Pharmacokinetics of imipenem in dogs

Corrie W. Barker, DVM; Weijiang Zhang, PhD; Susan Sanchez, PhD; Steven C. Budsberg, DVM, MS; F. Douglas Boudinot, PhD; M. A. McCrackin Stevenson, DVM, PhD

| Objective | To determine the plasma pharmacokinetics of imipenem (5 mg/kg) after single-dose IV, IM, and SC administrations in dogs and assess the ability of plasma samples to inhibit the growth of *Escherichia coli* in vitro. |
| Animals | 6 adult dogs. |
| Procedure | A 3-way crossover design was used. Plasma concentrations of imipenem were measured after IV, IM, and SC administration by use of high-performance liquid chromatography. An agar well antimicrobial assay was performed with 3 *E coli* isolates that included a reference strain and 2 multidrug-resistant clinical isolates. |
| Results | Plasma concentrations of imipenem remained above the reported minimum inhibitory concentration for *E coli* (0.06 to 0.25 µg/mL) for a minimum of 4 hours after IV, IM, and SC injections. Harmonic mean and pseudo-standard deviation half-life of imipenem was 0.80 ± 0.23, 0.92 ± 0.33, and 1.54 ± 1.02 hours after IV, IM, and SC administration, respectively. Maximum plasma concentrations (Cmax) of imipenem after IM and SC administration were 13.2 ± 4.06 and 8.8 ± 1.7 µg/mL, respectively. Time elapsed from drug administration until Cmax was 0.50 ± 0.16 hours after IM and 0.83 ± 0.13 hours after SC injection. Growth of all 3 *E coli* isolates was inhibited in the agar well antimicrobial assay for 2 hours after imipenem administration by all routes. |
| Conclusions and Clinical Relevance | Imipenem is rapidly and completely absorbed from intramuscular and subcutaneous tissues and effectively inhibits in vitro growth of certain multidrug-resistant clinical isolates of *E coli*. (Am J Vet Res 2003;64:694–699) |

Imipenem is a carbapenem β-lactam antimicrobial that has been used widely in human and veterinary medicine. It has a broad spectrum of antimicrobial activity and is effective against most gram-positive, gram-negative, aerobic, and anaerobic organisms.1-3 It has been used to treat various infections in humans, including soft tissue, abdominal, respiratory tract, urinary tract, and bone infections,4,13 and to provide perioperative antimicrobial prophylaxis in abdominal and colorectal surgeries.14,15 In addition, imipenem is used commonly in humans to treat various infections in humans, including soft issue, abdominal, respiratory tract, urinary tract, and bone infections,4-13 and to provide perioperative antimicrobial prophylaxis in abdominal and colorectal surgeries.14,15 

The pharmacokinetics of imipenem have been well established in human medicine.1-3,9-13,20 Both IV and IM administration routes are used in humans for treatment of nosocomial and serious infections.7,9,10,21 The use of imipenem in dogs has largely been extrapolated from the established dosage and pharmacokinetic data of humans. The pharmacokinetics of imipenem have been determined after IV administration in dogs,9,20 but neither the IM nor SC route has been thoroughly investigated.

The purpose of the study reported here was to characterize the pharmacokinetics of imipenem by use of high-performance liquid chromatography (HPLC) after single-dose IV, IM, or SC administrations (5 mg/kg) in dogs and to assess the ability of plasma samples to inhibit growth of *Escherichia coli* in vitro.

Materials and Methods

Dogs—Six 3- to 9-year-old sexually intact male cross-bred dogs weighing 19.6 to 38 kg (mean ± SD, 27.1 ± 7.1 kg) were used in this study.16 Dogs were housed at the University of Georgia Veterinary Medical Animal Care Facility. Animal studies were approved by the University of Georgia Animal Care and Use Committee and conducted in accordance with guidelines established by the National Institutes of Health Guide for the Care and Use of Laboratory Animals. A physical examination, CBC, serum biochemical profile, and urinalysis obtained by cystocentesis were performed on all dogs before starting the study. Urinalyses were repeated on the day of the first imipenem administration and the day after each additional administration. The CBC and serum biochemical profile were repeated 12 to 13 days after final administration.

Study design—The study was done in accordance with a randomized 3-way crossover design, with 3 dosing periods and a 1-week washout period between administrations. Imipenem (5 mg/kg)15 was administered IV, IM, and SC. The day prior to administration and sampling, the dogs were sedated with medetomidine (0.025 mg/kg, IV) for aseptic placement of an 18.5-gauge jugular catheter, which was used for blood sampling. An 18-gauge catheter was also placed in the cephalic vein of dogs scheduled for IV drug administration. Jugular catheters and cephalic catheters were flushed with 1 and 0.5 mL of saline (0.9% NaCl) solution, respectively, containing 5 U of sodium penicillin. 

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From the Departments of Small Animal Medicine and Surgery (Barker, Budsberg, McCrackin Stevenson) and Medical Microbiology and Parasitology (Sanchez, McCrackin Stevenson), College of Veterinary Medicine; the Department of Pharmaceutical and Biomedical Sciences, College of Pharmacy (Zhang, Boudinot); and Athens Diagnostic Laboratory (Sanchez), University of Georgia, Athens, GA, 30602. Dr. Zhang’s present address is TAP Pharmaceutical Products Inc, 673 N Field Dr, Lake Forest, IL 60045. Dr. Boudinot’s present address is the Department of Pharmacaceutics, College of Pharmacy, Virginia Commonwealth University, Richmond, VA 23298. Dr. McCrackin Stevenson’s present address is Laboratory of Intracellular Parasites, Rocky Mountain Laboratories, NIAID, NIH, 903 S 4th St, Hamilton, MT 59840. Supported in part by a donation to the University of Georgia College of Veterinary Medicine by Dr. and Mrs. Clarence A. Rawlings. Presented in part at the 37th Annual Scientific Meeting of the American College of Veterinary Surgeons, San Diego, October 2002. The authors thank Kate Pennick and Lynn Reece for technical assistance. Address correspondence to Dr. McCrackin Stevenson.

Harmonic mean and pseudo-standard deviation half-life of imipenem was 0.80 ± 0.23, 0.92 ± 0.33, and 1.54 ± 1.02 hours after IV, IM, and SC administration, respectively. Maximum plasma concentrations (Cmax) of imipenem after IM and SC administration were 13.2 ± 4.06 and 8.8 ± 1.7 µg/mL, respectively. Time elapsed from drug administration until Cmax was 0.50 ± 0.16 hours after IM and 0.83 ± 0.13 hours after SC injection. Growth of all 3 *E coli* isolates was inhibited in the agar well antimicrobial assay for 2 hours after imipenem administration by all routes.
heparin/mL. The effect of medetomidine was reversed with atipamezole (0.1 mg/kg, IM) after catheters were placed.

Food was withheld for 24 hours before and 6 hours after imipenem administration. Water was not withheld at any time. Purified imipenem powder was reconstituted with sterile saline solution (11 mg/mL). In this study, imipenem was not combined with cilastatin. Imipenem was given as an IV bolus through the cephalic catheter for 30 to 60 seconds. The IM injection was divided into 4 equal volumes (2.0 to 3.5 mL/injection site) because of the large volume and administered in 4 sites (bilateral epaxial and gluteal musculature) to minimize discomfort. Subcutaneous injections were given in the interscapular region. Subcutaneous injections were not divided, because the total volume (8.0 to 14.0 mL/injection site) is well tolerated in adult dogs. Adverse reactions to injections were recorded.

Blood samples were collected from the jugular catheter immediately before and 0.08, 0.25, 0.50, 0.75, 1, 1.5, 2, 4, 6, 8, and 12 hours after drug administration. Sampling was a 4-step procedure. Blood (2.5 to 3.0 mL) was aspirated from the jugular catheter into a sterile 3-mL syringe. A 1.5-mL sample for drug analysis was collected and transferred into a heparinized polypropylene microcentrifuge tube. The initial blood sample was returned to the dog through the jugular catheter, and the catheter was flushed with 1 mL of heparinized saline solution. The heparinized blood samples were labeled and centrifuged immediately at 1000 × g for 10 minutes. Plasma was removed from the samples and combined with an equal volume of stabilizing solution (0.5 M 3-morpholinopropanesulfonic acid-water-ethylene glycol [2:1:1]). The samples containing plasma in a stabilizing solution were vortexed, divided, and stored at −80°C for < 8 weeks before HPLC, and bacteriologic assays were performed. The addition of a stabilizing buffer prevents degradation of imipenem in plasma for > 90 days when samples are stored at −80°C. HPLC assay—The HPLC assay was performed with an established method. Briefly, plasma samples were subjected to ultrafiltration for 10 minutes at 6,000 × g by use of filter units, and 20 µL of the filtered sample was injected onto the HPLC column. Chromatography was performed by use of an HPLC system with a photodiode array detector. Chromatographic separation of the compounds was accomplished with a reversed-phase column (4 µm, 39 × 150 mm) and a mobile phase of 0.2M borate buffer (pH, 7.2) prepared with boric acid in HPLC-grade water at a flow of 1.5 mL/min. Absorbance was detected at 300 nm. Standard curves were graphed for each bacterial isolate with known concentrations of imipenem reconstituted in either plasma or PBS solution (PBSS); concentrations varied from 0 (negative control) to 0.39 mg/mL. These concentrations were chosen on the basis of known minimum inhibitory concentrations (MICs) of imipenem for the reference strain of E coli (0.06 to 0.25 µg/mL). Plates were incubated for 18 hours at 37°C. Each zone of inhibition was measured twice with standard calipers, and the mean of these 2 results was recorded. To test plasma samples from dogs, plates were prepared in duplicate as described, and the wells were filled with the designated test samples collected from the dogs after imipenem administration. Each of the E coli isolates was tested separately. Two negative control wells were included on each plate: 1 with PBSS alone and the other with plasma from a dog that had not been treated with imipenem. Positive control wells contained imipenem, which directly dissolved in dog plasma and PBSS at concentrations of 0.01 and 0.03 mg/mL for each solution. Plates were incubated, and the zones of inhibition were measured.

Data analyses—The plasma concentrations of imipenem detected at various collection times were analyzed with non-compartmental methods by use of computer software. The terminal elimination rate constant (λ) was obtained from the slope of the least squares linear regression fit of the logarithms of measurable concentrations versus time in the log-linear terminal phase of the curve. The area under the plasma concentration-time curve (AUC) from time zero to the last observed concentration (4 hours) was determined by the linear trapezoidal rule, and the AUC was extrapolated to time infinite by dividing the last measurable plasma concentration by λ. Harmonic means and pseudo-standard deviations were calculated for half-life (t½). The maximum plasma concentrations (Cmax) of imipenem and times to reach maximum plasma concentration (Tmax) were the observed values. Bioavailabilities after IM and SC administration were calculated by the ratio of the AUC from IM and SC administration to AUC from IV administration. Statistical analysis comparing pharmacokinetic values determined for various administration routes were performed by use of a 2-way ANOVA to test the effect of administration route, individual dog, and study period. Significance was defined as P < 0.05. Standard curves were graphed for each bacterial isolate by use of known imipenem concentration versus zone of inhibition. Nonlinear regression of these curves was performed with commercially available computer software. The resulting equations describing the standard curves were used to convert zone of inhibition measurements from the dog plasma samples to imipenem concentrations. Times after
injection versus imipenem concentrations were graphed. Paired t-tests were performed at each time point to compare the imipenem concentration for the reference strain of *E coli* with clinical isolates 2-2225 and 2-2559. Significance was defined as *P* < 0.05.

**Results**

**Pharmacokinetics**—Pharmacokinetic parameters of imipenem after single IV, IM, and SC administrations were determined (Table 1). Plasma imipenem concentrations after IV administration declined rapidly with a terminal harmonic mean t$_{1/2}$ of 0.80 ± 0.23 hours (Fig 1). Peak plasma concentrations were achieved rapidly after IM or SC administration, and terminal t$_{1/2}$ was lengthened, compared with IV administration. There were no significant differences in AUC values among IV, IM, and SC administration. The MIC of imipenem for *E coli* (0.06 to 0.25 mg/L) was exceeded for at least 4 hours after IV, IM, and SC administration.

**In vitro inhibition of bacterial growth**—The shapes of the standard curves were consistent with a 2-parameter hyperbola. Nonlinear regression of these curves described the line of best fit as:

\[ y = \frac{a}{b + x} \]

where *y* is zone of inhibition, *a* is maximum inhibition, *x* is imipenem concentration, and *b* is the concentration of imipenem inhibiting 50% of bacterial growth. Correlation coefficients for the standard curves of the 3 bacterial isolates ranged from 0.98 to 0.99. The constants (*a*, *b*) predicted for each standard curve were similar to the actual calculated values (with a probability of this similarity being due to chance alone of *<* 0.002), and the SE around the constants was low.

**Zones of inhibition** generated by agar well antimicrobial assays were converted to imipenem concentrations by use of the standard curves. Graphs of times after injection versus imipenem concentrations for all administration routes were similar to those drawn from HPLC data (Fig 1). The last plasma collection time point after IV or IM administration that yielded serum and inhibited bacterial replication was 2.0 hours. A small zone of inhibition was evident from samples obtained 4 hours after SC administration for several dogs: 3 with the reference strain, 3 with the 2-2225 isolate, and 2 with the 2-2559 isolate. No significant differences were detected between the reference strain of *E coli* and either of the multidrug-resistant clinical isolates when comparing concentrations of imipenem determined by the agar well antimicrobial assay.

**Response to injection**—An adverse effect associated with imipenem administration was vocalization (IM route, 2/6 dogs; SC route, 1/6 dogs) during injection. Hypersalivation (2/6 dogs) was observed during rapid IV administration. No other adverse reactions, either immediate or delayed, were noticed during imipenem administration or at the local injection sites.

**Blood analyses and urinalyses**—For 5 of the 6 dogs, no abnormalities were detected via the original

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**Table 1**—Pharmacokinetic parameters (mean ± SD) of imipenem after single-dose IV, IM, or SC administration (5 mg/kg) to 6 healthy dogs

<table>
<thead>
<tr>
<th>Pharmacokinetic parameters</th>
<th>Administration route</th>
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<tbody>
<tr>
<td></td>
<td>IV</td>
</tr>
<tr>
<td>AUC (mg·h/L)</td>
<td>20.2 ± 3.4</td>
</tr>
<tr>
<td>t$_{1/2}$ (h)</td>
<td>0.80 ± 0.23</td>
</tr>
<tr>
<td>CL (L/h/kg)</td>
<td>0.26 ± 0.06</td>
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<tr>
<td>Vss (L/kg)</td>
<td>0.32 ± 0.04</td>
</tr>
<tr>
<td>C$_{max}$ (mg/L)</td>
<td>NA</td>
</tr>
<tr>
<td>T$_{max}$ (h)</td>
<td>NA</td>
</tr>
<tr>
<td>F (%)</td>
<td>NA</td>
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*Harmonic mean ± pseudostandard deviation.

AUC = Area under the curve. t$_{1/2}$ = Half-life. CL = Total body clearance. Vss = Steady state volume of distribution. C$_{max}$ = Maximum plasma concentration. T$_{max}$ = Time until C$_{max}$ reached after drug administration. F = Bioavailability. NA = Not applicable.
observed in which the terminal t1/2 reflected the eliminations of the drug declined rapidly. However, after both administration of imipenem in our study, plasma concentrations of imipenem were similar to published values. The AUCs for imipenem administered by IM and SC routes studied. To the authors' knowledge, this is the first report of pharmacokinetic parameters of SC administration of imipenem to any species. In veterinary medicine, the ability to give continued injectable antimicrobials effectively by the SC route can make the difference between dogs remaining hospitalized or receiving continued care at home.

The t1/2 and AUC calculated after IV administration of imipenem were similar to published values. The t1/2 of imipenem after an IV bolus of 5 mg/kg to dogs was reported to be 0.51 hours.29 and the AUC was found to be 13.4 mg/L × hours.29 Following IV administration of imipenem in our study, plasma concentrations of the drug declined rapidly. However, after both IM and SC administrations, 'flip-flop' kinetics were observed in which the terminal t1/2 reflected the absorption rate rather than the elimination rate. Absorption of imipenem after IM or SC administration was characterized by a rapid burst effect followed by a sustained release of the drug, resulting in fairly constant plasma concentrations during a 2-hour period. The AUCs for imipenem administered by IM and SC routes were comparable to that for IV administration, indicating complete absorption of the drug. Thus, IM or SC administration of imipenem yields more desirable plasma concentration profiles than does IV administration.

The AUCs for imipenem administered by IM and SC administration are available for humans. The cost of these products is approximately $31/500-mg vial. Once reconstituted, commercial preparations of imipenem-cilastatin can be stored for no longer than 24 hours at 4°C.40 The combination product is a safer option for dogs with renal insufficiency, because the cilastatin should protect against renal tubular damage. A dose reduction is recommended in humans with renal failure, and this should apply to treatment of dogs as well.24,29

Mild changes in blood parameters reported in humans receiving imipenem include transient increases in liver enzyme activities, eosinophils, and platelets.38,39 In our study, 1 dog with preexisting high ALT activity had a persistent increase in ALT activity 12 days after administration of imipenem. It is unlikely that the increased liver enzyme activity in that dog was due to imipenem administration,
because the abnormality was present before and remained for 1 month after the study. Another dog had increased ALT and ALP activities 12 days after the third dose of imipenem, but liver enzyme activities returned to reference ranges 1 month after completion of the study. Based on the transient nature of the increased liver enzyme activity and its occurrence immediately after the study, the most likely cause was imipenem, metedomidine, or atipamezole. To the authors’ knowledge, there have been no reports of metedomidine or atipamezole causing increased liver enzyme activities. In humans, increased ALT and ALP activities associated with imipenem-cilastatin are usually transient and not associated with clinical abnormalities. However, hyperbilirubinemia and clinical icterus have been documented in humans after administration of imipenem-cilastatin.38,39 None of the dogs in our study developed clinical icterus.

Two of 6 dogs had eosinophilia 12 days after the last dose of imipenem administration. A CBC was not repeated to determine whether this abnormality resolved, and the importance of this delayed finding is not known. Transient eosinophilia has been reported in humans after administration of imipenem-cilastatin and is not associated with clinical problems.38,39 Other causes of eosinophilia could have been present in these dogs. All dogs were tested for heartworm disease and were receiving heartworm preventative medication. A fecal examination was not performed on the 2 dogs that developed eosinophilia. Although no abnormal skin lesions were seen, allergic skin disease was also not ruled out. Increases in platelet counts were not detected in any of the dogs in this study.

Mild abnormalities were also detected in some urine samples. Casts and WBCs in urine samples collected 24 hours after administration of imipenem may indicate mild renal inflammation secondary to imipenem administration. It is possible that this was seen because of using imipenem without cilastatin. Alternatively, the casts were low in number and may not be clinically important. One dog developed a UTI during the study. Clavamox administration successfully treated the Enterococcus infection. Although the UTI may have been related to imipenem-induced renal inflammation, it seems unlikely because few WBCs were consistently seen on urinalysis, and clinical signs and results of laboratory analyses never suggested systemic involvement or involvement of the upper portion of the urinary tract. An exhaustive workup to determine cause of the UTI was not done. The development of the UTI may have been related to a number of other factors as well. Some transient urinary retention occurred during the study in dogs hesitant to urinate while walked on a leash. Also, all dogs in the study were 3- to 9-year-old sexually intact male dogs, so subclinical prostatic infection would have to be considered.

Imipenem is an antimicrobial that has many indications for use in veterinary medicine. As nosocomial and multidrug-resistant infections in dogs become more common, the need for antimicrobials with a wide spectrum of activity, such as imipenem, increases. The tissue distribution of imipenem in dogs has not been reported and would be helpful for prescribing adequate dosages for achieving MICs in various tissues.

References