Flucloxacillin in bone

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SUMMARY Ten patients undergoing total hip replacement for osteoarthritis were each given intramuscular flucloxacillin about two hours preoperatively; bone and serum were sampled simultaneously at operation. Trabecular and compact bone were separated, partly dried, reduced to powders, and then extracted with buffer. The concentration of flucloxacillin in bone washings and serum was determined by well-diffusion assay. The mean concentration of flucloxacillin in serum was 8-9 mg/l, in trabecular bone washings, 1.3 mg/l, and in compact bone washings 0.9 mg/l. The amount of blood contaminating the bone washings was measured, and was calculated to account for at most 26% of the flucloxacillin present. The significance of these findings is discussed in relation to the prophylactic use of flucloxacillin in hip replacement surgery.

Although deep infection rates after total hip replacement operations are of the order of only 2 to 6% (Benson and Hughes, 1975) deep infection is a serious complication as it usually necessitates removal of the prosthesis.

The commonest organisms causing deep infection are coagulase-positive and -negative staphylococci (Visuri et al., 1976), and a number of studies have been made with various prophylactic antibiotics (Ericson et al., 1973; Parsons, 1976). Lincomycin and clindamycin have the drawback of causing pseudomembranous colitis in some patients, and fucidin, when used alone, may cause resistant organisms to emerge (Garrod et al., 1973a). The cephalosporins and gentamicin are broad-spectrum antibiotics, and gentamicin must be injected and is toxic at high concentrations. The penicillinase-resistant penicillins are active against most staphylococci, have a narrow spectrum of antibacterial activity, and are safe.

Some of the penicillinase-resistant penicillins have been studied in hip replacement surgery, and concentrations of oxacillin (Kolczun et al., 1974) and methicillin (Schurman et al., 1975) in bone have been measured. Little work, however, has been done on flucloxacillin; in the present investigation we measured bone concentrations of flucloxacillin and attempted to determine whether antibiotic found in bone was due to blood contamination.

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Excess surface blood was wiped off with tissue paper. A band saw was used to cut the bone, thus removing the entire surface layer, separating trabecular bone from a plate of compact bone in the upper aspect of the femoral neck, and reducing the bone to pieces of about 1 cm³. These were further reduced with pliers to about 3 mm³. Relatively sclerotic (‘hard’) and more normal (‘soft’) parts of the trabecular bone of patient 10 were treated separately. Cysts in bone were not sampled.

Bone fragments were exposed overnight to dried silica gel at 5°C and −70 cm mercury to render them more brittle; the reduction in weight was measured for one bone. Next the fragments were pulsed in a mortar and pestle, with paper between the mortar and the fragments of compact bone to prevent richochets. The size of bone particles after pestling was roughly measured by ruler. Phosphate buffer, pH 7, was added to the ground bone, 1 ml/g of bone. The suspension was shaken for 38-70 (mean 54) min at 5°C and then centrifuged at 2000 rpm for 10 min through a 13 mm antibiotic assay disc (Whatman Ltd) in a 5-ml syringe barrel (Gillette Surgical Ltd). The bone washings obtained were assayed for flucloxacillin concentration; protein and haemoglobin concentrations were measured (see below). The bone processing was based on a method of D. S. Reeves and his colleagues (personal communication).

Washings from patients 1, 9, and 10 were further filtered through a 0.2 µm membrane (Millipore Ltd) before being used for haemoglobin estimation. Washings from patient 10 were tested for the estimation of haemoglobin and flucloxacillin before and after membrane filtration.

**HAEMOGLOBIN IN BONE WASHINGS**

To assess contamination with blood, the haemoglobin concentration in bone washings from three patients was determined by measuring the maximum height of the peak of absorbance in the Soret band (400-420 nm) using an SP800A scanning spectrophotometer (Unicam Ltd). A control sequestrene specimen of normal venous blood of known haemoglobin concentration was treated as were the bones: that is, it was frozen at −20°C for 2 h, partially dried over silica gel, and shaken with 1 ml of buffer; doubling dilutions in buffer were made up to 1/10240 (0.01%, v/v). The maximum absorbance in the Soret band was measured for each dilution and plotted against haemoglobin concentration. This graph was used to convert maximum absorbances to concentrations of blood in the bone washings. To correct for any differences in haemoglobin concentration of the control blood and the patient’s blood, the concentration of blood in the bone washings was multiplied by the ratio of the haemoglobin concentrations of the control blood and the patient’s blood.

Protein in the bone washings of two patients was measured by means of Albustix (Ames Ltd) and by precipitation with 3% sulphosalicylic acid.

**ASSAY METHOD**

Concentrations of flucloxacillin in serum, marrow samples, erythrocytes, and bone washings were assayed by well diffusion. The medium used was Diagnostic Sensitivity Agar (Oxoid Ltd) and the test organism a clinical isolate of *Staphylococcus aureus* sensitive to methicillin. Plates were flooded with a 1/100 dilution of overnight broth culture and then drained for 3 min to yield semiconfluent growth. Standard solutions of flucloxacillin in phosphate buffer, pH 7, were used to assay bone washings, and standards were made up in horse serum to assay sera, marrow specimens, and erythrocytes. We compared flucloxacillin standards made up in phosphate buffer, pH 7, with and without the addition of horse serum, 1 g/l, which was equivalent to the concentration of protein found in bone washings. A control of flucloxacillin, 5 mg/l, was made separately from the standards. Standards and controls were stored in liquid nitrogen (−198°C) and compared with fresh standards after storage. Assay plates were kept at 5°C for 2 h to allow prediffusion and then incubated at 37°C for 18 h.

**AMOUNT OF ANTIBIOTIC IN BONE WASHINGS ATTRIBUTABLE TO CONTAMINATION WITH BLOOD**

The amount (c mg/l) of flucloxacillin in bone washings due to contamination with blood was calculated from the formula:

\[
c = \frac{b}{100} \times \left( \frac{100 - h}{100} \right) \times s
\]

where \(b\) is the concentration of blood in the bone washings (%), \(h\) the patient’s blood haematocrit (%), and \(s\) the concentration (mg/l) of flucloxacillin in the patient’s serum.

**Results**

**PATIENTS’ AGES, SEX, AND DRUG DOSAGES**

These are shown in Table 1. Ages ranged from 61 to 87 (mean 69-4) years. There were five men and five women. The weight-related doses of flucloxacillin for seven patients ranged from 6-4 to 15-9 (mean 10-4) mg/kg. The dose given to each is shown.

**DOSE SAMPLING INTERVAL**

The mean intervals between giving the injection of
Flucloxacillin in bone

Table 1 Patients' data and dosage

<table>
<thead>
<tr>
<th>Patient</th>
<th>Age (yr)</th>
<th>Sex (M/F)</th>
<th>Weight (kg)</th>
<th>Dose of flucloxacillin (g) (mg/kg)</th>
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<td>M</td>
<td>75</td>
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<td>F</td>
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<td>0.5</td>
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<td>3</td>
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<td>M</td>
<td>80</td>
<td>1.0</td>
</tr>
<tr>
<td>4</td>
<td>66</td>
<td>M</td>
<td>78</td>
<td>0.5</td>
</tr>
<tr>
<td>5</td>
<td>74</td>
<td>M</td>
<td>74</td>
<td>1.0</td>
</tr>
<tr>
<td>6</td>
<td>61</td>
<td>F</td>
<td>—</td>
<td>0.5</td>
</tr>
<tr>
<td>7</td>
<td>70</td>
<td>F</td>
<td>—</td>
<td>0.5</td>
</tr>
<tr>
<td>8</td>
<td>87</td>
<td>F</td>
<td>45*</td>
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</tr>
<tr>
<td>9</td>
<td>70</td>
<td>F</td>
<td>63</td>
<td>1.0</td>
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<tr>
<td>10</td>
<td>62</td>
<td>M</td>
<td>73</td>
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</tbody>
</table>

*Approximate weight.

Flucloxacillin and sampling bone and serum were 2 h 12 min and 2 h 13 min (ranges 1 h 45 min and 1 h 51 min-3 h 20 min) respectively. The maximum delay between sampling bone and serum in any patient was 15 min and the mean delay was 4.8 min. Marrow aspirates were taken from patients 1, 4, 9, and 10 within about two to three hours of the injection of flucloxacillin.

Storage of samples

Sera and bones were exposed to room temperature (c 20°C) for a mean of 3-5 h and a maximum of 9 h and 12 h respectively, to 5°C for up to 18 h and to −20°C for up to 10 days (mean 2.3 and 1.2 days). The ranges and means are thus similar for sera and bones. Marrow and erythrocyte specimens were stored for up to a few days at 5°C.

Character of the processed bone

Grinding of bone by mortar and pestle resulted in a fine bone-particle size of approximately 0.1 mm or less. Trabecular bone was reduced to a wet paste and compact bone to a damp powder (Fig. 1). The trabecular bone of patient 6 was unusually soft and degenerate. That of patient 10 was especially sclerotic in parts. On exposure to silica gel, the trabecular bone from patient 8 was reduced in weight by 17%, and compact bone by 15%. The protein content of bone washings from two patients was found to be about 1 g/l.

Contamination with blood

Table 3 shows the maximum absorbance in the Soret band for each sample of bone washings examined. In all cases the absorption peak was symmetrical, and maximum absorption occurred between 404 and 409 nm. The control normal blood specimen contained haemoglobin 13.7 g/dl. Dilutions of this blood in buffer gave similar symmetrical absorption peaks, with maxima between 405 and 407 nm. The maximum absorption was directly proportional to the concentration of haemoglobin. The slope of the line obtained by plotting concentration (v/v) of blood (%) against maximum absorbance (units) was 0.176, and multiplication by this factor converts maximum absorbance to concentration of blood in the washings. The haemoglobin concentrations of the patients, as shown in Table 3, were close to 13.7 g/dl. The corrected concentration of blood in the washings is shown. In none of the bone washings was this more than 2.6%. In the case of patient 10, soft and hard trabecular bone was tested separately, and the concentration of haemoglobin was higher in washings of the softer bone. Washings of compact bone in most cases contained less blood than trabecular bone washings from the same patient. After filtration through a 0.2 μm membrane the maximum absorbance of the washings of hard trabecular bone of patient 10 was unchanged by more than ± 10%.

Flucloxacillin standards

The accuracy of the flucloxacillin standards was such that the 5 mg/l control varied from the expected value by a maximum of ± 20%. Control solutions of flucloxacillin continued to meet this criterion after storage at −198°C for 30 days or at −20°C for five days. Standards made up in phosphate buffer with and without the addition of 1 g/l of serum gave zones of equal diameter. The lower limits of detection of flucloxacillin in buffer and in serum were 0.3 and 0.8 mg/l respectively. Fine growth occurred within the zone around standards and test fluids containing 3.1 mg/l or less of flucloxacillin, with more growth the lower the concentration. Standards and tests were identical in this.

Flucloxacillin assays

The sera contained 4.6-17.0 (mean 8.9) mg flucloxacillin per l (Table 2). Washings of trabecular bone contained flucloxacillin 0.3-2.2 (mean 1.3) mg/l, that is, 5.9-26.4 (mean 15.6)% of the corresponding serum concentrations. Nine of the washings of compact bone contained flucloxacillin 0.3-2.0 (mean 0.9) mg/l, that is, 5.2-22.6 (mean 11.6)% of the corresponding serum concentrations. The washings of the compact bone from patient 4 contained no detectable flucloxacillin. Filtration of the washings of hard trabecular bone of patient 10 through a 0.2 μm filter made no difference to the flucloxacillin concentration. Figure 2 shows the mean concentrations of flucloxacillin in sera and bone washings and the 95% upper and lower confidence limits of the means.

The marrow supernatants of patients 4 and 9 looked the same as serum, and contained 5.9 and 13 mg flucloxacillin per l, 128.3 and 76.5% of the corresponding serum concentrations. The marrow from patient 1 contained approximately
50% cellular material, and assay of the whole aspirate showed the flucloxacillin concentration to be 2.5 mg/l, 47.2% of the serum concentration. Marrow fat was aspirated from patient 10; it contained less than 0.8 mg flucloxacillin per l. Packed erythrocytes of patient 7 contained flucloxacillin 0.4 mg/l and lysed erythrocytes 0.8 mg flucloxacillin per l, 8.5 and 17% of the serum level.

Serum concentrations were directly proportional to the weight-related doses of flucloxacillin. Trabecular bone washings consistently had concentrations of flucloxacillin greater than or about the same as those in compact bone washings. Nineteen of 20 bone washings contained detectable flucloxacillin.

**ANTIBIOTIC DUE TO CONTAMINATION**

Table 3 shows the blood haematocrits and the concentrations of flucloxacillin found in the sera and bone washings of patients 1, 9, and 10. From these data the concentration of flucloxacillin in the bone.

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**Fig. 1 (a) Trabecular bone pieces.** 1 (b) Compact bone pieces. 1(c) Powdered bone: compact (left) and trabecular.
Flucloxacillin in bone

Table 2  Results of flucloxacillin assay of sera, bone washings, and marrow specimens

<table>
<thead>
<tr>
<th>Patient</th>
<th>Serum concentration of flucloxacillin (mg/l)</th>
<th>Concentration (mg/l) (and percentage of serum concentration) of flucloxacillin in:</th>
<th>Washings of</th>
<th>Compact bone</th>
<th>Marrow aspirate or supernatant from marrow aspirate</th>
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<tr>
<td></td>
<td></td>
<td></td>
<td>Trabecular bone</td>
<td></td>
<td></td>
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<tr>
<td></td>
<td></td>
<td></td>
<td>mg/l</td>
<td>%</td>
<td>mg/l</td>
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<tr>
<td>1</td>
<td>5·3</td>
<td>1·4 26·4</td>
<td>1·2 22·6</td>
<td>2·3 47·2</td>
<td></td>
</tr>
<tr>
<td>2</td>
<td>5·9</td>
<td>1·3 22·0</td>
<td>0·7 11·8</td>
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</tr>
<tr>
<td>3</td>
<td>12·0</td>
<td>1·6 13·7</td>
<td>1·2 10·0</td>
<td></td>
<td></td>
</tr>
<tr>
<td>4</td>
<td>4·6</td>
<td>1·0 21·7</td>
<td>nd</td>
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<tr>
<td>5</td>
<td>7·6</td>
<td>0·9 11·8</td>
<td>0·4 5·3</td>
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</tr>
<tr>
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<td>7·8</td>
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<tr>
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<td>10·0</td>
<td>2·2 22·0</td>
<td>2·0 20·0</td>
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<td>9</td>
<td>17·0</td>
<td>2·0 11·7</td>
<td>0·9 5·3</td>
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</tr>
<tr>
<td>10</td>
<td>13·5</td>
<td>0·8 5·9</td>
<td>0·7 5·2</td>
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</tr>
</tbody>
</table>

Table 3  Blood contamination of bone washings and contribution of contamination to total flucloxacillin in bone washings

<table>
<thead>
<tr>
<th>Patient</th>
<th>Nature of bone</th>
<th>Maximum absorbance of bone washings (units)</th>
<th>Patient's venous blood Haemoglobin Haematocrit Concentration of blood (v/v) in bone washings</th>
<th>Concentration of flucloxacillin in bone washings:</th>
<th>Patient's serum</th>
<th>Bone washings</th>
<th>Total Due to contamination</th>
<th>Corrected</th>
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<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>mg/l</td>
<td>% of total</td>
<td>mg/l</td>
<td>%</td>
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<tr>
<td>1</td>
<td>T</td>
<td>4·7</td>
<td>12·0 35</td>
<td>5·3 1·4</td>
<td>0·02</td>
<td>1·7</td>
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</tr>
<tr>
<td>1</td>
<td>C</td>
<td>2·1</td>
<td>12·0 35</td>
<td>5·3 1·2</td>
<td>0·01</td>
<td>0·8</td>
<td>1·19</td>
<td></td>
</tr>
<tr>
<td>9</td>
<td>T</td>
<td>11·8</td>
<td>13·7 45</td>
<td>2·0 17·0</td>
<td>0·2</td>
<td>9·5</td>
<td>1·9</td>
<td></td>
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<tr>
<td>9</td>
<td>C</td>
<td>6·0</td>
<td>13·7 45</td>
<td>1·0 17·0</td>
<td>0·9</td>
<td>10</td>
<td>0·8</td>
<td></td>
</tr>
<tr>
<td>10</td>
<td>T (H)</td>
<td>9·2</td>
<td>13·6 40</td>
<td>1·6 13·5</td>
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<td>16</td>
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</tr>
<tr>
<td>10</td>
<td>T (S)</td>
<td>15·5</td>
<td>13·6 40</td>
<td>2·6 13·5</td>
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<td>13·6 40</td>
<td>2·2 13·5</td>
<td>0·7</td>
<td>26</td>
<td>0·5</td>
<td></td>
</tr>
</tbody>
</table>

T = trabecular; (H) = hard; (S) = soft; C = compact.

washes due to blood contamination was calculated. It ranged from 0·01 to 0·2 mg/l, and the concentration was similar in washings of compact and trabecular bone from the same patient. Expressed as a percentage of total flucloxacillin in the washings, flucloxacillin due to blood contamination ranged from 0·8 to 26%. The corrected concentrations of flucloxacillin in these washings range from 0·5 to 1·9 mg/l.

Discussion

The results obtained were with uninfected osteoarthritis bone—and osteoarthritis is a common indication for total hip replacement. Little deterioration of flucloxacillin is likely to have taken place during storage of the bone, blood, or marrow samples before assay. Sutherland et al. (1970) showed that 10% w/v solutions of flucloxacillin are stable for 14 days at 5°C. Our samples were exposed to room temperature only briefly.

Parsons (1976) discusses two methods of measuring the concentration of antibiotic in bone: that of grinding bone to a paste and then placing the paste in the well of an assay plate, and that of agitating 0·2 g particles of bone in buffer and then assaying antibiotic in the buffer. We have used combines these elements. Bone particles, 1 μm in diameter, can be produced by use of a bone mill (Hansen et al., 1975), but this may not be necessary to extract antibiotic, and heat production in the mill may be a disadvantage. The concentrations we found in bone washings may have tended to underestimate the concentration of flucloxacillin in the bone, which may have been higher by two fold or more, according to how complete was the extraction of flucloxacillin from bone and to the in vivo distribution of flucloxacillin between different sorts of bone tissue. The ease with which we extracted flucloxacillin from bone, and the higher concentrations in washings of trabecular than of compact bone, suggest that flucloxacillin may be present predominantly in the extracellular fluid of bone.

The assay was accurate to within ± 20%; this
flucloxacillin in bone washings due to contamination with blood ignores any flucloxacillin contained in the erythrocytes. We found this to be little, and Korn-guth and Kunin (1976) observed that erythrocytes incubated in serum for three hours took up only 13-5% of the original serum concentration of di-
cloxacillin. Also, the concentrations of flucloxacillin we found in marrow preparations were in roughly inverse proportion to the cellularity of the preparations. Thus, contamination with blood appears to account for at most 26% of the flucloxacillin in the bone washings, and often for considerably less.

The timing of the dose of flucloxacillin was intended to produce the maximum concentration of antibiotic in the blood and tissues at about the time when organisms might be introduced to the tissues at operation. The mean concentration of flucloxacil-
in in serum was 8-9 mg/l. Trabecular bone washings had a mean flucloxacillin concentration of 1-3 mg/l and compact bone washings a mean concentration of 0-9 mg/l. These represent respectively 15-6 and 11-6% of the corresponding serum concentrations. Work with penicillins related to flucloxacillin has given similar results: methicillin (Schurman et al., 1975) and oxacillin (Kolczun et al., 1974) were detected in bone at, respectively, 23% of a mean serum concentration of 11-6 mg/l (after a dose of 1 g intramuscularly) and 5-9% of a mean serum concentration of 86 mg/l (after a dose of 2 g intra-
venously).

Lacey and Stokes (1977) presented evidence that, with heavy inocula of growing organisms, flucloxa-
cillin is susceptible in vitro to the penicillinase of Staph. aureus, and more so than cloxacin. How-
ever, the significance of this is doubtful when these agents are used prophylactically in hip surgery, where small inocula of organisms in a resting state are likely.

The commonest organisms causing deep infection after total hip replacement are coagulase-positive and -negative staphylococci (Visuri et al., 1976). The minimum inhibitory concentrations of flucloxacillin for most strains of these organisms are in the range 0-12-0-5 mg/l (Garrod et al., 1973b; Sutherland et al., 1970). The mean concentrations of flucloxacillin in trabecular and compact bone washings exceed these minimum inhibitory concentrations by about two- to 10-fold.

Methicillin resistance now occurs infrequently in Staph. aureus in Britain; less than 1% of consecutive isolates from septic lesions in seven London hospitals in 1973-75 were resistant (M. T. Parker, personal communication). It is a little more common among coagulase-negative staphylococci; a study of 820

strains of Staph. epidermidis (Baird-Parker groups SII and SV), isolated from the skin of babies and

![Graph](Fig. 2) Mean concentrations, and upper and lower 95% confidence limits of the means, of flucloxacillin in serum and washings of trabecular and compact bone.

compares favourably with the accuracy of plate assays of gentamicin in a series of clinical labora-
tories (Reeves, 1974). Fine growth occurred within zones of inhibition around wells containing only the lowest detectable concentrations of flucloxacillin. This phenomenon occurred equally with flucloxacil-
in standards and bone washings, so that assay was possible down to flucloxacillin levels in bone washings of 0-3 mg/l.

Individual bones varied in vascularity, which may account for differences in haemoglobin concentration in washings. The concentrations of haemoglobin and of flucloxacillin in washings of one bone were little affected by filtration through a 0-2 μm mem-
brane. The haemoglobin in the control blood and in bone washings was treated in exactly the same way; in both cases most of it was probably converted to methaemoglobin. The wavelengths in the Soret band at which oxyhaemoglobin and methaemoglobin absorb maximally are 413 and 406 nm (Varley, 1967). Haemoglobin and these related compounds have similar molecular extinction coefficients in the Soret band (Lemberg and Legge, 1949). Hence dilu-
tions of the control blood provided a valid means of measuring the blood contamination of the bone washings.

The formula used to calculate the amount of
adults in hospital, and of school children, showed 4-6% to be resistant to methicillin (J. F. Richardson, personal communication). Patients undergoing hip-replacement surgery have usually been recently admitted to hospital, and their skin flora may have changed little before the operation.

Only a small proportion of the cells of a methicillin-resistant strain of _Staph. aureus_ capable of growing in the presence of high concentrations of methicillin at 30°C can do so at 37°C (Parker and Hewitt, 1970). Over two-thirds (26 of 38) of methicillin-resistant strains of _Staph. epidermidis_ (SII) similarly showed only minimal resistance when tested at 37°C (J. F. Richardson, personal communication). Thus a small inoculum of a coagulase-positive or -negative staphylococcus into a wound is unlikely to contain cells capable of growing in the presence of methicillin in deep tissues.

The isoxazolyl penicillins have clinical advantages over various other antibiotics that have been studied in prophylaxis for hip-replacement surgery. It appears that flucloxacillin will reach concentrations in bone at the time of operation sufficient to inhibit the most likely pathogens causing deep post-operative infections, when given intramuscularly about two hours preoperatively, in a dose of 1 g, or more, in persons of above average weight.

We thank Beechams Ltd for supplying flucloxacillin powder, Mr R. Sutherland for helpful discussions on assay and bone processing technique, and Drs G. C. E. McAll and N. J. Riddell for great help in obtaining specimens and patients' data.

References


