DISPOSITION AND METABOLISM OF RIFAPENTINE, A RIFAMYCIN ANTIBIOTIC, IN MICE, RATS, AND MONKEYS

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ABSTRACT:

Rifapentine is a semisynthetic, rifamycin-class antibiotic (Arioli et al., 1981) under development for the treatment of pulmonary tuberculosis and for the prophylaxis of Mycobacterium tuberculosis and Mycobacterium avium complex infections. The antimicrobial spectrum of rifapentine is similar to that of its homologues, rifampin and rifabutin (Arioli et al., 1981; Dickinson and Mitchison, 1987; Heifets et al., 1990; Yates and Collins, 1982); however, rifapentine has demonstrated greater therapeutic efficacy in experimental mycobacterium infections, compared with rifampin (Arioli et al., 1981; Pattyn, 1987; TrufUfl-Pernot et al., 1983).

The structure of rifapentine is shown in fig. 1. Rifapentine differs from rifampin by the presence of a cyclopentyl ring instead of a methyl group at the piperazinyl moiety, which makes rifapentine more lipophilic. Pharmacokinetic studies of rats (Assandri et al., 1978) and healthy volunteers (Birmingham et al., 1978, Buniva et al., 1983) dosed with rifapentine indicate differences in the pharmacokinetic profiles between it and rifabutin. Peak serum concentrations obtained after oral or iv doses of rifapentine were comparable to those produced by rifampin, but the elimination half-lives of the two drugs were markedly different, with rifapentine persisting in serum about 4–5 times longer.

Previous studies in mice, rats, and monkeys have identified the pharmacokinetic profile of rifapentine and its metabolite, 25-desacetyl-rifapentine. In a study designed to assess disposition of radiolabeled-rifapentine in rats, Assandri and colleagues (Assandri et al., 1978) identified feces as the major route of elimination of intact rifapentine and metabolites. In a separate study (Assandri et al., 1984), the pharmacokinetics of 14C-rifapentine in rat, mouse, and rabbit were investigated. The mean bioavailability of rifapentine after a single oral dose of 3 mg/kg in rats was 84%. The primary route of elimination of radioactivity was through the feces (about 90%), with less than 10% of the radioactivity recovered in urine. The mean terminal elimination half-life of rifapentine was 14–21 hr in rats, 17–23 hr in mice, and only about 2 hr in rabbits. In these previous rifapentine studies, no attempts were made to determine mass balance or identify compounds.

The purpose of our studies was to determine the disposition and biotransformation of 14C-rifapentine in mice, bile duct–cannulated and uncannulated rats, and monkeys after either a single oral or iv dose. Mass balance studies included 14C analysis of urine, feces, bile, cage wash, carcasses, and cage air collected for up to 120 hr postdose. Radiochromatograms were similar for fecal samples from animals dosed by iv or orally. Ten regions of radioactivity were observed in mouse and rat fecal sample radiochromatograms, and seven regions of radioactivity were observed in monkey fecal sample radiochromatograms. The most abundant compound identified in feces was usually intact rifapentine (27%–41% of dose in mouse, 3%–35% of dose in rat, and 17%–29% of dose in monkey). Other peaks identified or characterized in feces based on liquid chromatography/ultraviolet/14C and/or liquid chromatography/mass spectrometry methods included 25-desacetyl-rifapentine, 3-formyl-25-desacetyl-rifapentine, and 3-formyl-rifapentine. The compounds rifapentine, 25-desacetyl-rifapentine, and 3-formyl-rifapentine were present in rat bile samples. These studies show that the metabolism and disposition of rifapentine in mice, rats, and monkeys were similar.
Materials and Methods

Chemicals and Dose Formulations. Standards of rifapentine, 25-desacetyl-rifapentine, and 3-formyl-rifapentine were synthesized at Hoechst Marion Roussel (Cincinnati, OH). The labeled drug was prepared with the imine carbon containing the $^{14}$C label (fig. 1). Unlabeled drug was coprecipitated with labeled drug to produce $^{14}$C-rifapentine and to limit the radioactivity exposure to less than 40 mCi/animal. The radiochemical purity was 98.41% and the chemical purity was 97.71%.

The iv dose (2 mg/ml solution for mice and 10 mg/ml solution for rats and monkeys) was prepared at 10°C by dissolving $^{14}$C-rifapentine in 0.1% sodium ascorbate/2% alcohol (95% ethanol)/20 mM sodium hydroxide. The specific activities of the oral doses were 24.0, 23.0, and 0.9 mCi/mg for mouse, rat, and monkey, respectively. After all of the test compound was dissolved, a small amount (about 3%) of 0.1 N HCl was added to adjust the pH to 10.0 ± 0.1. Immediately before administration, the iv dose was warmed to room temperature. The oral dose was prepared as a suspension (1 mg/ml for mice and rats,

Fig. 1. Structures of compounds either identified (standard exists) or characterized (MS data only) in mouse, rat, and monkey feces after iv or oral doses or $^{14}$C-rifapentine.

All molecular weights are the monoisotope mass. The asterisk (*) denotes the position of the $^{14}$C label.
and 5 and 20 mg/ml for the 10 and 40 mg/kg doses for monkeys) by slowly adding 0.5% aqueous methylcellulose to 14C-rifapentine while triturating.

Animals. Mice. Two groups, each consisting of six male NMRI mice weighing between 25 and 30 g, received either a single iv or oral 10 mg/kg dose of 14C-rifapentine. The mice were housed in glass Roth-type cages in a light- and temperature-controlled room. The animals fasted overnight before dosing, then had free access to food and water after dosing. The iv dose was administered into the tail vein and the oral dose was delivered by gavage.

Rats. Three parallel groups of four male Wistar rats weighing 250–300 g were administered 10 mg/kg 14C-rifapentine either orally (two groups) or by iv (one group). The rats were housed in glass Roth-type cages in a light- and temperature-controlled room. The animals fasted overnight before dosing, then had free access to food and water after dosing. The iv dose was administered into the tail vein and the oral doses were delivered by gavage.

The bile ducts of rats from two groups (oral and iv) were cannulated. Rats were anesthetized with isoflurane. A midline incision was made, and the common bile duct was isolated. A cannula, consisting of PE10 tubing attached to PE50 tubing, was inserted (PE10 end) into the bile duct slightly proximal to the small intestine. The cannula was exteriorized at the nape of the neck and was channeled through a rodent jacket and spring tether. The rats were housed in Roth-type metabolism cages. The rats were allowed to recover from the effects of the anesthetic before dosing.

Monkeys. One group of four male Cynomolgus monkeys weighing 3.0–5.0 kg was administered one of three treatments: 10 mg/kg 14C-rifapentine administered orally or by iv, or 40 mg/kg 14C-rifapentine administered orally. Monkeys were housed in individual stainless steel cages designed for the separation and collection of urine and feces. The treatments were administered sequentially, with a 3-week washout period between treatments. Before each dose, the animals fasted overnight and until about 4 hr postdose. The iv dose was administered via the saphenous vein, and the oral doses were administered via intragastric intubation.

All animals were housed according to the National Institutes of Health Guide for the Care and Use of Laboratory Animals, and all studies were conducted under protocols approved by the Hoechst Marion Roussel Animal Care and Use Committee. Anesthetic was not used during the dosing or sample collection for any of these studies.

Sample Collection. The collection times for urine, bile (rats only), and fecal samples were intervals of 0–6, 6–12, 12–24, 24–48, 48–72, 72–96, and 96–120 hr after the dose was administered. The samples were collected in tared glass containers for mice and rats and in plastic containers for monkeys. Cage washes were conducted at the end of each urine-collection interval (mice and rats) or after each 24-hr fecal collection (monkeys). In addition, for mice and the uncannulated rats, cage air was scrubbed through a CO2 trapping solution (5 M ethanolamine in 2-methoxyethanol, 3:7 v/v) to trap any 14C-labeled CO2 at 12-hr intervals. At the end of the mass balance study, carcasses of mice and rats were analyzed for radioactive content.

Rifapentine is known to degrade chemically by hydrolysis and oxidation (G. Beck, unpublished data, 1994). Therefore, all samples were collected over dry ice to minimize degradation, frozen, and stored at –20°C until analysis. In addition, 10 µl of 250 µg/ml ascorbic acid was added to all urine and bile samples immediately upon collection to stabilize the samples. Fecal samples were homogenized in methanol:water (1:1, v/v) containing 2.5 µg/ml ascorbic acid.

Fecal Sample Preparation for LC/UV/14C or LC/MS. Fecal samples were thawed and mixed with a methanol solution (1 µg/ml ascorbic acid) in a ratio of about 2 ml/g of fecal homogenate. Samples were centrifuged at 3500 rpm for 2 min after mixing; the supernatant was then transferred to a test tube. This extraction procedure was repeated two more times and the extracts were combined and dried in a vacuum centrifuge. The dry residue was reconstituted in a methanol/water solution, filtered, and injected onto the HPLC system.

Radioactivity Analysis. Concentrations of 14C in urine, bile, cage wash, carcass, and expired air were measured by direct liquid scintillation counting (LSC) with external standardization for quench correction. Weighted aliquots of urine, bile, and cage wash were transferred to vials, and 5–15 ml of Ultima Gold® (Packard, Meriden, CT) scintillant added. After incubation of carcass with Soluene-350 (Packard), 10 ml of Hionic Fluor® (Packard) was added as scintillant. The CO2 trapping solution from expired air was aliquoted into LSC vials and 10–15 ml of Permafluor V® (Packard) was added as scintillant. Activity of 14C in feces was determined after combustion of the dried samples in a Packard 307 oxidizer. The CO2 generated by the combustion was trapped in 9 ml of Carbo-Sorb E® (CO2 absorber; Packard), then 11 ml of Permafluor was added as scintillant. Each sample was counted by liquid scintillation analysis on a Tri-Carb 2300TR Liquid Scintillation Analyzer (Packard). All samples were counted for at least 5 min or 100,000 counts.

Metabolite Characterization. HPLC radiochromatograms and mass spectra were obtained for samples with radioactivity greater than 5% of the total dose. The HPLC system used for metabolite profiling included a Hitachi AS-4000 autosampler with a Hitachi L-6200A pump (San Jose, CA), Hitachi L-4250 UV detector set at 480 nm, and an INUS Systems RAM radioactivity detector. A Mac-Mod Zorbax RX-C8 column (250 × 4.6 mm, 5 µm; Chadds Ford, PA) with a gradient mobile phase was used to separate 14C-rifapentine-related compounds in urine, bile, and feces. Mobile phase A was 95% water/5% acetonitrile/0.1% heptafluorobutyric acid, and mobile phase B was 100% acetonitrile/0.1% heptafluorobutyric acid. A linear gradient was used to separate compounds in the extracts. Initially, 100% A flowed through the column for 1 min after the injection, followed by a linear ramp to 100% B in 35 min, where it remained for 10 more min. The flow rate was set to 1 ml/min and a 200 µl aliquot sample was injected onto the HPLC system. The LC/MS system used to analyze the fecal extracts and bile samples consisted of a Gilson Model 231 autosampler (Middleton, WI), HPLC pump (Microm BioResources Inc., Auburn, CA), Waters Model 490-MS UV detector with the Programmable Detector set at 480 nm (Milford, MA), and a Finnigan MAT TSQ 710 triple quadrupole mass spectrometer (San Jose, CA). The column and the mobile phase were identical to those used for the LC/UV/14C profiling analyses. The mass spectrometer was set in the positive ion mode to scan from 400 to 1400 atomic mass units in 2 sec. The standard Finnigan electrospray ionization source was used to produce ions. The heated capillary temperature was 150°C and spray voltage set at 4500 volts. The 1 ml/min flow from the LC column was split 10:1 before entering the mass spectrometer.

Determination of Extraction Efficiency and Column Recovery of 14C-Labeled Components. To determine extraction efficiency, mouse, rat, and monkey fecal samples were accurately weighed and processed as described above. A small amount of methanol was used to redisolve the dry residue, and the liquid was transferred to a scintillation vial along with scintillant. The sample was counted using a liquid scintillation counter. The extraction efficiency was calculated by comparing the radioactivity of a processed sample with the theoretical maximum value, based upon combustion data. To determine column recovery, two injections from the same processed solution of selected feces samples were made, one with the LC column in place and one without the LC column (assumed to be 100% recovery). Three-minute fractions of eluate were collected from the column and mixed with 15 ml of scintillant. The sample was counted using LSC. For the 100% recovery sample, the column was removed and the LC eluate was collected for 3 min and mixed with 15 ml of scintillant.

Results

Extraction Efficiency and Column Recovery. The extraction efficiencies for mouse and rat fecal samples were relatively high, ranging from 89% to 93% (mean, 90%) in mice, and from 82% to 105% (mean, 93%) in rats. Since the extraction efficiencies were high, the assumption was that no significant metabolite was unaccounted for in fecal homogenates in mice or rats. In monkeys, the extraction efficiencies ranged from 67% to 71% (mean, 69%). Because the extraction efficiencies were relatively low in this species, predose monkey feces was spiked with 14C-rifapentine to determine if a difference between spiked and study samples was observed. The extraction efficiencies for the spiked predose fecal samples ranged from 78% to 88% (mean, 83%). Thus a matrix effect may account for in the lower yield. The mean column recovery of duplicate measurements was 102% in mice, 102% in rats, and 103% in monkeys. No significant amount of radioactivity was retained on the column.

Mass Balance Studies. Mice. Mass balance data of excreted compounds after iv and oral administration of 14C-rifapentine in mice are
summarized in Table 1. After iv administration, radioactivity excreted in urine and feces during the 120-hr collection periods was 4.9% and 73.0%, respectively. The cage wash contained 3.7% of the dose, and 3.4% of the dose remained in the carcass. Expired air did not contain any radioactivity over the first 24 hr postdose. The mass balance after 144 hr postdose was 84.2%. After oral administration in uncannulated rats, radioactivity excreted in urine, feces, and bile over the first 24 hr postdose was 3.1%. Expired air did not contain any radioactivity greater than 5% of dose were submitted to LC/UV/14C. For all three species, the oral and iv routes of rifapentine administration produced similar metabolite profiles in feces samples.

Metabolites were identified by comparison of the LC/UV/14C chromatographic peak retention time of an authentic standard to that resulting from injection of a fecal or bile extract. Supporting identification data was produced using LC/MS. Standards of rifapentine and analogs gave a poor response by either atmospheric pressure chemical ionization or electrospray ionization LC/MS. No \((\text{M}+\text{H})^+\) ions were observed after injection of standards using positive atmospheric pressure chemical ionization, and the electrospray ionization \((\text{M}+\text{H})^+\) ion response of rifapentine was approximately 100 times less than the typical pharmaceutical compounds we study. This is possibly the result of factors including poor ionization efficiency and poorer transmission of higher mass ions through the quadrupole. In an effort to improve MS response, different mobile phase buffers and pH conditions were tested but did not yield any improvements. Thus, LC/MS could only be utilized to confirm the presence of some of the metabolites that were identified with the LC/UV/14C system.

**Mice.** A summary of identified metabolites and their abundance in fecal samples is presented in Table 2. Representative LC/14C chromatograms (fig. 2) resulting from injection of a mouse fecal extract contained 10 regions of significant radioactivity. The radiochromatograms were similar after either intravenous or oral dosing. One of the abundant regions of radioactivity (8–18% of dose) eluted form the column as a broad peak with retention time from 9 to 17 min. Occasionally, small peaks were observed on top of the broad peak. Unsuccessful attempts were made to resolve the broad peaks into abundant regions of radioactivity (8–18% of dose) eluted form the column as a broad peak with retention time from 9 to 17 min. Occasionally, small peaks were observed on top of the broad peak. Unsuccessful attempts were made to resolve the broad peaks into individual components using other columns and mobile phase components and conditions. The entire region was summed for the purpose of calculating percentage of dose.

Four of the peaks in each radiochromatogram were assigned as 25-desacetyl-rifapentine, rifapentine, 3-formyl-25-desacetyl-rifapentine, and 3-formyl-rifapentine. Initial assignments were made by matching LC/14C/UV retention times with standards. Separate LC/MS analyses discussed below were subsequently performed to verify some of these assignments.
The most abundant compound identified in mouse feces was usu-
ally rifapentine (27–41% of the dose). The other three compounds
identified, 25-desacetyl-rifapentine, 3-formyl-25-desacetyl-rifap-
entine, and 3-formyl-rifapentine, accounted for 7–11%, 0–1%, and
3–6% of the dose in mice, respectively. Together, these four com-
 pounds constituted approximately 65% of the radioactivity in feces.

3-Formyl-rifapentine is a known hydrolysis product of rifapentine in
an aqueous environment (G. Beck, unpublished data, 1994). To ensure
that processing did not degrade rifapentine in samples, an aliquot of
predose fecal homogenate was spiked with 14C-rifapentine and the
sample was prepared as previously described. No peaks other than
rifapentine were observed in these spiked radiochromatograms.

Two compounds, rifapentine and 3-formyl-rifapentine, in mouse
feces could be confirmed by LC/MS. Injection of 48-hr mouse fecal
extracts into the LC/MS resulted in reconstructed ion chromato-
graphic peaks with retention times and mass spectra consistent with
authentic standards. No mass spectra (unique from predose fecal
extracts) were observed for any of the other radioactive compounds
eluting from the LC column, presumably because of low concentra-
tions and/or low ionization yields. The structures for compounds
found in mouse, rat, and monkey urine, bile, and fecal extracts based
on LC/UV/14C and/or LC/MS data are shown in fig. 1.

FIG. 2. Radiochromatogram resulting from injection of a mouse sample (72-hr)
fecal extract after the animal received a 10 mg/kg iv dose of 14C-rifapentine.

Rat Bile. A summary of identified metabolites and their abundance
in bile samples is presented in table 2. A representative radiochro-
matogram from a rat bile extract is shown in fig. 4. Radiochromato-
grams from rat bile extracts were similar after either iv or oral dosing.
Three chromatographic peaks observed in the LC/14C chromatograms
were assigned as 25-desacetyl-rifapentine, rifapentine, and 3-formyl-
rifapentine, respectively, based upon a comparison of the retention
time and mass spectrum to an authentic standard. The other identified metabolites, based upon LC/UV/14C data, were not
observed by LC/MS, probably because of the relatively poor MS
response of rifapentine and analogs combined with their low concen-
tration.
The other identified metabolites, based upon LC/UV/\(^{14}\)C data, were not observed by LC/MS probably because of the relatively poor MS response of rifapentine and analogs combined with their low concentration. No mass spectra (unique from predose fecal extracts) were observed for any of the other radioactive compounds eluting from the LC column, presumably because of low concentrations and/or low ionization yields.

**Monkeys.** A representative LC/\(^{14}\)C chromatogram for the 10 mg/kg iv dose in monkeys that was obtained 72 hr postdose is presented in fig. 5. Most radiochromatograms obtained after the iv and oral doses were similar, with seven regions of significant radioactivity observed. As with the other species, one of the abundant regions of radioactivity (12%–19% of the dose) eluted from the column as a peak with retention time from 6 to 15 min. Four peaks in each radiochromatogram were assigned as 25-desacetyl-rifapentine, rifapentine, 3-formyl-25-desacetyl-rifapentine, and 3-formyl-rifapentine. In general, rifapentine represented 17–29% of the dose in a sample, while 25-desacetyl-rifapentine represented 14–20% of the dose. These were the largest amounts of radioactivity eluting from the column.

Identification or characterization by LC/MS was successful for four of the radioactive compounds in monkey feces eluting from the column. Fig. 6 illustrates a typical reconstructed ion chromatogram obtained after the injection of 200 \(\mu\)l of reconstituted fecal extract (72 hr postdose) from a monkey dosed with rifapentine 10 mg/kg iv. Four peaks were observed and their mass spectra were consistent with the assignment of 25-desacetyl-rifapentine, rifapentine, 3-formyl-25-desacetyl-rifapentine, and 3-formyl-rifapentine, respectively. As each compound eluted from the column, the \((M+H)^{+}\) ions were prominent in the respective mass spectra at \(m/z\) 684.6, 726.6, 835.7, and 877.7 for 3-formyl-25-desacetyl-rifapentine, 3-formyl-rifapentine, 25-desacetyl-rifapentine, and rifapentine, respectively. As each compound eluted from the column, the \((M+H)^{+}\) ion for the labeled analyte. User trace is UV wavelength 480 nm.

**Discussion**

The results of the rifapentine mass balance studies for mouse, rat, and monkey were strikingly similar. During the 120-hr collection period, less than 5% of radioactive drug was excreted in the mouse, rat, and monkey urine, indicating that renal excretion is a minor route of elimination of rifapentine in these species. The major route of elimination of radioactivity was into the feces, where more than 75% of the radioactive dose was recovered. In contrast to rifapentine, mass balance studies of \(^{14}\)C-rifabutin in rats and monkeys revealed that the amounts of radioactivity excreted in urine and feces after oral and iv doses were similar (about 44% of the dose each) (Battaglia et al., 1990, 1991).

Virtually no differences were found between the percentages of dose excreted in urine and feces after oral and iv administration of rifapentine in mouse, uncannulated rats, and monkeys. The relative amounts of radioactivity excreted into bile, feces, and urine after oral administration of rifapentine in bile duct–cannulated rats were also similar to that after iv administration. The ratio of radioactivity excreted in rat bile to feces was approximately 3:2 after both oral and iv.
iv dosing. These data support other reports that absorption of rifapentine is nearly complete (Weber et al., 1983).

Biliary excretion was a major route of elimination of radioactivity (approximately 51% of excreted radioactivity) after iv administration of rifapentine in bile–cannulated rats. Approximately 40% of the excreted radioactivity (about 23% of the dose) was recovered in feces after iv administration in bile duct–cannulated rats. This finding suggests that rifapentine may be cleared by another route of elimination, such as direct excretion through the intestinal wall. The role of gastrointestinal secretion in the clearance of rifapentine in rats is currently under investigation. The ratio of rifapentine to 25-desacetyl-rifapentine was approximately 10:1 in fecal samples of the cannulated rats, but for bile, the ratio was approximately 1:3. Since previous studies found no detectable amount of 25-desacetyl-rifapentine in circulating plasma (Weber et al., 1983), it appears that the 25-desacetyl metabolite is rapidly excreted into bile after being formed in the liver.

The total radioactivity recovered in urine and feces over 96 hr postdose in orally dosed uncanaluated rats (about 104.6%) was much greater than the recovery in urine, bile, and feces found in orally dosed bile duct–cannulated rats (about 41.1%). The balance of the radioactivity was primarily found in the carcass (table 1). This difference was most likely due to changes in the physiology of the rats, stemming from the stress of cannulation and/or surgical trauma. Further investigation of dose recovery after bile duct cannulation in rats is planned.

In general, examination of urine samples using LC/UV/14C was not done in any of the species because the total and individual sample recoveries of radioactivity in urine was less than 5% and 0.7% of dose, respectively. Only the urine sample with the highest level of radioactivity in each species was examined using LC/14C. The radiochromatogram contained small peaks corresponding to rifapentine and its hydrolysis product, 3-formyl-rifapentine. The radiochromatograms from rat fecal and bile samples after oral and iv administration of 14C-rifapentine were similar. By comparing the HPLC retention times of existing standards and/or mass spectral analysis of chromatographic peaks, components identified in the fecal and bile samples included rifapentine, 25-desacetyl-rifapentine, 3-formyl-rifapentine, and 3-formyl-25-desacetyl-rifapentine. The primary peak observed in fecal samples was rifapentine in all three species. Rifapentine is known to be unstable in solution and one major degradation product is 3-formyl-rifapentine. Similar to rifampin, the 3-formyl derivatives found in fecal samples are probably formed by nonenzymatic hydrolysis (Battaglia et al., 1990). Since these metabolites have not been observed in plasma, formation most likely occurs as the excreta resides in the gut.

In all three species, one region of radioactivity occurred as a broad rise in the background with few distinct peaks. The approximate retention time of this area was 9−17 min in mice (accounting for 8%−18% of dose), 7−17 min in rats (accounting for 1%−19% of dose), and 6−15 min in monkeys (accounting for approximately 12%−19% of dose). Attempts to resolve the broad peaks into individual components using other columns and mobile phase conditions were unsuccessful; therefore, structures could not be assigned. No unique mass spectra (from predose fecal extracts) were observed for any of the radioactive compounds eluting from the LC column other than rifapentine, 25-desacetyl-rifapentine, 3-formyl-rifapentine, or 3-formyl-25-desacetyl-rifapentine, presumably because of low concentrations and/or because the ionization efficiency was insufficient to yield discernible mass spectra.

A search of the published literature failed to identify any studies that evaluated the metabolism and excretion of rifampin in animals. The mass balance and metabolic profile of another rifamycin analog, rifabutin, has been reported in animals and humans (Battaglia et al., 1990, 1991; Utkin et al., 1997; Koudriakova et al., 1996). Parent drug accounted for only 8.5% of the radioactive rifabutin dose in urine of rats, and less than 0.5% of the dose in rabbits and monkeys (Battaglia et al., 1990). Most of the urinary radioactivity (more than 93%) after a single oral dose of 14C-rifabutin in rats, rabbits, and monkeys was constituted by unidentified peaks of polar compounds (Battaglia et al., 1990). The most abundant polar metabolite identified was N-isobutyl-4-hydroxy-piperidine. Lipophilic metabolites, which accounted for less than 20% of urinary radioactivity in rats and humans, included 25-O-desacetyl-rifabutin, 27-O-demethyl-rifabutin, and 31-hydroxy-rifabutin (Battaglia et al., 1990; Utkin et al., 1997; Koudriakova et al., 1996). Only small amounts of the 25-desacetyl derivative were found in the urine of rats (2.1% of the dose) (Battaglia et al., 1990). In bile duct–cannulated rats, approximately 24% of the rifabutin dose was excreted in bile, almost exclusively (98%) as metabolites (Koudriakova et al., 1996). These data suggest rifabutin is more extensively metabolized in animals than rifapentine is, with metabolic pathways that include oxidation.

In summary, the disposition and biotransformation of rifapentine in mouse, rat, and monkey are quite similar. Rifapentine is excreted primarily as intact drug in feces; less than 5% of rifapentine and its metabolites are excreted in the urine. The primary metabolite in bile and feces is 25-desacetyl-rifapentine, with smaller amounts of the degradation byproducts, 3-formyl-rifapentine and 3-formyl-25-desacetyl-rifapentine, formed in the gut.

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References

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