Assay of gentamicin in cerebrospinal fluid

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SYNOPSIS A comparison of standard curves obtained from a conventional plate diffusion assay method revealed significant differences when gentamicin standards were made up in different media. Standards made up in distilled water resulted in a curve which differed from that of standards made up in pooled human cerebrospinal fluid by a factor of up to 4. When the assay medium was supplemented with 0.5% sodium chloride, the difference between the two standard curves was reduced to a factor of about 1.5. The curve obtained from standards made up in 150 mM sodium chloride/4.5 mM calcium chloride correlated well with that from standards made up in cerebrospinal fluid. There was no evidence of gentamicin being bound to protein in the cerebrospinal fluid.

Over 50% of the instances of meningitis in neonates are caused by Gram-negative enteric bacilli (Mathies et al., 1971). Gentamicin, being effective against these organisms, is a useful agent in the treatment of meningitis in the newborn.

Systemic administration of gentamicin does not give rise to adequate therapeutic levels in the cerebrospinal fluid. Riff and Jackson (1971) could not detect measurable levels in the CSF after intramuscular injection unless the serum level exceeded 4 mg/l. McCracken et al. (1971) reported a mean CSF gentamicin level of 1.2 mg/l (range 0.2-2.9) after intramuscular administration of 1.5-2.5 mg/kg. The same authors measured 2.6 mg gentamicin/l (range 2.5-2.7) in CSF 24 hours after an intrathecal dose of 0.5 mg, and 15.8 mg/l (range 3.3-60) after an intraventricular dose of 0.5-1 mg. CSF levels of gentamicin up to 7 mg/l were obtained by Newman and Holt (1967) after 1-2 mg/kg per day intramuscularly and 0.2-1 mg/day intraventricularly.

It is well documented that the action of gentamicin is particularly susceptible to changes in the salt concentration of the medium (Rubenis et al., 1963; Schoutens and Yourassowsky, 1972). When it is necessary to assay the drug in body fluids this is usually done against reference standards which are made up in the same fluid in order to minimize any discrepancies due to variation in the constituents of the medium. However, this principle appears to have been neglected in assays of gentamicin in CSF, probably due, at least in part, to the fact that pooled human CSF is not readily available in many laboratories.

Reported levels of gentamicin in the CSF are often the result of assays against reference standards made up in water (Newman and Holt, 1967). In other cases the assay method is not described in sufficient detail, and this applies particularly to assays performed by a reference laboratory.

The purpose of this work was to show the extent of variation in results from standards made up in different media and to test the possibility of using a 'synthetic' substitute for pooled human CSF.

Material and methods

PLATE DIFFUSION ASSAY

Klebsiella Edwardsii (NCTC 10896) was used as the test organism. Overnight cultures were grown in Todd Hewitt broth (Oxoid). Bacto antibiotic medium 11 (Difco) was used as the assay medium. This was supplemented where appropriate with 0.5% sodium chloride. Fifty millilitre volumes of assay medium were dispensed into 13.5 cm diameter plastic Petri dishes. Each plate was surface-seeded with approximately 4 x 10^6 organisms. Twelve wells, 9 mm in diameter, were cut in each plate. Gentamicin sulphate (Gentamicin Injectable, Nicholas Laboratories) standards were prepared at concentrations of 1.25, 2.5, 5, 10, and 20 mg of gentamicin base/litre; 0.15 ml of standard was added to each well. Zone sizes were measured after 5-18 hours' incubation to the nearest 0.5 mm.

SOURCE OF CEREBROSPINAL FLUID

CSF was pooled from specimens obtained from patients in the hospital. Before use it was tested for
sterility and checked to ensure that no antibacterial agents were present.

**Protein Binding Assay**

The radioactive gentamicin used in protein-binding experiments was methyl-\(^{14}\)C-gentamicin sulphate having an activity of 0·44 mCi/l and a specific activity of 0·4 Ci/M of free base. Radioactive samples were added to 8 ml of 0·3% PPO/0·01% POPOP/toluene:ethanol (5:1) and counted in an Intertechnique model SL30 scintillation spectrometer.

Protein was estimated according to the method of Warburg and Christian (1942).

**Results**

Curves were obtained from standards made up in distilled water, pooled human CSF, 159 mM sodium chloride, and 150 mM sodium chloride/4·5 mM calcium chloride. Each standard was measured in triplicate. Figure 1 shows the results obtained using assay medium with no added salt. The resultant curve from standards made up in distilled water differs from the curve from standards in pooled CSF by a factor of 4. Standards made up in 150 mM NaCl/4·5 mM CaCl\(_2\) give a curve which correlates well with the CSF curve. The curve obtained from standards in 159 mM NaCl differs from the distilled water curve by a factor of 2.

Figure 2 shows the results obtained by the same procedure as above except that the assay medium was supplemented with 0·5% NaCl. The difference in the curves from standards made up in water and in CSF is reduced to a factor of about 1·5. Both these curves from standards made up in 159 mM NaCl and in 150 mM NaCl/4·5 mM CaCl\(_2\) correlate well with the CSF curve.

Results are shown for a single experiment but this was repeated four times. In the unsupplemented medium the CSF curve always differed from the distilled water curve by a factor greater than 3. When the medium was supplemented with 0·5% NaCl the difference always exceeded a factor of 1·3. These results were still reproducible when different batches of pooled CSF were used.

To test for protein binding 0·2 ml of \(^{14}\)C-gentamicin was added to 0·8 ml pooled human CSF to give a final concentration of 100 mg free base/l. 0·2 ml of \(^{14}\)C-gentamicin was also added to 0·8 ml of 150 mM NaCl/4·5 mM CaCl\(_2\) as a control. The samples were incubated for four hours at 37°C and

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**Table Distribution of \(^{14}\)C-gentamicin in the CSF**

<table>
<thead>
<tr>
<th></th>
<th>Total Radioactivity in DPM</th>
<th>Radioactivity in DPM associated with TCA Precipitate</th>
<th>% Activity associated with TCA Precipitate</th>
</tr>
</thead>
<tbody>
<tr>
<td>Test</td>
<td>2·5 × 10(^4)</td>
<td>80</td>
<td>0·03</td>
</tr>
<tr>
<td>Control</td>
<td>2·6 × 10(^4)</td>
<td>101</td>
<td>0·04</td>
</tr>
</tbody>
</table>

DPM = disintegrations per minute.

TCA = trichloroacetic acid.
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then added to an equal volume of 10% trichloroacetic acid. After centrifugation a sample was taken from the supernatant for counting and the precipitate was collected on to a 0.45 µ pore size membrane filter and counted. The results are shown in the table.

The protein content of the CSF was estimated at 890 mg/l.

Discussion

These results further emphasize the need for standardization in antibiotic assays. Gentamicin levels reported using reference standards made up in water may be four times as high as the actual levels. Thus, a level which appears to be above the MIC for the infecting organism may well be subtherapeutic.

To date there have been no reports of toxic effects due to intrathecal or intraventricular administration of gentamicin. However, since a large number of cases described have involved neonates, there probably has not been sufficient follow-up completely to rule out this possibility. It is also important therefore that assays are carried out accurately to ensure that potentially toxic levels are not reached. A level reported as being safe by one assay method could be toxic when assayed by a different method.

The salt solutions were prepared to correspond with the ionic strength of CSF. In salt-free medium the curves obtained from standards made up in 150 mM NaCl/4.5 CaCl₂ and 159 mM NaCl differed despite these solutions having the same ionic strength. This supports the theory that the effect of divalent cations such as Mg²⁺ and Ca²⁺ on the action of gentamicin is not solely related to their influence on the ionic strength of the medium (Gilbert et al., 1971). It is not certain why this effect of calcium ions is not apparent when the medium is supplemented with 0.5% NaCl.

There is also an increase in the range of zone diameters in the presence of 0.5% NaCl, and I have discussed this elsewhere (Deacon, 1976).

There was no association of gentamicin with CSF proteins in the specimen tested, which at 890 mg protein/l had a protein concentration considerably higher than normal. However, in bacterial meningitis the protein content of the CSF may increase to 10 g or more per litre and it is not known whether such high levels would affect protein-binding of the drug.

It is clear from these results that gentamicin in CSF should not be assayed against reference standards made up in water. If pooled CSF is not available then 150 mM NaCl/4.5 mM CaCl₂ would seem to be an adequate substitute.

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References


