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Synthesis and in Vitro Cytotoxicity Profile of the *R*- Enantiomer of 3,4-Dihydroxymethamphetamine (*R*-(-)- HHMA) : Comparison with Related Catecholamines.

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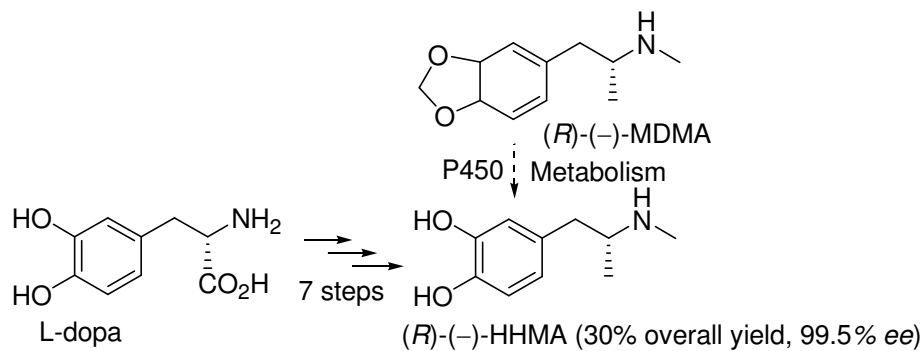
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1
2
3 **Abstract.** (±)-3,4-methylenedioxymethamphetamine (MDMA, also known as “ecstasy”) is a chiral
4 drug that is essentially metabolized in humans through *O*-demethylenation into 3,4-
5 dihydroxymethamphetamine (HHMA). There has recently been a resurgence of interest in the possibility
6 that MDMA metabolites, especially 5-(*N*-acetylcystein-S-yl)-*N*-methyl- α -methyldopamine,
7 (designated as 5-NAC-HHMA), might play a role in MDMA neurotoxicity. However, chirality of
8 MDMA was not considered in previously reported in vivo studies, because HHMA, the precursor of 5-
9 NAC-HHMA metabolite, was used as the racemate. Since the stereochemistry of this chiral drug needs
10 to be considered, the first total synthesis of *R*-(-)-HHMA is reported. Using L-dopa as the chiral source,
11 the preparation of *R*-(-)-HHMA is achieved through seven steps, in 30% overall yield and 99.5%
12 enantiomeric excess. The cytotoxicity of *R*-(-)-HHMA and related catecholamines has been further
13 determined by flow cytometric analysis of propidium iodide uptake in human dopaminergic
14 neuroblastoma SH-SY5Y cells, and by an *Escherichia coli* plate assay, specific for the detection of
15 oxidative toxicity. The good correlation between the toxicities observed in both systems suggests that
16 SH-SY5Y cells are sensitive to oxidative toxicity and that cell death (necrosis) would be mediated by
17 reactive oxygen species mainly generated from redox active quinonoid centers. In contrast, apoptosis
18 was detected for 3,4-dimethoxymethamphetamine (MMMA), the precursor of HHMA possessing a
19 protected catechol group. MMMA was not toxic in the bacterial assay, indicating that its toxicity is not
20 related to increased oxidative stress. Finally, we can conclude there is a need to distinguish the toxicity
21 ascribed to MDMA itself, also bearing a protected catechol moiety, from that depending on MDMA
22 biotransformation leading to catechol metabolites such as HHMA and the thioether conjugates.
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Introduction

The alarming increase in the recreational use of (\pm)-3,4-methylenedioxyamphetamine (MDMA, also known as “ecstasy”) and the multitude of adverse effects resulting from its misuse (1-5) require a complete understanding of the pharmacology and toxicology of this synthetic psychoactive drug. However, the precise mechanisms by which MDMA induces neurotoxic effects, both in laboratory animals (rodents, non-human primates) and humans, remain to be fully elucidated (6-11). MDMA-mediated neurotoxicity could be related to some of the adverse effects such as hyperthermia, the serotonin (5-HT) transporter action, 5-HT_{2A} agonism, glutamate excitotoxicity, the monoamine oxidase metabolism of dopamine and serotonin, nitric oxide together with the formation of damaging peroxynitrite and, importantly, the formation of MDMA neurotoxic metabolites (12). Among these factors, which may act collectively in a synergistic manner conducting to neurotoxicity, systemic metabolism of MDMA seems to be a crucial step (13-15).

In humans, MDMA is essentially metabolized through *O*-demethylation into 3,4-dihydroxymethamphetamine (HHMA) (16). Because of its catechol moiety, HHMA can be easily oxidized to the corresponding *o*-quinone species which exhibits a double reactivity. First, it can undergo redox-cycling which produces semiquinone radical and leads to the generation of reactive oxygen species (ROS). Second, it behaves as a highly electrophilic compound which can be conjugated with thiol nucleophiles such as glutathione or *N*-acetylcysteine (NAC), to form redox active catechol-thioether mono- and bis-conjugates (17) able to induce cytotoxicity mediated by ROS (18).

The potential role of metabolites in MDMA neurotoxicity has been a topic of recent interest (12, 17, 19). Among them, catechol-thioether metabolite 5-(*N*-acetylcystein-S-yl)-*N*-methyl- α -methyldopamine (here designated as 5-NAC-HHMA) has been the most strongly implicated in MDMA neurotoxicity (20, 21). Studies have been recently performed to further assess the 5-HT neurotoxic potential of 5-NAC-HHMA, but the results are at odds with those previously reported and argue against a pivotal role for 5-NAC-HHMA in MDMA-induced 5-HT neurotoxicity (19). The reasons for this

1 discrepancy is not clear, but it is important to note that 5-NAC-HHMA used in these studies was
2 prepared through a biomimetic electrochemical oxidation process (18), whereas the procedure
3 previously utilized for synthesizing this metabolite involved the oxidation of the catechol with
4 mushroom tyrosinase (20), which yields a different ratio of 5-NAC-HHMA diastereoisomers (22).
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MDMA is consumed as a racemate that is a 1:1 mixture of *R*(-)-MDMA and *S*(+)-MDMA enantiomers (Figure 1). Both enantiomers show different pharmacological and pharmacokinetic profiles (24-30). For most of the known psychomimetic agents having at least one chiral center, the most potent isomer exhibits the absolute *R* configuration (24). In the case of MDMA, the potency assignment is reversed since *S*(+)-MDMA has been shown to display effective psychomimetic properties in humans. Furthermore, although *R*(-)-MDMA has the higher affinity for the serotonin receptor 5-HT₂, *S*(+)-MDMA isomer produces the most lasting 5-HT deficits that probably reflects its neurotoxicity (25). Metabolism could also contribute to differences between both enantiomers. Especially, the difference in their pharmacokinetic properties may be caused by enantioselective metabolism by cytochrome P450 isoenzymes, in particular CYP2D6 and CYP2C19 (23). Therefore, HHMA, which is a major MDMA metabolite in humans, exists also as a pair of enantiomers (30) (Figure 1). HHMA being the precursor of the thioether metabolites, these conjugates exist as a mixture of diastereoisomers as they possess several chiral centers: that of HHMA present in the alkylamino side chain (carbon atom α to the amine function), together with those present in the thiol moieties. Because MDMA enantiomers have different biological activity, it is of a great interest to study whether the catechol-thioether diastereoisomers of HHMA also exhibit dissimilar biological properties. The preparation of such diastereoisomers requires the previous synthesis of enantiomerically pure precursors *R*(-)-HHMA and *S*(+)-HHMA, which are more difficult to synthesize than the racemate. To circumvent this problem, analytical and semipreparative methodologies for the diastereoisomeric separation of MDMA thioether conjugates

1 have been recently reported, but this separation method only furnished small quantities of both
2 diastereoisomers (22).
3

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5 This prompted us to envision the total synthesis of enantiomerically pure precursors, *R*-(-)-HHMA
6 and *S*-(+)-HHMA. Starting from these compounds, it would be then easy to prepare the corresponding
7 diastereomerically pure forms of thioether metabolites, either through the straightforward one-pot
8 electrochemical synthesis we have recently reported for preparing diverse catechol-thioether mono- and
9 bis-conjugates starting from (\pm)-HHMA racemate (18, 31), or by using the enzymatic procedure
10 described by other investigators (20, 21, 32).
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14 In this paper, we report the first total synthesis of *R*-(-)-HHMA in 30% overall yield and 99.5%
15 enantiomeric excess (e.e), using L-dopa as the inducer of chirality. To the best of our knowledge, a sole
16 synthesis of enantiomerically enriched *S*-(+)-HHMA (80% e.e.) is known (33). Consequently, as long as
17 the synthesis of optically pure *S*-enantiomer is not reported, the enantioselective toxicity of both
18 enantiomers of HHMA can not be addressed. Nevertheless, we have decided to examine the in vitro
19 toxicity profile of *R*-(-)-HHMA in a broader context, using different assays which could be predictive of
20 the toxicological effects associated with catecholamine-type structures. The cytotoxicity of *R*-(-)-
21 HHMA, and related catecholamines, (Figure 2) has been assessed in human dopaminergic
22 neuroblastoma SH-SY5Y cells. Flow cytometry has been used to discriminate between live, apoptotic
23 and necrotic cells, and to measure the activity of caspase-3, an effector of apoptosis. In addition, the
24 cytotoxicity observed in SH-SY5Y cells has been compared with that evaluated in an *Escherichia coli*
25 plate assay which is specific for the detection of oxidative toxicity (34).
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Experimental Procedures

Chemistry. All reagents and solvents (HPLC grade) were commercial products of the highest available purity and were used as supplied. Dopamine hydrochloride (DA, HCl) was purchased from Aldrich.

Analytical thin-layer chromatography was carried out on silica gel Macherey-Nagel Polygam SIL G/UV 254 (0.25 mm). Column chromatography was performed on Macherey-Nagel Si 60M silica gel (40-63 μm). Optical rotations were measured at 25 °C using a Perkin-Elmer 341 polarimeter. Melting points were measured on a Köfler apparatus.

HPLC was carried out using a Waters system consisting of a 600E multisolvent delivery system, a Rheodyne-type loop injector, and a 2487 dual-channels UV-visible detector set at 254 and 278 nm. A mixture of two solvents constituted the mobile phase: acetonitrile (MeCN) and solvent A. Solvent A was prepared by adding 1% concentrated trifluoroacetic acid (TFA) to deionized water. Semi-preparative reversed-phase HPLC was performed using a 250 \times 20 mm, 5 μm Kromasil C18 column and a 2 mL loop injector, whereas for the analytical reversed-phase HPLC, a 250 \times 4.6 mm, 5 μm Kromasil C18 column, together with a 50 μL loop injector, were used.

^1H NMR and ^{13}C NMR spectra were performed on a Bruker AC-300 spectrometer operating at 300 and 75 MHz, respectively. Chemical shifts are expressed as δ units (part per million) downfield from TMS (tetramethylsilane). The measurements were carried out using the standard pulse sequences. The carbon type (methyl, methylene, methine, or quaternary) was determined by DEPT experiments. ^1H and ^{13}C NMR spectra of all compounds are included in the supporting information as a proof of their identity.

High-resolution mass spectra (HRMS) were recorded on a LTQ-Orbitrap spectrometer operating in positive ion mode.

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(±)-3,4-dihydroxymethamphetamine hydrobromide (HHMA, HBr) was synthesized in two steps from commercially available 3,4-dimethoxyphenylacetone using an adaptation of previously reported procedures with slight modifications (35-38).

(±)-3,4-Dimethoxymethamphetamine [(±)-MMMA]. [previously described as the hydrochloride salt (37,38)] 6.45 mL of a 40% aqueous solution of methylamine (8 equiv) were added dropwise to a solution of 3,4-dimethoxyphenylacetone (3.0 g, 15.45 mmol) in MeOH (50 mL) at 0 °C. The resulting solution was stirred under nitrogen at 0 °C for 30 min, and sodium borohydride (NaBH₄) (632 mg, 16.7 mmol) was added in small portions for 30 min. Then, the reaction mixture was allowed to warm at room temperature and stirred for 1h. After addition of K₂CO₃ (2.2 g), the solvent was evaporated under reduced pressure. H₂O (15 mL) was added to the residue, and the reaction mixture was extracted with diethyl ether (40 mL). The organic phase was then washed with HCl 1N (15 mL). The resulting acidic aqueous phase was alkalized with K₂CO₃, and extracted with diethyl ether (50 mL). The organic phase was then dried over MgSO₄, filtered off and the solvent was evaporated. Column chromatography (dichloromethane-MeOH 80/20 v/v) afforded (±)-MMMA as a yellow oil (2.23g, 10.65 mmol) in 69% yield. ¹H NMR (CDCl₃) δ 1.30 (d, *J* = 6.5 Hz, 3H), 2.70 (m, 4H), 3.17 (dd, *J* = 13.4 Hz, 1H), 3.33 (m, 1H), 3.87 (s, 3H), 3.88 (s, 3H), 5.55 (broad s, 1H), 6.73 (m, 2H), 6.82 (d, *J* = 8.1 Hz, 1H). ¹³C NMR (CDCl₃) δ 15.2, 30.4, 39.2, 55.8, 55.9, 57.0, 111.4, 112.1, 121.3, 128.0, 148.0, 149.2.

(±)-3,4-Dihydroxymethamphetamine hydrobromide (HHMA, HBr). 3.3 mL of a 48% aqueous solution of HBr (18.7 mmol) were added to (±)-MMMA (523 mg, 2.5 mmol). The resulting solution was heated to reflux for 1.5h, under inert atmosphere. After addition of 0.5 mL of methanol and evaporation under reduced pressure, the purple oil was purified by column chromatography (dichloromethane-MeOH 90:10 v/v) affording HHMA, HBr as a colorless oil (524 mg, 2.0 mmol) in 80% yield. Spectroscopic data were identical to those previously reported (38).

Methyl (S)-2-Amino-3-(3,4-dihydroxyphenyl)propanoate Hydrochloride (1). Thionyl chloride (6.57 mL, 90 mmol) was added dropwise to a solution of L-dopa (1.97g, 10 mmol) in MeOH (100 mL) at 0°C. After 18h at 20°C, the solvent was evaporated giving amine chlorohydrate **1** in 100% yield

(2.48g) as a white solid; m.p. 95-96 °C (surprisingly, this value markedly differs from that previously reported - lit.(39) m.p. 174-175 °C); $[\alpha]_D^{25} + 9.4$ (*c* 0.03, MeOH); lit.(39) $[\alpha]_D^{23} + 7.9$ (*c* 1.0, MeOH); Spectroscopic data were identical to those previously reported (39,40).

Methyl (R)-2-(tert-Butoxycarbonylamino)-3-(3,4-dihydroxyphenyl)-propanoate (2). Saturated aqueous NaHCO₃ (7.5 mL) and di-*tert*-butyl dicarbonate (Boc₂O) (1.25 g, 5.41 mmol) in THF (5.5 mL) were added to ester **1** (0.96 g, 3.87 mmol) in THF (7.5 mL), at 0 °C, under nitrogen. The resulting solution was allowed to warm to room temperature and stirred for 1h. After acidification with 1N HCl aqueous solution (12 mL), the solvent was evaporated and the aqueous phase was extracted with ethyl acetate (40 mL). The organic phase was then dried over MgSO₄, filtered off and the solvent was evaporated under reduced pressure. Column chromatography (petroleum ether/ethyl acetate 60/40 v/v) afforded product **2** as a white solid (1.1g, 3.53 mmol) in 92% yield. m.p. 138-139 °C ; (lit.(39) m.p. 134-135 °C); $[\alpha]_D^{25} + 6.0$ (*c* 0.1, MeOH); lit.(39) $[\alpha]_D^{23} + 6.9$ (*c* 1.0, MeOH). Spectroscopic data were identical to those previously reported (39,41).

Methyl (S)-2-(tert-Butoxycarbonylamino)-3-(3,4-dimethoxyphenyl)-propanoate (3). To compound **2** (1.3 g, 4.2 mmol) in acetone (12 mL) were added MeI (2.1 mL, 33.6 mmol) and K₂CO₃ (5.8g, 42.1 mmol). The reaction mixture was heated to reflux for 6h. After filtration of the precipitate, the solution was concentrated and the resulting residue was dissolved in diethyl ether (40 ml). The organic phase was then washed with water (20 mL), dried over MgSO₄, filtered off and the solvent evaporated under reduced pressure. Column chromatography (petroleum ether/ethyl acetate 60/40 v/v) afforded product **3** as a white solid (1.41 g, 4.15 mmol) in 99% yield. m.p. 78-79 °C ; (lit.(39) m.p. 77-78 °C); $[\alpha]_D^{25} + 4.6$ (*c* 0.05, MeOH); lit.(39) $[\alpha]_D^{23} + 6.0$ (*c* 1.0, MeOH). Spectroscopic data were identical to those previously reported (39,42,43)

tert-Butyl [(S)-2-(3,4-Dimethoxyphenyl)-1-(hydroxymethyl)ethyl]-carbamate (4). Lithium aluminum hydride (LiAlH₄) solution in THF (1 M, 35.4 mL) was added dropwise to a solution of compound **3** (3 g, 8.84 mmol) in dry THF (40 mL). The reaction mixture was stirred under nitrogen at room temperature for 1h. Then, 0.5 mL ice-water and 0.5 mL NaOH 0.5 M were successively added.

1 The solid Al(OH)₃ was filtered off and washed with dichloromethane (2 × 40 mL). The filtrate was
2 concentrated to dryness, and the residue was dissolved in dichloromethane (40 mL). The resulting
3 organic phase was then washed with water (20 mL), dried over MgSO₄, filtered off and then
4 concentrated. Column chromatography (petroleum ether/ethyl acetate 60/40 v/v) afforded N-protected
5 α-aminoalcohol **4** as a white solid (2.5 g, 8.0 mmol) in 90% yield. m.p. 94-95 °C ; (lit.(39) m.p. 91-
6 92°C); [α]_D²⁵ – 19.0 (c 0.01, MeOH); lit.(39) [α]_D²³ – 19.6 (c 0.3, MeOH). Spectroscopic data were
7 identical to those previously reported (39,42,43).
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10 **(S)-2-(tert-butoxycarbonylamino)-3-(3,4-dimethoxyphenyl)-1-(4-methylbenzenesulfonyloxy)**
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12 **propane (5)**. To a solution of α-aminoalcohol **4** (2.34 g, 7.51 mmol) in dichloromethane (15.5 mL)
13 dried over CaCl₂, were added triethylamine (1.57 mL, 11.27 mmol), *p*-toluenesulfonyl chloride (1.72 g,
14 9.01 mmol), and dimethylaminopyridine (DMAP) (918 mg, 0.750 mmol). After stirring for 18 h at room
15 temperature, under nitrogen, the reaction mixture was diluted with dichloromethane (40 mL). The
16 organic phase was then washed with water (20 mL), dried over MgSO₄, filtered off and the solvent was
17 evaporated under reduced pressure. The crude residue was recrystallized from diethyl oxide/
18 dichloromethane mixture affording compound **5** as a white solid (2.94g, 6.3 mmol) in 84% yield. m.p.
19 155-156°C ; (lit.(44) m.p. 146-148 °C); [α]_D²⁵ – 16.2 (c 0.09, MeOH); lit.(44) [α]_D²⁵ – 15.4 (c 0.36,
20 CHCl₃). ¹H NMR (CDCl₃) δ 1.41 (s, 9H), 2.47 (s, 3H), 2.80 (m, 2H), 3.87 (s, 6H), 3.95 (m, 2H), 4.73
21 (d, *J* = 7.0 Hz, 1H), 6.64 (d, *J* = 7.9 Hz, 1H), 6.71 (s, 1H), 6.74 (d, *J* = 7.9 Hz, 1H), 7.36 (d, *J* = 8.2 Hz,
22 2H), 7.78 (d, *J* = 8.2 Hz, 2H). ¹³C NMR (CDCl₃) δ 21.7 , 28.3 , 36.,7 , 50.8, 55.9 (× 2), 69.9, 79.8,
23 111.2, 112.3, 121.3, 128.0, 129.1, 130.0, 132.5, 145.1, 147.8, 149.0, 155.0.
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48 **(R)-3,4-Dimethoxymethamphetamine (6) and (S)-3-(3,4-dimethoxyphenyl)-2-(N-methylamino)-**
49 **propan-1-ol (7)**. LiAlH₄ solution in THF (1 M, 4.45 mL) was added dropwise to a solution of
50 compound **5** (501 mg, 1.08 mmol) in dry dichloroethane (13.5 mL) at 0 °C. The resulting solution was
51 allowed to warm to room temperature, under nitrogen, and stirred for 45 min. Then, the reaction mixture
52 was heated to reflux for 1.5h. After hydrolysis with 0.5 mL ice-water and 0.5 mL NaOH 0.5 M, the solid
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Al(OH)₃ was filtered off and washed with dichloromethane (2 × 40 mL). The filtrate was concentrated to dryness. Column chromatography of the residue (dichloromethane/methanol 95/5 v/v with 1.5% triethylamine) afforded compounds **6** (134.5 mg, 0.64 mmol) and **7** (31 mg, 0.15 mmol) as colorless oils in 60% and 13% yields, respectively.

Compound **6**: ¹H NMR (CDCl₃) δ 1.02 (d, *J* = 6.2 Hz, 3H), 2.02 (broad s, 1H), 2.34 (s, 3H), 2.57 (m, 2H), 2.70 (m, 1H), 3.81 (s, 3H), 3.82 (s, 3H), 6.69 (m, 2H), 6.76 (d, *J* = 8.7 Hz, 1H). ¹³C NMR (CDCl₃) δ 19.5, 33.9, 43.0, 55.8 (× 2), 56.3, 111.1, 112.3, 121.1, 131.9, 147.4, 148.7. HRMS (ESI) *m/z* calcd for [M + H]⁺ 210.2955, found 210.1489. [α]_D²⁵ – 1.9 (*c* 0.06, MeOH).

Compound **7**: ¹H NMR (CDCl₃) δ 2.42 (s, 3H), 2.3 (m, 3H), 3.37 (dd, *J* = 10.8 Hz, 1H), 3.67 (dd, *J* = 10.8 Hz, 1H), 3.85 (s, 3H), 3.86 (s, 3H), 6.74 (m, 2H), 6.82 (d, *J* = 8.1 Hz, 1H). ¹³C NMR (CDCl₃) δ 33.7, 37.1, 55.9 (× 2), 61.9, 62.0, 111.3, 112.2, 121.1, 131.0, 147.5, 148.9. HRMS (ESI) *m/z* calcd for [M + H]⁺ 226.2949, found 226.1438. [α]_D²⁵ + 10.1 (*c* 0.01, MeOH).

(R)-3,4-Dihydroxymethamphetamine hydrobromide HHMA, HBr (8). 2.6 mL of a 48% aqueous solution of HBr (15 mmol) was added to compound **6** (420 mg, 2 mmol). The resulting solution was heated to reflux for 1.5h, under nitrogen. After addition of 0.5 mL of methanol and evaporation under reduced pressure, the orange oil was purified by column chromatography (dichloromethane/methanol 80/20) affording compound **8** as a colorless oil (376.5 mg, 1.44 mmol) in 72% yield. [α]_D²⁵ – 2.4 (*c* 0.05, MeOH); ¹H NMR (D₂O) δ 1.02 (d, *J* = 6.6 Hz, 3H), 2.50 (m, 4H), 2.66 (dd, *J* = 13.9 Hz, 1H), 3.18 (m, 1H), 6.42 (dd, *J* = 1.9 and 8.1 Hz, 1H), 6.58 (s, *J* = 1.9 Hz), 6.67 (d, 1H, *J* = 8.1 Hz). ¹³C NMR (D₂O) δ 14.8, 29.9, 37.9, 56.4, 116.2, 116.9, 121.7, 128.1, 143.0, 143.9. HRMS (ESI) *m/z* calcd for [M + H]⁺ 182.2419, found 182.1175.

(S)-3-(3,4-dihydroxyphenyl)-2-(N-methylamino)-propan-1-ol hydrobromide (9). Previous method, replacing compound **6** by compound **7** (414 mg, 2 mmol), afforded after column chromatography, compound **9** as a colorless oil (400.5 mg, 1.44 mmol) in 72% yield. [α]_D²⁵ + 2.1 (*c* 0.016, MeOH); ¹H NMR (D₂O) δ 2.62 (s, 3H), 2.74 (dd, *J* = 14.2 Hz, 1H), 2.83 (dd, *J* = 14.2 Hz,

1H), 3.37 (m, 1H), 3.53 (dd, $J = 12.3$ Hz, 1H), 3.73 (dd, $J = 13.6$ Hz, 1H), 6.64 (dd, $J = 1.9$ and 8.1 Hz, 1H), 6.73 (s, $J = 1.9$ Hz, 1H), 6.77 (d, $J = 8.1$ Hz, 1H). ^{13}C NMR (D_2O) δ 30.0, 32.5, 57.8, 61.0, 116.4, 116.9, 121.7, 127.8, 143.1, 144.1. HRMS (ESI) m/z calcd for $[\text{M} + \text{H}]^+$ 198.2413, found 198.1122.

Cell culture. Human neuroblastoma SH-SY5Y cells were obtained from the European Collection Cell Cultures (ECACC no. 94030304; Salisbury, UK). Cells were cultured in Dulbecco's modified Eagle's medium (DMEM) containing 15% fetal bovine serum (FBS), 1% non essential amino acids, 2 mM L-glutamine, 100 IU/mL penicillin and 100 $\mu\text{g}/\text{mL}$ streptomycin.

Flow cytometry. For cell analysis we used a Cytomics FC500 MPL multiwell-plate flow cytometer (Beckman-Coulter, Brea, CA), equipped with an air-cooled argon-ion laser tuned at 488 nm (blue light). To assay cell death, SH-SY5Y cells were seeded in plastic 96-well plates (final concentration in well 2.7×10^4 cells) and incubated in 10% FBS containing medium for 24 h (50% confluence) in CO_2 incubator at 37 °C. Then, the indicated concentrations of the compounds in 5% FBS containing medium were added and the plates kept for further 24 h. Cells were then trypsinized and resuspended in their original wells in 5% FBS containing medium, in the presence of 2.5 $\mu\text{g}/\text{mL}$ propidium iodide (PI). To identify PI uptake by dead cells (PI positive) by flow cytometry, cells were excited at 488 nm and the emission was detected at 625 nm. From dose-response curves of at least three experiments, TC_{50} , the molar concentration producing 50% of cell death, was calculated. SD, at doses in the vicinity of the TC_{50} values were lower than 10%. The percentage of viable (PI negative), apoptotic (PI intermediate) and necrotic (PI positive) cells was calculated by means of a plot correlating cell size (forward-angle light scatter) and PI uptake (45).

To measure the mitochondrial generation of superoxide, trypsinized cells were resuspended in their original wells in 5% FBS containing medium in the presence of 1.25 μM MitoSOX Red probe (Invitrogen), a mitochondrial superoxide indicator having excitation/emission maxima of approximately 510/580 nm. The mean fluorescence intensity was calculated for each sample. The results were expressed as Fluorescence Arbitrary Units (F.A.U). As a positive control we used the superoxide

1 generator plumbagin (2-methyl-5-hydroxy-1,4-naphthoquinone), inducing a mean intensity of 20 F.A.U.
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3 at the dose of 0.02 mM.
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5 Caspase-3 activity was measured by flow cytometry, using Cleaved Caspase-3 (Asp-175) Antibody
6 (Alexa Fluor[®] 488 Conjugate) in accordance with the protocol supplied by the manufacturer (Cell
7 Signaling Technology[®]). As a positive control, we used camptothecin inducing 54% of caspase-3
8 activity in SH-SY5Y cells at the dose of 0.5 μ M (24h).
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16 ***Escherichia coli* cytotoxicity plate assays.** The *E. coli* strains were IC5282 (OxyR⁺ strain), IC5204,
17 deficient in the OxyR function (OxyR⁻ strain), and IC2880, a tryptophan-requiring strain (34). Overnight
18 cultures were grown in Oxoid Nutrient Broth No. 2. Solid minimal E4 medium contained 15 g Difco
19 agar and 4 g glucose per liter of Vogel-Bonner E buffer (containing MgSO₄ · 7 H₂O (10 g), citric acid
20 monohydrate (100 g), K₂HPO₄ (500 g) and NaH₂PO₄ · 7 H₂O (175 g) per liter of distilled water). Top
21 agar contained 6 g Difco agar and 5 g NaCl per liter of distilled water. In the assays, 100 μ L of a suitable
22 dilution of the test compound were added to 100 μ L of the bacterial mix, containing approximately 800
23 test cells (from IC5282 or IC5204 strains) and 3x10⁷ filler cells (from IC2880 strain) unable to grow in
24 minimal E4 medium due to their requirement for tryptophan and used to obtain a background lawn.
25 Where indicated, mushroom tyrosinase (EC 1.14.18.1; Sigma) was added to the assay (100 μ L of a
26 suitable dilution from a starting solution in 0.1 M sodium phosphate pH 7.0). After the addition of 2.5
27 mL of molten top agar, the mixture was poured on plates containing solid E4 medium (about 35 mL),
28 which were incubated at 37 °C, for 24 or 48 h. Data were the mean of at least three experiments. From
29 the dose-response curves, the TC₅₀ value was calculated as the concentration (μ mol/plate) needed to
30 reduce the survival to 50%. SD at dose close to the TC₅₀ values were lower than 10%.
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55 Results and Discussion

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1 **Chemistry. Synthesis of (±)-HHMA.** (±)-HHMA was prepared in a straightforward manner (Scheme
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3 1), starting from commercially available 3,4-dimethoxyphenylacetone by reductive amination with
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5 sodium borohydride leading to (±)-MMMA in 69% yield. Complete demethylation of (±)-MMMA using
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7 hydrobromic acid, heated at reflux, afforded (±)-HHMA in 80% yield. Interestingly, this two step
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9 reaction sequence produced (±)-HHMA in a markedly improved overall yield (55%) when compared
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11 with earlier reported procedures starting from 3,4-dimethoxybenzaldehyde and nitroethane, for which
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13 the overall yields did not exceed 20% (35-38).
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17 **Synthesis of R(-)-HHMA.** We prepared R(-)-HHMA from commercially available L-dopa, chosen
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19 as the chiral source, through a seven step sequence which involved protection and deprotection reactions
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21 as reported in Scheme 2. This strategy required the intermediary synthesis of *N*-protected α -amino
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23 alcohol **4**, which has been previously described in four steps and 74.5% overall yield (39). Minor
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25 modifications of this procedure (solvent especially) allowed us to improve the yield in *N*-protected α -
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27 amino alcohol **4**, which was prepared in 82% overall yield. Quantitative esterification of L-dopa in
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29 MeOH, followed by protection of the resulting amino group by treatment with di-*tert*-butyl-dicarbonate
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31 (Boc₂O) in THF, gave access to compound **2** in 92% yield. Methylation of the aromatic hydroxyl groups
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33 in acetone led, in quantitative yield, to compound **3** whose the ester function was finally reduced with
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35 LiAlH₄ in THF, affording *N*-protected α -amino alcohol **4** in 90% yield. Treatment of **4** with *para*-
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37 toluenesulfonyl chloride (TsCl), in the presence of dimethylaminopyridine (DMAP) as the catalyst,
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39 provided the rather unstable tosylate **5** in 84% yield. Because **5** spontaneously converted through
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41 intramolecular cyclisation between its alcohol group and its carbamate function to oxazolidinone on
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43 silica gel (39), tosylate **5** had to be purified by recrystallisation. Concomittant reduction of the tosyl and
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45 BOC groups was achieved using LiAlH₄ in THF/dichloroethane 25/75 v/v, leading to R(-)-MMMA **6**
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47 as the major product (60%), with high optical purity (>99%) as determined by HPLC analysis of the
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49 mixture of Mosher amides prepared from racemate (±)-MMMA and from enantiomerically pure R(-)-
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51 MMMA **6** (Figures 3 and 4) (See the supporting information for the synthesis of the Mosher amides).
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1 Aminoalcohol **7** was also obtained as the minor product in 13% yield. Obviously, this reduction reaction
2 proved to be the more difficult step of the synthetic procedure, due to the occurrence of both
3 spontaneous cyclization of **5** to oxazolidinone and partial reduction of the tosyl group to aminoalcohol **7**.
4 So, a thorough control of temperature was necessary to obtain good yields in *R*-(-)-MMMA **6**. Finally,
5 complete demethylation of *R*-(-)-MMMA **6**, using hydrobromic acid heated at reflux, afforded *R*-(-)-
6 HHMA **8** in 72% yield with 99.5% ee, whereas that of aminoalcohol **7** led to the deprotected α -
7 aminoalcohol **9** in 72% yield.
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10 **Toxicological study.** The in vitro toxicological study was performed using the human dopaminergic
11 neuroblastoma SH-SY5Y cell line, which is commonly used for studying the toxicity of catecholamine
12 derivatives (46,47). Loss of cell viability was monitored by flow cytometric analysis of propidium iodide
13 (PI) uptake in dead cells (PI positive cells) having defective plasma membranes. In addition, as we
14 previously determined the cytotoxicity profile of (\pm)-HHMA in *Escherichia coli* assays, specific for the
15 detection of oxidative toxicity (18,34), these bacterial tests were also included in the present study.
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18 The results of the flow cytometric analysis of acute (24 h) cytotoxicity in SH-SY5Y cells are shown
19 in Table 1. *R*-(-)-HHMA enantiomer **8**, induced a decrease in cell viability similar to that resulting from
20 exposure to (\pm)-HHMA racemate, with a TC_{50} value of 0.30 mM. This decrease was also close to that
21 found for the control catecholamine dopamine (DA) (TC_{50} = 0.25 mM), whereas a lower toxicity (TC_{50}
22 = 1.0 mM) was observed for aminoalcohol **9** isolated as a byproduct from the synthesis of *R*-(-)-HHMA
23 **8** (Scheme 2). For (\pm)-MMMA racemate, an intermediate in the synthesis of (\pm)-HHMA possessing a
24 protected catechol group (Scheme 1) utilized for comparative purposes, toxicity was detected at high
25 doses (TC_{50} = 11.3 mM). Similarly, MDMA metabolite 4-hydroxy-3-methoxymethamphetamine [(\pm)-
26 HMMA)] has been described as considerably less toxic than catechol-containing metabolites to PC12
27 cells, in the MTT reduction assay (38). Our results on cytotoxicity obtained from the PI uptake assay
28 directly correlated with the extent of cell death evaluated by the MTT reduction assay (data not shown).
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E. coli based assays allow a mechanistic characterization of toxicity, so that they can be used as a model system. In these assays, toxicity mediated by ROS (ROS-TOX) can be detected in OxyR⁻ cells deficient in OxyR function, because they are unable to exert antioxidant defenses (34,48). In contrast, ROS-TOX is inhibited in OxyR⁺ cells due to the elimination of hydrogen peroxide by catalase and alkylhydroperoxide reductase, both enzymes encoded by OxyR-regulated genes, so that the toxicity arising from the binding of quinones with cellular nucleophiles could then be shown (48,49). The results of the *E. coli* plate assays are shown in Table 1. *R*-(-)-HHMA **8** and (±)-HHMA gave similar positive responses in the OxyR⁻ assay, with TC₅₀ values of 0.8 and 0.75 μmol/plate respectively, indicating high ROS-TOX induction. Furthermore, an increase in ROS-TOX (TC₅₀ = 0.6 and 0.5 μmol/plate, respectively) was observed in the presence of tyrosinase, which catalyzed the oxidation to *o*-quinone. For (±)-MMMA, ROS-TOX induction was abolished whereas, for aminoalcohol **9** and dopamine, it was poor (TC₅₀ ≈ 4 μmol/plate) but greatly increased following catalytic oxidation by tyrosinase (TC₅₀ ≈ 2 μmol/plate). The enhancement of toxicity promoted by tyrosinase suggests that the oxidation of these catecholamines to the corresponding *o*-quinone opens the way for redox cycling, leading to the formation of ROS (18). In contrast, the negative response in the OxyR⁺ assay (Table 1), even in the presence of tyrosinase (data not shown), was indicative of the absence of quinone-promoted toxicity, which is resistant to inhibition by antioxidant defenses (48,49).

The good correlation found between the toxicity observed in SH-SY5Y cells and in *E. coli* OxyR⁻ cells would indicate a susceptibility of the SH-SY5Y cell line to the oxidative toxicity induced by the catecholamine derivatives. Consistent with this possibility is the mitochondrial generation of superoxide anion by these compounds in SH-SY5Y cells, detected by flow cytometric analysis using the MitoSOX Red fluorochrome (Table 2). Note, however, that HHMA and DA induced similar toxicities in SH-SY5Y cells, whereas, converse to HHMA, DA was found a poor inducer of ROS-TOX in *E. coli* OxyR⁻ cells (Table 1).

Flow cytometry offers the possibility of multiparametric assays providing simultaneously several endpoints to characterize different modes of cell death at a single-cell level (45). In addition to the assay

1 of plasma membrane permeability (PI uptake) for cell death evaluation (TC₅₀ value), we used flow
2 cytometry to discriminate between live (PI negative), apoptotic (PI intermediate), and necrotic (PI
3 positive) subpopulations, by means of a plot correlating cell size (forward-angle light scatter) and PI
4 uptake (Table 3). For *R*-(–)-HHMA, (±)-HHMA, aminoalcohol **9** and DA induction of necrotic cells but
5 not that of apoptotic cells was detected. In contrast, induction of apoptotic cells (48% at 10 mM) was
6 clearly observed for (±)-MMMA. These results were complemented by a flow cytometric measurement
7 of cleaved (activated) caspase-3, a critical effector of most of the apoptotic pathways (Table 3). For *R*-
8 (–)-HHMA and (±)-HHMA, the activity of caspase-3 increased to a maximal value after exposure to 0.1
9 mM, a dose below the TC₅₀ value (0.3 mM). For aminoalcohol **9** and DA, maximal induction of
10 caspase-3 activity required a higher dose (1 mM), whereas for (±)-MMMA, a significant increase in
11 caspase-3 activity was observed at 10 mM, in the absence of any detectable cytotoxicity. These results
12 suggest that induction of necrosis is dominant over that of apoptosis in ROS-sensitive SH-SY5Y cells.
13 In these cells, induction of proapoptotic events such as caspase-3 activation would not be followed by
14 the detection of apoptosis.

35 Conclusions

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40 Despite several years of intensive efforts, the mechanisms by which MDMA destroys brain serotonin
41 (5-HT) axon terminals remains to be fully elucidated. In particular, the sequence order of events and the
42 identity of specific neurotoxic entities have yet to be determined. There has recently been a resurgence
43 of interest in the possibility that MDMA metabolites (especially 5-NAC-HHMA) might play a role in
44 MDMA neurotoxicity. The discrepant findings between the recently published work (19) with those
45 previously reported (20, 21), leave open the possibility that 5-NAC-HHMA may require the presence of
46 MDMA and/or elevated body temperature to be toxic, but also raise the question of the chirality of
47 MDMA which was not considered in previously reported in vivo studies, because HHMA, the precursor
48 of 5-NAC-HHMA metabolite, was used as the racemic mixture. In vitro metabolism of MDMA is
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1 enantioselective, with a preference for the *S*-enantiomer caused in part by CYP2C19 and CYP2D6-
2 mediated demethylenation (23). However, correlation with the in vivo situation is not straightforward
3 because in vivo catecholic phase I metabolites such as HHMA are further metabolized by *O*-methylation
4 and/or glucuronidation/sulfation. Recent investigations have revealed that the *S*-enantiomer of catecholic
5 phase I metabolites was preferably *O*-methylated by catechol-*O*-methyltransferase (COMT) (50),
6 resulting in the accumulation of the *R*-enantiomer. So, the synthesis of *R*-(-)-HHMA, by allowing the
7 synthesis of noticeable amounts of the corresponding diastereomerically pure form of 5-NAC-HHMA
8 metabolite, will permit a comprehensive examination of its participation in the in vivo neurotoxic
9 effects of ecstasy.

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21 The in vitro toxicological study, performed in human neuroblastoma SH-SY5Y cells, revealed similar
22 toxicities for HHMA racemate and its *R*-enantiomer. The inclusion in this work of an *E. coli* based plate
23 assay, specific for the detection of oxidative toxicity, showed a good correlation between the toxic
24 effects observed both in human and bacterial cells. This correlation led us to conclude that SH-SY5Y
25 cells exhibit a low protection against oxidative toxicity and that the observed cell death in these cells
26 would be mediated by ROS, mainly generated through the redox activity of the generated quinone
27 species. The toxicophore (i.e. the reactive chemical moiety) would be essentially the catechol group and
28 therefore the problem of the toxicity of the phase I catecholic metabolites of MDMA would be related to
29 the toxicity induced by other catecholamine compounds, as is the case with DA. Although to a lesser
30 extent, the nature of the alkylamino side chain of catecholamine derivatives also plays a role. As a result,
31 *R*-(-)-HHMA and aminoalcohol **9**, two enantiomeric pure forms, exhibit different cytotoxicity in *E. coli*
32 plate assays. Probably, a physical parameter such as lipophilicity may contribute to the exhaustive
33 toxicity. Accordingly, *R*-(-)-HHMA is at once more toxic ($TC_{50} = 0.75 \mu\text{mol per plate}$) and more
34 lipophilic ($\log P = 0.90$) than aminoalcohol **9**, for which $TC_{50} = 5 \mu\text{mol per plate}$ and $\log P = 0.04$.

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55 **Importantly, although HHMA racemate and its *R*-enantiomer exhibited close toxicological profiles, no**
56 **conclusion could be drawn about the enantioselective toxicity. For a complete comparison and**
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1 evaluation of enantioselective toxicity of both enantiomers, the synthesis of optically pure *S*-(+)-
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3 enantiomer is required. This synthesis is envisioned in our laboratory and will be reported in due course.
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5 The catechol toxicophore is neither present in MDMA, the precursor of HHMA, nor in MMMA for
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7 which the two phenolic groups are protected. For MMMA, no toxicity was detected in the bacterial plate
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9 assay whereas, at high doses, toxic effects could be detected in the flow cytometric assay measuring PI
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11 uptake. We can then speculate that the toxicity observed with MMMA is not related to increased
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13 oxidative stress. Consequently, caution is advised when comparing this toxicity with that induced by the
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15 catecholamine derivatives (38). This conclusion can be extended to MDMA, also bearing a protected
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17 catechol, so that in studies dealing with MDMA toxicity, there is a need to distinguish the toxicity
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19 ascribed to MDMA itself from that depending on MDMA biotransformation leading to catechol
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21 metabolites such as HHMA and the thioether conjugates.
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26 A general view, including qualitative as well as quantitative changes in toxicity, can be drawn. The
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28 parent drug (MDMA) associated to some adverse effects (12), is metabolized to redox active forms such
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30 as HHMA, which would induce, in cells endowed with low antioxidant defenses (Table 1), an oxidative
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32 toxicity resulting in necrosis (Table 3). These active forms could be further metabolized (e.g. by phase II
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34 conjugation reactions) to compounds such as MMMA able to induce a different kind of toxicity
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36 resulting in apoptosis (Table 3). This point of view might be considered when evaluating the *in vivo*
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38 situation, in which enantioselective metabolic reactions play a pivotal role.
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1 **Supporting Information Available.** Experimental procedures for the determination of enantiomeric
2 excess of compounds **6** and **8**; ^1H and ^{13}C NMR spectra of (\pm)-HHMA, (\pm)-MMMA, *R*-(-)-HHMA and
3 its synthetic intermediates, together with aminoalcohol derivatives **7** and **9**; analytical HPLC
4 chromatograms of (\pm)-HHMA, (\pm)-MMMA, *R*-(-)-HHMA and of aminoalcohol **9**. This information is
5 available free of charge via the Web at <http://pubs.acs.org>.
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References

- (1) Milroy, C. M., Clark, J. C., and Forrest, A. R. W. (1996) Pathology of deaths associated with "ecstasy" and "eve" misuse. *J. Clin. Pathol.* *49*, 149-153.
- (2) Walubo, A., and Seger, D. (1999) Fatal multi-organ failure after suicidal overdose with MDMA, "ecstasy": case report and review of the literature. *Hum. Exp. Toxicol.* *18*, 119-125.
- (3) Fineschi, V., Centini, F., Mazzeo, E., and Turillazzi, E. Adam (MDMA) and Eve (MDA) misuse: an immunohistochemical study on three fatal cases. (1999) *Forensic Sci. Int.* *104*, 65-74.
- (4) Cole, J. C., and Sumnall H. R. (2003) Altered states: the clinical effects of ecstasy. *Pharmacol. Ther.* *98*, 35-58.
- (5) Ricaurte, G. A., and McCann, U. D. (2005) Recognition and management of complications of new recreational drug use. *Lancet*, *365*, 2137-2145.
- (6) Green, A. R., Mehan, A. O., Elliot, J. M., O'Shea, E., and Colado, M. I. (2003) The pharmacology and clinical pharmacology of 3,4-methylenedioxymethamphetamine (MDMA, "ecstasy"). *Pharmacol. Rev.* *55*, 463-508 and references therein.
- (7) Lyles, J., and Cadet, J. L. (2003) Methylenedioxymethamphetamine (MDMA, Ecstasy) neurotoxicity: cellular and molecular mechanisms. *Brain Res. Rev.* *42*, 155-168 and references therein.
- (8) Morton, J. (2005) Ecstasy: pharmacology and neurotoxicity. *Curr. Opinion Pharmacol.* *5*, 79-86.
- (9) Baumann, M. H., Wang, X., and Rothman, R. B. (2007) 3,4-methylenedioxymethamphetamine (MDMA) neurotoxicity in rats: a reappraisal of past and present findings. *Psychopharmacology*, *189*, 407-424.

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- (10) De Win, M.M.L., Jager, G., Booij, J., Reneman, L., Schilt, T., Lavini, C., Olabarriaga, S.D., den Heeten, G.J., and van den Brink W. (2008) Sustained effects of ecstasy on the human brain: a prospective neuroimaging study in novel users. *Brain*, *131*, 2936-2945.
- (11) Gudelsky, G. A., and Yamamoto, B. K. (2008) Action of 3, 4-methylenedioxymethamphetamine (MDMA) on cerebral dopaminergic, serotonergic and cholinergic neurons. *Pharmacol. Biochem. Behav.* *90*, 198-207.
- (12) Capela, J. P., Carmo, H., Remiao, F. Lourdes Bastos, M., Meisel, A., and Carvalho, F. (2009) Molecular and cellular mechanisms of ecstasy-induced neurotoxicity: an overview. *Mol. Neurobiol.* *39*, 210-271 and references therein.
- (13) Goni-Allo, B., Puerta, E., Mathuna, B. O., Hervias, I., Lasheras, B., de la Torre, R., Aguirre, N. (2008) On the role of tyrosine and peripheral metabolism in 3,4-methylenedioxymethamphetamine-induced serotonin neurotoxicity in rats. *Neuropsychopharmacology* *54*, 885-900.
- (14) Esteban, B., O'Shea, E., Camarero, J., Sanchez, V., Green, A. R., and Colado, M. I. (2001) 3,4-Methylenedioxymehamphetamine induces monoamine release, but not toxicity, when administered centrally at a concentration occurring following a peripherally injected neurotoxic dose. *Psychopharmacology* *154*, 251-260.
- (15) Monks, T. J., Jones, D. C., Bai, F., and Lau, S. S. (2004) The role of metabolism in 3, 4-(±)-methylenedioxyamphetamine and 3,4-(±)- methylenedioxymethamphetamine (Ecstasy) toxicity. *Ther. Drug Monit.* *26*, 132-136.
- (16) Segura, M., Jordi, O., Farré, M., McLure, J.A., Pujadas, M., Pizarro, N., Llebaria, A., Joglar, J., Roset, P.N., Segura, J., and de la Torre, R. (2001) 3,4-Dihydroxymethamphetamine (HHMA). A major in vivo 3, 4-methylenedioxymethamphetamine (MDMA) metabolite in humans. *Chem. Res. Toxicol.* *14*, 1203-1208.

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- (17) Perfetti, X., O'Mathuna, B., Pizarro, N., Cuyas, E., Khymenets, O., Almeida, B., Pellegrini, M., Pichini, S., Lau, S. S., Monks, T. J., Farré, M., Pascual, A., Joglar, J., and de La Torre, R. (2009) Neurotoxic thioether adducts of MDMA identified in human urine after ecstasy ingestion. *Drug Metab. Dispos.* 37, 1448-1455.
- (18) Felim, A., Urios, A., Neudörffer, A., Herrera, G., Blanco, M., and Langeron, M. (2007) Bacterial plate assays and electrochemical methods: an efficient tandem for evaluating the ability of catechol-thioether metabolites of MDMA ("ecstasy") to induce toxic effects through redox-cycling. *Chem. Res. Toxicol.* 20, 685-693.
- (19) Mueller, M., Yuan, J., Felim, A., Neudörffer, A., Peters F.T., Maurer, H.H., McCann U.D., Langeron, M., Ricaurte, G.A. (2009) Further studies on the role of metabolites in MDMA-induced serotonergic neurotoxicity. *Drug Metab. Dispos.* 37, 2079-2086.
- (20) Jones, D.C., Duvauchelle, C., Ikegami, A., Olsen, C.M., Lau, S.S., de la Torre, R. and Monks, T.J. (2005) Serotonergic neurotoxic metabolites of ecstasy identified in rat brain. *J. Pharmacol. Exp. Ther.* 313, 422-431.
- (21) Erives, G. V., Lau, S. S., and Monks, T. J. (2008) Accumulation of neurotoxic thioether metabolites of 3,4-(±)-methylenedioxymethamphetamine in rat brain. *J. Pharmacol. Exp. Ther.* 324, 284-291.
- (22) Pizarro, N., de la Torre, R., Joglar, J., Okumura, N., Perfetti, X., Lau, S. S., and Monks, T. J. (2008) Serotonergic neurotoxic thioether metabolites of 3,4-methylenedioxymethamphetamine (MDMA, « ecstasy ») : synthesis, isolation and characterization of diastereoisomers. *Chem. Res. Toxicol.* 21, 2272-2279.
- (23) Meyer, M.R., Peters, F.T., and Maurer, H.H. (2008) The role of human hepatic cytochrome P450 isoenzymes in the metabolism of racemic 3, 4-methylenedioxymethamphetamine and its enantiomers. *Drug Metab. Dispos.* 36, 2345-2354.

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- (24) Anderson, G.M., Braun, G., Braun, U., Nichols, D.E., and Shulgin, A.T. (1978) Absolute configuration and psychotomimetic activity. In *Quasar Research Monograph 22* (Barnett, G., Trisc, M., and Willette, R., Eds) pp 8-15, National Institute on Drug Abuse, Washington, DC.
- (25) Hiramatsu, M., Nabeshima, T., Kameyama, T., Maeda, Y., and Cho, A.K. (1989) The effect of optical isomers of 3,4-methylenedioxyamphetamine (MDMA) on stereotyped behaviour in rats. *Pharmacol. Biochem. Behav.* *33*, 343-347.
- (26) Johnson, M. P., Hoffman, A. J., and Nichols, D. E. (1986) Effects of the enantiomers of MDA, MDMA and related analogues on [3H]serotonin and [3H]dopamine release from superfused rat brain slices. *Eur. J. Pharmacol.* *132*, 269-276.
- (27) Fantegrossi, W. E., Godlewski, T., Karabenick, R. L., Stephens, M. J., Ullrich, T., Rice, K. C., Woods, J. H. (2003) Pharmacological characterization of the effects of 3,4-methylenedioxyamphetamine (“ecstasy”) and its enantiomers on lethality, core temperature, and locomotor activity in singly housed and crowded mice. *Psychopharmacology* *166*, 202-211.
- (28) Pizarro, N., Farré, M., Pujadas, M., Peiro, A. M., Poset, P. N., Joglar, J., and de la Torre, R. (2004) Stereochemical analysis of 3,4-methylenedioxyamphetamine and its main metabolites in human samples including the catechol-type metabolite (3,4-dihydroxymphetamine). *Drug Metab. Dispos.* *32*, 1001-1007.
- (29) Fantegrossi, W. E. (2008) In vivo pharmacology of MDMA and its enantiomers in rhesus monkey. *Exp. Clin. Psychopharmacol.* *16*, 1-12 and references therein.
- (30) Fantegrossi, W. E., Murai, N., Mathuna, B. O., Pizarro, N., de la Torre, R. (2009) Discriminative stimulus effects of MDMA and its enantiomers in mice : pharmacokinetic considerations. *J. Pharmacol. Exp. Ther.* *329*, 1006-1015.

- 1 (31) Felim, A., Neudörffer, A., Monnet, F. P., and LARGERON, M. (2008) Environmentally friendly
2 expeditious one-pot electrochemical synthesis of bis-catechol-thioether metabolites of ecstasy: in
3 vitro neurotoxic effects in the rat hippocampus. *Int. J. Electrochem. Sci.* 3, 266-281.
4
5
6
7
8 (32) Macedo, C., Branco, P.S., Ferreira, L. M., Lobo, A. M., Capela, J. P., Fernandes, E., de Lourdes
9 Bastos, M., and Carvalho, F. (2007) Synthesis and cyclic voltammetry studies of 3,4-
10 methylenedioxymethamphetamine (MDMA) human metabolites. *J. Health Sci.* 53, 31-42.
11
12
13
14
15
16 (33) Pizarro, N., de la Torre, R., Farré, M., Segura, J., Llebaria, A., and Joglar, J. (2002) Synthesis and
17 capillary electrophoretic analysis of enantiomerically enriched references standard of MDMA and
18 its main metabolites. *Bioorg. Med. Chem.* 10, 1085-1092.
19
20
21
22
23
24 (34) Martínez, A., Urios, A., and Blanco, M. (2000) Mutagenicity of 80 chemicals in *Escherichia coli*
25 tester strains IC203, deficient in OxyR, and its *oxyR*⁺ parent WP2 *uvrA*/pKM101: detection of 31
26 oxidative mutagens. *Mutat. Res.* 467, 41-53.
27
28
29
30
31
32 (35) Morgan, P.H., and Beckett, A.H. (1975) Synthesis of some N-oxygenated products of 3,4-
33 dimethoxyamphetamine and its N-alkyl derivatives. *Tetrahedron*, 31, 2595-2601.
34
35
36
37 (36) Borgman, R.J. Baylor, M.R., McPhillips, J.J., and Stitzel, R.E. (1974) α -Methyldopamine
38 derivatives. Synthesis and pharmacology. *J. Med. Chem.* 17, 427-430.
39
40
41
42
43 (37) Cannon, J.G., Perez, Z., Long, J.P., Rusterholtz, D.B., Flynn, J.R., Costall, B., Fortune, D.H., and
44 Naylor, R.J. (1979) N-alkyl derivatives of (\pm)- α -methyldopamine. *J. Med. Chem.* 22, 901-907.
45
46
47
48
49 (38) Milhazes, N., Cunha-Oliveira, T., Martins, P., Garrido, J., Oliveira, C., Rego, A. C., and Borges,
50 F. (2006) Synthesis and cytotoxicity profile of 3,4-methylenedioxymethamphetamine (“ecstasy”)
51 and its metabolites on undifferentiated PC12 cells: a putative structure-toxicity relationship.
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60
- (39) Aubry, S., Pellet-Rostaing, S., and Lemaire, M. (2007) Oxidative nucleophilic substitution (S_NO_X) of the benzylic position as a tunable synthesis of tetrahydroisoquinoline natural alkaloid analogues. *Eur. J. Org. Chem.* 5212-5225.
- (40) Wells, G. J., Tao, M., Josef, K. A., and Bihovsky, R. (2001) 1,2-Benzothiazine 1,1-dioxide P_2 - P_3 peptide mimetic aldehyde calpain I inhibitors. *J. Med. Chem.* 44, 3488-3503.
- (41) Tang, S., Martinez, L. J., Sharma, A., and Chai, M. (2006) Synthesis and characterization of water-soluble and photostable L-DOPA dendrimers. *Org. Lett.* 8, 4421-4424.
- (42) Bloom, J. D., Dutia, M. D., Johnson, B. D., Wissner, A., Burns, M. G., Largia, E. E., Dolan, J. A., and Claus, T. H. (1992) (*R,R*)-5-[2-[[2-(3-chlorophenyl)-2-hydroxyethyl]-amino]propyl]-1,3-benzodioxole-2,2-dicarboxylate (CL 316, 243). A potent β -adrenergic agonist virtually specific for β_3 receptors. A promising antidiabetic and antiobesity agent. *J. Med. Chem.* 35, 3081-3084.
- (43) Garcia, E., Arrasate, S., Lete, E. and Sotomayor, N. (2005) Diastereoselective intramolecular α -amidoalkylation reactions of L-DOPA derivatives. Asymmetric synthesis of pyrrolo[2,1-*a*]isoquinolines. *J. Org. Chem.* 70, 10368-10374.
- (44) Clark, R.D., Berger, J., Lee, C.-H., and Muchowski, J.M. (1987) Synthesis and determination of the absolute stereochemistry of the enantiomers of 3-substituted 1,2,3,4-tetrahydroisoquinolines related to the calcium antagonist verapamil. *Heterocycles*, 26, 1290-1302.
- (45) Darzynkiewicz, Z., Juan, G., Li, X., Gorczyca, W., Murakami, T., and Traganos, F. (1997) Cytometry in cell necrobiology: analysis of apoptosis and accidental cell death (necrosis). *Cytometry* 27, 1-20.
- (46) Gomez-Santos, C., Ferrer, I., Santidrian, A.F., Barrachina, M., Gil, J., Ambrioso, S. (2003) Dopamine induces autophagic cell death and α -synuclein increase in human neuroblastoma cells. *J. Neurosci. Res.* 73, 341-350.

- 1 (47) Haque, M.E., Asanuma, M., Higashi, Y., Miyazaki, I., Tanaka, K.-I., Ogawa, N. (2003)
2 Apoptosis-inducing neurotoxicity of dopamine and its metabolites via reactive quinone
3 generation in neuroblastoma cells. *Biochim. Biophys. Acta* 1619, 39-52.
4
5
6
7
8 (48) Urios, A., Llargeron, M., Fleury, M.-B., and Blanco, M. (2006) A convenient approach for
9 evaluating the toxicity profile of in vitro neuroprotective alkylaminophenol derivatives. *Free*
10 *Radical Biol. Med.* 40, 791-800.
11
12
13
14
15
16 (49) Urios, A., López-Gresa, M.P., González, M.C., Primo, J., Martínez, A., Herrera, G., Escubedo,
17 J.C., O'Connor, J.E., Blanco, M. (2003) Nitric oxide promotes strong toxicity of phenolic
18 compounds against *Escherichia coli*: the influence of antioxidant defenses. *Free Radical Biol.*
19 *Med.* 35, 1373-1381.
20
21
22
23
24
25
26 (50) Meyer, M.R., and Maurer, H.H. (2009) Enantioselectivity in the methylation of the catecholic
27 phase I metabolites of methylenedioxy designer drugs and their capability to inhibit catechol-O-
28 methyltransferase-catalyzed dopamine 3-methylation. *Chem. Res. Toxicol.* 22, 1205-1211.
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Table 1. In Vitro Cytotoxicity Profile of *R*-(-)-HHMA in comparison with related catecholamines

compound	TC ₅₀ in assay			
	SH-SY5Y ^a		<i>E. coli</i> ^b	
	PI	OxyR ⁻	OxyR ⁻ /Tyr ^c	OxyR ⁺
<i>R</i> -(-)-HHMA 8	0.3	0.8	0.6	– ^d
(±)-HHMA	0.3	0.75	0.5	–
Aminoalcohol 9	1.0	5	2	–
(±)-MMMA	11.3	–	–	–
DA	0.25	4	1.8	–

^a SH-SY5Y = human dopaminergic neuroblastoma cells.

PI = propidium iodide assay: live cells, PI negative; dead cells, PI positive. TC₅₀ in mM.

^b TC₅₀ in μmol/plate.

^c Tyr = 50 units of tyrosinase per plate.

^d (-), survival >50% at the highest dose tested.

Table 2. Mitochondrial generation of superoxide anion by *R*-(-)-HHMA in comparison with related catecholamines

compound	dose (mM)	F.A.U. ^a
–	–	8.0
<i>R</i> -(-)-HHMA 8	0.1	8.0
	0.5	14.8
(±)-HHMA	0.1	8.6
	0.5	13.3
Aminoalcohol 9	0.25	11.5
	0.5	13.3
DA	0.1	9.1
	0.31	10.1

^a Mitochondrial generation of superoxide anion was determined by flow cytometry using the MitoSOX Red fluorochrome. F.A.U. = Fluorescence Arbitrary Units

Table 3. Flow cytometric characterization of cell death and of caspase-3 activity in treated SH-SY5Y cells

compound	dose (mM)	apoptotic cells (%) ^a	necrotic cells (%) ^b	caspase-3 activity (%) ^c
–	–	12	23	9
<i>R</i> -(-)-HHMA 8	0.1	10	34	36
	1	14	80	37
(±)-HHMA	0.1	9	30	42
	1	10	82	37
Aminoalcohol 9	0.5	9	29	15
	1	15	51	27
(±)-MMMA	10	48	25	18
DA	0.1	13	35	16
	1	11	83	22

^aPI intermediate cells; ^bPI positive cells.

^c Caspase-3 activity was determined by flow cytometry using cleaved caspase-3 (Asp175) antibody.

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3 **Figure Legends**
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7 **Figure 1.** Stereochemistry of the hepatic metabolism leading to *R*-(-)-HHMA and *S*-(+)-HHMA
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12 **Figure 2.** Chemical Structures of Studied Compounds
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17 **Figure 3.** HPLC chromatograms for the analytical resolution of Mosher amide diastereoisomers
18 prepared from (±)-MMMA racemate [(H₂O + 1 %TFA) / MeCN 53/47; flow rate: 0.9 mL.min⁻¹]:
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20 a) Mosher amide of *R*-(-)-MMMA isolated after semi-preparative HPLC; b) Mosher amide of *S*-(+)-
21 MMMA isolated after semi-preparative HPLC; c) mixture of the Mosher amides obtained from
22 (±)-MMMA racemate.
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32 **Figure 4.** HPLC chromatograms for the analytical determination of the enantiomeric excess of *R*-(-)-
33 MMMA **6** [(H₂O + 1 %TFA) / MeCN 53/47; flow rate: 0.9 mL.min⁻¹]: a) Mosher amide prepared from
34 enantiomerically pure *R*-(-)-MMMA **6**; b) mixture of the Mosher amides obtained from (±)-MMMA
35 racemate. c) Mosher amide prepared from enantiomerically pure *R*-(-)-MMMA **6** together with the
36 mixture of Mosher amides obtained from (±)-MMMA racemate.
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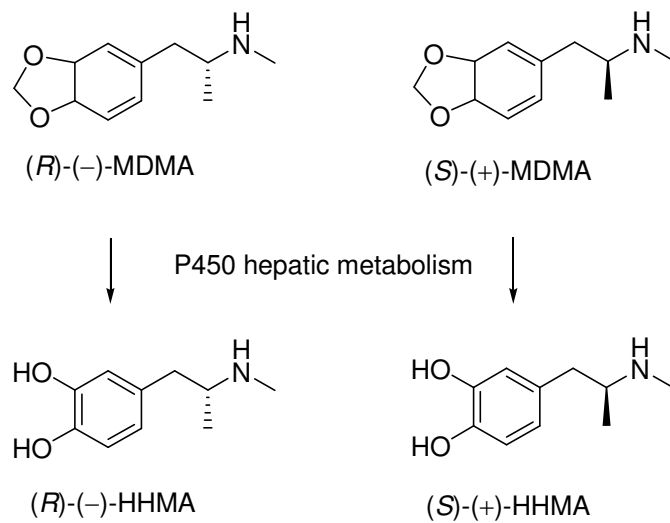
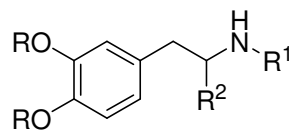


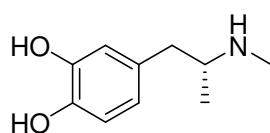
Figure 1



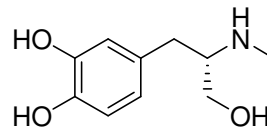
R = H, R¹ = R² = Me : HHMA

R = R¹ = R² = H : dopamine (DA)

R = R¹ = R² = Me : MMMA



R-(-)-HHMA **8**



Aminoalcohol **9**

Figure 2

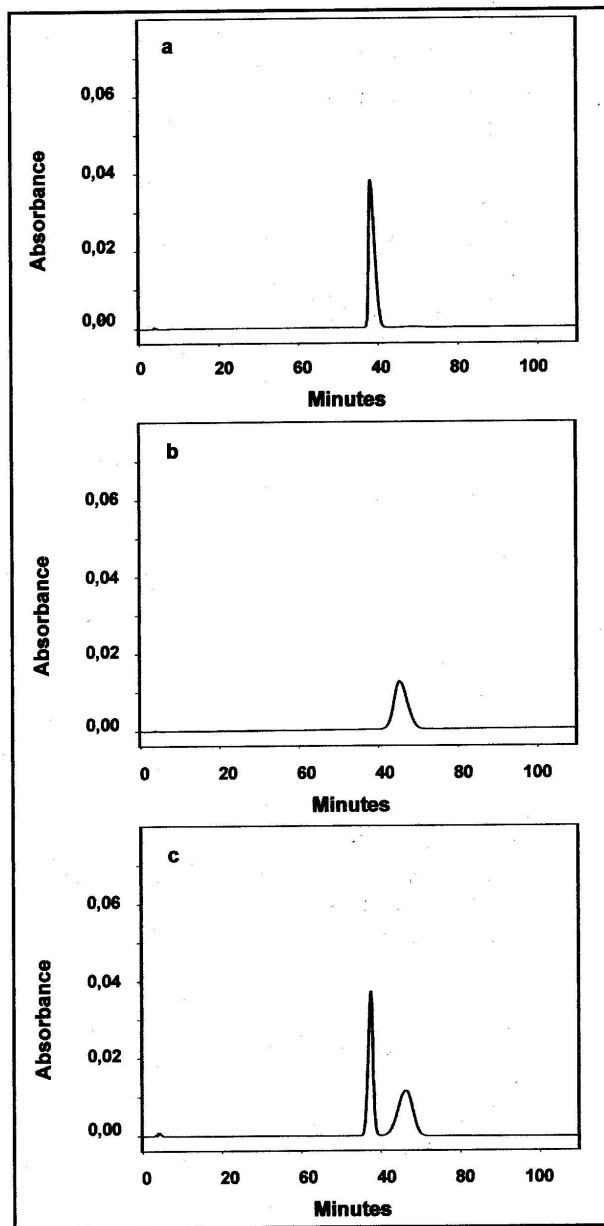


Figure 3

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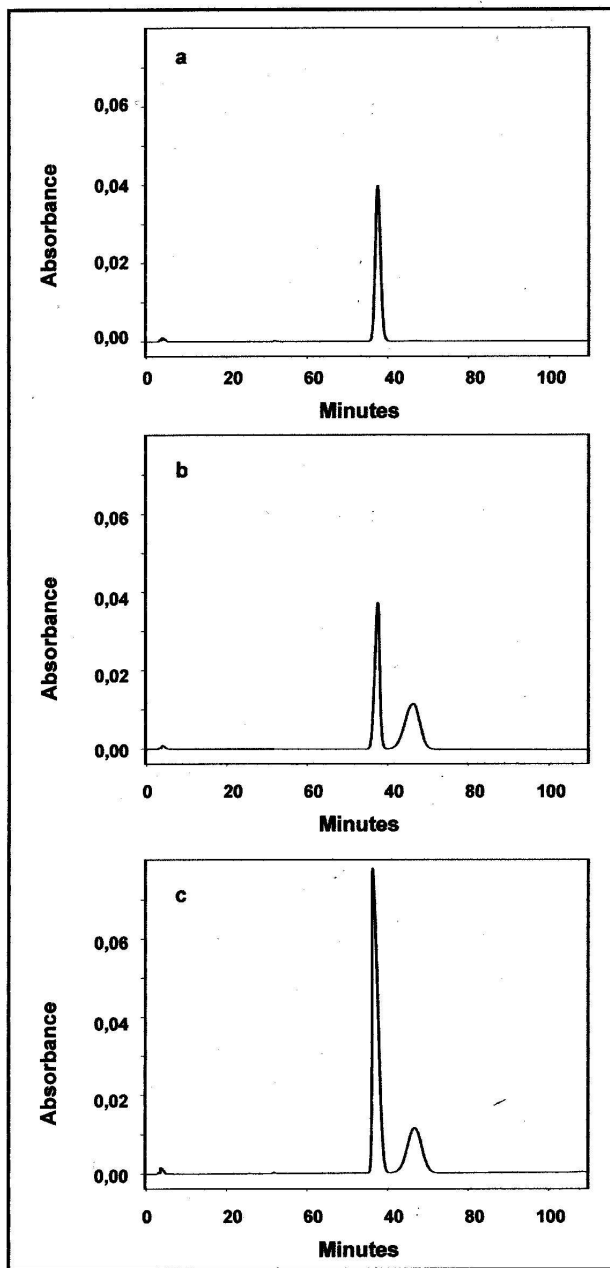


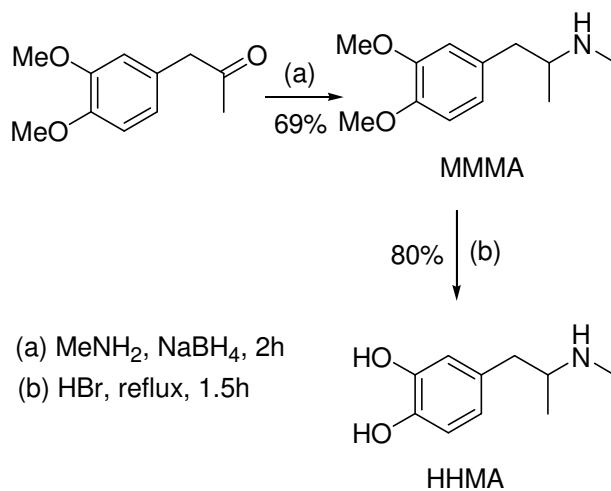
Figure 4

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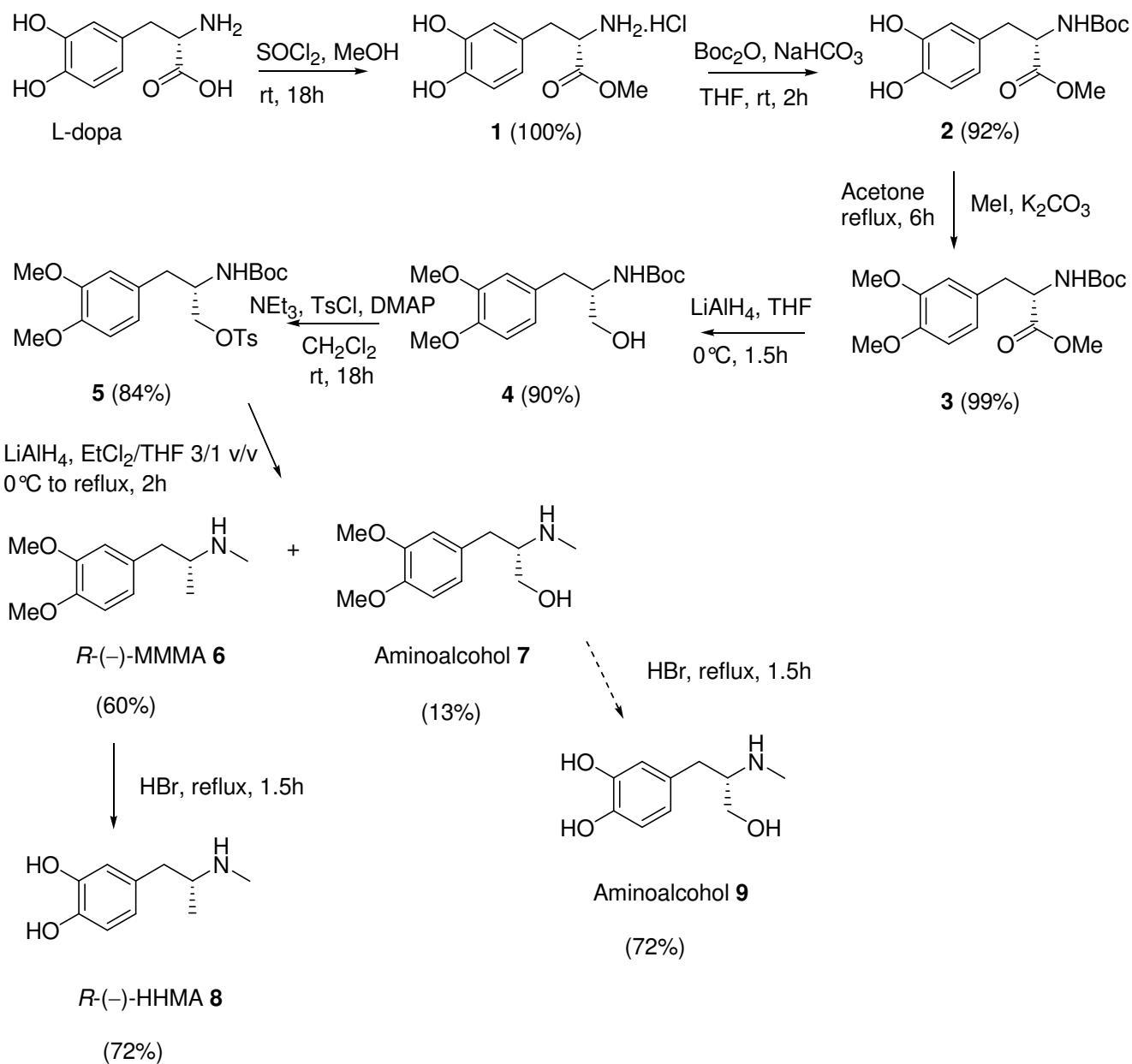
Scheme Legends

Scheme 1. Two step synthesis of (\pm)-HHMA

Scheme 2. Synthetic procedure for the total synthesis of *R*-(-)-HHMA



Scheme 1



Scheme 2