Local symptoms at the site of pacemaker implantation indicate latent systemic infection

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Background: To determine whether local complications at the site of pacemaker implantation indicate infection of the intravascular part of the leads as well as of the pacemaker pocket.

Methods: 105 patients admitted for local inflammatory findings, impending pacemaker or lead exteriorisation, frank pacemaker or lead exteriorisation, or overt infection were studied prospectively. After systematic lead extraction, the initial clinical presentation was related to the results of lead cultures.

Results: Regardless of the initial presentation, the intravascular parts of the leads gave positive cultures in 79.3% of patients. Additionally, 91.6% of the cultures of the extravascular lead segments were positive, in contrast to 38.1% positivity for wound swab cultures. No clinical observations or laboratory investigations permitted identification of patients with negative lead cultures. In a subgroup of 50 patients with manifestations strictly limited to the pacemaker implantation site, cultures of intravascular lead segments were positive in 72%. Infection recurred in 4/8 patients without complete lead body extraction (50%) vs 1/97 patients (1.0%) whose leads were totally extracted ($p < 0.001$).

Conclusions: Local complications at the site of pacemaker implantation are usually associated with infection of the intravascular part of the leads, with a risk of progressing to systemic infection. Such local symptoms should prompt the extraction of leads even in the absence of other infectious manifestations.

Avaliable studies suggest that pacemaker lead related systemic infection is caused by progression of infection from the pacemaker pocket.1 If that concept is valid, local infection of the pocket should in many patients be associated with contamination of the leads and a subsequent risk of systemic infection. The poor prognosis of this complication would make it mandatory to remove the pacing system (leads included) in its entirety. As extraction of the complication would make it mandatory to remove the pacing system being 2.8 (1.8) months.

Inflammation, defined as local pain and redness of the skin, was present in eight patients; impending exteriorisation—manifested as adherence of the lead or pulse generator to the skin without cutaneous breakthrough—was observed in 16; and frank exteriorisation had occurred in 31. An infection with fistula, abscess, or purulent collection was present in the remaining 50 patients.

All patients underwent the following workup: complete clinical examination, three series of blood cultures, serial complete blood counts, creatinine and C reactive protein assays, and erythrocyte sedimentation rate (ESR). Imaging consisted of chest x ray, transthoracic echocardiography (TTE), transoesophageal echocardiography (TOE), and ventilation/perfusion pulmonary scintigraphy, before any extraction attempt. The characteristics of patients are detailed in table 1 according to their initial presentation.

Lead extraction
All pacemakers and leads were systematically extracted except in five patients who refused lead extraction. Percutaneous extraction through the femoral vein was done as previously described.2 To prevent contamination of the lead by its passage through the pacemaker pocket, extraction from the femoral vein was undertaken by two operators, one liberating the proximal end of the lead and recovering the extravascular segment, the other recovering the intravascular segment of the lead from the femoral site.

Blood and lead cultures
Blood cultures were processed with an automated blood culture system (Bio Argos, Sanofi Diagnostics Pasteur, Marnes la Coquette, France).

Leads were cultured after extraction, and swab cultures from the pacemaker pocket were also obtained systematically. Bacteriological tests were done on the extravascular segment and two intravascular segments of the lead. These three parts were placed in sterile phosphate buffered saline and vortexed for one minute. An aliquot of the suspension was then plated on sheep blood agar under O₂ and on chocolate agar under CO₂. The plates were examined after 24 hours at 37°C. In addition, a brain–heart infusion medium was inoculated with 0.1 ml of the suspension for 15 days. An...
Patients were examined at the outpatient clinic 12 weeks after discharge and yearly thereafter. Additional follow up information was obtained by telephone contacts with patients and their primary physicians.

Table 1  Patient characteristics and results of cultures

<table>
<thead>
<tr>
<th></th>
<th>Inflammation (n = 8)</th>
<th>Impending ext (n = 16)</th>
<th>Exteriorisation (n = 31)</th>
<th>Local infection (n = 50)</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>63 (24)</td>
<td>74 (9)</td>
<td>74 (9)</td>
<td>68 (19)</td>
<td>NS</td>
</tr>
<tr>
<td>Sex (F/M)</td>
<td>1/7</td>
<td>5/11</td>
<td>5/26</td>
<td>12/38</td>
<td>NS</td>
</tr>
<tr>
<td>Procedures of pacemaker site (n)</td>
<td>2.8 (1.1)</td>
<td>3.3 (1.9)</td>
<td>2.4 (1.4)</td>
<td>3.0 (2.1)</td>
<td>NS</td>
</tr>
<tr>
<td>Time from last procedure to symptoms onset (months)</td>
<td>20 (20)</td>
<td>20 (20)</td>
<td>29 (31)</td>
<td>32 (30)</td>
<td>NS</td>
</tr>
<tr>
<td>Last procedure as initial implant</td>
<td>1 (12.5%)</td>
<td>3 (18.7%)</td>
<td>9 (29.0%)</td>
<td>12 (24.0%)</td>
<td>NS</td>
</tr>
<tr>
<td>History of local complications</td>
<td>3 (37.5%)</td>
<td>8 (50.0%)</td>
<td>6 (19.3%)</td>
<td>11 (22.0%)</td>
<td>NS</td>
</tr>
<tr>
<td>Intermittent fever</td>
<td>1 (12.5%)</td>
<td>2 (26.3%)</td>
<td>4 (12.9%)</td>
<td>8 (16.0%)</td>
<td>NS</td>
</tr>
<tr>
<td>Clinical pulmonary manifestations</td>
<td>0</td>
<td>1 (6.2%)</td>
<td>1 (3.2%)</td>
<td>4 (8.0%)</td>
<td>NS</td>
</tr>
<tr>
<td>Raised WBC count</td>
<td>2 (2.5%)</td>
<td>1 (6.2%)</td>
<td>5 (16.1%)</td>
<td>7 (14.0%)</td>
<td>NS</td>
</tr>
<tr>
<td>Laboratory findings of inflammation</td>
<td>2 (25%)</td>
<td>5 (31.2%)</td>
<td>12 (38.7%)</td>
<td>20 (40.0%)</td>
<td>NS</td>
</tr>
<tr>
<td>Pulmonary embolism</td>
<td>3 (37.5%)</td>
<td>1 (6.2%)</td>
<td>3 (9.7%)</td>
<td>10 (20.0%)</td>
<td>NS</td>
</tr>
<tr>
<td>Positive blood culture (by Duke criteria)</td>
<td>1 (12.5%)</td>
<td>1 (6.2%)</td>
<td>1 (3.2%)</td>
<td>2 (4.0%)</td>
<td>NS</td>
</tr>
<tr>
<td>Lead extraction refused or failed</td>
<td>3</td>
<td>3</td>
<td>2</td>
<td>2</td>
<td>NS</td>
</tr>
<tr>
<td>Wound swab culture</td>
<td>3 (37.5%)</td>
<td>8 (50.0%)</td>
<td>11 (35.5%)</td>
<td>18 (36.0%)</td>
<td>NS</td>
</tr>
<tr>
<td>Positive extravascular lead cultures</td>
<td>8/8 (100%)</td>
<td>13/14 (92.8%)</td>
<td>25/26 (96.1%)</td>
<td>41/47 (87.2%)</td>
<td>NS</td>
</tr>
<tr>
<td>Positive intravascular lead cultures</td>
<td>5/6 (83.3%)</td>
<td>10/13 (76.9%)</td>
<td>22/27 (77.7%)</td>
<td>36/46 (78.3%)</td>
<td>NS</td>
</tr>
<tr>
<td>S. epidermidis</td>
<td>5 (100%)</td>
<td>8 (80.0%)</td>
<td>16 (76.1%)</td>
<td>22 (61.1%)</td>
<td>NS</td>
</tr>
<tr>
<td>Other coagulase negative staphylococcus</td>
<td>1 (20%)</td>
<td>2 (20.0%)</td>
<td>4 (19.0%)</td>
<td>10 (27.8%)</td>
<td>NS</td>
</tr>
<tr>
<td>S. aureus or other microorganism</td>
<td>0</td>
<td>1 (10.0%)</td>
<td>6 (28.6%)</td>
<td>13 (35.6%)</td>
<td>NS</td>
</tr>
<tr>
<td>&gt;1 microorganism</td>
<td>2 (40%)</td>
<td>3 (30.0%)</td>
<td>2 (9.5%)</td>
<td>5 (13.9%)</td>
<td>NS</td>
</tr>
</tbody>
</table>

Values are mean (SD) or n (%). Details of the microorganisms refer to intravascular lead cultures only. Ext, exteriorisation; F, female; M, male; WBC, white blood cells.

**Statistical analysis**

Measurements between groups were compared by analysis of variance (ANOVA) followed by the appropriate post test or by Student's t test according to the case. Comparisons were two tailed and statistical analyses were done using the Prism software package (GraphPad, USA). A probability value of p < 0.05 was considered significant.

**RESULTS**

**Blood cultures**

A community acquired *Staphylococcus aureus* species was isolated from two separate blood cultures in two patients. Persistently positive blood cultures with a coagulase negative staphylococcus, consistent with endocarditis according to the Duke criteria, were found in one of eight patients with inflammation, one of 16 with impending exteriorisation, one of 31 with frank exteriorisation, and none of 50 with local infection (NS). A single positive blood culture identifying a coagulase negative staphylococcus was obtained in one of eight, one of 16, four of 31, and three of 50 patients with inflammation, impending exteriorisation, frank exteriorisation, and local infection, respectively (NS). Blood cultures were negative in the remaining 91 patients.

**Lead cultures**

Lead culture results are summarised in tables 1 and 2. In seven patients who underwent prolonged and technically complicated procedures, lead cultures were considered unreliable and were discarded from the analysis. Additionally, data pertaining to the intravascular segment of the lead were not available in the five patients mentioned above who refused extraction, and in one patient in whom we were unable to remove any segment of the intravascular part of the leads. Overall, cultures of the whole leads or of a segment of their intravascular parts were obtained in 92 patients. Cultures of these segments were positive in 73 of these (79.3%). Coagulase negative staphylococcal species, and *S. epidermidis* specifically, were isolated from 73.9% and 55.4% of these segments, respectively. The characteristics of patients with and without intravascular lead infections are compared in table 2. Fever, pulmonary embolism, increase white cell count, increased C reactive protein, and positive blood cultures predicted positive cultures from the intravascular...
part of the leads with a very high specificity (0.95, 1, 1, 0.74, and 1, respectively), but with a very low sensitivity (0.16, 0.20, 0.17, 0.45, and 0.07, respectively).

Each of the five species of microorganism that had been isolated from at least two separate blood cultures was always found on the intravascular lead culture. Additionally, when the culture of an intravascular lead segment was positive, the same microorganism was found on the culture of the extravascular portion of the lead. There were no patients with a positive culture on the proximal intravascular segment and a negative culture on the distal intravascular segment. Cultures of the extravascular lead segment were available in 95 patients and were positive in 87 of these (91.6%). In contrast, wound swab cultures were positive in only 38.1% of patients.

Validation of lead cultures
All 14 control leads tested in the operating room were free of infection related findings on direct examination, and a single culture of their 42 segments was positive for S epidermidis.

Cultures from the 11 leads extracted for dysfunction in 95 patients whose leads were fully removed (1.0%) were positive in 87 of these (91.6%). In contrast, wound swab cultures were positive in only 38.1% of patients.

Significance of isolated local symptoms
After detailed investigations, a subgroup of 50 patients was defined as follows: clinical findings strictly limited to the pacemaker implantation site; fever, laboratory markers of inflammation, and blood cultures negative; and pulmonary and cardiac imaging (TOE included) normal. In 36 of these (72%) the cultures of the intravascular and extravascular parts of the leads were positive. In nine additional patients, only the extravascular part of the lead gave a positive culture.

Outcomes
There were two procedural deaths (1.9%). These deaths occurred more than 20 minutes after completion of lead extraction; the patients collapsed suddenly and died in septic shock. Normal necropsy examinations confirmed this diagnosis. Thus these procedural deaths were not a result of mechanical problems but were directly related to the infective process.

Duration of follow up was 24 (17) months. Six patients (5.7%) died after discharge, one presumably from septic shock after having declined lead removal. The five remaining deaths were unrelated to pacemaker infection. Infection recurred in five patients (4.7%). In four of these, recurrence occurred with systemic symptoms including one case of septic shock, and tricuspid endocarditis in another case. Overall, full removal of pacing leads was not achieved or not attempted (consent not given) in eight patients, among whom four recurrences of infection (50.0%) were observed. In contrast, a single recurrence was observed among the 97 patients whose leads were fully removed (1.0%) (p < 0.0001).

### DISCUSSION
The main result of our study is that local symptoms at the site of pacemaker implantation are highly correlated (79.3%) with a positive culture of the intravascular part of the leads. This strong and striking correlation is present whatever the nature of these symptoms, and even in cases where the symptoms were an isolated finding after an extensive work up for infective endocarditis. Lead removal is associated with a favourable outcome; the converse was true when this removal was not achieved.

Our results confirm the mechanism of intravascular infection originating from the pacemaker pocket with subsequent risk of progressing to systemic infection or right heart endocarditis. Da Costa and colleagues have shown that pacing system infections are mainly caused by local contamination by local bacteriological flora during implantation. Skin microorganisms migrate from the insertion site along the surface of the lead, colonising its distal intravascular part and ultimately infecting the blood stream. Studies by transmission and scanning electron microscopy have shown that nearly all indwelling vascular catheters are colonised by microorganisms. There is agreement that all the implantable material should be extracted in patients with septicaemia related to an infected pacing system. On the other hand, there is no clear consensus with respect to the management of pathological findings limited to the pacemaker implantation site, though the trend is toward extraction of the entire system when the cause is infectious, rather than conservative management for purely mechanical complications. Patients with class I indications for lead removal according to the NASPE policy were excluded from our study, as clearly stated in the Methods section. Our study dealt with class II
indications in order to confirm or invalidate the policy of hardware removal. The study highlights the difficulty of excluding the presence of an infection on the grounds of clinical observation or detailed investigations. Griffith and colleagues have proposed a two-step approach primarily based on the results of wound swab cultures, reporting the need for a new contralateral pacemaker in fewer than 40% of patients after a mean follow up of 21.3 months. However, the lack of reliability of their wound swab cultures was illustrated by negative results in nearly 30% of patients who ultimately developed recurrences. In our study, 38.1% of wound swab cultures were positive, contrasting with 91.6% of positive lead cultures; one may question the appropriateness of exposing patients to an approach based on a test with such a low sensitivity.

In this series, some patients without complete lead removal had recurrences with systemic infection not present at the time of recruitment. As failure rates of percutaneous lead extraction procedures increase with the age of the lead, postponement of the procedure may not only allow the growth of large vegetations, with a risk of massive pulmonary embolism at the time of extraction, but also renders extraction more difficult. On the other hand, lead extraction procedures are technically challenging and carry a risk of death. In this study, though procedural mortality was 1.9%, this risk has to be compared with the risk of death in cases of uncontrolled infection—infection incurred in half the patients without complete extraction of the lead body, and this recurrence was fatal in one of these patients (12.5%).

Our studies were not designed to provide recommendations about antibiotic treatment. However, staphylococci adherent to polymer surfaces and their capacity for biofilm formation contribute to the pathogenesis of infections with implanted medical devices. The sterilisation of infected implanted devices by antibiotics is extremely difficult. This is why prolonged use of intravenous antibiotics is not the recommended form of treatment for infected pacemakers if the sterilisation of the implanted material has not been removed. The very small proportion of patients with antibiotic treatment before referral in this series could explain the 38.1% of positive wound swab cultures in our series in comparison with the 22.5% in the series by Griffith and colleagues. However, antibiotic treatment before lead extraction would probably not modify the results of cultures obtained directly from the implanted material, especially lead cultures. Several investigators have proposed conservative treatment and conservative management is required if lead extraction is impossible, unsuccessful, judged too risky, or refused. In cases where infected leads remain in place, prolonged follow-up is required and perhaps prolonged antibiotic treatment.

Almost all patients were referred from secondary care centres, and then admitted to our tertiary centre, specialising in pacemaker lead extractions. It should be emphasised that 28 patients (table 1, history of local complication) had undergone several previous therapeutic attempts because of their local symptoms before referral and inclusion in this study. This may have introduced a referral bias, with a selection of a larger proportion of infectious complications resistant to the previous conservative management usually undertaken in the referral centres. This bias, however, seems to have had a limited impact because the nature of the last procedure preceding inclusion in the study, and the percentage of patients with previous conservative surgery for local complications, were not statistically different in those with and without positive lead cultures.

Conclusions

Regardless of the initial clinical presentation, complications at the pacemaker implantation site are associated with an infectious process in the vast majority of cases. Furthermore, even in the absence of signs or symptoms of systemic infection, the process is rarely limited to the extravascular components of the pacing system. Consequently, the management of local symptoms—even when apparently confined to the pulse generator implantation site—should prompt the extraction of the entire system. A recurrence rate of infection in 50% of patients in whom the leads were not fully removed strongly supports this recommendation.

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REFERENCES

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Electrocardiographic gated multislice computed tomography of Uhl’s anomaly

Uhl’s anomaly of the right ventricle is an unusual cardiac disorder with almost complete absence of right ventricular myocardium, normal tricuspid valve, and preserved septal and left ventricular myocardium. A 12 year old boy was admitted with the complaint of fatigue with effort. A holosystolic murmur was heard on left lower sternal border. Electrocardiography showed right atrial dilation and low voltage in all leads. Cardiomegaly was apparent on chest x ray. Echocardiography revealed a greatly enlarged right atrium and ventricle. The right ventricular free wall seemed very thin.

Electrocardiographic gated multislice computed tomography (CT) showed an extremely thin walled right ventricle with almost complete absence of right ventricular free wall myocardium (panels A and B below). The interventricular septum and left ventricular myocardium were normal. Although magnetic resonance imaging can also show the absence of right ventricular myocardium, three dimensional reconstructed multislice CT imaging (right panel) makes it easier to see the difference between the normal left ventricular myocardium and the thin walled right ventricle.

This is the first case of Uhl’s anomaly that has been demonstrated by multislice CT, and the images suggest that multislice CT can be used for definition of many cardiac pathologies.

Reconstructed multislice CT images in sagittal (A) and axial (B) views show the extremely thin walled right ventricle (thick arrows), normal interventricular septum (arrowheads) and left ventricular myocardium (thin arrows).

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Electrocardiographic gated multislice computed tomography of Uhl's anomaly

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