Biodistribution and Plasma Protein Binding of Zoledronic Acid

H. Markus Weiss, Ulrike Pfaar, Alain Schweitzer, Hansjörg Wiegand, Andrej Skerjanec, and Horst Schran

Novartis Pharma AG, Basel, Switzerland (H.M.W., U.P., A.Sc., H.W.); and Novartis Oncology, Florham Park, New Jersey (A.Sk., H.S.)

Received February 15, 2008; accepted July 10, 2008

ABSTRACT:
The bisphosphonate zoledronic acid is a potent inhibitor of osteoclast-mediated bone resorption. To investigate drug biodistribution and elimination, 14C-zoledronic acid was administered intravenously to rats and dogs in single or multiple doses and assessed for its in vitro blood distribution and plasma protein binding in rat, dog, and human. Drug exposure in plasma, bones, and noncalcified tissues was investigated up to 240 days in rats and 96 h in dogs using radiometry after dissection. Drug biodistribution in the rat and within selected bones from dog was assessed by autoradiography. Concentrations of radioactivity showed a rapid decline in plasma and noncalcified tissue but only a slow decline in bone, to ~50% of peak at 240 days post dose, whereas the terminal half-lives (50–200 days) were similar in bone and noncalcified tissues, suggesting redistribution of drug from the former rather than prolonged retention in the latter. Uptake was highest in cancellous bone and axial skeleton. At 96 h after dose, the fraction of dose excreted was 36% in rat and 60% in dog; 94 to 96% of the excreted radioactivity was found in urine. Blood/plasma concentration ratios were 0.52 to 0.59, and plasma protein binding of zoledronic acid was moderate to low in all species. The results suggest that a fraction of zoledronic acid is reversibly taken up by the skeleton, the elimination of drug is mainly by renal excretion, and the disposition in blood and noncalcified tissue is governed by extensive uptake into and slow release from bone.

Bisphosphonates are therapeutic agents for the treatment of bone resorative diseases. This class of drugs possesses a strong affinity to bone tissue and markedly inhibits osteoclast activity (Cremers et al., 2005). The more potent nitrogen-containing bisphosphonates interfere with cellular signal transduction pathways in osteoclasts by inhibiting protein prenylation (Luckman et al., 1998). Following intravenous dosing, bisphosphonates show initially a rapid disappearance from the systemic circulation with several short elimination phases that are followed by long elimination phases (t1/2 months to years) (Lin, 1996). Elimination occurs almost exclusively by the kidney through glomerular filtration (Singer and Minoofar, 1995), and drug clearance is decreased in patients with renal impairment (Saha et al., 1994; Berenson et al., 1997). Bisphosphonates show negligible biotransformation (Lin et al., 1994). The rate and extent of skeletal uptake and release of drug is dependent on the osseous turnover rate as well as the intrinsic bone affinity of the bisphosphonate (Kasting and Francis, 1992). The distribution of bisphosphonates within the skeleton is not homogeneous. Bisphosphonate plasma concentrations during the early period after administration generally are dose proportional, but the very low concentrations long term postdose mainly reflect bone remodeling and do therefore not directly relate to the applied dose level or clearance processes (Gertz et al., 1993). Accordingly, an accurate determination of bisphosphonate terminal half-lives is difficult, and AUC0–∞ and clearance parameters extrapolated using apparent terminal half-lives derived from a short observation period are open to question.

The protein binding of bisphosphonates is typically low to moderate and may be calcium and pH dependent (Lin et al., 1993; Lin, 1996). Bisphosphonates are known to produce renal effects ranging from transient proteinuria and alterations in creatinine clearance (Pecherstorfer et al., 1996) to acute renal failure (Bounameaux et al., 1983). Reducing dose, rate of infusion, and dosing frequency can mitigate these effects (Kanis et al., 1983).

One of the newer and highly efficacious nitrogen containing bisphosphonate drugs is zoledronic acid (Fig. 1A). It ranks highest in its inhibitory potency in in vitro assays of bone resorption and calcium turnover (Green et al., 1994) as well as in assays of tumor cell invasion (Boissier et al., 2000). Following intravenous administration, zoledronic acid concentrations in patients’ plasma show a rapid decline from the observed end-of-infusion peak to approximately 1% of peak by 24 h postdose, followed by prolonged, very low drug plasma concentrations declining to below the limit of bioanalytical methodology over a period of days to weeks (Chen et al., 2002). The urinary excretion of zoledronic acid, consisting exclusively of unchanged drug and representing approximately 40% of the administered dose, is essentially complete over the first 24 h after dose. This suggests that approximately 60% of dose is retained in the skeleton. Bone remodeling processes will slowly release retained drug back into the systemic circulation from where it is renally excreted. Zoledronic acid renal clearance is approximately 50% of the glomerular filtration rate in patients with normal renal function (Skerjanec et al., 2003). Tran-
sient rises in serum creatinine have been observed in a small fraction (<1%) of patients with postmenopausal osteoporosis receiving once-yearly 5-mg doses of intravenous zoledronic acid (Black et al., 2007). The reported risk of adverse renal events is higher in cancer patients because of their disease state and associated comorbidities, exposure to other potentially nephrotoxic agents, and higher doses and more frequent dosing (Lewiecki and Miller, 2007). In a trial of monthly intravenous doses of zoledronic acid to treat skeletal metastases in patients with lung cancer and other solid tumors, the proportion of patients with decreased renal function was not significantly different between the 4-mg zoledronic acid (15-min infusion) and placebo groups (10.9 versus 6.7%). (Rosen et al., 2003).

 Differences in kidney retention and plasma protein binding have been proposed as potentially contributing to perceived differences in the renal safety of bisphosphonates currently in clinical use (Bergen et al., 2007; Lewiecki and Miller, 2007). For a better understanding of the biodistribution and excretion of zoledronic acid, including exposure of the kidney to the drug, studies in rat and dog models were performed. The study in the rat (intravenous doses on 16 consecutive days) was designed to approximate the treatment regimen of cancer patients (multiple myeloma or bone metastases) who receive a single intravenous administration of zoledronic acid every 3 to 4 weeks (12 to 16 doses per year). The in vitro blood distribution and plasma protein binding of zoledronic acid were determined for rat, dog, and human. Differences in plasma protein binding between zoledronic acid and ibandronate have been proposed as an additional potential contributing factor to the perceived differences in the renal safety of these two drugs (Body et al., 2005). Therefore, plasma protein binding of ibandronate and zoledronic acid was tested side-by-side under controlled assay conditions to obtain robust comparative data.

Materials and Methods

Radiolabels and Stock Solutions. Zoledronic acid and ibandronate in 14C-labeled form (Fig. 1) were prepared by the Isotope Laboratory of Novartis Pharma AG (Basel, Switzerland). The specific radioactivity of zoledronic acid was 1.5 to 1.9 MBq/mg for the study in dog and the blood distribution investigations and 3.9 to 6.1 MBq/mg for the studies in rat. In the plasma protein binding study, the specific radioactivity was 6.9 and 4.3 MBq/mg for zoledronic acid and ibandronate, respectively. For intravenous dosing, 14C-labeled zoledronic acid was dissolved in 0.9% sodium chloride to a concentration of 0.06 mg/g (rat) and 0.15 mg/g (dog). The pH was adjusted to 7.0 using a 1.5-MM solution of NaOH. For the in vitro investigations, aqueous stock solutions were prepared by serial dilution.

Animal Studies. All animal studies were performed in accordance with Swiss animal welfare regulations. The individual intravenous doses of zoledronic acid were 0.15 mg/kg for all animals. This dose is similar to the maximum dose tested in the oncology phase 3 clinical trials of 8 mg i.v. per patient (0.13 mg/kg for a 60-kg patient).

Excretion studies. For the excretion experiments, rats were housed singly on steel grids in metabolism cages. Urine (ice-cooled vials) and feces were collected quantitatively and separately in daily intervals up to 96 h after a single dose. Day 1 urine was collected in two fractions (0 to 8 and 8 to 24 h).

Studies in Dogs. The distribution and excretion of zoledronic acid was studied in three 91-month-old male beagle dogs. Old dogs were used since they, in contrast to rats, achieve skeletal maturity, the growth plates close, and bone growth ceases, resulting in a bone metabolism corresponding to that in adult humans. The average weight of the dogs was 13.1 kg. Prior to and after administration of zoledronic acid, the dogs had free access to food (pellets 3353; Provimi Kliba SA, Kaiseraugst, Switzerland) and tap water. Each dog received 0.15 mg/kg 14C-zoledronic acid as a 15-min intravenous infusion.

Distribution by dissection and autoradiography. At 96 h after the infusion of 14C-zoledronic acid, the dogs were anesthetized by an injection of 10 ml of a 10% solution of Pentothal (Abbott AG, Zug, Switzerland) in distilled water into the cephalic vein and then bled by severing the carotid artery. Tissues were harvested and assayed by radiometry and by quantitative autoradioluminography (QAL) (Schweitzer et al., 1987; Johnston et al., 1990). Radiometry samples were prepared as described by Botta et al. (1985). The LOQ for the determination of total radioactivity was defined as 1.8-fold the total background count (AUC). The LOQ for the determination of total radioactivity was defined as 1.8-fold the total background count. The area under the tissue or plasma concentration-time curve was corrected for organ growth during the observation period.

Studies in Rats. Male albino rats [Tif: RAII(SPF)] 6–8-weeks old (190–250 g) were given a diet of NAFAG pellets 890 (Nahr und Futtermittel AG, Gossau, Switzerland) and had free access to tap water. Zoledronic acid was administered via a single bolus injection into the tail vein. For the distribution studies, a single 0.15 mg/kg dose, and, for the multiple dose studies, 0.15 mg/kg daily on 16 consecutive days, was administered. The short term (4 days) excretion of zoledronic acid in urine and feces was studied after the single dose administration.

Distribution by dissection. At each sampling time three rats were anesthetized with ether and bled out after cardiac puncture. The tissues were harvested by dissection at 5 min, 4 h, and 24 h after the 1st dose, 24 h after the 8th dose, and then 1, 16, 31, 64, 128, and 240 days after the 16th daily dose (n = 3 rats/time point). For radiometry samples prepared as described by Botta et al. (1985), bones were dissolved in 25% HCl. The limit of quantification (LOQ) for the determination of total radioactivity was defined as 1.8-fold the total background count. The area under the tissue or plasma concentration-time curve was corrected for organ growth during the observation period.

Distribution by autoradiography. Tissue distribution was qualitatively assessed by whole-body autoradiography up to 12 months after a single 0.15 mg/kg intravenous dose. Immediately after sacrifice, the rats were deep-frozen in a mixture of dry ice and hexane at approximately −75°C and then embedded in a 2% precooled semisolid gel of sodium carboxymethylcellulose. Multiple sections of 40-µm thickness were obtained at varying depths at approximately −20°C in a Cryomacrocut (Reichert-Jung, Nussloch, Germany) according to the method of Ullberg (1977). After freeze-drying, the sections were fixed onto transparent adhesive tape, and autoradiographs were obtained after 2 days of exposure of selected sections to imaging plates as described for the dog (Studies in Dogs).

Distribution by autoradiography. Tissue distribution was qualitatively assessed by whole-body autoradiography up to 12 months after a single 0.15 mg/kg intravenous dose. Immediately after sacrifice, the rats were deep-frozen in a mixture of dry ice and hexane at approximately −75°C and then embedded in a 2% precooled semisolid gel of sodium carboxymethylcellulose. Multiple sections of 40-µm thickness were obtained at varying depths at approximately −20°C in a Cryomacrocut (Reichert-Jung, Nussloch, Germany) according to the method of Ullberg (1977). After freeze-drying, the sections were fixed onto transparent adhesive tape, and autoradiographs were obtained after 2 days of exposure of selected sections to imaging plates as described for the dog (Studies in Dogs).
order to minimize the background signal. The duration of exposure was chosen to allow detection of approximately 1 dpm/mg. At the end of the exposure, the imaging plates were first left at room temperature in the dark for 3 min, then transferred into a Fuji BAS 5000 confocal phosphorimager and scanned in steps of 25 μm. The resulting photostimulated light data files were corrected by background subtraction and processed electronically with a MCID/Elite (7.0) image analyzer (Imaging Research, St. Catharines, ON, Canada). The limit of detection (LOD) was taken as the sum of the mean of the background (n = 10 measurements) and 3 standard deviations on this mean; the LOQ was taken as 3× the LOD. Under the conditions of this study, the LOD and LOQ were 0.038 and 0.11 nmol/g, respectively, in the central cavity of the bones, and 0.24 and 0.74 nmol/g, respectively, in the compact bone, spongy bone, and periosteum. The image files were processed using the Adobe Photoshop Elements 2.0 software (Adobe Systems Inc., San Jose, CA).

**Radioactivity.** Radioactivity in the rat and the dog blood, plasma, tissues, urine, and feces samples was determined by liquid scintillation counting using a TriCarb liquid scintillation system (PerkinElmer Life and Analytical Sciences, Boston, MA) according to Botta et al. (1985). The LOQ was defined as 1.8-fold the background radioactivity.

**In Vitro Bioavailability.** Blood and plasma were obtained from male albino rats of the Hanover Wistar strain and from male beagle dogs. Human blood and plasma were from healthy male volunteer donors. Lithium heparin was used as anticoagulant. Animal blood and plasma were used as pools (n = 3). For human blood and plasma, three individual samples were used.

**Blood distribution.** Whole blood was spiked with zoledronic acid to concentrations of 30 to 5000 ng/ml, incubated for 30 min at 37°C, and centrifuged for 15 min at 1000 g for plasma separation. Drug-related radioactivity in blood (C_b) and plasma (C_p) was determined in triplicate by the combustion method. The hematocrit (H) was determined in triplicate. The fraction in plasma was determined as (C_b/C_p) · (1/H).

**Plasma protein binding.** Separation of plasma protein bound and unbound drug was performed by ultrafiltration. Initially, the recovery and free permeation of zoledronic acid and ibandronate in phosphate-buffered saline were analyzed: phosphate-buffered saline was spiked and 3×1-ml aliquots were introduced into Centrifree devices (molecular cut-off 30 kDa; Amicon Inc., Beverly, MA) and spun at 2000 g for 1 min (filtrate ≤500 μl). Samples were taken from the spiked solution, the filtrate, and retentate and analyzed by liquid scintillation counting, and filtrate and retentate weights were determined. The radioactivity recovery in the device was above 90%, and the free permeation through the membrane (concentration in filtrate/concentration in retentate) was ≥0.95, suggesting no relevant bias due to the separation technique. The plasma pH was adjusted to 7.4 ± 0.1 (using 5% lactic acid or 1 N NaOH) before spiking to concentrations of 2 to 2000 ng/ml zoledronic acid or ibandronate. After incubation for 30 min at 37°C under constant gentle agitation, the spiked plasma samples (n = 3) were centrifuged at 2000 g for 10 min in the prewarmed Centrifree devices. Total radioactivity was determined by liquid scintillation counting in the ultrafiltrate [concentration of unbound drug (C_u)] and in the sample introduced into the reservoir before ultrafiltration [concentration in plasma (C_p)]. The unbound fraction in plasma was calculated as C_u/C_p. Values of 30 dpm above background (14C) were defined as the LOQ for radioactivity analysis; all reported values were above this limit.

**Results**

**Tissue Distribution in Rats.** Five minutes after a single intravenous dose of 14C-zoledronic acid (0.15 mg/kg) to adolescent rats, the highest concentration of drug-related radioactivity was detected in plasma (5383 pmol/g), which declined very rapidly (29 pmol/g at 4 h and 3.9 pmol/g at 24 h) because of extensive distribution and renal elimination. The highest tissue concentration was initially observed in the kidney (3218 pmol/g at 5 min after dosing). This also dropped rapidly within 4 h to 403 and 183 pmol/g at 24 h postdose. At 4 h, the highest concentrations were detected in the calcified tissues, which, in contrast to noncalcified tissue, continued to show uptake of zoledronic acid-related radioactivity up to 24 h postdose (Fig. 2A, inset). This preferential and extensive uptake of zoledronic acid into bones was more pronounced after multiple intravenous doses of 0.15 mg/kg 14C-zoledronic acid (daily on 16 consecutive days) to adolescent rats. There was no indication of saturation of available binding capacity in bones: zoledronic acid concentrations in the calcified tissues were 6- to 7-fold higher at 24 h after the 8th dose and 10- to 12-fold higher at 24 h after the 16th dose compared with the single dose. 14C-Zoledronic acid-related radioactivity in the calcified tissues declined very slowly over the observation period of 240 days (Fig. 2A). The AUC_{0–256d} values of the bone tissues were 100- to 1500-fold higher than in the nonmineralized tissues and at least 4000-fold higher than in plasma (Table 1; Fig. 2). Sixteen days after the last dose, the drug concentrations in rat plasma and all nonmineralized tissues (listed in Materials and Methods) dropped below 0.4 nmol/g, whereas concentrations in bone remained high at >10 nmol/g up to 240 days after dosing. Considering the entire 240-day observation period, the con-
centrations in the analyzed bones (cranium, vertebrae thoracales, and tibia) decreased by a factor of approximately 2. Based on this observation, approximate half-lives of 150 to 200 days were estimated. When the observed tibia concentrations were corrected for bone growth, no elimination was discernible within the 240 day post-treatment period. Qualitatively similar results were obtained by whole body autoradiography after a single intravenous dose of 0.15 mg/kg 14C-zoledronic acid (Fig. 3). Five minutes after dosing, high concentrations of drug-related radioactivity were found in the blood and the highly perfused organs such as liver, kidney, and bone, whereas, at 1 month after dosing (Fig. 3B) and up to the final autoradiogram taken at 12 months postdose, visible exposure was confined to bones.

**Tissue Distribution in Dogs.** In skeletally mature dogs, 96 h after administration of a single intravenous infusion of zoledronic acid, the plasma and blood radioactivity concentrations as well as concentrations in brain and muscle were below the LOQ (Fig. 4). Measurable concentrations were found in all of the other tissue specimens assayed, suggesting a high volume of distribution. As observed for the rat, there was a striking difference between the concentrations measured in noncalcified tissues versus bone tissues. Exposure was higher in bones of the axial skeleton compared with the appendicular bones or the head (including teeth). Analysis of the distribution of radioactivity within selected bones of the dog by quantitative autoradiography revealed high labeling in the cancellous bones. Distribution was mainly into the more dynamic portions of bone and areas with a high surface area, such as spongy bone. Uptake of drug in compact bone was lower, and only little labeling was detected in the central cavity of bone. Teeth and jaws showed no exceptional differences compared with noncalcified (cancellous bone) compared with compact bone or the noncalcified (non-cancellous) bone.

**Excretion of Zoledronic Acid in Rats and Dogs.** After administration of a single intravenous dose of 0.15 mg/kg zoledronic acid to adolescent rats, 36.0% of the dose was recovered in urine and feces within 96 h (Fig. 6A). Of this, 96% was present in the urine (34.6% of the dose) and 4% in the feces (1.4% of the dose). In skeletally mature dogs, the total cumulative mean recovery was 60.5% within 96 h of dosing, with 94% of this being excreted in the urine (Fig. 6B).

**In Vitro Blood Distribution and Plasma Protein Binding.** In blood of human, dog, and rat, zoledronic acid was mainly located in the plasma fraction. The mean fraction of drug in plasma was 90 ± 6%, 95 ± 5%, and 91 ± 9% for rat, dog, and human, respectively, corresponding to mean blood to plasma concentration ratios of 0.54, 0.52, and 0.59.

**Discussion**

The biodistribution studies in rat and dog demonstrate several orders of magnitude higher affinity of zoledronic acid for calcified compared with noncalcified tissue. As shown in rats, bone exhibits a large storage capacity for drug, which was not saturated after 16 daily doses, representing the cumulative annual dose of zoledronic acid in patients with metastasis to bone. In the skeleton of mature dogs, the uptake of drug was higher in the axial skeleton compared with appendicular bones or the head. It was most pronounced in the dynamic parts of bone and in regions with a high surface area (cancellous bone) compared with compact bone or the noncalcified central cavity of bone. Teeth and jaws showed no exceptional differences in drug uptake compared with other bones.

Bisphosphonates are known to complex with hydroxyapatite, thereby leading to a high sequestration of the drug into bone (Cremers et al., 2005). The prolonged but extremely low exposure level of zoledronic acid results in a limited accumulation of drug in the bone tissue.
Zoledronic acid-related radioactivity in plasma and noncalcified tissues seen in the rat 240 days after the last dose likely reflects the slow release from bone subsequent to its initial rapid uptake, rather than a very low systemic clearance (Lin, 1996). This parallels the observations with other bisphosphonates (Lin et al., 1991). The accuracy of reported terminal half-lives of bisphosphonates may be strongly influenced by detection limits, observation times, and sampling schedules as well as differences in bone turnover due to age and disease state. Thus, the reported 2.2-h half-life of zoledronic acid in the dog (Martin-Jimenez et al., 2007), which was derived from a 9-h observation period, does not represent the terminal elimination phase. The half-life of ibandronate in the rat kidney of 24 days, determined by Bauss and Russell (2004), has been cited as a potential reason for the purportedly better renal safety of this bisphosphonate (Body et al., 2005). However, when calculating the terminal kidney half-life of ibandronate by more appropriately using the last 2 (or 3) values rather than all 4 of the concentration values reported by Bauss and Russell (2004), it turns out to be 138 days (or 90 days), which is similar to the half-lives reported for other bisphosphonates, including zoledronic acid.

Bisphosphonates are generally not subject to hepatic metabolism (Lin, 1996; Cremers et al., 2005), which is in line with the observed relatively low concentrations of zoledronic acid in the liver compared with other nonosseous tissue in rats and dogs (Figs. 2 and 4) and negligible fecal excretion of drug following intravenous dosing (Fig. 6). Pamidronate and alendronate represent examples of other bisphosphonates where drug concentrations in rat kidney are higher than in the liver (Pongchaidecha and Daley-Yates, 1993; Lin, 1996).

Variability in the pharmacokinetic profiles of bisphosphonates between and within species is mainly attributable to differences in the rates of renal excretion and uptake into calcified tissues. Within 1 to 5 days after administration to humans, 30 to 60% of the bisphosphonate dose is renally excreted (Lin, 1996; Kino et al., 1999). This was also observed for zoledronic acid (Fig. 6). Since uptake into bone and renal elimination are competing processes, a faster bone uptake should result in a lower amount of drug excreted by the kidney. Therefore, predominance of bone formation in the adolescent rats could explain their lower degree of renal excretion of zoledronic acid as compared with the skeletally mature dogs (Fig. 6). In addition, Lin et al. (1994) reported a higher apparent uptake clearance of alendronate by bone for rat compared with dog. A similar difference for zoledronic acid may contribute to the observed difference in excretion in rat and dog.

Zoledronic acid did not partition strongly into blood cells. Binding to plasma proteins was moderate in rat plasma and low in dog and human plasma. In the absence of active transport processes, the extent of tissue distribution of a drug is driven by the ratio between tissue
binding and plasma protein binding. For zoledronic acid, and likely bisphosphonates in general, the balance is greatly in favor of the tissues due to the high affinity of the drug for bone. Plasma protein binding is low and clearly not capable of significantly restricting the uptake of these drugs by bone. Lin et al. (1991) reported that the apparent uptake clearance of alendronate by bone was higher in rat compared with dog, despite the more than 10-fold higher plasma protein binding in rat.

Both rat and dog show the lowest exposure to zoledronic acid-related radioactivity in blood and plasma compared with all other tissues analyzed (Table 1; Figs. 2 and 4). Earlier reports have suggested that high bisphosphonate doses accompanied by high concentrations in plasma overload renal elimination mechanisms, and the retained compound can damage renal cells (Body et al., 2005). These reports also have speculated that a higher plasma protein binding of ibandronate as compared with zoledronic acid may contribute to differences in the renal safety of the two drugs (Body et al., 2005; Bergner et al., 2007). The reported concentration of 14C-ibandronate in the rat kidney, 0.3% of 0.10 mg/kg dose 24 h after drug administration (Bauss and Russell, 2004), or approximately 118 pmol/g assuming 250 g/rat and 2 g/rat kidney (Davies and Morris, 1993), is proportional to the observed zoledronic acid 24-h kidney concentration of 184 pmol/g per 0.15 mg/kg dose, indicating no difference in kidney retention between the two drugs. Plasma protein binding of zoledronic acid was observed to be dependent on plasma free calcium levels and pH, as has been reported previously for alendronate (Lin et al., 1993; Lin, 1996). Slight shifts in the pH of plasma during in vitro incubations (e.g., due to loss of carbon dioxide) may contribute, together with other factors, to interstudy variability of plasma protein binding, possibly leading to the wide differences in reported values for ibandronate (Barrett et al., 2004; Dooley and Balfour, 1999). Under rigorous testing conditions, the protein binding of ibandronate in human plasma was found to be slightly lower than that of zoledronic acid. Both drugs showed a qualitatively similar binding pattern in plasma of the tested species (rat > dog > human; Table 2) and a slight concentration dependence in dog and human plasma. These in vitro findings are in line with the available data reported for other bisphosphonates: for alendronate, plasma protein binding in the rat is higher than in dog and human (Lin et al., 1999), and alendronate and etidronate show a concentration dependence of plasma protein bind-
ing (Lin, 1996). The renal clearance of zoledronic acid, 60 ml/min (Skjerjane et al., 2003), is identical to that reported for ibandronate (Barrett et al., 2004), consistent with our finding of comparable plasma protein binding of the two drugs.

The biodistribution results suggest that zoledronic acid disposition in noncalcified tissue is governed by the extensive uptake into and slow release from bone, as generally observed for bisphosphonate drugs. The findings that zoledronic acid and ibandronate show similar dose normalized levels in the rat kidney, have comparable elimination half-lives in rat kidney, and do not appreciably differ in their plasma protein binding across rat, dog, and human counter the reported claims of pharmacokinetic and biodistribution differences providing the basis for potential renal safety differences in animals and humans.

Acknowledgments. We are grateful to Heidi Hügli, Marcel Frensenau, Barbara Handisch, and Lothar Dillo for technical assistance, Thomas Mönius and Ines Rodriguez for preparation of 14C labeled compounds, and Alessandro Probst, Helmut Schütz, and Jonathan Green for valuable advice and comments.

References


Address correspondence to: H. Markus Weiss, Novartis Pharma AG, DMPK, WSJ-210.4.25, CH-4002 Basel, Switzerland. E-mail: markus.weiss@novartis.com