METABOLISM OF THE PSYCHOTOMIMETIC TRYP TAMINE DERIVATIVE 5-METHOXY-N,N-DIISOPROPYLTRYPTAMINE IN HUMANS: IDENTIFICATION AND QUANTIFICATION OF ITS URINARY METABOLITES

Tooru Kamata, Munehiro Katagi, Hiroe T. Kamata, Akihiro Miki, Noriaki Shima, Kei Zaitsu, Mayumi Nishikawa, Einosuke Tanaka, Katsuya Honda, and Hitoshi Tsuchihashi

Forensic Science Laboratory, Osaka Prefectural Police Headquarters, Osaka, Japan (T.K., M.K., H.T.K., A.M., N.S., K.Z., M.N., H.T.); and Department of Legal Medicine, University of Tsukuba, Tsukuba, Japan (E.T., K.H.)

Received June 15, 2005; accepted November 7, 2005

ABSTRACT:

The urinary metabolites of 5-methoxy-N,N-diisopropyltryptamine (5-MeO-DIPT) in humans have been investigated by analyzing urine specimens from its users. For the unequivocal identification and accurate quantification of its major metabolites, careful analyses were conducted by gas chromatography/mass spectrometry, liquid chromatography/mass spectrometry, and liquid chromatography/tandem mass spectrometry, using authentic standards of each metabolite synthesized. Three major metabolic pathways were revealed as follows: 1) side chain degradation by O-demethylation to form 5-hydroxy-N,N-diisopropyltryptamine (5-OH-DIPT), which would be partly conjugated to its sulfate and glucuronide; 2) direct hydroxylation on position 6 of the aromatic ring of 5-MeO-DIPT, and/or methylation of the hydroxyl group on position 5 after hydroxylation on position 6 of the aromatic ring of 5-OH-DIPT, to produce 6-hydroxy-5-methoxy-N,N-diisopropyltryptamine (6-OH-5-MeO-DIPT), followed by conjugation to its sulfate and glucuronide; and 3) side chain degradation by N-deisopropylation, to the corresponding secondary amine 5-methoxy-N-isopropyltryptamine (5-MeO-NIPT). Of these metabolites, which retain structural characteristics of the parent drug, 5-OH-DIPT and 6-OH-5-MeO-DIPT were found to be more abundant than 5-MeO-NIPT. Although the parent drug 5-MeO-DIPT was detectable even 35 h after dosing, no trace of its N-oxide was detected in any of the specimens examined.

5-Methoxy-N,N-diisopropyltryptamine (5-MeO-DIPT) is a tryptamine derivative that possesses, in its chemical structure, the critical features necessary for hallucinogenic/stimulant activity; substitutions on the indole ring and the ethylamine side chain of tryptamine result in a pharmacologically active substance (Fig. 1). Shulgin and Carter (1980) synthesized 5-MeO-DIPT and reported its hallucinogenic activity when administered orally. It produces pharmacological effects similar to those of other Schedule I hallucinogens such as dimethyltryptamine (DMT) (Shulgin and Carter, 1980). It is often used orally at a dosage ranging from 6 to 20 mg, and is sometimes taken through the routes of smoking and snorting. Oral administration of 5-MeO-DIPT produces subjective effects with an onset at about 20 to 30 min, a peak at about 1 to 1.5 h, and a duration of about 3 to 6 h (DEA, 2003). Also, 5-MeO-DIPT causes mydriasis, and high dosages produce nausea, jaw clenching, muscle tension, and overt hallucinations with both auditory and visual distortions.

The abuse problem of 5-MeO-DIPT first emerged in 2001 in the United States (DEA, 2001), and then in the same year in Japan (Katagi et al., 2002), and, since then, in various other countries. 5-MeO-DIPT has often been sold under the name “Foxy” or “Foxy Methoxy,” in tablet and capsule forms, which is often used in raves (lively parties involving dancing and drinking). Because this drug is easily available through the Internet, its use has been extensively and rapidly spreading, especially among teenagers and young adults. Because of its escalating popularity and potent physiological effects, increasing numbers of acute poisoning cases (e.g., its overdose) have been reported. Thus, 5-MeO-DIPT was amended into Schedule I of the Controlled Substances Act in 2003 in the United States (DEA, 2003) and was banned in Japan in April 2005.

Although research on the metabolism of 5-methoxy-N,N-dimethyltryptamine (5-MeO-DMT) in mammals (Sitaram et al., 1987) provides some useful information for analyzing 5-MeO-DIPT metabolites in human body fluids, only a few data have been reported for its metabolism in humans. Recently, “tentative identification” of several metabolites in humans, but without using authentic standards, was reported (Meatherall and Sharma, 2003; Wilson et al., 2005). In this report we have aimed for the careful investigation and indisputable identification of the metabolites of 5-MeO-DIPT in humans. The authors for the first time synthesized the authentic stan-
dards of its metabolites that were predicted based on previous studies (Meatherall and Sharma, 2003; Wilson et al., 2005; Tsutsumi et al., 2005a,b). Utilizing the authentic standards, several urine specimens from 5-MeO-DIPT users were analyzed by gas chromatography/mass spectrometry (GC/MS), liquid chromatography/mass spectrometry (LC/MS), and liquid chromatography-tandem mass spectrometry (LC-MS/MS). The excretion profiles are presented, and the metabolic pathways of 5-MeO-DIPT are discussed.

Materials and Methods

Materials. 5-MeO-DIPT hydrochloride, 5-methoxy-N-isopropyltryptamine (5-MeO-NIPT) hydrochloride, 5-hydroxy-N,N-diisopropyltryptamine (5-OH-5-MeO-DIPT) hydrochloride, 6-hydroxy-5-methoxy-N,N-diisopropyltryptamine (6-OH-5-MeO-DIPT), and 5-methoxy-N,N-diisopropyltryptamine-N-oxide (5-MeO-DIPT-NO) were synthesized in our laboratory according to previously published methods with partial modification, as detailed below (their chemical structures can be seen in Fig. 5). Every synthesized compound was ensured to be >95% pure based on high-resolution MS analysis by the flow-injection method. Stock standard solutions of these five compounds were prepared in methanol (1 mg/ml each) and then diluted to appropriate concentrations with distilled water or control human urine, immediately before use. The internal standard (I.S.) 5-methyletryptamine (5-MT) hydrochloride was purchased from Aldrich Chemical Co. (Milwaukee, WI), and an I.S. solution (200 ng/ml) was prepared in distilled water. Acetonitrile and methanol were of high-performance liquid chromatography grade, and other chemicals used were of analytical grade. Sulfatase/β-glucuronidase (Helix pomatia, type H-1) was obtained from Sigma-Aldrich (St. Louis, MO), N-Methyl-N-trimethylsilyltri-fluoroacetamide (MSTFA) used for trimethylsilyl (TMS) derivatization was purchased from Wako Pure Chemicals (Osaka, Japan).

Chemical Synthesis. 5-MeO-DIPT (I). I was synthesized from 5-methoxytryptamine (Aldrich Chemical Co.) and purified, according to the method of Shulgin and Shulgin (1997a).

5-MeO-NIPT (II). II was synthesized according to the above-mentioned procedure, with a slight modification: instead of excess 2-isopropanol, an equivalent amount of 2-isopropanol was added to 5-methoxytryptamine, and the resultant reaction mixture was purified on a column packed with Chromatorex NH, using chloroform as an eluent (Fuji Silysia Chemical, Ltd., Kasugai, Japan).

5-OH-5-MeO-DIPT (III). III was synthesized in analogy to the modified procedure for 5-hydroxy-N,N-dimethyltryptamine (Shulgin and Shulgin, 1997b): diisopropylamine was used instead of dimethylamine.

6-OH-5-MeO-DIPT (IV). IV was synthesized according to the procedures of Shulgin and Shulgin (1997c,d), with modifications as follows. To concentrated nitric acid, which was stirred and cooled in an external ice-bath, finely powdered 4-benzyloxy-3-methoxybenzaldehyde (Avocado Research Chemicals, Ltd., Heysham, UK) was added. (The temperature must not be allowed to rise above 0°C). After 2 h of additional stirring, the reaction mixture was poured over chipped ice, and the product was extracted with ethyl acetate. The extract was washed with a saturated aqueous sodium bicarbonate solution and water to remove remaining acid. After evaporation, the residue was subjected to column chromatography, using a silica gel column and an ethyl acetate/n-hexane mixture (1:1 v/v) as a developing solvent, to obtain 2-nitro-4-benzyloxy-3-methoxybenzaldehyde. (The yield in this step was 70%).

A solution of 2-nitro-4-benzyloxy-3-methoxybenzaldehyde in glacial acetic acid was treated with nitromethane (Wako Pure Chemicals), followed with anhydrous ammonium acetate. After being held at reflux for 2 h, the reaction mixture was poured into ice water. The product was extracted with ethyl acetate, and the extract was evaporated to dryness after washing with water. The residue was subjected to column chromatography, using a silica gel column and an ethyl acetate/n-hexane mixture (1:2 v/v) as a developing solvent, to isolate 2,2′-dinitro-4-benzyloxy-3-methoxyxylene (78%).

To glacial acetic acid, 2,2′-dinitro-4-benzyloxy-3-methoxyxylene and electrolytic powdered iron were added, stirred, and heated gently until an exothermic reaction set in, and this was maintained at a controlled pace with external cooling. When the spontaneous reaction had subsided, the reaction mixture was refluxed for 15 min, cooled, neutralized with sodium hydroxide, and extracted with ethyl acetate. The extract was evaporated. The residue was subjected to column chromatography, using a Chromatorex NH column and an ethyl acetate/n-hexane mixture (1:3, v/v) as a developing solvent, to obtain 6-benzyloxy-5-methoxyindole (30%).

To a well stirred, cold solution of 6-benzyloxy-5-methoxyindole in anhydrous diethyl ether, a solution of oxalyl chloride (Wako Pure Chemicals) in diethyl ether was added dropwise with protection from atmospheric moisture. The reaction mixture was stirred for an additional 20 min, and the intermediate indoleglyoxylic acid was separated as a crystalline solid but was not isolated. This was treated with 40% solution of diisopropyl amine in anhydrous diethyl ether, dropwise, until the pH reached 8 to 9. The reaction mixture was diluted with chloroform and shaken with 5% aqueous potassium bisulfate (Wako Pure Chemicals) solution, followed by a saturated aqueous sodium bicarbonate solution. After drying, the organic solvent was removed. The residue was subjected to column chromatography, using a Chromatorex NH column and an ethyl acetate/n-hexane mixture (1:5 v/v) as a developing solvent, to isolate 6-benzyloxy-5-methoxyindol-3-yl- N,N-diisopropylglyoxylamide (55% in these two steps).

To a well stirred suspension of lithium aluminum hydride in dry tetrahydrofuran (THF), a solution of 6-benzyloxy-5-methoxyindol-3-yl-N,N-diisopropylglyoxylamide in anhydrous THF was added dropwise. The mixture was brought to a reflux temperature, held there for 15 min, and allowed to return to room temperature. The excess hydride was destroyed by adding ethyl acetate and water. The solids were removed by filtration, the filter cake was washed with THF, and the pooled filtrate and washings were evaporated under vacuum. The residue was subjected to column chromatography, using a Chromatorex NH column and an ethyl acetate/n-hexane mixture (1:5, v/v) as a developing solvent, to obtain 6-benzyloxy-5-methoxy-N,N-diisopropyltryptamine (35%).

A solution of 6-benzyloxy-5-methoxy-N,N-diisopropyltryptamine in methanol containing 10% palladium-activated carbon catalyst was shaken under 3 atm of hydrogen for 6 h, and the solids were removed by filtration. Evaporation of the solvent under vacuum gave a residue. The residue was subjected to column chromatography using a Chromatorex NH column and an ethyl acetate/n-hexane mixture (1:1 v/v) as a developing solvent, to isolate 6-hydroxy-5-methoxy-N,N-diisopropyltryptamine (88%). The overall isolated yield was 28%.

5-MeO-DIPT-NO (V). V was synthesized from I, according to the procedure of Cymerman Craig and Purushothaman (1970).

The identities of all synthesized compounds were confirmed by high-resolution MS and MS/MS in the flow-injection method, in addition to GC/ electrospray ionization (EI) MS with and without TMS derivatization and LC/ESI MS. The GC/EI MS and LC/ESI MS data are described under Results. The high-resolution mass spectral data (calculated exact mass is given in parentheses) are as follows: I: m/z 275.2102 [M + H]+ (275.2133), 174.0896 [M – (C3H7)2N]+ (174.0919), 114.1314 (C3H7)2NCH2]+ (114.1283). II: m/z 233.1659 [M + H]+ (233.1654), 190.0854 [M – (C3H7)NH]+ (190.0919). III: m/z 261.1967 [M + H]+ (261.1967), 160.0756 [M – (C3H7)2]+ (160.0762), 113.1312 (C3H7)2NCH2]+ (113.1283). IV: m/z 291.2079 [M + H]+ (291.2072), 190.0855 [M – (C3H7)2N]+ (190.0852). V: m/z 291.2099 [M + H]+ (291.2072), 174.0897 [M – (C3H7)2NO]+ (174.0919).

Apparatus. GC/MS was carried out on a GCMS QP-2010 (Shimadzu, Kyoto, Japan). A fused-silica capillary column, DB-5MS (30 m × 0.25 mm i.d.; 0.25 μm; J&W Scientific, Rancho Cordova, CA) was used for separation. Injections were effected automatically in the splitless mode at 260°C. The column oven temperature was maintained at 80°C for 2 min and then raised at 20°C/min to 310°C. The transfer line temperature was set at 260°C. High-purity helium, at a flow rate of 3 ml/min, was used as the carrier gas. The EI
After adding 60 M ascorbic acid, and the specimen was adjusted to pH 5 with 10% acetic acid. At 40°C. The residue was dissolved in 100 ml of 0.5 M acetate buffer (pH 5.0) and after hydrolysis were adjusted to pH 8 with 2.8% ammonium hydroxide.

Extraction for LC/MS and LC-MS/MS of Conjugates. Urine specimens before and after hydrolysis were each mixed vigorously with 2 volumes of methanol. After centrifugation, the supernatant was recovered and evaporated to dryness. The residue was dissolved in 100 µl of distilled water and filtered. Aliquots of 5 µl were automatically injected into the LC/MS and LC-MS/MS systems.

Validation of the LC/MS Procedure. To quantify 5-MeO-DIPT and its metabolites in urine, the LC/MS procedure optimized was validated. A 300-µl drug-free urine spiked with the synthesized standards at 1 µg/ml each was processed as described above. The residue was dissolved in 100 µl of distilled water and added to 100 µl of 200 ng/ml I.S. (5-MT) solution. A 5-µl aliquot was injected into the LC/MS system in the selected ion monitoring mode, where the protonated molecules of each analyte were selected as the monitoring ions, and the peak area ratios to I.S. were calculated. The recoveries at 1 µg/ml were 93.1%, 92.3%, 95.7%, and 105% for 5-MeO-DIPT, 5-MeO-NP, 5-OH-5-MeO-DIPT, and 6-OH-5-MeO-DIPT, respectively (n = 5). The detection limits were 0.03 µg/ml for 6-OH-5-MeO-DIPT and 0.003 µg/ml for the others. Calibration curves constructed by the I.S. method showed good linearity over the ranges from 0.1 to 10 µg/ml for 6-OH-5-MeO-DIPT and from 0.01 to 10 µg/ml for the others. The within-day relative standard deviations (evaluated at 1 µg/ml, n = 5) ranged from 3.35 to 5.17% for all of the analytes. These results guaranteed the reliability of the present procedure for the analysis of urine specimens from 5-MeO-DIPT users.

Results

GC/MS and LC/MS of Predicted Metabolite Standards. The EI mass spectra of the authentic standards, except for 5-MeO-DIPT-NO, indicated their characteristic structures: predominant ions due to the α-cleavage of amine moieties (m/z 72 for 5-MeO-NP; m/z 114 for 5-MeO-DIPT, 5-OH-5-MeO-DIPT, and 6-OH-5-MeO-DIPT) and some other ions, including a very small molecular ion (m/z 274 for 5-MeO-DIPT, m/z 232 for 5-MeO-NP, m/z 260 for 5-OH-DIPT, and m/z 290 for 6-OH-5-MeO-DIPT) (data not shown). 5-MeO-DIPT-NO was not detectable by GC/MS because N-oxide metabolites readily degrade in the injection port of the gas chromatograph.

GC/MS was also carried out after TMS derivatization because of its higher sensitivity and clearer identification of amines and phenols. The EI mass spectra of their TMS derivatives also had a molecular ion (m/z 346 for 5-MeO-DIPT-TMS, m/z 304 for 5-MeO-NP-TMS, m/z 404 for 5-OH-DIPT-di-TMS, and m/z 434 for 6-OH-5-MeO-DIPT-di-TMS), but their relative intensities were still very low, as in the case without derivatization (data not shown).

Unlike GC/MS spectra, the ESI mass spectra of the authentic standards taken by LC/MS (data not shown) were characterized by the predominant protonated molecules at m/z 275 for 5-MeO-DIPT, m/z 233 for 5-MeO-NP, m/z 261 for 5-OH-DIPT, m/z 291 for 6-OH-5-MeO-DIPT, and 5-MeO-DIPT-NO. It should be noted that the ESI mass spectrum of 5-MeO-DIPT-NO had a weak dimerization ion [2M + H]+, which often appears specifically for amine N-oxides, at m/z 581.

Identification of the Metabolites in Urine. The extracts of the unhydrolyzed urine specimens were first analyzed by GC/MS with and without TMS derivatization, and the retention times and mass spectra of compounds detected were compared with those of the standards. As a result, free-form 5-MeO-DIPT, 5-MeO-NP, 5-OH-DIPT, and 6-OH-5-MeO-DIPT were detected in the extracts without derivatization, and TMS derivatives of these four compounds were also clearly detected in the derivatized extracts.

LC/ESI MS was next carried out under the optimized conditions, and the results were compared with those of the authentic standards. Figure 2 shows the extracted ion chromatograms obtained from User A’s urine (11 h post-intake). Although 5-MeO-DIPT, 5-MeO-NP,
5-MeO-DIPT, and 6-OH-5-MeO-DIPT were confirmed, no trace of 5-MeO-DIPT-NO was detected in this specimen.

To examine the stability of the predicted N-oxide metabolite, six control urine samples from drug-free volunteers were spiked with 5-MeO-DIPT-NO at 1 μg/ml, and the samples were stored at −20°C for 1 month. No noticeable decrease of 5-MeO-DIPT-NO was detected in any of the samples tested. Thus, the absence of 5-MeO-DIPT-NO was not attributed to its denaturation or decomposition.

**Excretion of 5-MeO-DIPT and Its Metabolites.** The concentrations of 5-MeO-DIPT and its three metabolites identified in the urine specimens were quantified by the validated LC/MS procedure, using the calibration curves constructed, and the excretion profiles were investigated. Also, the concentrations before and after enzymatic hydrolysis were compared. The results are summarized in Table 1. Although 5-MeO-DIPT, 5-MeO-NIPT, 5-OH-DIPT, and 6-OH-5-MeO-DIPT were detected in all of the samples, no trace of 5-MeO-DIPT-NO was detected in any of the specimens examined.

Because excretion of the glucuronides and sulfates was expected for 5-OH-DIPT and 6-OH-5-MeO-DIPT from their increased levels after hydrolysis, their direct detection was attempted by LC-MS/MS. Thus, we concluded that the detection of "N-oxide" (it should be named "hydroxylamine," rather than "N-oxide") in their reports required reexamination by LC/MS, based on our previous studies that dealt with N-oxide metabolites of several other drugs (Katagi et al., 2000, 2001).

Previous studies on the rat reported metabolic routes of DMT and 5-MeO-DMT, which include N-oxidation, N-demethylation, O-demethylation, and oxidative deamination (Sitaram et al., 1987). Based on that previous report, 5-MeO-NIPT, 5-OH-DIPT, and 5-MeO-DIPT-NO were predicted as the major urinary metabolites of 5-MeO-DIPT. In addition, Morano et al. (1993) proposed that etryptamine (ethyltryptamine) is metabolized mainly by 6-hydroxylation, like other indole derivatives, which suggested the possibility of hydroxylation at the 6-position of 5-MeO-DIPT into 6-OH-5-MeO-DIPT-oxide. Thus, the present authors selected 6-OH-5-MeO-DIPT, in addition to the above-mentioned three metabolites, as the expected major metabolites that retain the structural characteristics of the parent drug 5-MeO-DIPT. Based on the GC/MS and LC/MS analyses in this study, 5-MeO-DIPT, 5-MeO-NIPT, 5-OH-DIPT, and 6-OH-5-MeO-DIPT were indisputably identified in human urine from 5-MeO-DIPT users.

In this study, no trace of 5-MeO-DIPT-NO was detected in any of the urine specimens from 5-MeO-DIPT users. In the previous study by
Sitaram et al. (1987) on the metabolism of 5-MeO-DMT, an analog of 5-MeO-DIPT, 5-MeO-DMT-N-oxide was identified as the abundant and characteristic urinary metabolite in the rat. This difference is probably attributed to steric interference with N-oxidation on the tertiary nitrogen atom by the two bulky isopropyl moieties, although the possibility of interspecies variations should not be excluded. Also, no noticeable decrease of 5-MeO-DIPT-NO during sample storage was detected, as mentioned above. Thus, there is probably no metabolic transformation of 5-MeO-DIPT into 5-MeO-DIPT-NO in humans.

Our studies confirm that 5-MeO-NIPT and 5-OH-DIPT are metabolites found in human urine as predicted by Meatherall and Sharma (2003) and Wilson et al. (2005). However, 6-OH-5-MeO-DIPT was also identified, rather than 5-MeO-DIPT-N-oxide, as the GC/MS product close to the 5-OH-DIPT peak. Because 5-MeO-DIPT-N-oxide and 6-OH-5-MeO-DIPT have the same molecular weight, we concluded that the metabolite that Meatherall’s team expected to be 5-MeO-DIPT-N-oxide was probably the 6-OH-5-MeO-DIPT identified here.

Excretion of 5-MeO-DIPT and Its Metabolites. As summarized in Table 1, enzymatic hydrolysis increased 6-OH-5-MeO-DIPT concentration to almost the same level as that of 5-OH-DIPT, although no notable increase was observed for 5-OH-DIPT. These facts indicate that 5-OH-DIPT and 6-OH-5-MeO-DIPT are abundant and characteristic metabolites and that N-deisopropylation is a minor metabolic pathway for 5-MeO-DIPT.

Based on the quantitative results listed in Table 1, it should be noted that the conjugation rate of 5-OH-DIPT seemed to be very low, although most of the 6-OH-5-MeO-DIPT was excreted as its conjugates. For determining exact rates of sulfates and glucuronides, anal-
ysis should be done using authentic standards of conjugates, or by performing selective hydrolysis using sulfatase and β-glucuronidase inhibitors. However, in the direct analysis of the conjugates of hydroxylated metabolites of phenethylamine-type drugs by LC/ESI MS and LC/ESI MS/MS, we found that the sensitivities in detecting glucuronides were 1.2 to 2.3 times higher than those of sulfates in the LC/ESI MS of p-hydroxymethamphetamine and 4-hydroxy-3-methoxymethamphetamine, the major metabolites of methamphetamine and 3,4-methylenedioxymethamphetamine. Based on these facts, the present data in Fig. 3 suggest that 5-OH-DIPT is excreted in urine mainly in the unconjugated form, whereas 6-OH-5-MeO-DIPT is excreted mostly as its sulfate.

These results revealed three metabolic pathways of considerable quantitative significance for 5-MeO-DIPT in humans. As shown in Fig. 5, the first pathway leads, via side chain degradation by O-demethylation, to 5-OH-DIPT, partly followed by conjugation to form its sulfate and glucuronide. The second pathway leads, via direct hydroxylation on position 6 of the aromatic ring of 5-MeO-DIPT, and/or methylation of the hydroxyl group on position 5, after hydroxylation on position 6 of 5-OH-DIPT, to produce 6-OH-5-MeO-DIPT, followed by conjugation to its sulfate and glucuronide. Although the hydroxylated 5-OH-DIPT (Fig. 5) is also expected to be methylated to form 5-hydroxy-6-methoxy-diisopropyltryptamine, the positional isomer of 6-OH-5-MeO-DIPT and 5-hydroxy-6-methoxy-diisopropyltryptamine should be distinguished by their retention times on GC/MS and LC/MS according to our previous studies. The positional isomers of N-benzylpiperazine metabolites (Tsutsumi et al., 2005b) and methylendioxymethamphetamine (Ecstasy) metabolites (unpublished data) had been well separated by GC/MS and LC/MS, and the urinary hydroxy-methoxy-diisopropyltryptamine detected in this study completely agreed with 6-OH-5-MeO-DIPT. We, therefore, concluded that the hydroxylated metabolite is 6-OH-5-MeO-DIPT. The third pathway leads, via side chain degradation by N-deisopropylation, to the corresponding secondary amine, 5-MeO-NIPT. Of these, 5-OH-DIPT and 6-OH-5-MeO-DIPT were of the major metabolites. However, further investigation of hydrolysis conditions is necessary to exactly evaluate the contributions of demethylation and hydroxylation to 5-MeO-DIPT metabolism, because the hydrolysis of 5-OH-DIPT-sulfate and 6-OH-5-MeO-DIPT-sulfate could not be completed under the current conditions (Fig. 3). Also, a more quantitative experiment using radiolabeled 5-MeO-DIPT in an animal model might be very useful for more detailed survey of the metabolism of 5-MeO-DIPT. It is well known that oxidative N-deamination is a major metabolic pathway for 5-MeO-DMT in the rat (Sitaram et al., 1987) and, thus, it is expected to be an important metabolic pathway for 5-MeO-DIPT in humans. However, possible metabolites like 5-methoxyindoleacetic acid and 5-hydroxyindoleacetic acid, formed by the oxidative deamination, retain few structural characteristics of the parent compound. In fact, 5-hydroxyindoleacetic acid, which is commercially available, is known to be present at low μg/ml levels in the urine of healthy humans. Thus, the identification of characteristic urinary metabolites of 5-MeO-DIPT reported here will be of importance in forensic and clinical urine analysis. In addition, the present study will provide useful information in the urine analysis for unknown tryptamine-type drugs that may be encountered in the future.

References


Address correspondence to: Tooru Kamata, Forensic Science Laboratory, Osaka Prefectural Police Headquarters, 1-3-18, Hommachi, Chuo-ku, Osaka 541-0053, Japan. E-mail: t-kamata@mahoroba.ne.jp