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EXPERIMENTAL EPIDEMIOLOGY
OF COCCIDIOIDOMYCOSIS:
1. EPIZOOTIOLOGY OF
NATURALLY EXPOSED
MONKEYS AND DOGS

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OCTOBER 1965

UNITED STATES ARMY
BIOLOGICAL LABORATORIES
FORT DETRICK
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October 1965
In conducting the research reported here, the investigators adhered to "Principles of Laboratory Animal Care" as established by the National Society for Medical Research.
FOREWORD

The work reported here was performed under contract with the University of Arizona. Mr. Converse, Mr. Snyder, and Dr. Ray are employees of the U.S. Army Biological Laboratories; Drs. Reed and Kuller, and Mr. Trautman are employed by the Department of Animal Pathology, College of Agriculture and Agricultural Experiment Station, University of Arizona.
ACKNOWLEDGMENT

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ABSTRACT

Animals from constant populations of monkeys and dogs (24 of each divided among 3 exposure sites) housed for 1 year in the open in a known endemic area for coccidioidomycosis (Tucson, Arizona) were removed, upon contracting infection with Coccidioides immitis, to air-conditioned quarters for further observation and were immediately replaced at the exposure sites with other susceptible animals. Periodic soil and air samples were obtained, and appropriate climatic data were recorded throughout the 1-year period. Clinical and laboratory observations were continuously recorded for all animals, and complete necropsies were performed at the termination of the experiment. Approximately 15% of the monkeys and 58% of the dogs became infected, the majority during the cooler months. Comparison of the pathogenesis of the disease in the naturally infected monkeys with that in experimentally infected monkeys indicated a natural airborne infectious dose of probably less than 10 arthrospores. The infection rate, as well as the extent of disease, in the naturally infected dogs was greater than in either the naturally infected monkeys or the experimentally infected dogs, and was attributed to their contact with the ground. The lack of mortality in the naturally infected animals of either species indicated very low natural infectious doses of Coccidioides immitis. The ecological and climatic parameters of this study were similar to those of other studies in the same general area.
I. INTRODUCTION

The endemic areas of coccidioidomycosis, and the climatic and geo-
physical conditions necessary for growth of Coccidioides immitis in the
soil in these areas, have been firmly established. Among the major con-
tributions to these findings have been the very thorough epidemiological
studies of Dr. C.E. Smith and his associates, correlating rainfall and
dry, dusty atmospheric conditions with season morbidty rates; Hugen-
holtz's study of the optimal climatic factors for growth of the organism
in the soil; the extensive studies of Dr. Roger Egeberg and co-workers,
associating salt content of the soil at various seasons with optimal
growth conditions for \textit{C. immitis}; and a demonstration of the close
association of the boundaries of the Lower Sonoran Life Zone with those
of the known endemic areas, by Dr. Keith Maddy.4

Although infections with \textit{C. immitis} result chiefly from inhalation
of arthrospores picked up by wind from the soil, it is extremely diffi-
cult to isolate the organism from air samples. Consequently, the size
of the infectious dose received in nature remains a matter of conjecture.
The purpose of this study was an attempt to determine the approximate
dose received by man in natural infections, by the use of laboratory
animals as "biological air-samplers."5

We have at our disposal a large amount of data on the pathogenesis
of coccidioidomycosis in monkeys and dogs exposed to graded respiratory
doses of \textit{C. immitis} of 10 to 80,000 arthrospores. In our experience, the
dog was as susceptible to the disease as the monkey, but more resistant
to its effects, through its ability to maintain a blood supply to the
lesions for a longer period of time and by a faster and more prolific
collagen response. It was postulated that the monkey was most susceptible
to the ravages of the disease; the dog least affected; with man somewhere
on the scale between the species, probably much closer to the dog than to
the monkey. It seemed logical that a comparison of the pathogenesis
(clinical and laboratory observations) of the disease in animals exposed
in the open, in an endemic area, with that of animals receiving known
experimental doses might lead to a valid estimation of the infectious
dose received by man in nature.
II. MATERIALS AND METHODS

A. SELECTION OF EXPOSURE AREA

The Tucson area in southern Arizona, lying in the heart of the endemic area for coccidioidomycosis, was chosen as the exposure site. The infectivity for man in this area approaches 70\% in long-time residents, and for cattle is closer to 80\%.

Dr. Raymond E. Reed of the Animal Pathology Department of the University of Arizona, a recognized authority on both natural and experimental coccidioidomycosis in dogs, agreed to supervise this study. The facilities of the College of Agriculture of the University of Arizona were available for the project.

B. PHYSICAL SET-UP OF EXPOSURE SITES

Three chain-link fenced-in areas, approximately 100 to 200 feet apart, were arranged in a shallow arc, affording each pen exposure to the prevailing wind (Fig. 1 and 2). The pens were located at the University's Casa Grande Farm in the Santa Cruz River basin (a venturi-like geographical site, which funnels the prevailing winds through the area). This farm contains feed-lots, in which practically all cattle imported from non-endemic areas eventually become infected with C. immitis.

In each of the three enclosures (Fig. 3), 8 dogs were allowed free run in the area (36 x 40 ft), and 8 monkeys (Macaca mulatta) were confined in open cages (26 inches above ground) under appropriate shelter (Fig. 4). All animals remained at the open exposure sites for a period exceeding 1 year unless they became infected with C. immitis, at which time they were immediately removed to air-conditioned quarters at the University's Campbell Avenue Farm, for further clinical and laboratory studies, and were replaced at the exposure site with reserve, susceptible animals.

C. PROCEDURES FOR DETERMINATION OF INFECTION

Two dogs and two monkeys from each pen were subjected to coccidioidin dermal sensitivity tests, immunodiffusion precipitin tests, and thoracic radiographs each week, thus providing a population observation turnover once every month. Tests were immediately repeated on all animals each time an infection was noted. All animals were critically observed, daily, for clinical symptoms of infection.

Controls for the study consisted of 10 monkeys and 10 dogs, inoculated intratracheally with either 10 or 100 C. immitis arthrospores and isolated from the area under study.
Figure 1. Aerial View of Casa Grande Exposure Area. The three exposure sites (location of pens) are indicated by arrows. Note dry bed of Santa Cruz River.

Figure 2. View of Same Area as in Figure 1. Note Santa Cruz River meandering through center of photograph. Exposure pens are at upper left, near horizon.
Figure 3. View of One of the Three Exposure Pens. The solid-appearing structure extending up the fence from the ground is made of louvered aluminum and permits the entry of wind and dust. The roofed structure visible inside the fence is the monkey shelter.

Figure 4. Close-Up View of Monkey Shelter. Note automatic bottled-gas heater, extending up through center of cage battery, that furnishes heat during severe weather conditions. The battery consists of 4 cages (housing 2 monkeys each) completely open on 4 sides and partially open on a 5th side. Note one of the oil-drum dog shelters in the background. The dogs have free run of the fenced area (36 x 40 ft.).
D. METEOROLOGICAL DATA

Climatic factors such as maximum and minimum temperature, relative humidity, wind speed and direction, and rainfall were recorded daily.

E. SOIL AND AIR SAMPLING

Four to eight surface and subsurface soil samples were collected, bimonthly, in and around the exposure pens. These were plated directly on Mycobiotic agar, and injected (1:10 dilution of soil suspension) into 6 mice (intranasally and intraperitoneally), for recovery of \textit{C. immitis}.

Mycobiotic agar plates were exposed, daily, in and around the exposure pens, for recovery of \textit{C. immitis} from the air.

F. PATHOLOGY STUDIES

At termination of the experiment (approximately 52-54 weeks), all animals were subjected to complete necropsies; gross pathology was recorded, impression smears and cultures made from the lungs and any suspicious lesions, and histological sections of all tissues were stained with routine and specific fungal stains.

III. RESULTS AND DISCUSSION

The meteorological factors (Table 1) in the area under study (throughout the period Oct. 1963 to Oct. 1964) were similar to those of former studies in the same general region. Maddy\textsuperscript{4} and Hugenholtz\textsuperscript{2} reported mean July temperatures of 80 to 90 F over a period of years at various locations, as compared with 88 for the present study, with mean maximum, and peak temperatures of 105 and 110 degrees, respectively, as compared with 101 and 110 in our study. Mean January temperatures of 50 to 55 degrees in the former studies were in accord with the 47 F temperature at Casa Grande Farm. The average yearly rainfall of 9 inches reported by Maddy, and 6 to 10 inches by Hugenholtz, compared favorably with the 12.5 inches recorded by us. The only factors that might have affected this study adversely were mean winter temperatures about 10 degrees below normal, with a very late, cold spring.

Prevailing winds varied alternately from the southwest and from the southeast. All attempts to isolate \textit{C. immitis} from the air failed, although two soil samples collected in August 1964 and two in Oct. 1964 were positive for the fungus.
TABLE 1. COMPARISON OF WEATHER CONDITIONS IN THREE EPIDEMIOLOGICAL STUDIES

<table>
<thead>
<tr>
<th>Observation</th>
<th>Maddy a</th>
<th>Hugenholtz b</th>
<th>Present Study</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mean July Temperature</td>
<td>90</td>
<td>80-90</td>
<td>88</td>
</tr>
<tr>
<td>Mean Maximum Temperature</td>
<td>105</td>
<td>101</td>
<td></td>
</tr>
<tr>
<td>Peak Temperature</td>
<td></td>
<td>110</td>
<td></td>
</tr>
<tr>
<td>Mean January Temperature</td>
<td>50</td>
<td>50-55</td>
<td>47</td>
</tr>
<tr>
<td>Minimum Temperature</td>
<td></td>
<td>14</td>
<td></td>
</tr>
<tr>
<td>Rainfall (inches per year)</td>
<td>5-20</td>
<td>6-10</td>
<td>12.5</td>
</tr>
<tr>
<td>(Avg. 9)</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Five of 34 monkeys and 29 of 50 dogs became infected during the 12-month exposure period (Fig. 5 and Table 2), the majority of the animals (25/34) during the cooler months of Nov. through March.

Each increase in infections occurred after a cycle of rainfall followed by a comparatively dry period, as has been previously reported. However, the low infectivity rate (unexpected) during July and August may have been due to excessive rainfall (4.5 and 2.5 inches, respectively).

Only three of the five infected monkeys demonstrated positive serological titers and only two monkeys exhibited histological lung changes indicative of coccidioidomycosis (and these of a very minor nature). Lung cultures of all monkeys were negative for C. immitis. The five infected monkeys remained in good health during the 12-month period, showing no clinical symptoms of disease; their serological titers were comparatively low (mean maximum of 1-8, with a range of negative to 1-64); and very little evidence of infection was noted in the X-rays.

A comparison of these data with those for experimentally infected monkeys (respiratory doses of from 10 to 300 arthrospores) in several former studies (Table 3) indicated a very small natural infectious dose (probably in the range of 10, or less than 10, arthrospores). This undoubtedly represented airborne infection, since the monkeys were housed 26 inches above ground level.
Figure 5. Correlation of Weather Conditions with Infection with C. jejuni.
Cross-hatched bar indicates monthly rainfall in inches; solid bar, number of infections; and solid lines, mean maximum and minimum temperatures for each month. The figures in parentheses represent relative humidity at the temperature indicated.
<table>
<thead>
<tr>
<th>Animal</th>
<th>Number Exposed</th>
<th>Number Infected</th>
<th>Number Positive Skin Test</th>
<th>Number Positive Serology</th>
<th>Number Showing Pathological Changes</th>
<th>Number Positive Lung Cultures</th>
<th>Average Days to Infection</th>
</tr>
</thead>
<tbody>
<tr>
<td>Dog</td>
<td>50</td>
<td>29</td>
<td>23</td>
<td>23</td>
<td>19</td>
<td>3</td>
<td>86/ (10-357)</td>
</tr>
<tr>
<td>Monkey</td>
<td>34</td>
<td>5</td>
<td>5</td>
<td>3</td>
<td>2</td>
<td>0</td>
<td>88</td>
</tr>
</tbody>
</table>

Very minor

a. 10-20 days in peak infection period (Jan. to March). Figures in parentheses indicate span.
<table>
<thead>
<tr>
<th>Dose /</th>
<th>Mean / Maximum Titer</th>
<th>% Mortality</th>
<th>Dose /</th>
<th>Mean / Maximum Titer</th>
<th>% Mortality</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>1 - 8 (Neg. - 64)</td>
<td>0</td>
<td></td>
<td>1 - 16 (Neg. - 512)</td>
<td>0</td>
</tr>
<tr>
<td></td>
<td>10 (64 - 256)</td>
<td>40</td>
<td></td>
<td>10</td>
<td>-/</td>
</tr>
<tr>
<td></td>
<td>50 - 100 (128 - 256)</td>
<td>30</td>
<td></td>
<td>100</td>
<td>-</td>
</tr>
<tr>
<td></td>
<td>300 (128 - 1024)</td>
<td>40</td>
<td></td>
<td>1000</td>
<td>47</td>
</tr>
</tbody>
</table>

a. Numbers refer to experimental aerosol arthrospore doses.\(^8, 7\)
b. Immunodiffusion precipitin test. Numbers in parentheses indicate spread.
c. Numbers refer to intratracheal arthrospore dose (Reed, Raymond K., personal communication).
d. Test not made.
The infection rate, as well as the extent of the disease, was much greater in the naturally infected dogs than in the naturally infected monkeys, indicating a larger infectious dose in the dogs. This was attributed to their contact with the soil, their habit of constantly digging and fighting, with the consequent stirring up of dust, and particularly the fact that the majority of dog infections (19/29) occurred in one of the three exposure pens; all of which indicated that the dogs were becoming infected from contact with the ground, rather than from normal airborne aerosols.

Of the 29 naturally infected dogs (Table 2), 23 exhibited positive skin tests, 23 positive precipitin titers, 19 showed histological lung changes due to C. immitis, and 2 had C. immitis isolated from the lungs by culture. Four of the 29 were diagnosed by histological studies only; all clinical and laboratory tests were negative.

No deaths resulting from coccidioidomycosis occurred in either species. Clinical signs and symptoms of the disease were lacking in the monkeys, and very mild in the dogs, all of which indicated low infectious doses received by the naturally exposed animals.
LITERATURE CITED


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