SEROLOGICAL EVIDENCE OF CANDIDA INFECTION 
AFTER OPEN-HEART SURGERY

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Candida endocarditis is a well-recognised complication of open-heart surgery (Jamshidi, Pope and Friedman, 1963; Climie and Rachmaninoff, 1965; McConnell and Roberts, 1967; Newsom, Lee and Rees, 1967; Conway et al., 1968), but its diagnosis presents difficulties. Even when infection is established, blood cultures may fail to yield candida. When, on the other hand, the diagnosis is in doubt, the isolation of Candida species may be of little significance without other evidence of infection, because of the frequency with which these organisms are found in mucocutaneous situations (Mackenzie, 1961; Marples, 1966) from which they may contaminate specimens. Clearly a reliable serological test would be of great value.

Winner (1955) concluded from a study of over 2500 subjects that a high agglutination titre was of little or no diagnostic significance in a case of suspected candida infection. However, Stallybrass (1964) and Taschdjian et al. (1964a and b; 1967) have produced evidence that precipitins to somatic antigens of Candida species seldom appear in the absence of infection. Our observations made on patients undergoing open-heart surgery suggest that the finding of high levels of precipitins to candida may be indicative either of candida endocarditis or inapparent candida infection.

METHODS

Sera

We have examined 409 sera for the presence of agglutinins and precipitins to Candida albicans, C. tropicalis, C. guilliermondii and C. parapsilosis. The study began when, between Nov. 1965 and May 1966, sera were received from four patients (M.B., C.B., J.C. and R.H.) in whom a clinical diagnosis of candida endocarditis had been made. When precipitins to Candida species were detected in them, the study was extended to another 64 patients who underwent open-heart surgery between June 1966 and Sept. 1967; two of these patients (D.S. and J.R.) developed candida endocarditis. As far as possible blood was taken immediately before operation and thereafter at weekly intervals. Agglutination and gel diffusion precipitin tests were carried out on all sera.

Agglutination tests

Agglutination tests were carried out with suspensions of whole yeast cells. Candida species were grown on glucose peptone agar for 48 hr at 37°C, washed off with isotonic saline and made into stock suspensions of approximately 50 per cent. packed-cell volume, which

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were kept frozen. When required, the stock suspension was diluted in saline to 2 per cent. packed-cell volume by haematocrit estimation, formaldehyde was added to a concentration of 0·08 per cent. and the suspension allowed to stand overnight. The following day, the yeasts were centrifuged out, washed once and resuspended in saline to a concentration of 2 per cent. (v/v). Doubling dilutions of the sera from 1 in 4 were made in saline in 0·6 ml volumes and to each dilution was added 0·06 ml of a 2 per cent. suspension of candida. The serum-yeast mixtures were incubated at 37°C for 3 hr and then stored at 4°C overnight. Agglutination of the yeast cells was readily seen with oblique lighting against a black background.

Precipitin tests

Antigens for precipitin tests were prepared by one or other of two methods according to the availability of apparatus. Washed cells of _C. albicans_ suspended in saline were forced through a hydraulic press (X-press, AB Bion, 25-ml model) and the other three species were disrupted overnight in a Mickle shaker. Suspended solids were removed by centrifugation and the liquid phases were freeze-dried. Working antigens were prepared by dissolving the dried material in a buffer at pH 8·6 (boric acid 6·7 g and sodium borate decahydrate 13·4 g per litre of distilled water) to concentrations of about 50 mg per ml. The optimal concentration for precipitation was determined by testing each antigen against serum from a rabbit hyperimmunised with the homologous species of _Candida_; this was done to try to obviate variations between the batches of antigen. Trial batches of _C. albicans_ antigen prepared by the Mickle shaker method gave much the same results in precipitin tests as those produced by the hydraulic press. Taschdjian _et al._ (1964a) used antigens prepared by a similar method but disintegrated the yeasts by ultrasonication. Stallybrass used a Mickle shaker.

Precipitin tests were carried out by double diffusion in agar buffered to pH 8·6 with the same borax-boric acid buffer as was used as antigen solvent; 2·5-ml quantities of agar were poured into plastic petri plates 5 cm in diameter. A pattern of wells was cut in the agar; a central well was surrounded by six equally spaced peripheral wells; two adjacent peripheral wells had a diameter of 2 mm and the other wells had one of 6 mm; the rims of the peripheral wells lay 5 mm from that of the central well. The patient's serum was placed in the central well and suitable positive control sera were placed in opposite peripheral 6-mm wells. Antigen from one _Candida_ species was placed in an opposing pair of wells, which included a small (2 mm) and a large one (6 mm), and antigen from another species was placed in the remaining pair of wells, one large and one small. Thus, two plates were required to test a serum with four antigens. The diameters of the small and large antigen wells were chosen so as to give a volume ratio of about 1:10 because it was found in preliminary experiments that weakly reacting sera gave the best precipitation reactions with the smaller volume of antigen and stronger sera reacted better with the larger volume. The plates containing antigens and antisera were kept for 5 days at 26°C and then washed in four changes of saline over a period of 24 hr. The agar was then lifted out of the petri dishes, placed on 3 × 1 in. (7·5 × 2·5 cm) glass slides, trimmed to fit, covered with filter paper and dried at 37°C. The dried slides were stained for 10 min. in naphthalene black (0·05 g dissolved in methanol 50 ml, distilled water 40 ml and glacial acetic acid 10 ml) and then differentiated in the same methanol-water-acetic acid solvent.

Results

The results of the precipitin tests, as shown in the table, placed the patients in three well-defined groups. (1) The precipitin test was negative with antigens of all four _Candida_ species in 35 patients (21 male and 14 female). Agglutination titres greater than 16 were obtained with _C. albicans_ in 27 of these patients but with _C. parapsilosis_ in only one. Two of the patients died, both from causes not connected with candida infection. (2) The precipitin test was positive with antigen from only one species of _Candida_ in 13 patients (8 male and
<table>
<thead>
<tr>
<th>Group of patients: type of valve replacement or other operation (number in group)</th>
<th>Number of patients in group giving a negative precipitin reaction with all four species</th>
<th>Number of patients in group giving a positive precipitin reaction with one species only</th>
<th>Number of patients in group giving a positive precipitin reaction with two or more species</th>
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<tbody>
<tr>
<td></td>
<td>C. albicans</td>
<td>C. parapsilosis</td>
<td>C. albicans</td>
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<tr>
<td><strong>Patients with endocarditis</strong></td>
<td>Total</td>
<td>and an agglutination titre &gt;16 with</td>
<td>Total</td>
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<tr>
<td>Homograft aortic (6)</td>
<td>0</td>
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<td>0</td>
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<tr>
<td><strong>Patients without endocarditis</strong></td>
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<tr>
<td>Homograft aortic (15)</td>
<td>8</td>
<td>6</td>
<td>0</td>
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<tr>
<td>Starr-Edwards aortic (18)</td>
<td>11</td>
<td>10</td>
<td>0</td>
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<td>Starr-Edwards aortic and mitral (5)</td>
<td>1</td>
<td>1</td>
<td>0</td>
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<tr>
<td>Starr-Edwards mitral (17)</td>
<td>9</td>
<td>7</td>
<td>1</td>
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<tr>
<td>Starr-Edwards mitral and tricuspid (3)</td>
<td>3</td>
<td>1</td>
<td>0</td>
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<td>University of Capetown mitral (2)</td>
<td>2</td>
<td>2</td>
<td>0</td>
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<tr>
<td>Repair, atrial septal defect (1)</td>
<td>1</td>
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<td>0</td>
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<td>Resection, coarctation aorta (1)</td>
<td>0</td>
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<td>0</td>
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<td><strong>All patients (68)</strong></td>
<td>35</td>
<td>27</td>
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5 female). The positive precipitin result was obtained with *C. albicans* in 12 patients and with *C. parapsilosis* in one. Agglutination titres greater than 16 were obtained with *C. albicans* in all 13 patients but with *C. parapsilosis* in only five of them. One patient died from failure of Starr-Edwards valve suture; he was the one patient in whom the precipitin test was positive with *C. parapsilosis*. (3) The precipitin test was positive with at least two species of *Candida* in 20 patients (18 male and 2 female). All 20 patients gave precipitation with *C. albicans* and 16 of them did so with *C. parapsilosis*. Agglutination titres greater than 16 were obtained with *C. albicans* in all 20 patients and with *C. parapsilosis* in 19 of them. This group included all six patients with candida endocarditis, of whom five died (*vide infra*). Of the other 14 patients in the group, one died and the cause of death was multiple myocardial infarcts.

In all three groups, sera that agglutinated *C. albicans* to titres greater than 16 also agglutinated *C. tropicalis* and *C. guilliermondii* to similar levels. Of the 20 sera in the third group, 17 precipitated with *C. tropicalis* antigen and 8 with *C. guilliermondii* antigen.

**Findings in six patients with endocarditis**

*Patients M.B., C.B. and J.C.* The pathological findings in these three patients were reported by McConnell and Roberts. In all three, *C. albicans* was isolated from vegetations on the aortic valve cusps at necropsy. In M.B., blood culture 3 mth after operation yielded *C. albicans* and, despite treatment with amphotericin B, he died 10 days later. Serum obtained at necropsy gave positive precipitin tests with *C. albicans* and *C. tropicalis*. In C.B., yeasts were seen in a large embolus removed from near the bifurcation of the aorta 2 mth after operation and treatment with amphotericin B was begun. Specimens of serum obtained 2 wk later and at necropsy after 5 wk were precipitin-positive with all four species of *Candida*. In J.C., serum obtained 5 wk after operation was precipitin-positive with *C. albicans*, *C. tropicalis* and *C. parapsilosis*. One week later, *C. albicans* was cultured from his urine, but he died soon afterwards despite treatment with amphotericin B.

*Patient D.S.* A specimen of serum obtained 11 days after operation from this patient was precipitin-negative, but agglutinated *C. parapsilosis* at a dilution of 1 in 16. Although he developed the infectious-mononucleosis-like illness known as post-cardiotomy syndrome (Wheeler, Turner and Scannell, 1962; Smith, 1964) and had a monocyte count of 7000 per mm$^3$, subsequent sera were also precipitin-negative and the *C. parapsilosis* agglutinin titre fell to less than 4. After the patient had been discharged from hospital no serum was examined until his return to hospital 7 mth later with suspected endocarditis; he then died suddenly as a result of a coronary embolus. At necropsy, *C. parapsilosis* was recovered from vegetations on the homograft valve. His serum was precipitin-positive and gave high agglutination titres with all four species of *Candida*.

*Patient R.H.* Specimens of serum obtained 4 and 8 wk after operation were precipitin-positive with all four species of *Candida*, but the patient remained well and 7 mth later his serum was precipitin-negative. Just over a year later when he was readmitted with a suspected embolus, his serum was strongly precipitin-positive and gave very high agglutination titres with all four species. Blood cultures yielded *C. parapsilosis*. Treatment was begun and subsequently the aortic valve homograft was removed and replaced by a Starr-Edwards prosthesis. *C. parapsilosis* was isolated from vegetations on the excised homograft valve. Subsequently the patient made a good recovery and over a period of 10 mth there was a steady diminution in precipitin reactions and agglutinin titres. A year after the second valve operation, precipitins could be detected only by concentrating the serum, although agglutinins were still moderately elevated. Thereafter no further sera were examined for 10 mth and this final specimen was completely negative by both precipitation and agglutination tests.
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Patient J.R. No specimen of serum was examined at the time of the original homograft operation, but when the patient was readmitted 18 mth later with suspected endocarditis, precipitin and agglutinin tests were negative on two sera. Blood cultures yielded a streptococcus and he was given penicillin for 6 wk. His pyrexia returned, however, and blood cultures now yielded C. albicans; three sera obtained at weekly intervals were strongly positive with all four candidas by precipitation and agglutination. After treatment with amphotericin B for 4 wk the homograft was replaced by a Starr-Edwards valve. He died 5 wk after this operation and at necropsy the Starr-Edwards valve was enshrouded with a membrane from which a penicillin-resistant strain of Staphylococcus aureus was isolated. No candida was isolated or seen in sections although in sera obtained at weekly intervals after operation the precipitin reactions and agglutinin titres had not declined significantly.

Times of appearance of precipitins and rise in agglutinin titres

In 47 patients, at least one specimen of serum was obtained before operation and none of these specimens was precipitin-positive. After operation the precipitin test became positive with one species of Candida in 9 of the patients and with two or more species in 10 of them. In the majority, the positive reaction was first detected in the 2nd wk after operation. The agglutination titre rose after operation in 36 patients. In only 4 patients was the rise detected in the serum taken during the 1st wk after operation. In 31 of them, however, there was a sharp rise in agglutinins in the serum between the 10th and 13th post-operative days. In one patient the rise was not observed till the 19th day. Appropriate investigations in these patients did not reveal any association between the appearance of candida antibodies and either the post-cardiotomy syndrome or heavy candida infection of mouth, throat or intestinal tract.

DISCUSSION

The 68 patients from whom specimens of serum were obtained included six in whom candida infection of the aortic valve had become established. In two of them, after initial treatment with amphotericin B, the infected valve was successfully removed and replaced by a Starr-Edwards prosthesis. If this method of treatment is to succeed it is clear that diagnosis should be made as early as possible. In both the successfully treated patients an early indication of the nature of the infection was obtained by examining the serum for precipitins and agglutinins to four species of Candida. In these, as in the other four patients with candida endocarditis, we found precipitins to two or more species of Candida and relatively high titre of agglutinins to C. albicans and C. parapsilosis. The presence of antibodies to species of Candida other than the infecting organism is doubtless due to antigenic sharing among the species and to the fact that cross-reactions become more evident as the titre of antibodies rises.

An unexpected and unexplained finding was the appearance of high antibody titres in some patients who showed no sign of candida endocarditis. Chew and Theus (1967) and Pepys et al. (1968) detected precipitins to C. albicans group-A mannann in 48 and 23 per cent. of healthy subjects respectively. Other workers have, however, found precipitins only in patients with deep-seated candida infections and Pepys et al. suggest that these differences may be attributable
to the use of antigens of different concentration and chemical composition. Pepys et al. noted that precipitins against mannan and, less frequently, precipitins against protein antigens of *C. albicans*, are common in asthma; these workers used purified mannan in low concentration as their precipitating antigen and found positive reactions in 4.5 per cent. of healthy subjects when unconcentrated serum was used. The antigens used in the present study contained little mannan. It is important to note that in our investigation, sera were obtained from 47 patients before operation and, by our methods, none of these sera was precipitin-positive.

The timing of the antibody response with the appearance of precipitins and the rise in agglutinin titre in the 2nd wk after operation suggests that infection began at about the time of operation. It is doubtful whether contaminated homograft valves can be incriminated as the source of the infection, since the antibody response often occurred after the insertion of a Starr-Edwards prosthesis. Such prostheses are also susceptible to candida infection (Watankunakorn et al., 1968). During the period under review, antibiotic treatment was given for a few days during and after operation and there was no evidence that heavy proliferation of candida in the upper respiratory tract or alimentary tract was the cause of the antibody response. Other possible causes include undetected candidaemia or deep-seated transient infection in, for example, the lungs. It is clearly important to discover whether similar candida antibody reactions can be demonstrated after other types of surgery, including other forms of chest surgery.

Although precipitins to several species of *Candida* were present in all our patients with candida endocarditis, it is clear that this finding is not by itself diagnostic of that infection. Nevertheless, patients who develop this serological pattern shortly after heart valve surgery should be watched carefully for infective complications, and special attention should be paid to patients with homograft valve replacements since these replacements are peculiarly susceptible to candida infection.

**SUMMARY**

Six patients with homograft aortic valves who developed endocarditis due to infection with *Candida albicans* or *C. parapsilosis* were found to have serum precipitins and high titres of agglutinins to several species of *Candida*. In 14 of 62 other patients undergoing open-heart surgery, a similar serological pattern developed in the 2nd wk after operation, suggesting the possibility that otherwise undetected candida infection was present. A further 13 of the 62 patients developed antibodies to a single species of *Candida*. Such patients, especially those with homograft valve replacements, should clearly be watched carefully for infective complications.

We wish to thank Dr E. Mavis McConnell and Dr F. Whitwell for much valuable help and advice on histological aspects of this problem; the physicians and surgeons of the Liverpool Regional Cardiac Centre for the opportunity to investigate their patients; and the staffs of the pathology departments of Sefton General and Broadgreen Hospitals for numerous cultures and sera.
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REFERENCES

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