G.P. 770	Died, 8 days; small abscess only.
8284	II.	Discrete colonies from spleen.	Rat 163	Killed, 12 days; no local lesion; grey foci in liver and spleen; no bipolar bacilli in spleen.
8439	II. (<i>found dead.</i>)	Discrete colonies from spleen.	Rat 162	Died, 2 days; slight necrosis at site of inoculation, many bipolar bacilli; a few in spleen. Colon-like colonies were isolated from local lesion and spleen of Rat 162. Rat 166, inoculated from local lesion of Rat 162, was killed in 9 days and found healthy.
10150	I. (<i>found dead.</i>)	Overgrown.	Rat 168	Died, 4 days; local necrosis and oedema, a few grey foci in liver; coliform organism isolated.
10549	II.	Discrete colonies from liver.	Rat 173	Killed, 7 days; no local lesion; grey foci in liver and spleen; no bipolar bacilli in spleen.
10678	III.	Discrete colonies from spleen and liver.	Rat 174	Died, 7 days; slight necrosis, but no congestion at site of inoculation; grey foci in liver and spleen. B. Gärtner isolated.
12194	II. (<i>bad.</i>)	Overgrown.	Rat 191	Died, 6 days; local abscess, few grey foci in liver. B. Gärtner isolated.

12520	III.	Discrete colonies from spleen.	Rat 193	Killed, 8 days; slight local reaction; grey foci in liver and spleen. B. Gärtner isolated.
12919	III. (<i>bad.</i>)	Discrete colonies from spleen.	Rat 195	Killed, 7 days; slight local reaction; grey foci in liver and spleen. B. Gärtner isolated.
14638	III. (<i>bad.</i>)	Discrete colonies from spleen.	Rat 214	Killed, 8 days; no local reaction; grey foci in spleen and liver.
14790	II.	Discrete colonies from spleen, yielding B. Gärtner.	Rat 217	Died, 6 days; local necrotic nodule, grey foci in liver and spleen; colon-like culture from spleen.
14792	II.	Overgrown.	Rat 218	Died, 7 days; necrosis at site of inoculation, congestion without oedema; culture from spleen unlike plague.
14793	II.	Overgrown.	Rat 219	Died, 5 days; small necrotic nodule at site of inoculation, no subcutaneous congestion, spleen firm, irregular grey foci in liver. Culture from liver coliform.
15064	III.	Discrete colonies from spleen.	Rat 224	Killed, 7 days; no local lesion, grey foci in liver and spleen.

Bacteria of some Special Interest.

Bacilli belonging to the Gärtner Group.

An organism belonging to this group was isolated sixteen times, thrice from the original material and in thirteen instances from rats inoculated with original material. A feature of these cases was the occurrence in the tissues of bipolar staining bacilli closely resembling the plague bacillus.

The organisms gave the following reactions. Acid and gas were produced in glucose, galactose, fructose, mannite, and dulcitol. In maltose acid was produced, sometimes with and sometimes without gas. In glycerin and iso-dulcitol acid was produced but no gas. No change of reaction occurred in lactose, saccharose, raffinose, inulin, or salicin. Litmus milk became acid at first, but afterwards changed to alkaline. The above test media were kept under observation in every instance for at least 18 days.

Several other cases, in addition to the above 16, produced in experimental rats lesions typical of Gärtner infection, but it was not thought necessary to investigate these culturally.

A Cocco-bacillus from a Rabbit.

In addition to the rats' tissues, specimens from one rabbit were received for examination as being under suspicion of plague.

The animal had been found dead. The post-mortem report stated that there was subcutaneous congestion, a small and granular spleen, a congested liver with necrotic patches, and copious pleural effusion; numerous bipolar bacilli were seen in the liver and the heart's blood, and a few in the spleen.

From the tissues forwarded for examination, the liver and spleen yielded on agar slants pure cultures of a cocco-bacillus; the colonies presented a resemblance to those of plague in their general appearance, but were more translucent; the cultures grew more readily than those of plague and were not sticky.

Rat 212 was killed 13 days after inoculation with an emulsion of the liver of the rabbit. There was an ulcer at the site of inoculation, but no glandular enlargement; with the exception of a few areas of congestion in the lungs, the organs were normal.

G.P. 900 died one day after inoculation with an emulsion of the rabbit's spleen. The local lesion was deeply congested and oedematous, and there was oedema of the adjacent subcutaneous tissue. There was slight excess of peritoneal fluid, but all the internal organs appeared normal. Swarms of very small bipolar bacilli were found in the local lesion, and a few short, thick bacilli in the spleen and blood. A culture from the local lesion proved identical with those obtained from the tissues of the original rabbit.

Rabbit 13 died 3 days after cutaneous inoculation in the ear with a minute dose of culture from the original rabbit's spleen. Post mortem, there was some thickening and bluish redness of the ear; the spleen and liver were both covered with a thin fibrinous layer which was easily removed; irregular, greyish

foci were then found in the substance of each organ; the lungs were congested and oedematous. In the spleen small bacilli were abundant; in the blood small bacilli with well-marked bipolar staining were numerous.

The cultural characteristics of the organism were as follows: it was non-motile, Gram-negative, and formed a deposit in peptone water without producing turbidity. It formed acid, but no gas, in media containing respectively—glucose, galactose, fructose, saccharose, mannite, and glycerin. It produced no change of reaction in media containing—maltose, lactose, raffinose, iso-dulcitol, dulcitol, adonitol, salicin, or inulin; no change was produced in litmus milk or malachite green.

From the above cultural and animal tests, the organism was evidently the bacillus of rabbit septicaemia.

Colonies bearing some Resemblance to Plague.

In 19 of the cases which proved to be negative, the primary cultures contained colonies which, during the first two days, bore some resemblance to plague colonies. They became visible within the first twenty-four hours after inoculation as small, almost transparent points. Subsequently their development was slow; they remained small and transparent, but gradually became slightly raised in the centre. Subcultures grew in the form of discrete colonies, and never produced more than a very thin and transparent layer. After the first two days, therefore, there was no difficulty in distinguishing these growths from plague. As a matter of subsidiary interest, however, they received further investigation.

In every case examined the colonies consisted of a Gram-positive coccus with a tendency to assume slightly elongated forms which might perhaps be called very short bacilli.

The organism, when tested after being in culture for a considerable period, was not pathogenic for mice, no more than a small, purulent focus being formed at the site of inoculation. One strain inoculated subcutaneously into a rat and a guinea-pig failed to produce any disease.

In seven cases the coccus produced acid and clot in litmus milk, and acid, without gas, in glucose, galactose, fructose, maltose, saccharose, lactose, mannite, dextrin, and salicin. No change of reaction was obtained in raffinose, dulcitol, or inulin. In five other cases the organism was a somewhat less active fermenter, saccharose being unattacked in one case, mannite and salicin in a second, mannite and lactose in a third, and mannite, lactose and salicin in a fourth. In the fifth case the coccus had no action on litmus milk, and failed to attack lactose, saccharose, mannite, and salicin.

It is doubtful whether these cocci were responsible for any pathogenic changes in the rats from which they had been obtained, and it is noteworthy, in this connection, that all the experimental rats inoculated with the original material remained healthy. They were all from cases showing marked pleural effusion, but there is no evidence that the cocci were responsible for this condition.

EVIDENCE AS TO DISTRIBUTION AND APPARENT LIMITATION OF INFECTION.

The Area from which Rats were Collected.

The area investigated is shown on the accompanying map, prepared from data furnished by Mr. Huddart.

At the beginning of the enquiry attention was confined to the borough of Ipswich and its environment, the rural districts of Samford and Woodbridge, and the urban districts of Woodbridge and of Felixstowe and Walton, this being the area where plague rats had previously been found.

Six rats obtained during the latter half of July from the parishes of Sutton, Bawdsey, and Trimley St. Mary, in Woodbridge rural district, and from the parish of Bentley, in Samford rural district, proved to be plague infected.

As soon as these cases were fully established, steps were taken to extend the area of enquiry, whilst continuing the search for infected localities in the original area. The enlarged area comprised a larger field in East Suffolk, a small portion of West Suffolk, and a strip of Essex adjacent to the south of Suffolk. Various reasons determined the actual demarcation of the area. To the north it included certain parishes from which reports had been received last year of rats being found dead in unusual numbers; to the north-west it included the whole of Bosmere and Claydon rural district, which had not been searched during the Board's enquiry in the beginning of the present year; to the west it extended far enough to include a parish in which a plague-infected hare had been found last year; and to the south it took in a fairly wide strip of land on the right bank of the Stour, including a parish where a plague-infected hare had been found last year.

Comparison with the Area previously Investigated.

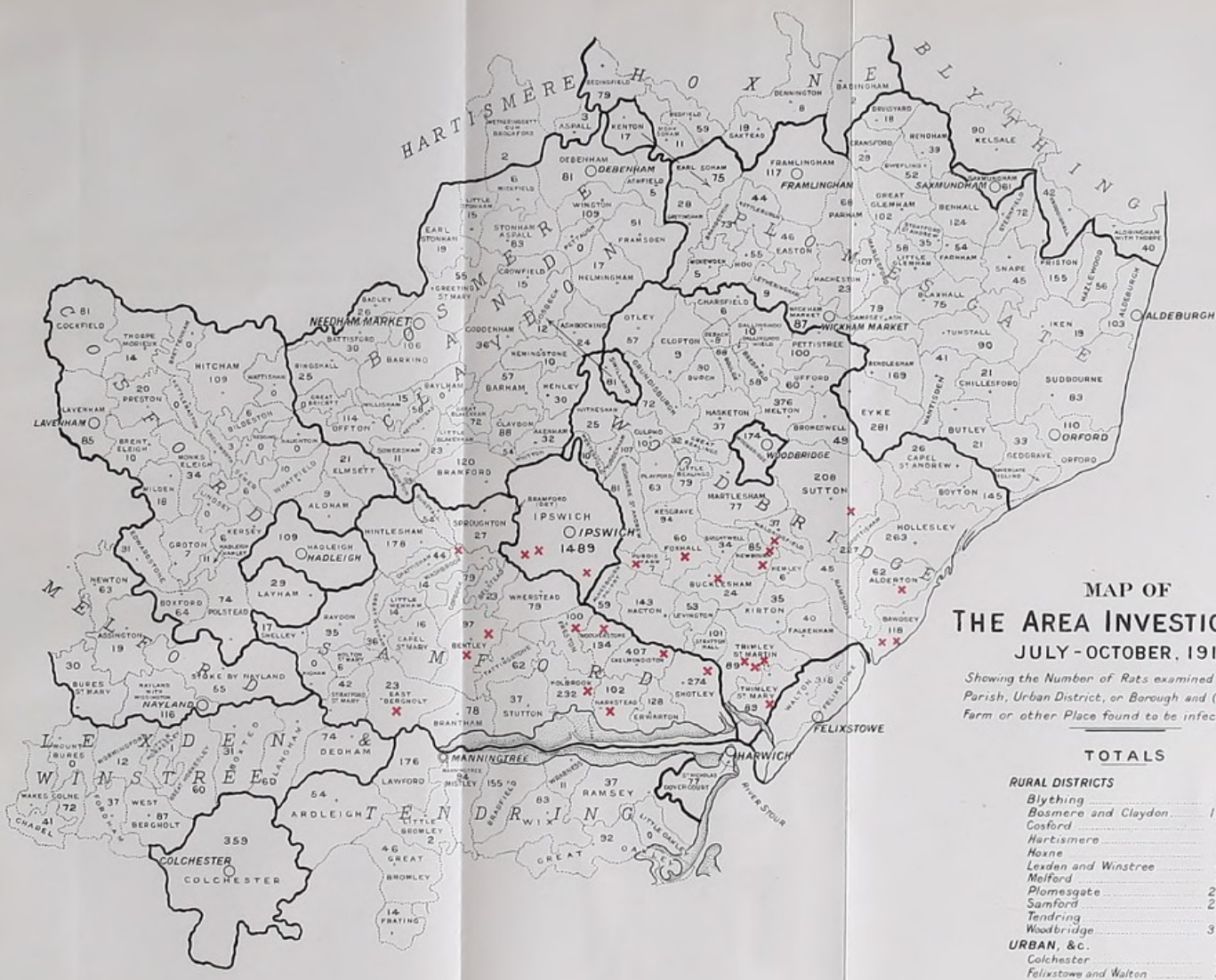
The present investigation has been especially directed to the whole of the area which was not examined during the rat enquiry conducted for the Board during the period January 16th to February 14th, 1911.*

The area referred to consists of the rural districts of Cosford, Samford, Bosmere and Claydon, and Woodbridge, the urban districts of Hadleigh, Woodbridge, and Felixstowe and Walton, and the borough of Ipswich.

In addition to comprising this area, the present enquiry has extended a short distance to the north and to the south. (*See map.*)

The result of the former enquiry was that no plague-infected rats were found in a wide peripheral zone surrounding the area omitted from that investigation. In the present enquiry plague-infected rats have been found only in this last-mentioned area, and only in a limited portion thereof, which corresponds closely

* Reports to Local Government Board on Public Health and Medical Subjects. New Series, No. 52.



**MAP OF
THE AREA INVESTIGATED
JULY-OCTOBER, 1911.**

Showing the Number of Rats examined from each Parish, Urban District, or Borough and (X) each Farm or other Place found to be infected.

TOTALS	
RURAL DISTRICTS	
Blything	172
Bosmere and Claydon	1519
Cosford	648
Hartismere	5
Hasne	178
Laxden and Winstree	475
Melford	283
Plomesgate	2718
Samford	2398
Tendring	764
Woodbridge	3585
URBAN, &c.	
Colchester	359
Felixstowe and Walton	318
Hadleigh	109
Harwich and Dovercourt	77
Ipswich	1489
Saxmundham	61
Woodbridge	174
Total	15,332

871. 3453/66. 126. 7. 12.

Malby & Sons, Lith.



to, but is slightly more limited than, the area proved to be infected with rat plague before the former enquiry was instituted.

The Supply of Rats.

During July and August, the first two months of the present enquiry, rats were, as was anticipated, particularly difficult to obtain. They were dispersed in the open country amongst the standing crops and densely overgrown ditches and hedgerows; and during harvest farmers and their assistants were too busy to spare time for ratting.

There was, however, an important reason for commencing the present enquiry at the beginning of July. Rat-plague usually tends to expand from sporadic outbreaks into epizootic proportions at the beginning of autumn; therefore it was particularly desirable to locate early cases, in order that preventive measures might be directed to the foci where these were found, before the time arrived when rapid spread of the infection was likely to occur.

This anticipation was justified by results. The first rat (No. 1,030) which proved to be plague-infected was caught on July 17th, and by the end of that month infected rats had been secured from five different foci. The Board at once advised the local authorities concerned, and the Board's investigators, as soon as a case was diagnosed, visited the farm or other premises where the case occurred, warned the occupier of the danger, and advised thorough and persistent rat destruction. This procedure was maintained throughout the enquiry, whenever fresh cases of rat-plague were detected.

The abnormally hot and dry weather experienced throughout July and August this year made rat-catching particularly difficult; at the same time it had the advantage of reducing to some extent the normal proliferation amongst rats, owing to their difficulties in obtaining the moisture requisite for their own subsistence and for suckling their young. Another reason for their scarcity was that a large number of farmers throughout the area investigated had been systematically killing rats throughout the year.

In September and October the area of enquiry had been extended, the harvest was over, the rats were beginning to come back to food supplies near farm buildings, and the farmers and their men had more time to interest themselves in ratting. For these reasons there was then no difficulty about obtaining a daily supply of as many rats as could be dealt with in the Ipswich laboratory; in fact, prompt measures had to be taken to reduce the inflow, which towards the end of September became excessive.

For many reasons the numbers of rats taken from each parish (*see* map) varied considerably. Parishes differ greatly in size and in the number of farms which they contain; the rat population is very irregularly distributed; and in some districts the farmers are much more enterprising in rat destruction than in others. Special attention was paid to parishes which were under suspicion last year owing either to the dis-

covery of rat-plague or to local reports of excessive mortality amongst rodents. In some of these parishes no plague could be discovered, although every effort was made to obtain for examination as many rats as could be found.

As soon as a positive case was diagnosed, further supplies were stopped from the parish whence the rat had been obtained, as it was the object of the enquiry to determine the limits of the infection rather than to ascertain how many cases of infection could be discovered in particular areas; and, owing to the migratory habits of rats, the discovery of one infected pocket was considered sufficient to stigmatise its environment, within a radius of several miles, as possibly plague-infected.

Although special search for dead rats was invited and the same price was paid for these as for rats which had been caught, out of the total 15,332 rats only 69 dead rats were brought in. During the September and October of the previous year dead rats were reported in large numbers, particularly within or near to localities where the existence of rat-plague had been proved. All the known pockets where rat-plague had previously been found were re-examined several times during the present enquiry, but in only a few cases were fresh cases of the disease discovered.

ACKNOWLEDGMENTS.

Dr. Klein, F.R.S., has given us valuable assistance in the diagnosis of rat-plague, and, before the commencement of the enquiry, Dr. C. J. Martin, F.R.S., and Dr. Rowland kindly allowed us to make post-mortem examinations of a large number of rats which had died from experimental inoculation with plague bacilli.

The Lister Institute kindly seconded Dr. G. H. Macalister, who had already gained extensive experience in the post-mortem examination of rats, to act as one of our assistants throughout the enquiry. He and his colleague, Dr. Brooks, though engaged upon work which was extremely unpleasant, tiring, and of long duration, always exhibited remarkable skill in the selection of the material which they submitted to us for diagnosis.

Dr. Pringle, Medical Officer of Health for Ipswich, was most helpful in facilitating arrangements at the Ipswich laboratory, the use of which for the purpose of the enquiry must, we fear, have caused him personal inconvenience in the prosecution of his own bacteriological work. Dr. Heath, then Medical Officer of Health for East Suffolk, gave us invaluable advice and assistance throughout the enquiry by placing at our disposal his experience of rat-plague in the district which we were investigating. Dr. Hollis, Medical Officer of Health for Woodbridge rural district, and Dr. Redpath, Medical Officer of Health for Woodbridge urban district, also took great interest in the enquiry and assisted us in the examination of areas likely to prove plague infected.

The Chief Constables of East Suffolk, West Suffolk, Essex, and Colchester gave us every possible facility for the collection of rats through the aid of the police; and the highly efficient

organisation of their forces is responsible in large measure for the maintenance of a satisfactory supply of rats throughout the enquiry. The individual constables undertook the extra work which devolved on them with willingness and enthusiasm, and spared neither time nor trouble in obtaining supplies of rats from the farms on their beats. Special thanks are due to Captain Mayne, Chief Constable of East Suffolk, and his assistant, Superintendent Staunton, for their keen personal interest in the enquiry and their skill in organisation.

Throughout the four months of the enquiry Mr. Gordon Merriman worked daily with the rat-catchers for the purpose of collecting and investigating the fleas from the freshly killed animals. His practical advice, based on the local knowledge thus gained and on his personal experience of ratting, added much to the efficiency of our control over the daily work of the rat-catchers.

SUMMARY.

During the period July 1st—October 31st, 1911, 15,332 rats were examined for plague infection. These rats were obtained (*see* map facing p. 26) partly from the area which in the previous year had been found to be infected with rat-plague and partly from the districts immediately adjacent to this area.

As a result of the present enquiry 27 farms or other premises were found to harbour plague-infected rats.

These 27 places are all within the area previously pronounced to be infected. No plague-infected rats have been discovered outside this area.

Experience of rat collection during the present enquiry showed that the rat destruction which had been maintained by local enterprise since the end of 1910 had in many localities appreciably diminished the rat population.

RECOMMENDATIONS.

At the close of the present enquiry the conclusion appeared to be justified that prompt and persistent rat destruction, systematically conducted throughout the infected area and in the districts peripheral to this area, if maintained for a period of at least 18 months, held out the best prospect of eliminating the disease in its present area of prevalence.

As there is an appreciable risk of rat plague being communicated, directly or indirectly, to human beings, and as a declaration of human plague might involve serious commercial loss, the expenditure requisite for efficient rat destruction in the affected area is strongly to be recommended.

ARTHUR EASTWOOD,
F. GRIFFITH.

APPENDIX.

Report upon the Post-Mortem Examination of Rats at Ipswich by G. H. Macalister, M.A., M.D., Assistant Bacteriologist at the Lister Institute of Preventive Medicine, and R. St. John Brooks, B A., M.B.

INTRODUCTION.

On June 30th, 1911, we proceeded to Ipswich, with instructions from the Local Government Board to assist in an enquiry, the purpose of which was to determine the extent of the plague epizootic amongst rats in the surrounding districts. The share of this work allotted to us consisted in examination, at the Municipal Laboratory, of the animals received, and in selection, for further investigation by the Board's pathologists in London, of material which appeared suggestive of plague. For this purpose, rats presenting features in any degree consistent with the presence of plague were regarded by us as suspicious.

PRELIMINARIES.

The rats were delivered at the laboratory in boxes. These were opened, and the rats taken out and affixed to dissecting boards. Dissections were made so as thoroughly to expose the principal lymphatic glands, thoracic and abdominal viscera. After inspection, rats obviously free from plague and in other respects devoid of interest, were removed to suitable receptacles, pending transference to the destructor. Those of pathological interest, either with regard to the possibility of plague-infection or from other cause, were set apart for further scrutiny.

For purposes of identification and reference, all rats were numbered.

Rats set aside on the strength of gross post-mortem appearances as possibly infected with plague were submitted to a complete microscopical examination. Films from heart-blood, liver, spleen and lymphatic glands, stained with carbol-thionin blue, were searched for the presence of organisms resembling plague bacilli. After this, the evidence, macroscopic and microscopic, was reviewed, and the rat was either rejected or reserved for cultural and inoculation tests. In the latter case, the liver, spleen and any other portions indicated by the post-mortem characters were packed and sent to London, accompanied by cards describing the pathological features presented by the animal from which they were taken. Cultivations were usually taken from the organs before despatch. Rats which had been so treated were then immersed in strong formalin, and removed separately in tins for destruction. A record was kept of other features of pathological interest.

NUMBER OF RATS, &C. EXAMINED.

Summary of Results.

Sexes and Pregnancies:—

Males	7,490
Females	7,342
Pregnant Females	1,213
Average number of Fœtuses	9.9

Species:—

<i>Mus decumanus</i>	15,167
<i>Mus rattus</i>	165*
Ferrets	2
Moles	3
Hares	3
Stoat	1
Weasel	1
Rabbit	1

Plague infection.

—	Killed.	Found dead.	Total.
Rats examined	15,263	69	15,332
Sent up as suspicious, and proved negative.	105	11	116
Proved plague infected by inoculation.	21	14	35

The work was performed by us with the help of two attendants, and we wish to record our appreciation of the valuable and conscientious service rendered by J. Yerrell, our chief assistant.

POST-MORTEM APPEARANCES.

The accompanying tables show diagrammatically the characters of the selected rats, those found upon further enquiry in London to be infected with plague being placed in Table I., whilst particulars regarding those which proved to be negative are set forth in like manner in Table II. A certain number of rats which were sent up and found to be negative are excluded from Table II. for reasons hereinafter stated. (*See* page 34.) The following paragraphs contain some amplification of these tabular statements.

Plague-Infected Rats.

Lymphatic Glands.—Enlargement of glands was present in most infected rats; three only failing to exhibit this feature. It was confined to the submaxillary region in five instances, whilst four rats showed congestion of these glands with no increase in size. On eight occasions, the lymphatic glands

* This number includes 134 rats caught on board ships in the port of Ipswich, and 31 caught in and around the docks.

showed general congestion, but no enlargement. In some of the rats the enlarged glands were associated with minute ecchymoses upon the surface or on section, and definite hæmorrhages were observed in a few cases. These appearances were accompanied in some instances by a peripheral zone of congested tissues surrounding the gland, and paling off into a less-marked degree of general subcutaneous congestion (2,876 and 3,430). In one rat (2,876) cross section of an enlarged and congested inguinal gland displayed a necrotic centre. In another (15,079) the left pelvic gland was necrotic. Involution forms, resembling those of *B. pestis*, were observed in some cases, upon microscopic examination of smears from enlarged glands (2,876, 11,724, 12,013, and 13,663). Bipolar bacilli were demonstrated in 23 cases.

Subcutaneous Congestion.—Congestion, distributed universally over the subcutaneous tissues, was seen in 24 cases. In two instances (2,064 and 2,522) it was described as being confined to the thoracic and axillary regions. In four cases only was it altogether absent. This feature was very generally present to a marked degree in the infected rats obtained during the concluding period of the investigation. In general, the congestion was manifested as a uniform subcutaneous blush due to distension of the capillaries, but in a few cases venous engorgement was most prominent, dilation of individual venules being obvious.

Hæmorrhages.—Small hæmorrhages were found in 10 rats, but in no case was this feature conspicuous. Four instances of retroperitoneal hæmorrhage, the pelvic glands being invaded in two cases, were observed. The lungs showed minute ecchymoses on two occasions. Effusions of blood into the bronchial glands and mesentery, and into the subcutaneous tissues of the axillary and submaxillary regions were also described.

Pleural Effusion.—In three plague-infected rats there was no effusion into the pleural cavity. In one putrid rat turbid fluid was found; four cases showed blood-stained effusion. In all the remaining cases a clear pleural effusion, usually considerable in amount, was observed.

Liver.—The pathological changes exhibited by the liver varied greatly in degree and in quality, variations in size, colour and consistency, surface and surface markings being observed. In two rats, both somewhat decomposed, the liver was not described as abnormal. Three rats (2,064, 5,381, and 13,817) presented livers which appeared at first sight to be normal, but scrutiny with a hand lens showed the presence of fine pitting distributed universally over the surface of the organ, indicative, possibly, of fibrotic change. In a fourth animal (13,663) this pitting was accompanied by signs of congestion. Enlargement, with congestion, was recorded on nine occasions. No register was kept of variations in the degree of consistency of the organ, as this depended to a large extent upon its state of preservation. In 17 cases, the liver is said to be mottled. The term "mottling," as applied to liver changes, is not intended to indicate any specific pathological conditions, but is used to indicate a mottled or marbled appearance of the organ associated

with abnormal definition of the liver lobules. This condition was often shown by livers undergoing putrefactive change (*e.g.*, 10,848).

Granular livers, in which were present necrotic foci appearing as scattered whitish points, were found in 18 cases. In two instances the liver is described as granular but no obvious areas of necrosis were observed. It was noticed in one rat (8,086), which had been found dead, that although the liver showed advanced putrefactive changes, whitish points of necrosis stood out clearly from the discoloured surface. The manner in which these various features—congestion, mottling, and necrotic change—were combined in different cases, is shown in the table; but it may here be stated that the complex to which has been applied the name of 'typical pest liver' (*i.e.*, an enlarged, pale, mottled liver covered with numerous necrotic points) was found only in three cases (2,876, 5,381, and 12,012).

In every animal except one (1,264) microscopical examination showed the presence of bipolar bacilli in liver smears, well-marked involution forms being observed in a number of cases. In two livers (2,522 and 11,407) bipolar bacilli were seen in relatively small numbers, together with a large variety of putrefactive forms. In Rat No. 8,086 referred to above, very large involution forms of the swollen and globular types were observed.

Spleen.—In eight cases the spleen was macroscopically normal. Fourteen rats showed spleens which presented signs of congestion, being dark in colour and firm in texture. Fifteen exhibited various degrees of granular change. Enlarged spleens showing no distinctive pathological change were not regarded as abnormal. In two cases (2,876 and 13,817) coarse punctate necrotic foci were observed. In one rat (1,030) the spleen showed a white infarction.

Microscopical examination showed the presence of bipolar bacilli in 26 cases, enormous numbers being observed in seven of these. Well-marked involution forms were found in several instances (*e.g.*, 8,086). Spleen smears from seven rats did not show any organisms, although with one exception (1,264) these were found in corresponding preparations from the liver. It appeared that there was little correlation between the degree of gross morbid change observed and the number of bacilli seen on microscopic examination. In some cases (3,430 and 14,855) spleens which appeared normal to naked-eye inspection yielded films found to be crowded with organisms. This condition was met with in the case of the liver in two cases (5,382 and 13,817).

Heart Blood.—Films prepared from the heart blood showed no organisms in six cases. The type of organism observed in the remainder was similar to that shown by the smears from liver and spleen. The number varied from very few (in 1,264) to enormous quantities (in 1,694). The small number which was found in 1,264 afforded in that case the only microscopical evidence of plague infection, as the smears from liver and spleen showed no organisms.

Oedema, &c.—Pelvic oedema was seen in one case (1,769). Another rat (10,848) showed the rare condition of general sub-

cutaneous oedema. In rat 13,817 the lungs were markedly engorged, and were found upon microscopical examination to contain large numbers of bipolar bacilli.

Negative Rats.

Particulars are shown in Table II. of 76 rats selected as possibly plague infected, which, however, were shown by cultural and inoculation experiments to be negative. Some of the salient features of these rats are summarised below. In addition to the negative rats included in Table II. 35 rats were sent up for further investigation: 19 of these exhibited abundant pleural effusion containing numerous bipolar bacilli, and are more fully described below; the remaining 16 were found to harbour bipolar bacilli in their accessory sexual glands. In most rats, even those apparently healthy, smears from accessory sexual glands showed large numbers of organisms, which often exhibited bipolar staining and bore a close resemblance to the plague bacillus. Cultural investigation and inoculations carried out in these cases proved the absence of any such infection. As soon as it was ascertained that neither of these conditions was in any way associated with infection suggestive of plague, such animals were regarded as negative.

Lymphatic Glands.—Glandular enlargement was a very common feature, and was particularly well marked in two cases. Bipolar bacilli were found in gland smears from 21 rats, and in some cases organisms resembling plague involution forms were observed. Enlargement of lymphatic glands, particularly those in the submaxillary region, was observed in a great majority of the rats received for examination, but in most cases, when microscopic examination was made, no organisms were found. The pelvic glands also were as a rule conspicuous in the rats examined during the course of this enquiry, although it had previously been observed by one of us who had the opportunity of examining a series of rats in January, that these glands were not usually obvious at that season. In the majority of cases enlargement of the glands was not accompanied by congestion.

Subcutaneous congestion.—Intense and general subcutaneous congestion was only observed twice. Localised congestion was seen in a number of cases. In 22 rats no signs of congestion were observed.

Hæmorrhages.—Hæmorrhages were described as occurring in 11 cases. The majority of these were strictly localised and were probably traumatic in origin.

Pleural Effusion.—Clear pleural effusion was observed in 39 cases, the amount of fluid being scanty in the majority of these. Effusion described as blood-stained occurred in 39 instances. In several decomposed rats turbid fluid was found in the pleural cavity. These numbers include 19 rats not included in Table II. In 12 of these the pleural effusion was blood-stained, and in seven instances it was clear or turbid. Microscopical examination of films prepared from such pleural effusions showed the presence of a great variety of organisms, some of which simulated

very closely both typical and pleomorphic forms of plague bacilli. Further, the groupings of organisms on some films bore a most striking resemblance to those characteristic of *B. pestis*. These organisms were subsequently regarded as being putrefactive.

Liver.—The livers showed punctate foci of necrosis in 16 cases, in five of which the feature was well marked. In four such rats infection with the Gaertner bacillus was subsequently demonstrated. Microscopically no organisms were found in 26 cases; 50 showed the presence of bipolar bacilli.

Spleen.—Granular changes in the spleen were frequently observed. In three animals the spleen showed necrotic foci. Bipolar bacilli were found in 42 rats.

Heart Blood.—Films prepared from the heart blood showed cocco-bacilli or bipolar bacilli in 36 rats.

Minor changes in the viscera were observed in a great number of rats which were rejected at once, as well as in those reserved for microscopic examination. Mottling or marbling of the liver due to commencing putrefaction was very common. Minor granular changes in the spleen, overgrowth of submaxillary glands, and effusions of blood into the pleural cavity were noticed with great frequency. No statistical record of such changes was kept, but, in about 600 such rats, microscopic examinations were made. These gave negative results.

OTHER PATHOLOGICAL CONDITIONS.

Gaertner Infection.

In 14 of the cases, where infection with plague was, upon preliminary examination, considered possible, further investigation demonstrated the presence of bacilli of the Gaertner group. The allocation of these organisms was based upon cultural and fermentation reactions.

Five other rats were noticed, which demonstrated so clearly the post-mortem character associated with Gaertner infection, that they were not included among the plague-suspects. Nineteen rats altogether were, therefore, regarded as being infected with this organism. It is probable, however, as the investigation was not directed definitively to the detection of such cases, that this number does not represent the degree of prevalence of this disease among the rats received.

The most prominent post-mortem features typically associated with Gaertner infection were displayed by the liver and spleen. The former was not as a rule granular, but thickly peppered with white points. It should be noted that although a typical plague-infected liver differs markedly from the appearances presented by the organ in a typical Gaertner infection, in certain cases the post-mortem appearances are of no great assistance in differential diagnosis, and further investigation is needed to determine the nature of the disease under consideration. The spleen, on the other hand, was much more markedly granular than that found in the majority of plague-infected rats. Microscopically the number of organisms seen was usually small, but these exhibited,

not infrequently, bipolar staining. Some slight amount of subcutaneous congestion was shown by a number of Gaertner-infected rats.

Trypanosomiasis.

Trypanosomiasis was very common, particularly during the first two months of the investigation. The most general and prominent indication of this infection was blood-stained pleural effusion consisting in many cases of pure blood. Rats exhibiting such a blood-stained effusion, whilst appearing fairly healthy in other respects, might with some degree of certainty be regarded as infected with trypanosomiasis. A continuous series of 2,000 rats received in August were investigated particularly with regard to this point. Of these, 145 showed pleural effusion which was decomposed and putrid in many cases, and clear in some others, while in 55 cases the fluid was blood-stained: 45 of these showed the presence of trypanosomes. These parasites were also found in five cases where the pleural effusion was turbid. Incidentally, three rats in this series of 2,000 were found to be plague-infected. In these three (2,522, 2,876, 3,346) the effusions were severally turbid, clear, and blood-stained. Other appearances, associated with trypanosomiasis, were congestion of the subcutaneous tissues and lymphatic glands, enlargement of glands, and congestion of the liver. In one case these features were particularly obvious, and the right inguinal gland, particularly, was enlarged and surrounded by a zone of deeply congested tissues. Microscopic examination of this gland showed the presence of many trypanosomes, the agglutination phenomenon being well demonstrated. Large numbers of trypanosomes were found also in the heart blood. This rat was interesting, as showing that the infection was introduced probably within the area drained by the right inguinal gland. It was also very important, in that the post-mortem appearances presented a picture very closely resembling that associated with plague. Agglutination of trypanosomes was, on another occasion, found in a film prepared from the turbid pleural effusion.

Other Infections.

Suppurative conditions of the lungs occurred with great frequency. This phenomenon is extremely common amongst rats, and has been observed on every occasion when large numbers have been examined. In one series, all stages of the condition were seen, from the earliest, presenting a few small foci containing glairy material, to advanced stages in which the lung tissue is almost entirely replaced by caseous matter, the healthy lung which remains being thereby compressed and displaced and the bony wall of the thorax deformed. Suppurations, general and localised, in the abdominal cavity were not infrequent, and abscesses in lymphatic glands were very common. Osteomyelitis of the hind-limb bones was twice found.

Twenty-five cases of rat-leprosy were observed.

DISCUSSION.

Some facts, observed in the study of both negative and positive cases, may illustrate certain difficulties of preliminary diagnosis, which assume their greatest importance in the investigation of outbreaks of rat-plague occurring in this country.

Infected rats presenting atypical appearances were relatively common, particularly during the early part of the enquiry, at the commencement, that is, of the seasonal epizootic. On the other hand, those found during the last two months of the investigation showed features typical of plague. Broadly speaking, preliminary diagnosis during the first period of two months was usually tentative, while during the second period it was, in the great majority of instances, certain. It must not be assumed, however, that this statement implies that negative cases could always be diagnosed as such without resort to animal experiment. Even in regard to cases in which the diagnosis appeared certain, a caveat must be entered. A study of the characters shown by some rats (4,843, 4,886, and 10,150), in Table II., will illustrate the danger of making positive diagnoses upon the evidence afforded by post-mortem features, not confirmed by inoculation experiments. Reference, on the other hand, to three cases (1,264, 2,064, and 11,407), in Table I., will illustrate the necessity of care in the investigation of epizootics where rats presenting such slender evidence of infection may occur. Reports of similar investigations carried out at Hamburg and elsewhere indicate that such atypical plague rats may always be encountered in the investigation of European epizootics.

The individual features of greatest service for diagnosis were subcutaneous congestion, clear pleural effusion, and granular, necrotic changes in the liver. All three were present, usually to a marked degree, in most of the plague-infected rats, but only in 9 rats found to be negative. Pleural effusion, either clear or blood-stained, and congestion was observed in a considerable number of cases. No rat found positive failed to display obviously one or more of these characters. In order of relative value, subcutaneous congestion and pleural effusion would come first, the liver changes being less important, but their collective significance should be considered rather than the individual prominence of each single factor.

Enlargement of lymphatic glands possesses, generally speaking, little weight as primary evidence. At the season in which the enquiry was carried out, pelvic glands were conspicuous in most rats. Submaxillary glands were enlarged as a rule. Congestion in regions where this feature was generally not obvious was of some value as a secondary piece of evidence.

In general the facts observed agree with those which emerge from the great bulk of work which has been published upon the question of plague epizootics.

Reference has already been made to the misleading character of gross morbid changes associated in one case with trypanosomiasis. Enlarged and congested glands, subcutaneous congestion, pleural effusion, engorgement of liver and spleen, all

features suggestive of plague, were shown, though microscopic examination demonstrated readily the real nature of the infection.

In one case a dual infection of pest and trypanosomiasis was present, and it is difficult to say how far each infection was responsible for the post-mortem appearances observed.

The peppery liver associated with Gaertner infections is a crisply defined pathological entity, but in many cases this organ presented appearances which caused trouble in diagnosis. Concurrent microscopic and macroscopic evidence gave in most cases clear indication of the true nature of the infection, but atypical livers and spleens showing signs of congestion with minor necrotic and granular changes were in a number of instances reserved for inoculation purposes. The result of inoculations with such material showed the importance of microscopic examination, for negative results were obtained in every case where no bipolar bacilli were found in the films.

Microscopical examination presents, however, a number of fallacies. Bipolar staining is at times shown in intestinal and putrefactive organisms, as well as by certain bacteria, probably of intestinal origin, which are found in the accessory sexual glands of otherwise healthy rats. Putrefactive organisms may also present forms resembling very closely pleomorphic plague bacilli, and assume groupings usually regarded as characteristic of pest. These were best seen in decomposed pleural effusions. Inoculation of such material from 16 cases—not recorded in Table II.—showed the fallacious nature of such appearances. As previously mentioned, many rats not infected with plague revealed organisms in the liver and spleen which could not be differentiated from plague bacilli on microscopical examination alone.

In the infected rats generally, plague bacilli were less numerous in blood films than in spleen smears, but in a few cases the reverse held true. In some rats received in October plague bacilli were present in enormous numbers in the blood. In one or two of these, the experiment was tried of cutting off an inch of tail and preparing a film from the cut surface of the stump. Films crowded with plague bacilli were thus obtained. This point has a practical application, indicating that the practice, prevalent amongst those who catch rats, of cutting off the tails is not without danger. The cut stump presents a sharp vertebral point which might readily inflict an inoculation scratch.

CONCLUSIONS.

(1.) Many plague-infected rats in the earlier period (July and August) of the epizootic presented atypical appearances.

(2.) The majority of those found in the later period were typical and could usually be diagnosed upon post-mortem examination with a considerable degree of certainty. A number of the negative cases presented difficulty in diagnosis throughout the whole enquiry.

(3.) Features suggestive of plague infection may be simulated by rats infected with trypanosomiasis or with bacilli of the Gaertner group.

(4.) The most valuable diagnostic points are the presence of subcutaneous congestion, pleural effusion, and necrotic changes in the liver. These should be considered collectively.

(5.) Minor granular and necrotic changes in liver and spleen uncorroborated by other suggestive pathological characters, macroscopic and microscopic, are in general not indicative of plague.

(6.) Major granular and necrotic changes in liver and spleen are very suggestive, but not necessarily indicative, of pest infection.

(7.) Fallacies in direct microscopic diagnosis are presented by certain intestinal and putrefactive organisms.

KDKDKKDKKDDKDKKKKKKKDKK

+ + + + +
+ + + + +
+ + + + +

+ + + + +
+ + + + +
+ + + + +

+ + + + +
+ + + + +
+ + + + +

+ + + + +
+ + + + +
+ + + + +

+ + + + +
+ + + + +
+ + + + +

+ + + + +
+ + + + +
+ + + + +

+ + + + +
+ + + + +
+ + + + +

+ + + + +
+ + + + +
+ + + + +

+ + + + +
+ + + + +
+ + + + +

+ + + + +
+ + + + +
+ + + + +

+ + + + +
+ + + + +
+ + + + +

+ + + + +
+ + + + +
+ + + + +

+ + + + +
+ + + + +
+ + + + +

+ + + + +
+ + + + +
+ + + + +

+ + + + +
+ + + + +
+ + + + +

7643
8086
10169
10840
10848
11407
11724
12012
12013
12245
13663
13712
13713
13817
14499
14816
14854
14855
14911
15079
15219

TABLE II.—NEGATIVE CASES.

Serial No.	Macroscopic Appearances.										Microscopic Appearances.				Killed or found dead.
	Glands.		Subcutaneous congestion.	Haemorrhages.	Pleural effusion.	Liver.			Spleen.		Glands.	Liver.	Spleen.	Heart blood.	
	Enlargement.	Congestion.				Necrosis.	Motting.	Congestion.	Necrosis.	Granulation.					
51	+	-	±	-	++*	-	+	+	-	-	+	+	-	+	K
92	+	-	-	-	-	-	+	+	-	-	+	+	-	+	K
102	+	-	-	-	-	-	+	+	-	-	+	+	-	+	K
334	+	-	-	-	-	-	+	+	-	-	+	+	-	+	K
444	+	-	-	-	-	-	+	+	-	-	+	+	-	+	K
544	+	-	-	-	-	-	+	+	-	-	+	+	-	+	K
550	+	-	-	-	-	-	+	+	-	-	+	+	-	+	K
669	-	-	+	+	++*	-	+	+	-	-	+	+	-	+	K
711	+	-	+	+	++*	-	+	+	-	-	+	+	-	+	K
723	+	-	+	+	++*	-	+	+	-	-	+	+	-	+	K
752	+	-	+	+	++*	-	+	+	-	-	+	+	-	+	K
818	+	-	+	+	++*	-	+	+	-	-	+	+	-	+	K
920	+	-	+	+	++*	-	+	+	-	-	+	+	-	+	K
1444	+	-	+	+	++*	-	+	+	-	-	+	+	-	+	K
2557	+	-	-	-	++*	-	+	+	-	-	+	+	-	+	K
2651	+	-	-	-	++*	-	+	+	-	-	+	+	-	+	K
2952	-	-	+	+	++*	-	+	+	-	-	+	+	-	+	D
2994	-	-	+	+	++*	-	+	+	-	-	+	+	-	+	K
3023	+	-	+	+	++*	-	+	+	-	-	+	+	-	+	K
3030	+	-	+	+	++*	-	+	+	-	-	+	+	-	+	K
3031	-	-	-	-	++*	-	+	+	-	-	+	+	-	+	K
3051	+	-	-	-	++*	-	+	+	-	-	+	+	-	+	K
3052	-	-	-	-	++*	-	+	+	-	-	+	+	-	+	K
3053	-	-	-	-	++*	-	+	+	-	-	+	+	-	+	K
3056	+	-	+	+	++*	-	+	+	-	-	+	+	-	+	K

Serial No.	Macroscopic Appearances.										Microscopic Appearances.				Killed or found dead.
	Glands.		Subcutaneous congestion.	Haemorrhages.	Pleural effusion.	Liver.		Spleen.		Glands.	Liver.	Spleen.	Heart blood.		
	Enlarge-ment.	Conges-tion.				Necrosis.	Mot-ting.	Conges-tion.	Necrosis.					Granu-lation.	
11805	+	-	++	-	-	+	-	+	+	+	-	+	+	D	
12005	+	-	++	-	+	+	-	+	+	+	-	+	+	K	
12194	+	-	-	-	-	-	-	+	+	+	-	+	+	K	
12520	+	-	-	-	-	-	-	-	+	+	-	+	+	K	
12919	+	-	-	-	-	-	-	-	-	-	-	+	+	K	
12932	+	-	+	-	-	-	-	-	-	-	-	+	+	K	
13832	+	-	++	-	-	-	-	-	-	-	-	+	+	K	
13835	+	-	+	-	-	-	-	-	-	-	-	+	+	K	
13891	-	-	-	-	-	-	-	-	-	-	-	+	+	K	
14594	-	-	+	-	-	-	-	-	-	-	-	+	+	D	
14628	+	+	+	-	-	-	-	-	-	-	-	+	+	K	
14688	+	-	-	-	-	-	-	-	-	-	-	+	+	K	
14641	+	-	-	-	-	-	-	-	-	-	-	+	+	K	
14704	+	-	-	-	-	-	-	-	-	-	-	+	+	K	

