The Pharmacokinetics of Once-Daily Dosing With Gentamicin in Women With Postpartum Endometritis

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ABSTRACT

Objective: To evaluate the pharmacokinetics and cost of once-daily dosing with gentamicin in women with postpartum endometritis.

Methods: Gentamicin in a single daily dose of 4.5 mg/kg was administered intravenously to 10 women with postpartum endometritis. Peak and trough gentamicin levels were measured, and nephrotoxicity and clinical ototoxicity were monitored. Pharmacokinetic data were analyzed, and a cost analysis of once-daily gentamicin administration was performed.

Results: The mean elimination constant was 0.105 ± 0.008 L/h, and the mean volume of distribution was 0.34 ± 0.07 L/kg. Mean peak gentamicin levels exceeded 11 mg/L, and all trough levels were <0.3 mg/L. Cost savings of 44% were achieved with once-daily dosing of gentamicin, compared with traditional thrice-daily dosing.


KEY WORDS

gentamicin; postpartum endometritis

Gentamicin remains a popular bacteriocidal antibiotic for the treatment of infections caused by gram-negative organisms. Postpartum endometritis, typically a polymicrobial infection of both aerobic and anaerobic organisms, is often treated with clindamycin and gentamicin for broad-spectrum coverage. While it is a highly effective antibiotic, gentamicin is associated with potentially serious toxicity, including nephrotoxicity and ototoxicity. A review of prospective clinical trials of adult patients treated with aminoglycosides between 1975 and 1982 indicated that the average frequencies of nephrotoxicity and ototoxicity of gentamicin are 14% and 8%, respectively.¹ Moreover, wide variations in dosage requirements of gentamicin have been demonstrated in the postpartum period, and subtherapeutic levels are seen in one third of women receiving standard dosing regimens.² ³

Aminoglycosides are commonly administered in either two or three divided doses per day. Two properties of aminoglycoside agents, the postantibiotic effect (PAE) and concentration-dependent killing, have raised interest in the idea of less frequent dosing of aminoglycosides. Postantibiotic effect is defined as ongoing bacterial inhibition after the antibiotic exposure has ceased. As such, the efficacy of aminoglycosides may not rely on maintaining serum concentrations above the minimum inhibitory concentration.⁴ The duration of the PAE

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correlates with the aminoglycoside concentration. Aminoglycoside antibiotics exhibit concentration-dependent bactericidal activity, with greater antibacterial activity seen in higher doses of the drug. In addition, a once-daily dosing regimen of aminoglycoside antibiotics may result in less drug accumulation in the renal cortex, thereby potentially decreasing nephrotoxicity. We report on the pharmacokinetics and cost of once-daily dosing with gentamicin in postpartum women with endometritis.

SUBJECTS AND METHODS

Women 18 years of age and older with postpartum endometritis were eligible for the study and were enrolled between July 1995 and January 1996. This study was approved by the institutional review board. The diagnosis of endometritis was based on temperature elevation greater than 38°C and uterine tenderness occurring at least 24 hours postpartum in the absence of any other source of infection. Exclusion criteria included aminoglycoside allergy, renal impairment (defined as serum creatinine >1.2 mg/dL or estimated creatinine clearance <60 mL/min), known hearing loss, or immunocompromised status. After obtaining informed consent, patients were administered a single dose of gentamicin at a dose of 4.5 mg per kilogram of actual body weight over 30 minutes every 24 hours. Gentamicin was used in combination with clindamycin to provide polymicrobial coverage. Blood urea nitrogen and creatinine were determined prior to initiation of therapy and every 48 hours during treatment. Nephrotoxicity was defined as an elevation of serum creatinine from baseline greater than 0.5 mg/dL. Gentamicin levels were drawn 30 minutes after completion of the first and second doses, 30 minutes prior to the second and third doses, and an 8-hour level was drawn after the second dose. Pharmacokinetic data were analyzed for each patient using standard pharmacokinetic principles. Oto-toxicity was evaluated clinically by regularly questioning patients about hearing loss, tinnitus, and vertigo. No audiometric evaluations were performed.

Cost analysis was based on 72 hours of treatment using 300 mg/d of gentamicin in the once-daily regimen compared with 100 mg every 8 hours in the traditional dosing regimen. Patient charges from the pharmacy for preparation and dispensing and nursing administration times were calculated. Nursing costs were based on hospital costs and were determined using an average registered nurse wage of $18/h. Charges for intravenous tubing and gentamicin level monitoring were excluded in the cost analysis, as intravenous tubing costs are minimal, and gentamicin level monitoring may be necessary with both single and thrice-daily treatment regimens.

Gentamicin serum concentrations were determined by fluorescence polarization immunoassay using an automated fluorescence polarization analyzer. For levels between 2 and 5 mg/L, the coefficient of variation was 5%. Levels between 10 and 15 mg/L had a coefficient of variation of 6%.

RESULTS

Ten women with postpartum endometritis were enrolled in this study. Table 1 summarizes demographic data as well as half life and estimated creatinine clearance. The mean (± standard deviation [SD]) elimination constant was 0.105 ± 0.008 L/h, with a range of 0.096–0.122 L/h. The mean volume of distribution (± SD) was 0.34 ± 0.07 L/kg, with a range of 0.27–0.51 L/kg. The mean first peak level (± SD) was 11.6 ± 2.3 mg/L. The mean second peak level (± SD) was 13.0 ± 2.5 mg/L. All first and second trough levels were less than 0.3 mg/L. A favorable clinical response was seen in nine patients. The median duration of therapy was 3 days. One patient was administered heparin therapy for presumed septic pelvic thrombophlebitis. There was no nephrotoxicity nor clinical ototoxicity detected.

Based on a 3-day course of gentamicin, the patient charge from our pharmacy for the once-daily regimen was $241, compared with $422 for the thrice-daily regimen. After adding nursing administration times, the total patient charges were $250 and $449 respectively. This represents a 44% re-

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Mean ± SD</th>
<th>Range</th>
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</thead>
<tbody>
<tr>
<td>Age (y)</td>
<td>26.2 ± 8.4</td>
<td>18–41</td>
</tr>
<tr>
<td>Weight (kg)</td>
<td>80.3 ± 23.9</td>
<td>56–136</td>
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<tr>
<td>Dose (mg)</td>
<td>360 ± 105</td>
<td>255–600</td>
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<tr>
<td>Half life (h)</td>
<td>4.36 ± 0.22</td>
<td>3.96–4.67</td>
</tr>
<tr>
<td>Estimated clearance</td>
<td>149 ± 53</td>
<td>84–271</td>
</tr>
</tbody>
</table>

*SD, standard deviation.
ONCE-DAILY DOSING OF GENTAMICIN FOR POSTPARTUM ENDOMETRITIS  SUNYECZ ET AL.

Once-daily dosing of gentamicin for the treatment of postpartum endometritis when compared with the thrice-daily regimen for 72 hours of treatment.

DISCUSSION

This study demonstrates several important points regarding a once-daily dose of gentamicin among postpartum women. Our data show that no drug accumulation occurred during our treatment period, as trough levels were consistently less than 0.3 mg/L. This is important clinically as data suggests that aminoglycoside-free periods during therapy are necessary to reduce the risk of nephrotoxicity and ototoxicity. We were also able to consistently achieve elevated peak levels with our dosing regimen, optimizing the PAE and concentration-dependent killing characteristics of aminoglycoside antibiotics. Our pharmacokinetic parameters were consistent with published data from Sangha et al. evaluating once-daily dosing with gentamicin in intensive care unit patients with open fractures. The numerous physiologic changes that occur during the postpartum period may cause wide fluctuations in the pharmacokinetic parameters. This may explain the differences in elimination rate constant, volume of distribution, and half life between our data and those reported recently by Del Priore et al. An additional factor that may reflect a higher half life is that our laboratory could not detect gentamicin levels less than 0.3 mg/L, the trough value used to calculate the half life. Higher trough levels lengthen the half life. Finally, we were able to demonstrate significant cost savings with a once-daily regimen in the postpartum patient with endometritis, reducing charges by nearly one half. The results of this study support the conclusions of two reports of once-daily gentamicin in the treatment of puerperal uterine infections. Once-daily dosing of gentamicin for the treatment of postpartum endometritis is efficacious and cost-effective, and it appears that this regimen is not associated with a high incidence of nephrotoxicity or ototoxicity.

REFERENCES

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