proteins such as plasma-albumin? The acidic non-steroidal anti-inflammatory drugs displace a number of compounds that are bound to albumin, so this factor might complicate the whole issue. I wonder if you have any evidence in relation to this?

**DR. BERRY** McCarty, Polcyn, Collins, and Gottschalk (1970) found that it was albumin-bound, but we were unable to prove this ourselves. There was no evidence that drugs displaced technetium from protein binding.

**PROF. J. J. R. DUTHIE** (Edinburgh) It would not be inappropriate to say that many of our young men have devoted a vast amount of energy and expertise to acquire information which might be obtained more readily by asking the patient a simple question, namely 'Do you feel better?'

**DR. HUSKISSON** It is because of the unreliability of this technique in the past that we are spending so much time and energy to find a better method.

**A. SPEAKER** We have been studying the metacarpal-phalangeal joints using technetium, and have found in some patients, who have been on large doses of salicylates, that, although there has been a clinical improvement, the technetium count rate has in fact increased. This may reflect displacement of technetium from its binding sites by the salicylates.

**References**


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**Search for Mycoplasma in Synovial Fluids from Patients with Rheumatoid Arthritis.** By G. D. WINDSOR, A. NICHOLLS, R. N. MAINI, R. LEMCKE, D. G. EDWARD, and D. C. DUMONDE (Kennedy Institute of Rheumatology, Wellcome Research Laboratories, and Lister Institute of Preventive Medicine, London)

This communication describes attempts to isolate mycoplasmas from the joints of patients with rheumatoid arthritis. Reports of culture isolations have included different strains of mycoplasmas (Bartholomew, 1965; Jansson and Wager, 1967; Williams, 1968) and tiny colonies suspected to be a 'mycoplasma' (Jansson, Mäkiisara, Vainio, Vainio, Snellmann, and Tuuri, 1971). One interpretation of the lack of uniformity of observations has been that the organisms were tissue culture contaminants or commensals of doubtful clinical significance. However, the findings of Williams and his colleagues of a particular species, *Mycoplasma fermentans*, in 40 per cent. of synovial fluids from patients with rheumatoid arthritis using sucrose gradient separation techniques, accompanied by an apparent specific in vitro cellular immunological reaction to the organism has led to speculation as to its possible aetiological significance (Williams, 1970; Williams, Brostoff, and Roitt, 1970).

73 samples of synovial fluids and nine membranes were obtained from patients with rheumatoid disease (including those with arthritis of recent onset) and with other conditions causing joint effusions. The samples were investigated for the possible presence of mycoplasmas:

(a) By direct culture in solid and liquid media. Some of the specimens were cultured in two independent laboratories.

(b) A sucrose gradient of a similar composition to that described by Williams (1968) was used to separate synovial fluid into visible bands by ultra-centrifugation, and 0.2 ml. volume of material from each band was cultured in liquid and solid medium.

No mycoplasmas were isolated by the direct technique from fourteen synovial fluids and homogenates from nine synovial membranes. The results of examining 69 synovial fluids on sucrose gradients are summarized below:

<table>
<thead>
<tr>
<th>Fluids</th>
<th>RA</th>
<th>Non-RA</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number synovial fluids</td>
<td>64</td>
<td>5</td>
</tr>
<tr>
<td>Visible bands cultured</td>
<td>141</td>
<td>7</td>
</tr>
<tr>
<td>Mycoplasma isolations</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Achromobacter isolations</td>
<td>13</td>
<td>0</td>
</tr>
</tbody>
</table>

A colour change to an acid pH was observed in the liquid medium of cultures of sixteen bands which on subculture yielded an organism belonging to the Achromobacter species. This was regarded as a contaminant as further isolation of these bacteria was prevented by suitable precautions. Mycoplasmas were not isolated from any of the samples examined.

**Discussion**

**DR. D. TAYLOR-ROBINSON** (London) What I have to say confirms what has just been said. I shall briefly summarize about 7 years' work; my colleagues and I have been looking for mycoplasmas in joints since 1965. We started in Salisbury obtaining specimens from a variety of places and after moving to Northwick Park Hospital in 1970 we continued the studies. Synovial membranes and synovial fluids have been examined, and although there were a range of diagnoses, rheumatoid arthritis comprised 109 of 146 cases. Most of these were acute exacerbations of chronic disease, and very few of them early cases. From one case of rheumatoid arthritis, we isolated a glucose-fermenting mycoplasma which was not identified further. In addition, we isolated T-mycoplasmas from the joints of two patients having polyarthritis and psoriasis. This was work done in collaboration with Dr. E. M. C. Dunlop and Prof. Barrie Jones. It is interesting that about 6 months ago Dr. P. Gill (McGill University) told me that he had isolated a T-mycoplasma from the joint of a boy suffering from Still's disease. In collaboration with clinicians at Taplow (Prof. Bywaters, Dr. Ansell, Dr. Gumpel, and Dr. Swannell) specimens have been obtained and divided, a portion being examined by us and a portion by Miss Auriol Hill at the Laboratory Animals Centre, Carshalton. The isolation of a mycoplasma from one patient was of particular interest, because there was a 16-year history of agammaglobulinaemia. Three years before the synovial membrane specimen was taken, a diagnosis of nodular rheumatoid arthritis had been made. At the time of admission to hospital the patient had a respiratory infection and an acute exacerbation of arthritis. The mycoplasma was identified as *Mycoplasma pneumoniae*. In view of the history of respiratory infection, the identification of the mycoplasma and its isolation in two separate laboratories, it is extremely unlikely to be a laboratory contaminant. Finally, my colleague Andrew Campbell has
examined specimens not only by conventional mycoplasma isolation methods but also by the sucrose density gradient technique. In no case was a mycoplasma isolated. In summary, it seems to me that it is possible very occasionally to isolate a mycoplasma from a joint. This does not seem unreasonable in a patient who has an immunity defect such as agammaglobulinemia. On the basis of these studies I cannot believe that Mycoplasma fermentans is present in a large proportion of joints. I think, however, it is worth remembering that, in mycoplasma-induced arthritis in animals and birds, the organisms disappear although the manifestations of the disease continue. Although my enthusiasm for this sort of study is waning, I firmly believe the continued studies should be directed at looking at very early cases of rheumatoid arthritis where these can be obtained and certainly not at the chronic disease.

DR. K. N. LLOYD (Cardiff) I am a clinician and worked with Dr. Williams for some years. I think that there may be a transport problem here. Our specimens came from the patient and were straightaway stored at —70°C. and examined at leisure, can I ask at what temperature your specimens were transported by post?

DR. WINDSOR As the synovial fluid has to be centrifuged for 2 hours at 4°C, we thought it reasonable to transport them on ice. Transport time was about 2 hours. There were some synovial fluids that had been stored at —70°C but the later ones were cultured direct.

DR. K. N. LLOYD (Cardiff) Concerning the band found on the sucrose density gradient, have you looked at it under the electron microscope to see whether there are any mycoplasma-like bodies?

DR. WINDSOR I should think that cultural techniques are more sensitive than electron microscopy; we can put Mycoplasma fermentans into synovial fluids and then recover one or two particles per ml. were the isolation techniques good enough.

DR. K. N. LLOYD (Cardiff) Dr. Taylor-Robinson and Dr. Mervyn Williams have stressed that the high isolation rate came from the early case. In only ten of your patients was disease duration 1 year or less. Finally, what was the clinical activity of these patients and in particular were they on gold therapy, because we know that gold is mycoplasmacidal?

DR. MAINI All the patients had active rheumatoid arthritis when the synovial fluid specimens were obtained, and they included ten whose arthritis was of very recent onset (a few months to one year). None of the patients was receiving drugs known to suppress mycoplasma growth, such as gold, chloroquine, or antibiotics.

DR. D. TAYLOR-ROBINSON (London) Dr. Dourmashkin has looked at some of these specimens by electron microscopy. If you look at the band from a gradient by negatively staining it, then you can see anything you wish to. What you have to do is to examine sectioned material. The best approach is to take a synovial fluid, centrifuge it, and section the deposit. When we isolated Mycoplasma pneumonia there was not the slightest doubt about it. The difficult problem is looking at specimens when mycoplasma have not isolated. Occasionally you can see things that strongly resemble mycoplasmas but there is no way of knowing whether they are or not.
Search for mycoplasma in synovial fluids from patients with rheumatoid arthritis.

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