METABOLISM OF SELEGILINE IN HUMANS
Identification, Excretion, and Stereochemistry of Urine Metabolites

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ABSTRACT:

Nine urinary metabolites of selegiline hydrochloride [N-methyl-N-propargyl[2-phenyl-1-methyl]ethylammonium chloride], a monoamine oxidase inhibitor, after administration to humans were identified. Their identities were confirmed by comparison of the spectra from GC/MS of peaks with those of authentic compounds. The following metabolites and unchanged drug (selegiline) were detected in urine: (R)-desmethylselegiline, (R)-methamphetamine, (R)-amphetamine, (1S,2R)-norephedrine, (1R,2R)-norpseudoephedrine, (1S,2R)-ephedrine, (1R,2R)-pseudoephedrine, (R)-p-hydroxyamphetamine, and (R)-p-hydroxymethamphetamine. The metabolites excreted 2 days after administration of 2.5–10 mg of selegiline hydrochloride amounted to 44–58% of the dose. Selegiline was metabolized by three distinct pathways: N-dealkylation, β-carbon hydroxylation, and ring-hydroxylation. The major metabolite was (R)-methamphetamine. During metabolism, no racemic transformation occurred and β-carbon hydroxylation showed apparently product stereoselectivity.

Selegiline is a selective, irreversible inhibitor of monoamine oxidase that has been used in combination with L-dopamine in the treatment of parkinsonism (1, 2).

There have been several reports of studies of selegiline metabolism either in vivo or in vitro. Reynolds et al. (3) showed that methamphetamine and amphetamine were excreted in human urine after oral administration of the drug. Later, Heinonen et al. (4) identified the three metabolites (methamphetamine, amphetamine, and desmethyl-selegiline) in human urine. Also, p-hydroxyamphetamine and p-hydroxymethamphetamine have been identified as additional urinary metabolites in the rat (5–7). However, no p-hydroxylated metabolite in humans has been identified, and neither unchanged drug nor β-hydroxylated metabolite was found in urine, after administration of selegiline to humans or rats.

Selegiline has a chiral center on the α-carbon atom of the phenylethylamine moiety. There are no detailed studies on the stereoselective metabolism of selegiline in the literature (8, 9).

Ten metabolites, together with unchanged drug, were identified and quantified from human urine after administration of selegiline. Herein we report on the stereochemical aspects of selegiline metabolism.

Materials and Methods

Chemicals. (R)-Amphetamine sulfate, (S)-amphetamine sulfate, (R)-methamphetamine, and (1R,2R)-norpseudoephedrine · HCl were purchased from Sigma (Deisenhofen, Germany). (1R,2S)-Norephedrine · HCl, (1S,2R)-norephedrine · HCl, (1R,2S)-ephrine · HCl, (1S,2R)-ephrine · HCl, and (1S,2R)-pseudoephedrine · HCl were purchased from Fluka (Neu-Ulm, Germany). Pure standards of p-hydroxyamphetamine, p-hydroxymethamphetamine, p-hydroxynorephedrine, and p-chlorphentermine (internal standard) were provided by Tropon (Köln, Germany). Arylsulfatase/β-glucuronidase was provided by Serva (Heidelberg, Germany). Reagents were purchased from various sources: the chiral derivatizing reagent MTPA · Cl was from JPS Chimie (Bevaiz, Swiss); and diethylether, methanol, acetonitrile, TFA, sodium bicarbonate, and potassium carbonate were from Merck (Darmstadt, Germany). MSTFA, MTESTFA, and MBTFA were purchased from Sigma.

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Fig. 1. GC-NPD chromatogram of the metabolites in urine 2–4 hr after administration of selegiline.

Peaks: A1, amphetamine; A2, methamphetamine; A3, desmethylselegiline; A4, selegiline; A5, nor(pseudo)ephedrine; and A6, (pseudo)ephedrine.
Drug Administration and Sample Collection. 10, 5, and 2.5 mg of selegiline HCl (2, 1, and 0.5 tablets of Movergan, respectively; ASTA Pharma AG, Frankfurt, Germany) were orally administered to male volunteers. Subjects were four healthy males, aged 25–34. Urine samples were collected at various times over 72 hr and stored at 4°C.

Isolation of Unconjugated Metabolites. To 3.0 ml of urine, 100 mg of sodium bicarbonate:potassium carbonate (2:1, g/g) and 50 ng of internal standard (p-chlorphentermine) were added. The metabolites were extracted with 8 ml of diethylether-tert-butanol (7:1, v/v). The organic layer was transferred into a 15-ml glass centrifuge tube, with 0.4 ml of 0.06 M hydrochloric acid added. Extraction was performed by mixing for 5 min at 1200 g, and the organic layer was aspirated and discarded. The aqueous layer was dried in a desiccator over phosphorus pentoxide:potassium hydroxide.

Isolation of Conjugated Metabolites. 3.0 ml of urine was adjusted to pH 5.2, with 1 ml of 0.2 M sodium acetate buffer, and incubated with 50 ml of arylsulfatase/b-glucuronidase from Helix pomatia (Serva) at 52°C for 5 hr. After cooling, the solution was neutralized with 5 M KOH and adjusted to pH 9.6 with 200 mg of sodium bicarbonate:potassium carbonate (2:1, g/g). Fifty nanograms of internal standard (p-chlorphentermine) was added. The procedure for the extraction of the hydrolyzed metabolites from urine is identical with that for the extraction of unconjugated metabolites.

Quantification of Selegiline and Its Metabolites (10). The dry residue was dissolved in 50 μl of a mixture of acetonitrile:TFA (60:40, v/v) that contained 200 ppm of methyl orange. The mixture was titrated with MSTFA until the color of the reaction mixture changes from red to yellow. The sample was heated for 10 min at 60°C. Two drops of MBTFA were added to the reaction mixture, and the sample was heated for an additional 10 min at 60°C. After cooling, 2 ml of the solution was injected into the GC/MS system.

Identification of the Stereoisomers (11). The dry residue was dissolved in 50 μl of a mixture of acetonitrile:TFA (60:40, v/v) that contained 200 ppm of methyl orange. The mixture was titrated with MSTFA until the color of the reaction mixture changed from red to yellow. The sample was heated for 10 min at 60°C. Two drops of MBTFA were added to the reaction mixture, and the sample was heated for an additional 10 min at 60°C. After cooling, 2 μl of the solution was injected into the GC/MS system.

GC/MS. All mass spectra were obtained with an HP 5890/5971A instrument, including an HP 9144 disk drive and an HP Think Jet printer. Separation was achieved with an HP fused-silica capillary column with cross-linked 5% phenylmethylsilicone (SE-54) that was ~17 min length, with a 0.2 mm internal diameter, and a 0.33 μm film thickness. The operating parameters of GC/MS were as follows: detector, mass selective detector in scan or SIM mode; ionization, electron impact mode; ionization potential, 70 eV; injector temperature, 280°C; interface temperature, 300°C; initial oven temperature, 100°C; ramp, 15°C/min; final oven temperature, 320°C for 2 min; carrier gas, helium at a flow of 1.0
ml/min; and split ratio, 1:8. The ions m/z 140 for amphetamine-N-TFA, m/z 154 for methamphetamine-N-TFA, m/z 179 for norephedrine-N-TFA-O-TMS, norpseudoephedrine-N-TFA-O-TMS, pseudoephedrine-N-TFA-O-TMS, m/z 206 for p-hydroxymethamphetamine-N-TFA-O-TMS and p-hydroxymethamphetamine-N-TFA-O-TMS, m/z 178 for desmethylselegiline, and m/z 96 for selegiline were selected for GC/MS (SIM) detection. Full-scan mass spectra (m/z 40–550) were recorded for analyte identification.

**Results and Discussion**

**Identification of Metabolites.** To detect the urinary metabolites of selegiline, selegiline was administered to humans. The extracts from human urine were examined by GC-NPD and GC/MS as free form or derivatives. Identification of metabolites was conducted by comparison of the EI mass spectra and the gas chromatographic retention times of the extracted metabolites with those of authentic compounds. In this study, nine urinary metabolites and unchanged drug were identified.

GC-NPD of the extract from human urine without derivatization gave six peaks (A1–A6) (fig. 1), that after N-trifluoroacetylation,

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**Fig. 4.** EI spectra of (B5 and B6) nor(pseudo)ephedrine, N-TFA, O-TMS; (B7 and B8) pseudoelehedrine, N-TFA, O-TMS; (B9) p-hydroxymethamphetamine, N-TFA, O-TMS; and (B10) p-hydroxymethamphetamine, N-TFA, O-TMS.

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**Fig. 5.** Gas chromatographic separation of N-MTPA(+), O-TES derivatives of the authentic standards on capillary column.


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**Fig. 6.** Metabolic pathways of selegiline in humans.
Peaks A4 and B4 showed the same retention time and mass spectra as authentic selegiline. Selegiline does not have any functional group to be derivatized by the described derivatization method, such as the amino, hydroxyl, or carboxylic group. The retention times and mass spectra before and after derivatization would be unchanged. The EI mass spectrum showed the base peak at m/z 96 and principal fragment ions at m/z 56, 91, 65, 97, and 115 (fig. 3).

Otherwise, peak A5 after selective derivatization gave two peaks (B5 and B6), and peak A6 gave two peaks (B7 and B8). Peak B6 had a longer retention time than peak B5, but no significant difference was between the mass spectra of the two peaks. The EI mass spectra of peaks B5 and B6 showed the base peak at m/z 179, and at the same retention times and diagnostic ions as the derivatives of norpseudoephedrine and norephedrine, respectively (fig. 4). Also, peak B8 had a longer retention time than peak B7, but no significant difference was between the mass spectra of the two peaks. In EI mass spectra, peaks B7 and B8 gave the base peak at m/z 179, and at the same retention times and diagnostic ions as the derivatives of ephedrine and pseudoephedrine, respectively (fig. 4).

Peak B9 gave EI mass spectrum with a molecular ion at m/z 319, and principal fragment ions at m/z 140, 179, and 206. The analytical data of B9 agreed with those of p-hydroxymethamphetamine (fig. 4).

The EI mass spectrum of B10 showed a molecular ion at m/z 333 and diagnostic ions m/z 154, 179, and 206, and was identical to that of the derivative of p-hydroxymethamphetamine (fig. 4).

**Identification of the Stereoisomers of the Selegiline Metabolites.**

For the identification of the stereoisomers of the selegiline metabolites, the method of Shin and Donike (11)—applicable to the resolution and quantification of trace optical isomers containing more than one reactive functional group—was used. Figure 5 shows the chromatogram of the N-MTPA, O-TES derivatives of the authentic standard. All enantiomers were well separated through conversion to the derivatives, but the derivatives of (1S,2R)-ephedrine and (1S,2S)-pseudoephedrine were overlapped. The stereochemical identities of the metabolites in urinary extracts were confirmed by comparison of the gas chromatographic retention times of the diastereomeric derivatives of the extracted metabolites with those of the authentic standard. It appeared from this work that the metabolites identified—amphetamine, methamphetamine, desmethylselegiline, p-hydroxymethamphetamine, and p-hydroxymethamphetamine—have the (R)-absolute configuration. We also identified the absolute configurations of the four β-hydroxylated metabolites. Their absolute configurations were (1S,2R)-norephedrine, (1R,2R)-norpseudoephedrine, (1S,2R)-ephedrine and (1R,2S)-pseudoephedrine. No peak of (1R,2S)-ephedrine or (1S,2S)-pseudoephedrine, which could not be sep-
FIG. 7. Excretion of selegiline and its metabolites after an oral dose of 10 mg of selegiline · HCl.
ated by this method, was found in the extracted metabolites. These facts indicated that it was during metabolism of selegiline that no racemic transformation at the 2-carbon position occurred.

The main routes of metabolism of selegiline in humans are shown in fig. 6.

**Urinary Excretion of Metabolites.** The urinary excretion of the metabolites after oral administration of selegiline · HCl for the time period of collection (0–48 h) is summarized in table 1. Selegiline was detected as free form. Determination of amphetamine, methamphetamine, and desmethylselegiline was conducted as trifluoroacetate. Determination in urine is therefore too low to be detected by the available analytical method (12, 13).

Identification and detection of unchanged drug and new metabolites in this study may give especially useful new information for the studies of selegiline treatment.

### References