Tetrafluorobenzyl derivatives and pharmaceutical composition for preventing and treating acute and chronic neurodegenerative diseases in central nervous system containing the same
TETRAFLUOROBENZYL DERIVATIVES AND PHARMACEUTICAL COMPOSITION FOR PREVENTING AND TREATING ACUTE AND CHRONIC NEURODEGENERATIVE DISEASES IN CENTRAL NERVOUS SYSTEM CONTAINING THE SAME

Title:

TETRAFLUOROBENZYL DERIVATIVES AND PHARMACEUTICAL COMPOSITION FOR PREVENTING AND TREATING ACUTE AND CHRONIC NEURODEGENERATIVE DISEASES IN CENTRAL NERVOUS SYSTEM CONTAINING THE SAME

Abstract:
The present invention relates to a tetrafluorobenzyl derivative and a pharmaceutical composition for prevention and treatment of acute and chronic neurodegenerative diseases in central nervous system and ophthalmic diseases containing the same. The tetrafluorobenzyl derivative of the present invention can effectively be used to prevent and treat chronic neurodegenerative diseases such as Alzheimer's disease, Parkinson's disease and Huntington's disease, degenerative brain disease such as epilepsy and ischemic brain disease such as stroke.

Inventors/Applicants:


Applicant (for designated States except US):


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For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.
TECHNICAL FIELD

The present invention relates to a tetrafluorobenzyl derivative and a pharmaceutical composition comprising the same as a pharmaceutically effective ingredient and, more particularly, to a novel tetrafluorobenzyl derivative therapeutically effective for the treatment and prevention of neurological diseases and ocular diseases.

BACKGROUND ART

Recent advances in medicine have extended the life span of human beings and as a result, age-related acute and chronic neurological diseases, such as Alzheimer's disease, stroke, Parkinson's disease etc. increase. These neurological diseases are characterized by the progress of degeneration of specific neurons over the course of diseases. As mature neurons do not regenerate once they die, neuronal death in neurological diseases above can
result in incurable loss of essential brain function including cognition, sensation, and movement and thus economic and social overload.

Excitotoxicity, oxidative stress, and apoptosis have been implicated as major routes of neuronal death occurring in various neurological diseases and propagate through distinctive signaling pathways for each route.

Glutamate is the excitatory neurotransmitter mediating slow excitatory synaptic transmission through N-methyl-D-aspartate (NMDA) receptors and fast excitatory synaptic transmission through kainate or α-amino-3-hydroxy-5-methyl-4-isoxazolepropionic (AMPA) receptors. In the resting state of neurons, Mg\(^{2+}\) blocks NMDA receptor channels in a voltage-dependent manner. With stimuli causing membrane depolarization, Mg\(^{2+}\) is liberated from the NMDA channels, rendering the channels permeable to Ca\(^{2+}\) and Na\(^{+}\). Activation of NMDA receptors plays an important role in physiological process including learning and memory [Siegel G. J. et al., Basic Neurochemistry, 6\(^{th}\) edition, Lippincott Williams & Wilkins, 315 – 333 (1999)]. Besides physiological roles, brief and excess activation of NMDA receptors can cause rapidly evolving neuronal death and mechanisms underlying NMDA receptor-mediated neurotoxicity have been extensively studied over the
last two decades.

accumulation of glutamate causes neuronal cell death primarily through activation of NMDA receptors. In fact, administration of NMDA receptor antagonists have been reported to reduce neuronal death following hypoxic-ischemic brain injury [Goldberg, M. P. et al., J. Pharmac. Exp. Ther., 243:784-791(1987); Simon et al., Science 226:850-852 (1984); Sheardown, M. J. et al., Science 247:571-574(1990)].

Extensive evidence supports that excitotoxicity also contributes to neuronal death in neurodegenerative diseases. The key pathological features of Huntington's disease (HD) include degeneration of GABAergic neurons and selective sparing of NADPH diaphorase-containing neurons in the striatal area. These pathological features of HD are observed following the intrastriatal injections of NMDA or quinolinic acid, an NMDA receptor agonist [Ferrante, R. J et al., Science, 230(4625):561-563(1985); Beal, M. F. et al., Nature, 321(6066):168-171(1986); Koh, J.Y. et al., Science, 234(4772):73-76(1986)]. Amyotrophic lateral sclerosis (ALS) is accompanied by degeneration of upper and lower motor neurons and marked by neurogenic atrophy, weakness, and fasciculation. While the pathogenesis of ALS remains to be resolved, excitotoxicity has been expected to participate in the process of the ALS. In particular, ALS patients show defects in synthesis and transport of

Although NMDA receptor-mediated excitotoxicity plays a causative role in stroke and neurodegenerative diseases, the therapeutic potential of NMDA receptor antagonists has been limited by unexpected side effects in brain. In particular, systemic administration of NMDA receptor antagonists impairs normal brain function and can cause widespread neuronal damage in adult rat brain [Olney et al., Science 244:1360-1362 (1989)]. The neuropsychopathological side effects are produced by high-affinity NMDA receptor antagonists such as phencyclidine and related NMDA receptor antagonists such as MK-801 (dizocilpine maltate), tiletamine and ketamine and may be overcome with administration of channel-blocking NMDA receptor antagonists with low affinity and rapid-kinetic response [Rogawski, Amino Acids 19:133-149 (2000)].


Zinc (Zn²⁺) is a transition metal which is highly present and plays a dynamic role in brain. Within cells, zinc is associated with metalloproteins to control the enzymatic activity and structural stability of the proteins. Also, zinc regulates gene expression by binding to various transcription factors. In the CNS, zinc is localized at the synaptic terminal of glutamatergic neurons, released in an activity-dependent manner, and regulates activity of various neurotransmitter receptors and ion channels.

Zn²⁺ mediates neurodegenerative process observed in seizure, ischemia, trauma, and Alzheimers disease (AD). The central administration of kainate, a seizure-inducing excitotoxin, causes the translocation of Zn²⁺ into postsynaptic degenerating neurons in several forebrain areas. Translocation of zinc into adjacent neurons was also observed following ischemic and traumatic brain disease, and the blockade of its transition inhibited neuronal cell death [Frederickson, C. J. and Bush, A. I., Biomems. 14:353-366(2001);

Zinc has been known to enter neurons through Ca\(^{2+}\) permeable NMDA and AMPA/KA receptors, voltage-gated Ca\(^{2+}\) channel, or zinc transporter protein, and to induce neuronal death by the activation of NADPH oxidase generating reactive oxygen species. Zn\(^{2+}\) is observed in the extracellular plaque and degenerating neurons in AD, which likely contributes to neuronal degeneration in AD [Suh et al., *Brain Res.* 852:274-278 (2000); Bush et al., *Science* 265:1464-1467 (1994); Lee et al., *Proc. Natl. acad. Sci. U.S.A.* 99:7705-7710 (2002)]. Therefore, the inhibition of release and toxicity of zinc has been suggested as new strategy of prevention and treatment for Alzheimer's disease [Fredrickson and Bush, *Biometals*; 14:353-66 (2001)].

As described above, NMDA receptor-mediated excitotoxicity, oxidative stress, and zinc can contribute to neuronal death in various acute and neurodegenerative diseases in the nervous system. Thus, efficient therapeutic drugs preventing each route of neuronal deaths should be developed to treat such catastrophic neurological diseases.

We have investigated to develop neuroprotective drugs with multiple neuroprotective effects against
excitotoxicity or oxidative stress and succeeded in inventing tetrafluorobenzyl derivatives that can be applied to treat stroke, trauma, and some neurodegenerative diseases.

DISCLOSURE OF THE INVENTION

Throughout this specification and the claims which follow, unless the context requires otherwise, the word "comprise", and variations such as "comprises" and "comprising", will be understood to imply the inclusion of a stated integer or step or group of integers or steps but not the exclusion of any other integer or step or group of integers or steps.

The reference in this specification to any prior publication (or information derived from it), or to any matter which is known, is not, and should not be taken as an acknowledgment or admission or any form of suggestion that that prior publication (or information derived from it) or known matter forms part of the common general knowledge in the field of endeavour to which this specification relates.

Leading to the present invention, the intensive and thorough research on the treatment of disorders in the central nervous system, conducted by the present inventors, results in the finding that novel tetrafluorobenzyl derivatives have potent neuroprotective activity against various types of neuronal death induced in cell culture and animal models of neurological diseases.
Accordingly, it is a preferred object of the present invention to provide a novel tetrafluorobenzyl derivative.

It is another preferred object of the present invention to provide a pharmaceutically effective composition for the treatment and prevention of neurological diseases and ocular diseases.

In one aspect of the present invention, there is provided a novel tetrafluorobenzyl derivative, represented by the following chemical formula 1,

![Chemical Formula 1]

wherein,

- $R_1$, $R_2$, and $R_3$ are hydrogen or halogen;
- $R_4$ is hydroxy, alkyl, alkoxy, halogen, alkoxy substituted with halogen, alkanoyloxy or nitro;
- $R_5$ is carboxylic acid, ester having $C_1$-$C_4$ alkyl, carboxyamide, sulfonic acid, halogen, or nitro;
- or a pharmaceutically acceptable salt thereof.
In another aspect of the invention there is provided a pharmaceutical composition comprising a tetrafluorobenzyl derivative or its pharmaceutically acceptable salt as referred to above, and a pharmaceutically acceptable carrier.

In another aspect of the present invention, there is provided a method for the treatment or prevention of neurological diseases and ocular diseases in a patient, comprising administering to the patient the tetrafluorobenzyl derivative or its pharmaceutically acceptable salt as an effective ingredient.

In a further aspect of the invention there is provided use of the tetrafluorobenzyl derivative or its pharmaceutically acceptable salt in the manufacture of a medicament for treatment or prevention of neurological diseases and ocular diseases.

The present invention provides a novel tetrafluorobenzyl derivative represented by the following chemical formula 1, or its pharmaceutically acceptable salt.

[Chemical Formula 1]
wherein, \( R_1 \), \( R_2 \), and \( R_3 \) are hydrogen or halogen

\( R_4 \) is hydroxy, alkyl, alkoxy, halogen, alkoxy substituted with halogen, alkanoyloxy or nitro;

\( R_5 \) is carboxylic acid, ester having \( C_1-C_4 \) alkyl, carboxyamide, sulfonic acid, halogen, or nitro;

In here, alkyl group is \( C_1-C_4 \) alkyl and more preferably \( C_1-C_2 \) alkyl. Alkyl described above definitely contains methyl, ethyl, propyl, isopropyl, n-butyl, sec-butyl, or tert-butyl.

Alkoxy group is \( C_1-C_4 \) and more preferably \( C_1-C_2 \) alkoxy. Alkoxy described above definitely contains methoxy, ethoxy, or propanoxy.

Halogen can be substituted with fluoride, chloride, bromide, or iodide.

Alkanoyloxy is \( C_2-C_{10} \) alkanoyloxy and more preferably \( C_3-C_5 \) alkanoyloxy. Alkanoyloxy described above definitely contains ethanoyloxy, propanoyloxy, or cyclohexanecarbonyloxy.

In ester having \( C_1-C_4 \) alkyl, \( C_1-C_4 \) alkyl can be methyl, ethyl, isopropyl, or butyl.
Specific compounds of interest within Formula I are as follows:

2-Hydroxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)-benzoic acid (hereinafter, referred to as '2-Hydroxy-TTBA'),

2-Nitro-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)benzoic acid,

2-Chloro-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)benzoic acid,

2-Bromo-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)benzoic acid,

2-Hydroxy-5-(2,3,5,6-tetrafluoro-4-methylbenzylamino)benzoic acid,

2-Methyl-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)benzoic acid,

2-Methoxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)benzoic acid,

5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)-2-trifluoromethoxy benzoic acid.

2-Nitro-4-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)phenol,

2-Chloro-4-(2,3,5,6-tetrafluoro-4-
trifluoromethylbenzylamino)phenol (hereinafter, referred to as '2-Chloro-TTP'),

2-Hydroxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)benzamide (hereinafter, referred to as '2-Hydroxy-TTA'),

2-Hydroxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)benzenesulfonic acid (hereinafter, referred to as '2-Hydroxy-TTS'),

Methyl 2-hydroxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)benzoate,

2-Ethanyloxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)benzoic acid (hereinafter, referred to as '2-Ethan-TTBA'),

2-Propanoyloxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)benzoic acid (hereinafter, referred to as '2-Propan-TTBA'), or

2-Cyclohexanecarbonyloxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)benzoic acid (hereinafter, referred to as '2-Cyclohexan-TTBA').

However, the compounds described above are just representative of the present invention, which could include more compounds.

The present invention provides tetrafluorobenzyl derivatives represented by formula (I) and
pharmaceutical composition for prevention and treatment of neurological and ocular diseases containing the effective component as a pharmaceutically acceptable salt.

As drugs for pharmaceutical use, the salt of the compound of formula (I) doesn't have toxicity, and should be pharmaceutically acceptable. Various kinds of salts can be used to prepare pharmaceutically acceptable salts, including non-toxic compound of the present invention.

The pharmaceutically acceptable salts of the compounds in the present invention include alkali metals, such as lithium, sodium or potassium, and alkaline earth metals, such as calcium or magnesium. Acid addition salts may be prepared by reacting the solution of pharmaceutically acceptable nontoxic salts such as hydrochloric acid, fumaric acid, maleic acid, succinic acid, acetic acid, citric acid, tartaric acid, carbonic acid, or phosphoric acid with the compound of the invention.

The compound of formula (I) in the present invention can be used for cure of normal and pathological neurodegenerative diseases among diseases or symptoms in cerebrovascular and neurological systems. In the concrete, the compound above represented by formula (I) is used for prevention or treatment of
thromboembolism, ischemic stroke, hemorrhagic stroke, cerebrovascular convulsion, brain aging, traumatic brain injury, traumatic spinal cord injury, cardiac arrest, arterial hypotention, hypoglycemia, anoxia, and hypoxia. Also, the compound of formula (I) in the present invention can be beneficially used for decreasing neurodegenerative diseases such as Huntington's disease, Alzheimer's disease, senile dementia, Pick's disease, Korsakov's syndrome, olivopontocerebellar degeneration, amyotrophic lateral sclerosis (ALS), Parkinson's disease, Down's syndrome, Glutaric acidemia, epilepsy, multi-infarct dementia, and brain inflammation. They have application to treatment of ocular diseases such as glaucoma, macular degeneration, diabetic retinopathy, uveitis. Moreover, they have application to the prevention and treatment of drug addiction, depression, and pain.

The composition of the present invention can be treated by oral administration, intravenous injection or non-oral administration, and treated by various forms such as tablet, capsule, powder, grain, sterilized solution, suspension or suppository for rectal administration. Major effective elements of the composition can be made as a solid tablet using pharmaceutical carriers, for example common tablet element such as corn dextrin, lactose, sucrose,
sorbitol, talc, stearic acid, magnesium stearate, decalcium phosphate or gums, and additional pharmaceutical diluted solution. Tablets or pillets of the pharmaceutical composition in the present invention can be manufactured for sustained release dosage form as facilitated forms for administration using well-known coating method etc. in the appropriate industry. For example, tablets or pillets can be composed with inner and outer administrative elements. The inner administrative elements of tablets or pillets can be manufactured as wrapped with outer administrative elements. Liquid forms of the composition in the present invention manufactured for oral administration or the injection include solution, appropriately flavored syrup, water-soluble suspension, water-insoluble suspension, emulsion made by edible oil such as cotton oil, sesame oil, coconut oil, or peanut oil, elixir, and similar pharmaceutical carriers. Tragacanth gum, acacia, alginic acid sodium salt, dextran, sodium carboxymethylcellulose, methylcellulose, polyvinylpyrrolidone, or synthesized or natural gums like gelatin etc can be used as appropriated aid to dispersion or suspension in making water-soluble suspension.

Quantity of medication can be determined by several related factors such as diseases, age, sex,
weight, and degrees of illness of patients etc. for the treatment of neurodegeneration.

The tetrafluorobenzyl derivatives related to the present invention may be synthesized from the following reaction schemes. However, the compounds described in the schemes are just representative of the present invention, which could include more compounds.

<Reaction scheme 1>

\[
\begin{array}{c}
\text{O}_2\text{N} - \text{R}_5 - \text{R}_4 \\
\downarrow^a \\
\text{R}_2 \text{F}_3 \text{F}_2 \text{CH}_2 \text{Br} + \text{NH}_2 - \text{R}_5 - \text{R}_4 \rightarrow \text{R}_3 \text{C}_2 \text{F}_2 \text{CH}_2 \text{NH} - \text{R}_5 - \text{R}_4
\end{array}
\]

The tetrafluorobenzyl derivatives were synthesized from the following reaction. First, the nitrobenzene compounds, where hydrogens at 3 and 4 positions were substituted with \( R_5 \) and \( R_4 \) respectively, were hydrogenated for 12 hours under 3 atm pressure (reaction condition \( a \)). The resulting aniline compounds were reacted with 2,3,5,6-tetrafluoro-4-methylbenzyl bromide in DMF in the presence of
triethylamine for 12 hours to give the desired tetrafluorobenzyl derivatives (reaction condition b).

In the above scheme, \( R_1, R_2, \) and \( R_3 \) represent hydrogen or halogen; \( R_4 \) represents hydroxy, alkyl, alkoxy, halogen, alkoxy substituted with halogen, alkanoyloxy, nitro; \( R_5 \) represents carboxylic acid, esters of carboxylic acid, carboxyamide, sulfonic acid, halogen, and nitro group.

For the synthesis of compound where \( R_4 \) is hydroxy group and \( R_5 \) is carboxyamide group, the hydroxy and amino group in 5-(2,3,5,6-tetrafluoro-4-trifluorobenzylamino)benzoic acid were first protected with trifluoro group by reacting with trifluoromethyl acetic anhydride in the catalytic amount of \( c-H_2SO_4 \) (reaction condition a). The resulting 2-carboxy-5-[2,3,5,6-tetrafluoro-4-trifluoromethyl-benzyl]-(2,2,2-trifluoroacetyl)amino)phenyl ester compound was then reacted with \( SOCl_2 \) followed by ammonium carbonate to
give 2-carbamoyl-4-[2,3,5,6-tetrafluoro-4-
trifluoromethyl-benzyl-(2,2,2-
trifluoroacetyl)amino]phenyl ester compound (reaction
condition b), which was then hydrolyzed with HCl
solution to give the desired carboxyamide compound.
(reaction condition c) In this scheme, \( R_1, R_2, R_3 \)
and \( R_4 \) are same groups, which are previously defined while \( R_5 \)
are \( C_1-C_4 \) substituted alkyl group.

<Synthesis Example 1>

Preparation of 2-hydroxy-5-(2,3,5,6-tetrafluoro-4-
trifluoromethylbenzylamino)benzoic acid (2-Hydroxy-
TTBA)

To a solution of 5-aminosalicylic acid (1.02 g,
6.66 mmole, purchased from Aldrich Chemical Company,
USA, A7, 980-9) and triethylamine (1 ml) in dried DMF
(30 ml) was added 2,3,5,6-tetrafluoro-4-
trifluoromethylbenzyl bromide (1.23 g, 7.18
mmole)(Aldrich, 40, 640-6) at room temperature under a
nitrogen atmosphere. The reaction mixture was stirred
for 2 hr at room temperature and then solvent was
removed in vacuo. The reaction mixture was diluted
with ethyl acetate and then extracted with ethyl
acetate. The organic layer was washed with water and
brine, and then dried over anhydrous MgSO\(_4\). After
evaporation of the solvent, the residue was
recrystallized from ether/hexane (1:10) to give 1.60 g (64% yield) of 2-hydroxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)benzoic acid as a white solid.

\[ \text{mp} \ 179 \degree C, \]

\[ {^1H-}\text{NMR} : \ 8.0 \ (s, 1H), \ 7.3 \ (d, 1H), \ 6.7 \ (t, 1H), \ 5.5 \ (s, 2H), \]

\[ \text{IR (KBr pellet)} : \ 3386, \ 1741, \ 1500 \text{ cm}^{-1} \]

Elemental analysis for C\textsubscript{15}H\textsubscript{14}F\textsubscript{7}N\textsubscript{2}O\textsubscript{3}

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<Synthesis Example 2>

Preparation of 2-nitro-5-(2,3,5,6-tetrafluoro-4-trifluoromethyl benzylamino)benzoic acid

According to the similar procedure in Synthesis Example 1, by using 5-amino-2-nitrobenzoic acid (1.03 g, 5.65 mmole) and 2,3,5,6-tetrafluoro-4-trifloromethylbenzyl bromide (1.01 g, 6.04 mmole) (ACROS, 33074-0010), 1.50 g (76.3% yield) of 2-nitro-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)benzoic acid was obtained as a pale yellow solid.

\[ \text{mp} \ 121\degree C, \]
**Synthesis Example 3**

Preparation of 2-chloro-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)benzoic acid

According to the similar procedure in Synthesis Example 1, by using 5-amino-2-chlorobenzoic acid (1.02 g, 5.94 mmole)(ACROS, 32525-5000), DMF (50 ml) and 2,3,5,6-tetrafluoro-4-trifluoromethylbenzyl bromide (1.16 g, 6.99 mmole), 2.04 g (85.5% yield) of 2-chloro-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)benzoic acid was obtained as a pale brown solid.

mp 58°C,

\[ ^1H\text{-NMR} : 7.26 (d, 1H), 7.24 (s, 1H), 6.7 (d, 1H), 4.12 (s, 2H), \]

IR(KBr pellet): 3402, 1720, 1494, 1434, 929 cm\(^{-1}\)

HPLC (0.01% TFA-ethyl acetate : 0.1 TFA-water = 80 : 20, Rt=4.2 mins): 97% purity

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<Synthesis Example 4>

Preparation of 2-bromo-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)benzoic acid

(4-1) Preparation of 5-amino-2-bromobenzoic acid

A mixture of 2-bromo-5-nitrobenzoic acid (1.10 g, 4.06 mmol)(Aldrich, 38, 184-5), activated Pd-C (43.62 mg, 0.41 mmol)(Aldrich, 20, 569-9) in methanol (30 ml) was hydrogenated for 4 hr under 30 psi of hydrogen pressure. After the mixture was filtered, the filtrate was concentrated to give 0.80 g (91.2% yield) of 5-amino-2-bromobenzoic acid as a pale yellow solid.

IR(KBr pellet): 3111, 2558, 2499, 1716, 1676 cm⁻¹

(4-2) Preparation of 2-bromo-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)benzoic acid

According to the similar procedure in Synthesis Example 1, by using 5-amino-2-bromobenzoic acid (1.05 g, 6.12 mmole) which was prepared from Synthesis Example (4-1), DMF (50 ml) and 2,3,5,6-tetrafluoro-4-trifloromethylbenzyl bromide (1.16 g, 6.99 mmole), 2.02 g (76.9% yield) of 2-bromo-5-(2,3,5,6-tetrafluoro-4-
trifluoromethylbenzylamino)benzoic acid was obtained as a pale yellow solid.

mp 70°C,

$^1$H-NMR : 7.42 (d, 1H), 7.3 (s, 1H), 6.9 (d, 1H), 5.5 (s, 2H),

IR (KBr pellet) : 3438, 1695, 1491, 1425, 939 cm$^{-1}$

HPLC (0.1% TFA-ethyl acetate : 0.1 % TFA-water = 80 : 20, Rt= 4.5 mins) 99% purity

Elemental analysis for C$_{15}$H$_{12}$BrF$_3$NO$_2$

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<Synthesis Example 5>

Preparation of 2-hydroxy-5-(2,3,5,6-tetrafluoro-4-methylbenzylamino)benzoic acid

According to the similar procedure in Synthesis Example 1, by using 4-methyl-2,3,5,6-tetrafluoro-benzyl bromide (1.23 g, 7.18 mmole) (Aldrich, 40, 646-6) instead of 2,3,5,6-tetrafluoro-4-trifluoromethyl benzyl bromide, 1.60 g (64.0% yield) of 2-hydroxy-5-(2,3,5,6-tetrafluoro-4-methylbenzylamino)benzoic acid was obtained as a white solid.

mp 212°C,

$^1$H-NMR : 8.0 (s, 1H), 7.3 (d, 1H), 6.7 (t, 1H), 5.5 (s, 2H), 2.2-2.3 (s, 3H),
IR(KBr pellet) : 3386, 1741, 1500 cm\(^{-1}\)

Elemental analysis for C\(_{14}\)H\(_{15}\)F\(_4\)N\(_8\)O\(_3\)

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<Synthesis Example 6>

Preparation of 2-methyl-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)benzoic acid

(6-1) Preparation of 2-methyl-5-nitrobenzoic acid

To a solution of 2-methylbenzoic acid (3.05 g, 22.3 mmole)(ACROS, 13904-0010) in c-HNO\(_3\) (20 ml) was carefully added Conc.H\(_2\)SO\(_4\) (15 ml) at 0 °C. The resulting solution was refluxed at 100 - 120 °C for 5 hours. After the reaction mixture was cooled to room temperature, 50 ml of ice chip was added. The resulting precipitate was filtered, washed with water and dried to give 3.90 g (97.5% yield) of 2-methyl-5-nitrobenzoic acid as a white solid.

IR(KBr pellet) : 1531, 1350 cm\(^{-1}\)

(6-2) Preparation of 5-Amino-2-methylbenzoic acid

According to the similar procedure in Synthesis Example (4-1), by using 2-methyl-5-nitrobenzoic acid (4.10 g, 22.6 mmole) which was prepared from Synthesis
Example (6-1), 2.01 g (60.0% yield) of 5-amino-2-methylbenzoic acid was obtained as a white solid.

\[ \text{IR (KBr pellet): } 3437, 3336, 1633, 1400 \text{ cm}^{-1} \]

(6-3) Preparation of 2-methyl-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)benzoic acid

According to the similar procedure in Synthesis Example 1, by using 5-amino-2-methylbenzoic acid (2.06 g, 15.0 mmole), DMF (60 ml), TEA (4 ml), and 2,3,5,6-tetrafluoro-4-trifluoromethylbenzyl bromide (5.52 ml, 17.7 mmole), 1.50 g (27.0% yield) of 2-methyl-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)benzoic acid was obtained as a yellow solid.

\[ \text{mp } 84^\circ C, \]

\[ \text{\textsuperscript{1}H-NMR: } 7.32(s, 1H), 7.3(d, 1H), 6.9(d, 1H), 5.5(s, 2H), 2.2(s, 3H), \]
\[ \text{IR (KBr pellet): } 3417, 1716, 1496 \text{ cm}^{-1} \]

Elemental analysis for C\textsubscript{15}H\textsubscript{10}F\textsubscript{5}NO\textsubscript{2}

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<Synthesis Example 7>

Preparation of 2-methoxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)benzoic acid
According to the similar procedure in Synthesis Example (6-1), by using 2-methoxybenzoic acid (2.10 g, 13.79 mmole) (ACROS, 17375-2000), 2.50 g (97.0% yield) of 2-methoxy-5-nitrobenzoic acid was obtained as a white solid.

IR (KBr pellet): 3099, 2986, 2965, 1736, 1547 cm\(^{-1}\)

According to the similar procedure in Synthesis Example (4-1), by using 2-methoxy-5-nitrobenzoic acid (4.10 g, 22.6 mmole) which was prepared from Synthesis Example (7-1), 1.90 g (98.0% yield) of 5-amino-2-methoxybenzoic acid was obtained as a brown solid.

IR (KBr pellet): 1394, 1220 cm\(^{-1}\)

According to the similar procedure in Synthesis Example 1, by using 5-amino-2-methoxy-benzoic acid (2.10 g, 12.7 mmole), DMF (50 ml), TEA (6 ml), and 2,3,5,6-tetrafluoro-4-trifluoromethylbenzyl bromide (2.00 ml, 14.0 mmole), 1.50 g (31.5% yield) of 2-methoxy-5-(2,3,5,6-tetrafluoro-4-
trifluoromethylbenzylamino)benzoic acid was obtained as a yellow solid:

mp 94°C,

'H-NMR 7.42(s, 1H), 6.4(d, 1H), 6.2(d, 1H), 5.5(s, 2H), 3.73(s, 3H),

IR(KBr pellet): 3429, 1730, 1496 cm⁻¹

Elemental analysis for C₁₆H₁₀F₇NO₃

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<Synthesis Example 8>

Preparation of 5-(2,3,5,6-tetrafluoro-4-trifluoromethyl-benzylamino)-2-trifluoromethoxybenzoic acid

(8-1) Preparation of 5-nitro-2-trifluoromethoxybenzoic acid

According to the similar procedure in Synthesis Example (6-1) (Lancaster, 15687), by using 2-trifluoromethoxybenzoic acid (2.10 g, 9.81 mmole), 1.40 g (55.0% yield) of 5-nitro-2-trifluoromethoxybenzoic acid was obtained as a white solid.

IR(KBr pellet): 1488, 1354 cm⁻¹
(3-2) Preparation of 5-amino-2-trifluoromethoxybenzoic acid

According to the similar procedure in Synthesis Example (4-1), by using 5-nitro-2-trifluoromethoxybenzoic acid (1.40 g, 5.34 mmole) which was prepared from Synthesis Example (8-1), 1.02 g (90.0% yield) of 5-amino-2-trifluoromethoxybenzoic acid was obtained as a pale yellow solid.

IR(KBr pellet) : 1627, 1369 cm⁻¹

(3-3) Preparation of 5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)-2-trifluoromethoxybenzoic acid

According to the similar procedure in Synthesis Example 1, by using 5-amino-2-trifluoromethoxybenzoic acid (1.35 g, 6.23 mmole), DMF (50 ml), TEA (6 ml), and 2,3,5,6-tetrafluoro-4-trifluoromethylbenzyl bromide (1.10 ml, 6.73 mmole), 2.25 g (81.0% yield) of 5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)-2-trifluoromethoxybenzoic acid was obtained as a yellow solid. :

mp 38°C,

¹H-NMR 7.3 (s, 1H), 7.12 (d, 1H), 6.8 (d, 1H), 5.5 (s, 2H),

IR(KBr pellet): 3383, 1712, 1504, 1446, 929 cm⁻¹
Elemental analysis for C₁₄H₁₂F₁₀N₂O₃

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<Synthesis Example 9>

Preparation of 2-nitro-4-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)phenol

According to the similar procedure in Synthesis Example 1, by using 4-amino-2-nitrophenol (1.00 g, 6.49 mmole), DMF (30 ml), TEA (0.5 ml), and 2,3,5,6-tetrafluoro-4-trifluoromethylbenzyl bromide (1.30 g, 7.78 mmole), 1.00 g (40.0% yield) of 2-nitro-4-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)phenol was obtained as a reddish solid:

- mp 126-128°C,
- $^1$H-NMR(DMSO-$d_6$) δ 4.23 (d, 2H), 6.93 (m, 2H), 7.12 (s, 1H),
- $^{13}$C NMR(DMSO-$d_6$) δ 36.36, 105.48, 120.63, 122.64, 123.32, 136.17, 140.86, 142.34, 144.06, 144.83, 146.44, 151.23,
- IR(neat): 3391, 3255, 1545, 1339 cm$^{-1}$

Elemental analysis for C₁₄H₁₂F₁₅N₂O₃

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<Synthesis Example 10>

Preparation of 2-chloro-4-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)phenol

According to the similar procedure in Synthesis Example 1, by using 4-amino-2-chlorophenol (3.00 g, 19.6 mmole), DMF (80 ml), TEA (0.5 ml), and 2,3,5,6-tetrafluoro-4-trifluoromethylbenzyl bromide (1.40 g, 8.36 mmole), 2.00 g (76.0% yield) of 2-chloro-5-(2,3,5,6-tetrafluoro-4-methylbenzylamino)phenol was obtained as a yellow solid.

mp 52°C,

\[ \text{H-NMR(CDC}_1\text{)} \ 6.9 \ (d, 1H), 6.7 \ (s, 1H), 6.5 \ (d, 1H), 4.4 \ (s, 2H), \]

IR(KBr pellet) 3382, 1687, 1617, 1586 cm\(^{-1}\)

Elemental analysis for C\(_{16}\)H\(_{18}\)F\(_7\)N\(_2\)O\(_3\)

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<Synthesis Example 11>

Preparation of 2-hydroxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)benzamido
Preparation of 2-carboxy-4-[2,3,5,6-tetrafluoro-4-trifluoromethyl-benzyl]-(2,2,2-trifluoroacetyl) amino]phenyl trifluoroacetate

To a solution of 2-hydroxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino) benzoic acid (2.00 g, 5.12 mmole) and trifluoroactic anhydride (15 ml) was added c-$\mathrm{H}_2\mathrm{SO}_4$ (0.50 ml) at 10°C under a nitrogen atmosphere. The reaction mixture was stirred for 20 min at room temperature and then quenched with an ice (10 g). After the solvent was removed in vacuo, the residue was dissolved in ethyl acetate (50 ml). The organic layer was washed with water (20 ml x 2), 10% NaHCO$_3$ (20 ml x 3), 0.5 N HCL (20 ml x 2), and water (20 ml). The organic layer was dried over anhydrous Na$_2$SO$_4$. After evaporation of the solvent, the residue was recrystallized from ethyl acetate/hexane (1:10) to give 1.40 g (47% yield) of 2-carboxy-4-[2,3,5,6-tetrafluoro-4-trifluoromethylbenzyl]-(2,2,2-trifluoroacetyl)amino]phenyl trifluoroacetate as an yellow solid.

mp 174 -180°C,

$^1$H-NMR(DMSO-d$_6$)  δ 5.16(d, 2H), 6.94(d, 1H), 7.44(d, 1H), 7.81(s, 1H),

$^{13}$C-NMR(DMSO-d$_6$)  δ  43.03, 113.2, 114.3, 117.2, 117.8, 118.9, 128.7, 130.2, 135.5, 141.8, 143.8, 144.3, 146.2, 146.3, 155.2, 155.5, 161.6, 170.5,
IR (neat): 1711, 1673, 1498, 1331, 724 cm⁻¹

(11-2) Preparation of 2-carbamoyl-4-[2,3,5,6-tetrafluoro-4-trifluoromethylbenzyl-(2,2,2-trifluoroacetyl)amino]phenyl trifluoroacetate

To a solution of 2-carboxy-4-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzyl-(2,2,2-trifluoroacetyl)amino)phenyl trifluoroacetate (600 mg, 1.05 mmole) in anhydrous methylene chloride (15 ml) was added SOCl₂ (1.18 ml, 21.0 mmole) at 40°C under a nitrogen atmosphere. After the reaction mixture was stirred for 1 hr, the solvent was removed in vacuo. The resulting residue was dissolved in anhydrous methylene chloride (60 ml) and added with ammonium carbonate (assay > 30%, 2.00 g). After the reaction mixture was stirred for 1 hr, it was filtered to remove the remained ammonium carbonate. The organic layer was washed with water (40 ml x 3) and then dried over anhydrous Na₂SO₄. After evaporation of the solvent, the residue was recrystallized from methylene chloride/hexane (1:10) to give 370 mg (61% yield) of 2-carbamoyl-4-[2,3,5,6-tetrafluoro-4-trifluoromethylbenzyl-(2,2,2-trifluoroacetyl)amino]phenyl trifluoroacetate as a white solid. mp 179 -180°C,
Preparation of 2-hydroxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino) benzamide

To a solution of 2-carbamoyl-4-[2,3,5,6-tetrafluoro-4-trifluoromethylbenzyl-(2,2,2-trifluororacetyl)amino]phenyl trifluoroacetate (300 mg, 0.52 mmole) in methanol (8 ml) and water (3 ml) was added c-HCl (2.0 ml) at 40°C under a nitrogen atmosphere. After the reaction mixture was stirred for 24 hr, the organic solvent was removed in vacuo. The residue was extracted with ethyl acetate (20 ml x 3). The organic layer was washed with water and then dried over anhydrous Na₂SO₄. After evaporation of the solvent, the residue was recrystallized from methylene chloride/hexane (1:10) to give 120 mg (60% yield) of 2-hydroxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)benzamide as a white solid. mp 143 -145°C,

\[^1\text{H-}\text{NMR(DMSO-}\text{d}_6\text{)} \delta \ 4.37(\text{s, 2H}), \ 6.63(\text{d, 1H}), \ 6.76(\text{d, 1H}), \ 7.14(\text{s, 1H}), \]
<Synthesis Example 12>
Preparation of 2-hydroxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)benzene sulfonic acid

According to the similar procedure in Synthesis Example 1, by using 4-amino-2-hydroxybenzene sulfonic acid (1.00 g, 5.30 mmole), DMF (10 ml), TEA (1.0 ml), and 2,3,5,6-tetrafluoro-4-trifluoromethylbenzyl bromide (0.88 g, 5.30 mmole), 0.61 g (28.0% yield) of 2-hydroxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)benzene sulfonic acid was obtained as a yellow solid:

mp: above 300°C,

$^1$H-NMR (DMSO-d$_6$) $\delta$ 4.28 (s, 2H), 5.58 (m, 2H), 6.82 (s, 1H),

$^{13}$C NMR (DMSO-d$_6$) $\delta$ 32.08, 106.41, 111.39, 112.10, 119.15, 119.33, 119.51, 126.11, 135.09, 137.17, 139.17, 139.70, 139.89, 140.40, 140.59, 141.61,

IR (neat) 3427, 3227, 1492, 1331, 1196, 1135, 628 cm$^{-1}$

<Synthesis Example 13>
Preparation of methyl 2-hydroxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino) benzoate

(13-1) Preparation of methyl 5-amino-2-hydroxybenzoate

To a solution of 5-aminosalicylic acid (3.00g, 19.6 mmole) in methanol (80 ml) was added c-H₂SO₄ (8 ml) at 0°C. After the reaction mixture was refluxed for 6 hr, the solvent was removed in vacuo. The resulting residue was partitioned with ethyl acetate and water. The organic layer was washed with water (40 ml x 3) and then dried over anhydrous MgSO₄. After evaporation of the solvent, the residue was recrystallized from ethyl acetate/hexane to give 2.50 g (76% yield) of methyl 5-amino-2-hydroxybenzoate as a yellow solid.

\[
\text{IR (KBr pellet)} 3406, 3327, 2950, 1672, 1616, 1492, 1440, 788 \text{ cm}^{-1}
\]

(13-2) Preparation of methyl 2-hydroxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino) benzoate

According to the similar procedure in Synthesis Example 1, by using methyl 5-amino-2-hydroxybenzoate (2.00 g, 11.9 mmole), DMF (60 ml), TEA (0.5 ml), and 2,3,5,6-tetrafluoro-4-methylbenzyl bromide (2.60 g, 14.3 mmole), 3.20 g (85.0% yield) of methyl 2-
hydroxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)benzoate was obtained as a pale yellow solid:

\[ \text{mp: 127}^\circ\text{C,} \]

\[ \text{H-NMR(CDC}_3\text{)}\delta 7.15(\text{s, 1H}), 6.9(\text{d, 1H}), 6.7(\text{d, 1H}), 4.5(\text{s, 2H}), 3.9(\text{s, 3H}), \]

\[ \text{IR(KBr pellet) 3382, 1687, 1617, 1586 cm}^{-1} \]

Elemental analysis for \( \text{C}_{16}\text{H}_3\text{F}_{7}\text{N}_0\text{O}_3 \)

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<Synthesis Example 14>

Preparation of 2-ethanoyloxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethyl-benzylamino)benzoic acid

(14-1) Preparation of 5-tert-butoxycarbonylamino-2-hydroxybenzoic acid

The mixture of 5-aminosalicylic acid (1.01 g, 6.59 mmole), Di-BOC (2.87 g, 13.1 mmol), TEA (1.0 ml) in DMF (20.0 ml) was stirred for 2 hr at room temperature. After the reaction mixture was concentrated, the residue was dissolved in ethyl acetate. The organic layer was washed with water and dried over anhydrous \( \text{Na}_2\text{SO}_4 \). After evaporation of the
solvent, the residue was recrystallized from ethyl acetate/hexane to give 1.42 g (84% yield) of 5-tert-butoxy-2-hydroxybenzoic acid as a white solid.

mp: 282°C,

\[ \text{H-NMR (DMSO-d}_6) \delta 7.9-8.0 (s, 1H), 7.4-7.5 (d, 1H), 6.8-6.9 (d, 1H), 1.4-1.6 (s, 9H), \]

\[ \text{C NMR (DMSO-d}_6) \delta 171.93, 156.50, 152.97, 131.06, 126.36, 119.39, 117.03, 113.13, 79.05, 28.42 \]

(14-2) Preparation of 2-ethanoyloxy-5-tert-butoxycarbonylaminoacetic acid

To a solution of 5-tert-butoxycarbonylamino-2-hydroxybenzoic acid (1.02 g, 4.02 mmole) in DMF (20.0 ml) was added with acetyl chloride (39 mg, 0.483 mmole) and potassium carbonate (555 mg, 4.02 mmole). The reaction mixture was stirred for 4 hr at room temperature. After the reaction mixture was concentrated, the residue was dissolved in ethyl acetate. The organic layer was washed with water and brine and dried over anhydrous Na\(_2\)SO\(_4\). After evaporation of the solvent, the residue was recrystallized from ethyl acetate/hexane to give 0.62 g (52% yield) of 2-ethanoyloxy-5-tert-butoxycarbonylaminoacetic acid as a white solid.

mp: 76-78°C,
**Preparation of 2-ethanoyloxy-5-aminobenzoic acid**

The mixture of 2-ethanoyloxy-5-tert-butoxycarbonylaminobenzoic acid (1.01 g, 3.42 mmole) in TFA/CH$_2$Cl$_2$ (20.0 ml, 1:1 v/v) was stirred for 30 min at room temperature. After the reaction mixture was concentrated, the residue was dissolved in ether. The resulting residue was recrystallized from ethyl acetate/hexane to give 0.65 g (97% yield) of 2-ethanoyloxy-5-aminobenzoic acid as a white solid.

mp: 133-136°C,

$^1$H-NMR(DMSO-d$_6$) δ 7.5-7.6 (s, 1H), 7.2-7.3 (d, 1H), 7.0-7.1 (d, 1H), 2.1-2.2 (s, 3H),

$^{13}$C NMR(DMSO-d$_6$) δ 170.83, 165.57, 143.40, 124.41, 123.45, 121.09, 118.64, 113.98, 30.98.

**Preparation of 2-ethanoyloxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino) benzoic acid**

According to the similar procedure in Synthesis Example 1, by using 2-ethanoyloxy-5-aminobenzoic acid (0.81 g, 4.10 mmole) which was prepared from Synthesis Example (14-3), DMF (15 ml), TEA (0.1 ml), tetrabutyl
ammonium iodide (5 mg), and 2,3,5,6-tetrafluoro-4-trifluoromethylbenzyl bromide (1.02 ml, 6.15 mmole), 0.92 g (53.0% yield) of was obtained 2-ethanoyloxy-5-(2,3,5,6-tetrafluoro-
4-trifluoromethylbenzylamino)benzoic acid as a white solid. :

mp: 185-187°C,

$^1$H-NMR(DMSO-d$_6$) δ 2.16(s, 3H), 4.46(s, 2H), 6.82(d, 1H), 6.88(d, 1H), 7.17(s, 1H),

$^{13}$C NMR(DMSO-d$_6$) δ 21.64, 36.37, 114.36, 116.97, 123.94, 127.78, 124.81, 141.40, 142.44, 144.44, 145.04, 145.84, 146.85, 166.35, 170.18,

IR(KBr pellet) 3410, 1747, 1705, 1489 cm$^{-1}$

Elemental analysis for C$_{17}$H$_{13}$F$_{11}$NO$_4$

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<Synthesis Example 15>

Preparation of 2-propanoyloxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)benzoic acid

Preparation of 5-tert-butoxycarbonylamino-2-propanoyloxybenzoic acid

According to the similar procedure in Synthesis Example (14-2), by using propanoyl chloride (446 mg,
4.82 mmole), 0.71 g (57.0% yield) of was obtained 5-tert-butoxycarbonylamino-2-propanoyloxybenzoic acid as a white solid.

mp: 137-142°C,

\[ ^1H-NMR(DMSO-d_6) \delta 7.9-8.0 (s, 1H), 7.4-7.5 (d, 1H), 6.8-6.9 (d, 1H), 2.5-2.6 (t, 2H), 1.4-1.6 (s, 9H), 1.0-1.2 (q, 3H), \]

\[ ^{13}C\text{ NMR}(DMSO-d_6) \delta 172.689, 165.598, 152.804, 144.674, 137.287, 124.045, 79.618, 28.358, 27.259, 9.080 \]

(15-2) **Preparation of 5-amino-2-propanoyloxybenzoic acid**

According to the similar procedure in Synthesis Example (14-3), by using 5-tert-butoxycarbonylamino-2-propanoyloxybenzoic acid (1.01 mg, 3.26 mmole), 0.67 g (97.0% yield) of was obtained 5-amino-2-propanoyloxybenzoic acid as a white solid.

mp: 213-220°C,

\[ ^1H-NMR(DMSO-d_6) \delta 7.5-7.6 (s, 1H), 7.2-7.3 (d, 1H), 7.0-7.1 (d, 1H), 2.4-2.6 (t, 2H), 1.0-1.2 (q, 3H), \]

\[ ^{13}C\text{ NMR}(DMSO-d_6) \delta 172.78, 165.35, 145.54, 137.100, 124.90, 124.76, 124.23, 121.72, 27.32, 9.10. \]

(15-3) **Preparation of 2-propanoyloxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino) benzoic acid**
According to the similar procedure in Synthesis Example 14-4, by using 5-amino-2-propanoyloxybenzoic acid (0.32 g, 1.53 mmole) which was prepared from Synthesis Example (15-2), DMF (10 ml), TEA (0.05 ml), tetrabutyl ammonium iodide (3 mg), and 2,3,5,6-tetrafluoro-4-trifluoromethylbenzyl bromide (0.38 ml, 2.30 mmole), 0.34 g (51.0% yield) of was obtained 2-propanoyloxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)benzoic acid as a white solid.

mp 188-191°C,
\[ ^1H-NMR(\text{acetone-d}_6) \delta 1.17 (t, 3H), 2.06 (q, 2H), 4.67 (s, 2H), 6.94 (m, 2H), 7.37 (s, 1H), \]
\[ ^13C \text{NMR}(\text{acetone-d}_6) \delta 8.65, 27.41, 36.34, 114.76, 117.27, 123.57, 124.16, 124.59, 141.56, 142.44, 145.29, 145.31, 165.29, 172.77, \]

IR(KBr pellet) 3414, 1745, 1702, 1491 cm\(^{-1}\)

Elemental analysis for C\(_{15}\)H\(_{17}\)F\(_7\)NO\(_4\)

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<Synthesis Example 16>

Preparation of 2-cyclohexanecarbonyloxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)benzoic acid
(16-1) Preparation of 5-tert-butoxycarbonylamino-2-cyclohexanecarbonyloxybenzoic acid

According to the similar procedure in Synthesis Example (14-2), by using cyclohexylcarbonyl chloride (707 mg, 4.82 mmole), 0.72 g (49.0% yield) of was obtained 5-tert-butoxycarbonylamino-2-cyclohexanecarbonyloxybenzoic acid as a white solid.

mp: 68-74°C,

$^{1}$H-NMR(DMSO-$d_6$) δ 7.9-8.0 (s, 1H), 7.4-7.5 (d, 1H), 6.8-6.9 (d, 1H), 2.4-2.6 (t, 1H), 1.0-2.0 (m, 19H)

(16-2) Preparation of 5-amino-2-cyclohexanecarbonyloxybenzoic acid

According to the similar procedure in Synthesis Example 14-3, by using 5-tert-butoxycarbonylamino-2-cyclohexanecarbonyloxybenzoic acid (1.01 mg, 2.78 mmole), 0.70 g (96.0% yield) of was obtained 5-amino-2-cyclohexanecarbonyloxybenzoic acid as a white solid.

mp: 116-121°C,

$^{1}$H-NMR(DMSO-$d_6$) δ 7.5-7.6 (s, 1H), 7.4-7.5 (d, 1H), 6.9-7.0 (d, 1H), 2.4-2.6 (t, 1H), 1.0-2.0 (m, 10H),

$^{13}$C NMR(DMSO-$d_6$) 173.87, 170.74, 165.60, 160.09, 158.61, 158.27, 143.14, 140.71, 130.03, 124.66, 123.44, 121.70, 119.12, 118.61, 113.81, 42.42, 31.25, 28.94, 25.65, 22.38, 14.29.
(16-3) Preparation of 2-cyclohexanecarbonyloxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)benzoic acid

According to the similar procedure in Synthesis Example (14-4), by using 5-amino-2-cyclohexanecarbonyloxybenzoic acid (1.03 g, 3.95 mmole) which was prepared from Synthesis Example (16-2), DMF (15 ml), TEA (0.10 ml), tetrabutyl ammonium iodide (10 mg), and 2,3,5,6-tetrafluoro-4-trifluoromethylbenzyl bromide (1.01 ml, 5.92 mmole), 0.95 g (49.0% yield) of was obtained 2-cyclohexanecarbonyloxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethylbenzylamino)benzoic acid as a white solid.

mp 190-193°C,

$^1$H-NMR(acetone-\textit{d}) $\delta$ 1.20-1.57 (m, 6H), 1.64-1.81 (m, 4H) 2.53 (m, 1H), 4.67 (s, 2H), 6.90 (d, 1H), 6.96 (d, 1H), 7.36(s, 1H),

$^{13}$C NMR(acetone-\textit{d}) $\delta$ 25.57, 26.07, 36.23, 43.13, 114.67, 117.21, 122.21, 122.36, 124.45, 124.56, 142.32, 142.74, 144.21, 145.24, 146.82, 165.29, 173.96, IR(KBr pellet) 3402, 1724, 1707, 1491 cm$^{-1}$

Elemental analysis for C$_{22}$H$_{18}$F$_7$NO$_4$

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BRIEF DESCRIPTION OF THE DRAWINGS

The above and other objects, features and other advantages of the present invention will be more clearly understood from the following detailed description taken in conjunction with the accompanying drawings, in which:

Fig. 1a. The effect of 2-Hydroxy-TTBA on NMDA-induced excitotoxicity.

Mouse cortical cell cultures (DIV 12 - 14) were exposed to 300 μM NMDA for 10 min, alone or with inclusion of 3 - 300 μM 2-Hydroxy-TTBA. Neuronal death was analyzed 24 hr later by measuring levels of LDH released into the bathing medium, mean ± SEM (n = 9 - 12 culture wells per condition), scaled to mean LDH efflux value 24 hr after sham wash (=0) and continuous exposure to 500 μM NMDA (=100) that causes near complete neuronal death. *, Significant difference from the vehicle control, p<0.05 using ANOVA and Student-Neuman-Keuls' test.

Fig. 1b. The effect of 2-Hydroxy-TTS, 2-HydroxyTTA, 2-Ethan-TTBA, 2-Propan-TTBA, or 2-Cyclohexan-TTBA on NMDA-induced excitotoxicity.

Mouse cortical cell cultures (DIV 12 - 14) were
exposed to 300 μM NMDA for 10 min, alone (□) or with inclusion of 3 – 300 μM 2-Hydroxy-TTS (●), 2-Hydroxy-TTA (○), 2-Ethan-TTBA (▼), 2-Propan-TTBA (▼), or 2-Cyclohexan-TTBA (■). Neuronal death was analyzed 24 hr later by measuring levels of LDH released into the bathing medium, mean ± SEM (n = 9 - 12 culture wells per condition), scaled to mean LDH efflux value 24 hr after sham wash (=0) and continuous exposure to 500 μM NMDA (=100). *, Significant difference from the vehicle control, p<0.05 using ANOVA and Student-Neuman-Keuls' test.

Fig. 2a. Blockade of NMDA currents by 2-Hydroxy-TTBA.

Typical NMDA-induced inward currents (NMDA currents) were evoked by applying 300 μM NMDA to cortical neurons which were held at -60 mV. Successive application of various concentrations of 2-Hydroxy-TTBA reduced the response elicited by 300 μM NMDA in a concentration-dependent manner (n=8). The graph shows a dose response relation of 2-Hydroxy-TTBA to NMDA currents; its IC₅₀ value was close to 35 μM and Hill coefficient 0.91.

Fig. 2b. Pretreatment with 2-Hydroxy-TTBA did not influence NMDA currents.

Cortical neurons were treated with 100 μM 2-Hydroxy-TTBA, washed thoroughly, and then applied with
300 μM NMDA. The pretreatment with 2-Hydroxy-TTBA did not influence NMDA-induced inward currents, suggesting that 2-Hydroxy-TTBA exert its blocking action only when the receptor has been activated by agonist (n=6).

Fig. 3a. The effect of 2-Hydroxy-TTBA on FeCl₂-induced oxidative stress.

Mouse cortical cell cultures (DIV 12 - 14) were exposed to continuously to 50 μM Fe²⁺, alone or with inclusion of 0.1 - 100 μM 2-Hydroxy-TTBA (●) or trolox (○), a membrane-permeable form of vitamin E). Neuronal death was analyzed 24 hr later by measuring levels of LDH released into the bathing medium, mean ± SEM (n = 9 - 12 culture wells per condition), scaled to mean LDH efflux value 24 hr after sham wash (=0) and continuous exposure to 500 μM NMDA (=100). *, Significant difference from the vehicle control, p<0.05 using ANOVA and Student-Neuman-Keuls’ test.

Fig. 3b. The effect of 2-Hydroxy-TTS, 2-Hydroxy-TTA, 2-Chloro-TTP, 2-Ethan-TTBA, 2-Propan-TTBA, or 2-Cyclohexan-TTBA on FeCl₂-induced oxidative stress.

Mouse cortical cell cultures (DIV 12 - 14) were exposed to continuously to 50 μM Fe²⁺, alone (◆) or with inclusion of 1 - 30 μM 2-Hydroxy-TTS (●), 2-Hydroxy-TTA (○), 2-Chloro-TTP (▼), 2-Ethan-TTBA (▼), 2-Propan-TTBA (■), or 2-Cyclohexan-TTBA (□). Neuronal death was analyzed 24 hr later by measuring levels of
LDH released into the bathing medium, mean ± SEM (n = 9 - 12 culture wells per condition), scaled to mean LDH efflux value 24 hr after sham wash (=0) and continuous exposure to 500 μM NMDA (=100). *, Significant difference from the vehicle control, p<0.05 using ANOVA and Student-Neuman-Keuls' test.

Fig. 4. The effect of 2-Hydroxy-TTBA on SNP-induced oxidative stress.

Mouse cortical cell cultures (DIV 12 - 14) were exposed to continuously 5 μM sodium nitroprusside (SNP), alone (●) or with inclusion of 0.1 - 10 mM aspirin (○), 1 - 100 μM trolox (▼), or 0.1 - 10 μM 2-Hydroxy-TTBA(▼). Neuronal death was analyzed 24 hr later by measuring levels of LDH released into the bathing medium, mean ± SEM (n = 9 - 12 culture wells per condition), scaled to mean LDH efflux value 24 hr after sham wash (=0) and continuous exposure to 500 μM NMDA (=100). *, Significant difference from the vehicle control, p<0.05 using ANOVA and Student-Neuman-Keuls' test.

Fig. 5. The effect of 2-Hydroxy-TTBA on zinc toxicity.

Mouse cortical cell cultures (DIV 12 - 14) were exposed to 300 μM Zn²⁺ for 30 min, alone or with inclusion of 3 - 300 μM 2-Hydroxy-TTBA. Neuronal death was analyzed 24 hr later by measuring levels of LDH
released into the bathing medium, mean ± SEM (n = 9 - 12 culture wells per condition), scaled to mean LDH efflux value 24 hr after sham wash (=0) and continuous exposure to 500 µM NMDA (=100).

Fig. 6a. Free radical scavenging activity of 2-Hydroxy-TTBA.

2-Hydroxy-TTBA (●) or trolox (○) was reacted with 100 µM 1,1-diphenyl-2-picrylhydrazil (DPPH, a stable free radical) dissolved in ethanol for 10 min. The radical scavenging activity was determined by measuring the decrease in DPPH levels at 517 nm. *, Significant difference from the vehicle control, p<0.05 using ANOVA and Student-Neuman-Keuls' test.

Fig. 6b. The DPPH assay of 2-Hydroxy-TTS, 2-Hydroxy-TTA, or 2-Chloro-TTP.

2-Hydroxy-TTS (●), 2-Hydroxy-TTA (○), 2-Chloro-TTP (▼) or control (▼) was reacted with 100 µM DPPH dissolved in ethanol. The radical scavenging activity was determined by measuring the decrease in DPPH levels at 517 nm. *, Significant difference from the vehicle control, p<0.05 using ANOVA and Student-Neuman-Keuls' test.

Fig. 7a. 2-Hydroxy-TTBA reduces neuronal death in the spinal cord in an animal model of ALS.

Bright field photomicrographs of spinal cord sections stained with eosin from amyotrophic lateral
sclerosis (ALS) mice overexpressing the mutant SOD1-G93A that received vehicle (A) or 2-Hydroxy-TTBA (B, 10 mg/kg/day through drinking water) for 8 weeks from the age of 2 months.

Fig. 7b. 2-Hydroxy-TTBA reduces neuronal death in the spinal cord in an animal model of ALS.

Degenerating neurons from Fig. 7a. were analyzed by counting viable neurons after staining with eosin in the dorsal and ventral horn of spinal cord from ALS mice treated with a vehicle (Control) or 2-Hydroxy-TTBA, mean ± S.E.M (n = 5 animals for each condition). *, Significant difference from the vehicle control, p<0.05 using the independent t-test.

Fig. 8a. Intraperitoneal administration of 2-Hydroxy-TTBA reduces ischemic injury in brain.

Adult rats received transient cerebral ischemia by occluding right middle cerebral artery and both common carotid arteries for 60 min with intraperitoneal injections of vehicle or 50 mg/kg 2-Hydroxy-TTBA at 5 min, 30 min, or 1 hr after reperfusion. Infarct volume was analyzed 24 hr later after staining brain slices with 2% 2,3,5-triphenyltetrazolium chloride (TTC), mean ± SEM (n = 8 - 11 rats per each condition). *, Significant difference from the vehicle control, p<0.05 using ANOVA and Student-Neuman-Keuls’ test.

Fig. 8b. Intraperitoneal administration of 2-
Hydroxy-TTBA reduces ischemic injury in brain.

Adult rats received transient cerebral ischemia by occluding right middle cerebral artery and both common carotid arteries for 60 min with intraperitoneal injections of vehicle (control) or 50 or 100 mg/kg 2-Hydroxy-TTBA at 5 min after reperfusion. Infarct volume was analyzed 24 hr later after staining brain slices with 2% 2,3,5-triphenyltetrazolium chloride (TTC), mean ± SEM (n = 7 - 8 rats per each condition).

*, Significant difference from the vehicle control, p<0.05 using ANOVA and Student-Neuman-Keuls' test.

Fig. 8c. Intravenous injections of 2-Hydroxy-TTBA reduces ischemic injury in brain.

Adult rats received transient cerebral ischemia by occluding right middle cerebral artery and both common carotid arteries for 60 min, with intravenous injections of vehicle (control) or 5 mg/kg 2-Hydroxy-TTBA at 30 min after occlusion or 5 min, 30 min, 1 hr, 2 hr, or 4 hr after reperfusion. Infarct volume was analyzed 24 hr later, mean ± SEM (n = 8 - 11 rats per each condition). *, Significant difference from the vehicle control, p<0.05 using ANOVA and Student-Neuman-Keuls' test.

Fig. 8d. Intravenous injections of 2-Hydroxy-TTBA reduces ischemic injury in brain.

Adult rats received transient cerebral ischemia
by occluding right middle cerebral artery and both common carotid arteries for 60 min, with intravenous injections of vehicle (control) or 1, 2.5, 5, 10 or 20 mg/kg 2-Hydroxy-TTBA at 5 min after reperfusion. Infarct volume was analyzed 24 hr later, mean ± SEM (n = 8 - 10 rats per each condition). *, Significant difference from the vehicle control, p<0.05 using ANOVA and Student-Neuman-Keuls' test.

Fig. 8e. Oral administration of 2-Hydroxy-TTBA reduces ischemic injury in brain.

Adult rats received transient cerebral ischemia by occluding right middle cerebral artery and both common carotid arteries for 60 min, with oral administrations of vehicle (control) or 10 or 20 mg/kg 2-Hydroxy-TTBA at 5 min after reperfusion. Infarct volume was analyzed 24 hr later, mean ± SEM (n = 8 - 10 rats per each condition). *, Significant difference from the vehicle control, p<0.05 using ANOVA and Student-Neuman-Keuls' test.

Fig. 9a. Effects of 2-Hydroxy-TTBA on mitochondrial ROS generation at 72 hr of recirculation after global ischemia.

Mitotracker CM-H$_2$X ROS was injected into the lateral ventricle of rat brain at 24 hr before ischemic surgery, and then animals received transient forebrain ischemia for 10 min, with intraperitoneal injections of
vehicle (control) or 2-Hydroxy-TTBA (50 mg/kg) at 5 min after reperfusion. The levels of mitochondrial ROS were analyzed 2 hr later, mean ± S.E.M. (n = 50 – 70 neurons randomly chosen from the CA1 sector of the hippocampal formation per each condition). *, Significant difference from the vehicle control, p<0.05 using the independent t-test.

Fig. 9b. 2-Hydroxy-TTBA prevents neuronal death in the CA1 sector at 72 hr of recirculation after global ischemia.

Adult rats received transient global ischemia for 10 min, with intraperitoneal injections of vehicle (control) or 2-Hydroxy-TTBA (50 mg/kg) at 5 min after reperfusion. The number of degenerating neurons in the CA1 was analyzed 3 d later by counting viable neurons after staining with cresyl violet, mean ± S.E.M. *, Significant difference from the vehicle control, p<0.05 using the independent t-test.

Fig. 10a. 2-Hydroxy-TTBA reduces death of dopaminergic neurons in the substantia nigra 3 d following MPTP injection.

Bright field photomicrographs showing dopaminergic neurons in the substantia nigra immunostained with anti-tyrosine hydroxylase (TH) antibody 3 d following sham control (A) or the single daily injection of 1-methyl-4-phenyl-1,2,3,6-tetra-
hydropyridine (MPTP, 40mg/kg, SC) in C57/BL6 mice, alone (B) or with intraperitoneal injections (2 times per day) of 2-Hydroxy-TTBA (C, 25 mg/kg; D, 50 mg/kg) prior to MPTP administration.

Fig. 10b. 2-Hydroxy-TTBA reduces death of dopaminergic neurons in the substantia nigra 3 d following MPTP injection.

Quantification of TH-positive dopaminergic neurons in the substantia nigral sections 3 d following the single daily injection of (MPTP, 40mg/kg, SC), alone or with pretreatment with 2-Hydroxy-TTBA as described above. *, Significant difference from the control, p<0.05 using ANOVA and Student-Neuman-Keuls' test.

Fig. 10c. 2-Hydroxy-TTBA reduces death of dopaminergic neurons in the substantia nigra 7 d following MPTP injection.

Bright field photomicrographs showing dopaminergic neurons in the substantia nigra immunostained with anti-tyrosine hydroxylase (TH) antibody 7 d following sham control (A) or the single daily injection of 1-methyl-4-phenyl-1,2,3,6-tetrahydropyridine (MPTP, 40mg/kg, SC) in C57/BL6 mice, alone (B) or with intraperitoneal injections (2 times per day) of 2-Hydroxy-TTBA (C, 25 mg/kg; D, 50 mg/kg) prior to MPTP administration.
Fig. 11. 2-Hydroxy-TTBA prevents generation of mitochondrial ROS in the dorsal horn neurons of the spinal cord following traumatic injury.

Adult rats received compression injury at the level of T8 spinal cord, alone (control) or with injections of 2-Hydroxy-TTBA (50 mg/kg, ip) and Mitotracker CM-H$_2$X ROS immediately after injury. Animals were euthanized 2 d later, spinal cord sections sections immunostained with an antibody for NeuN (a neuronal marker protein), and levels of mitochondrial ROS in the dorsal horn neurons (DHN) analyzed by measuring fluorescence intensity of oxidized Mitotracker CM-H$_2$X, mean ± S.E.M. [n = 12 rats for each condition (5 spinal cord sections for each rat)]: *, Significant difference from the vehicle control, $p<0.05$ using the independent t-test.

This present invention is described particularly in experimental examples using 2-Hydroxy-TTBA, 2-Hydroxy-TTS, 2-Hydroxy-TTA, 2-Chloro-TTP, 2-Ethan-TTBA, 2-Propan-TTBA and 2-Cyclohexan-TTBA manufactured in synthesis example as follows.

However, the examples described below are just representative of the present invention, which could include more examples.
mixed cortical cell cultures of neurons and glia

For mixed neuron-glia culture, mouse cerebral cortices were removed from brains of the 14 - 16 day-old-fetal mice (E14-16), gently triturated and plated on 24 well plates (2X10^5 cells/plate) precoated with 100 µg/ml poly-D-lysine and 4 µg/ml laminine. Cultures were maintained at 37°C in a humidified 5% CO_2 atmosphere. Plating media consist of Eagles minimal essential media (MEM, Earles salts, supplied glutamine-free) supplemented with 5% horse serum, 5% fetal bovine serum, 26.5 mM bicarbonate, 2 mM glutamine, and 21 mM glucose.

After 7 - 8 days in vitro (DIV 7-8), 10 µM cytosine arabinofuranoside (Ara-C) was included to halt overgrowth of glia. The drug treatment was carried on DIV 12 - 15 cortical cell culture. Overall neuronal cell injury was assessed by measuring amount of lactate dehydrogenase (LDH) released into the bathing medium 24 hr after neurotoxic insults as previously described [Koh and Choi, J Neurosci Methods 20:83-90, 1987].

Inhibitory effects of excitotoxicity by 2-hydroxy-TTBA, 2-hydroxy-TTA, 2-Hydroxy-TTS, 2-Ethan-TTBA, 2-Prppan-TTBA or 2-Cyclohexan-TTBA
At DIV 13-15 in mixed neuron-glia culture as shown in Example 1, cortical cell cultures were exposed to 300 μM NMDA for 10 min, alone or with 3 - 300 μM 2-Hydroxy-TTBA, 10 - 300 μM 2-Hydroxy-TTS, 30 - 300 μM 2-Hydroxy-TTA, 30 - 300 μM 2-Ethan-TTBA, 10 - 300 μM 2-Propan-TTBA, or 10 - 300 μM 2-Cyclohexan-TTBA. Neuronal death was assessed 24 hr later by measuring amount of LDH released into the bathing medium. *

Significant difference from the vehicle control, p<0.05 using ANOVA and Student-Neuman-Keuls’ test.

Cortical cell cultures exposed to 300 μM NMDA for 10 min underwent widespread neuronal death (approximately 75 - 80% neurons died) over the next 24 hr. NMDA-induced neuronal death was blocked by co-treatment with 2-Hydroxy-TTBA in a dose-dependent manner at doses of 10 - 300 μM (Fig. 1a).

Concurrent administration of 2-Hydroxy-TTS, 2-Hydroxy-TTA, 2-Ethan-TTBA, 2-Propan-TTBA, or 2-Cyclohexan-TTBA also prevented NMDA-induced neuronal death at doses of 30 - 300 μM. (Fig. 1b).

Example 3

Blocking effect of 2-Hydroxy-TTBA on NMDA-induced inward currents

Whole cell recordings were performed on cortical cell cultures at room temperature as described [Seo et

Typical NMDA-induced inward currents (NMDA currents) were evoked immediately after applying 300 μM NMDA to cultured cortical neurons which were held at -70 mV. Bath application of 2-Hydroxy-TTBA immediately depressed NMDA-evoked currents in a dose dependent manner (n=7-15 neurons/condition, IC₅₀ value = 30.55 ± 2.96 μM) (Fig 2a). Application of 100 μM 2-Hydroxy-TTBA alone had no effect on the holding currents and had little effect on the response of the cell to a subsequent treatment of 300 μM NMDA (Fig. 2b), suggesting that 2-Hydroxy-TTBA exerts antagonistic effect only when the NMDA receptor is activated (n=6).

<Example 4>

Blockade of oxidative neuronal death by 2-Hydroxy-TTBA, 2-Hydroxy-TTA, 2-Hydroxy-TTS, 2-Chloro-TTP, 2-Ethan-TTBA, 2-Propan-TTBA or 2-Cyclohexan-TTBA

(4-1) Inhibition of FeCl₂-induced free radical toxicity

Mixed cortical cell cultures (DIV 13 - 15) were continuously exposed to 50 μM FeCl₂, which produces hydroxyl radical via a Fenton reaction, alone or with inclusion of 2-Hydroxy-TTBA, 2-Hydroxy-TTA, 2-Hydroxy-TTS, 2-Chloro-TTP, 2-Ethan-TTBA, 2-Propan-TTBA, 2-
Cyclohexan-TTBA or trolox (a vitamin E analogue) at indicated doses. Neuronal cell death was analyzed 24 hr later by LDH assay as described above. *, Significant difference from the vehicle control (FeCl₂), p<0.05 using ANOVA and Student-Neuman-Keuls’ test.

Cortical cell cultures exposed to FeCl₂ underwent widespread neuronal death over the next 24 hr. 2-Hydroxy-TTBA and trolox prevented FeCl₂-induced free radical neurotoxicity in a dose-dependent manner. However, 2-Hydroxy-TTBA was 30-fold stronger than trolox in preventing free radical neurotoxicity (Fig. 3a). Moreover, synthetic derivatives of 2-Hydroxy-TTBA (2-Hydroxy-TTS, 2-Hydroxy-TTA, 2-Chloro-TTP, 2-Ethan-TTBA, 2-Propan-TTBA, or 2-Cyclohexan-TTBA) showed much stronger neuroprotective effects than trolox in preventing FeCl₂-induced free radical neurotoxicity (Fig. 3b).

(4-2) Inhibition of SNP (sodium nitroprusside) cytotoxicity

Mixed cortical cell cultures were continuously exposed to 5 μM SNP, a nitric oxide (NO) donor, alone or with inclusion of 0.1 - 10 μM 2-Hydroxy-TTBA, 1 -10 μM trolox or 100 - 10,000 μM aspirin. Neuronal cell death was analyzed 24 hr later by LDH assay. *, Significant difference from the vehicle control (SNP),
Administration of SNP resulted in complete neuronal cell death and some glial cell death. SNP toxicity was completely blocked in the presence of 2-Hydroxy-TTBA and trolox. The former was 100-fold stronger than the latter in preventing NO toxicity. Aspirin, a structural component of 2-Hydroxy-TTBA, did not reduce SNP cytotoxicity (Fig. 4).

(4-3) Inhibition of zinc neurotoxicity

To induce zinc (Zn²⁺) neurotoxicity, mixed cortical cell cultures were exposed to 100 μM ZnCl₂ for 30 min in a HEPES-buffered control salt solution (HCSS) : (in mM) 120 NaCl, 5 KCl, 1.6 MgCl₂, 2.3 CaCl₂, 15 glucose, 20 HEPES and 10 NaOH, alone or with 3 - 300 μM 2-Hydroxy-TTBA. After exposure, cultures were washed out 3 times and exchanged with MEM adjusted to 25 mM glucose and 26.2 mM sodium bicarbonate. Neuronal cell death was analyzed 24 hr later by LDH assay. *, Significant difference from the vehicle control (SNP), p<0.05 using ANOVA and Student-Neuman-Keuls' test.

Concurrent treatment with 2-Hydroxy-TTBA prevented zinc-induced neuronal death in a dose-dependent manner at doses of 10 - 300 μM (Fig. 5).

(4-4) DPPH assay of 2-Hydroxy-TTBA, 2-Hydroxy-TTA, 2-
Hydroxy-TTS or 2-Chloro-TTP

To examine the free radical scavenging effects, 1 - 100 μM 2-Hydroxy-TTBA, 2-Hydroxy-TTA, 2-Hydroxy-TTS, 2-Chloro-TTP or trolox was reacted with 100 μM DPPH (2,2-diphenyl-1-picryl-hydrazyl radical), a stable free radical, dissolved in ethanol. After incubation for 30 min, relative decrease in DPPH absorption at 517 nm was measured by a spectrophotometer, mean ± SEM (n = 3 test tubes per condition). *, Significant difference from the vehicle control (DPPH alone), p<0.05 using ANOVA and Student-Neuman-Keuls’ test.

Compared to the anti-oxidant trolox, 2-Hydroxy-TTBA reduced levels of DPPH at lower doses, suggesting that 2-Hydroxy-TTBA is a direct anti-oxidant stronger than trolox (Fig. 6a). Other synthetic derivatives also reduced levels of DPPH (Fig. 6b).

<Example 5>
Prevention of neuronal cell death in spinal cord of ALS mouse by 2-hydroxy-TTBA

Transgenic mice with the G93A human SOD1 mutation (B6SJL-TgN(SOD1-G93A)1Gur), animal models of ALS (ALS mice), were obtained from Jackson Laboratories (ME, USA). 2-hydroxy-TTBA was administered to 2 month-old wild type and ALS transgenic mice through drinking water bottle (10 mg/kg per day) for 8 weeks. Animals
were then euthanized and spinal cords processed for histological examination by staining with hematoxylin and eosin. Spinal motor neurons from ALS mice without being treated with 2-hydroxy-TTBA (control) underwent degeneration as evident by eosinophilic staining, which was reduced in ALS mice treated with 2-hydroxy-TTBA (Fig. 7a).

The number of degenerating neurons was analyzed by counting eosinophilic neurons in ventral horn (VH) and dorsal horn (DH). Administration of 2-hydroxy-TTBA significantly prevented degeneration of dorsal and ventral horn neurons in the spinal cord from ALS mice (Fig. 7b). *, Significant difference from the control, p<0.05 using independence t test.

<Example 6>
Prevention of hypoxic-ischemic brain injury by 2-Hydroxy-TTBA

(6-1) Induction of transient focal cerebral ischemia in rat

Sprague-Dawley rats were anesthetized with chloral hydrate, and subjected to focal cerebral ischemia by occlusion of middle cerebral artery (MCAO) as previously described [Tamura et al., J. Cerebr. Blood Flow Metab. 1:53-60(1981)]. Both common carotid
arteries (CCAs) were exposed and right middle cerebral artery (rMCA) was exposed under the surgical microscope by making a 3 mm diameter craniotomy rostral to the foramen ovale. CCAs and rMCA were occluded with microclips. The clips were released 60 min later and restoration of blood flow in rMCA was observed under the microscope.

(6-2) 2-Hydroxy-TTBA reduces infarct volume after 60 min MCAO.

Animals received MCAO for 60 min, alone or with administration of 2-Hydroxy-TTBA (50 mg/kg, i.p.) at indicated points of time after reperfusion. Saline was injected as a control. Animals were euthanized 24 hr later and brains removed and sectioned coronally into seven 2-mm slices in a brain matrix. Brain slices were placed in 2% 2,3,5,-triphenyltetrazolium chloride solution, followed by 10% formalin overnight. The infarction area, outline in white, was measured (TINA image analysis system) and infarction volume was calculated by summing the infarct volume showing white of sequential 2-mm-thick section, mean ± SEM (n = 8-12 rats/condition) (Fig. 8a). Note that delayed administration (i.p) of 2-Hydroxy-TTBA up to 1 hr after reperfusion significantly reduced infarct volume. Higher doses of 2-Hydroxy-TTBA (100 mg/kg, i.p) also
showed similar protection against 60 min MCAO (Fig. 8b).

Additional experiments were performed to examine effects of intravenous injections of 2-Hydroxy-TTBA (5 mg/kg) against 60 min MCAO. 2-Hydroxy-TTBA was administered at indicated points of time after reperfusion. The delayed injections of 2-Hydroxy-TTBA up to 4 hr after reperfusion significantly attenuated infarct volume evolving 24 hr after 60 min MCAO, mean ± SEM (n = 8 - 12 rats/condition) (Fig. 8c).

Dose-response experiments of the intravenous injections of 2-Hydroxy-TTBA showed that 2-Hydroxy-TTBA attenuated infarct volume at doses as low as 1 mg/kg. The protective effects of 2-Hydroxy-TTBA were observed at doses higher than 2.5 mg/kg, mean ± SEM (n = 8 - 12 rats/condition) (Fig. 8d).

Protective effects of 2-Hydroxy-TTBA were also verified through oral administration. In particular, administration of 2-Hydroxy-TTBA (10 or 20 mg/kg, p.o.) at 5 min after reperfusion reduced infarct volume 24 hr after 60 min MCAO, mean ± SEM (n = 8 - 12 rats/condition) (Fig. 8e).

<Example 7>

Prevention of the CA1 neuronal cell death after transient forebrain ischemia by 2-Hydroxy-TTBA
(7-1) Induction of global ischemia in rat

Male adult Sprague-dawley rats (250 - 300 g) were anesthetized with chloral hydrate and subjected to four-vessel occlusion model (occlusion of both common carotid arteries and vertebral arteries) as previously described [Pulsinelli and Brierley, stroke 10: 267-272 (1979)].

(7-2) Inhibition of mitochondrial ROS generation after global ischemia by 2-Hydroxy-TTBA

Rats received the intracerebroventricular injections of 0.4 nmol Mitotracker CM-H₂X Ros. After 24 hr, rats received four-vessel occlusion for 10 min. Immediately after reperfusion, rats received the intraperitoneal injections of saline (control) or 2-Hydroxy-TTBA (50 mg/kg). Levels of mitochondrial ROS were analyzed 2 hr later by measuring the fluorescence intensity of oxidized Mitotracker CM-H₂X Ros in mitochondria. Administration of 2-Hydroxy-TTBA reduced production of mitochondrial ROS 2 hr after transient global ischemia (Fig. 9a).

(7-3) Inhibition of neuronal cell death in the CA1 field following transient forebrain ischemia by 2-Hydroxy-TTBA

Rats received four-vessel occlusion for
10 min followed by administration of saline (control) or 2-Hydroxy-TTBA (50 mg/kg, ip). Animals were euthanized 3 d later and processed for analysis of neuronal death in the CA1 field after staining with hematoxylin and eosin (mean ± SEM, N=12 rats/condition). Administration of 2-Hydroxy-TTBA prevented delayed neuronal cell death in the CA1 areas evolving after transient global ischemia (Fig. 9b). *, Significant difference from the control, p<0.05 using independence t test.

<Example 8>
Prevention of dopaminergic neuronal cell death in the substantial nigra following the injection of MPTP (1-methyl-4-phenyl-1,2,3,6-tetrahydropyridine) by 2-Hydroxy-TTBA

(8-1) Inhibition of dopaminergic neurons by 2-Hydroxy-TTBA

C57/BL6 mice (male, 8 weeks) received the injections of 40 mg/kg MPTP (s.c.), alone or with 25 or 50 mg/kg 2-Hydroxy-TTBA (ip) every 12 hr per day beginning 30 min before MPTP injection. After 3 or 7 days, all animals were anesthetized with chloral hydrate and perfused transcardially with PBS followed by 4% paraformaldehyde. Brains were immediately
removed and immersed in the fixative for 8 - 10 h. The fixative was replaced with 30% sucrose, incubated at 4°C for 2 d, and stored at -70°C. The brains were sectioned at a thickness of 30 μm on a sliding microtome (TPI, Inc., MO). Sections were then stored in 0.1 M phosphate buffer (pH 7.4, 30% (v/v) glycerol, 30% ethylene glycol) at 4°C until use. The sections were immunostained with anti-TH (tyrosine hydroxylase) antibody, colored by DAB (diaminobenzidine), and then observed under light microscope to analyze the degeneration of dopaminergic neurons.

Mice treated with MPTP showed marked degeneration of the dopaminergic neurons in the substantia nigra over the next 3 d. The intraperitoneal injections of 2-Hydroxy-TTBA significantly attenuated degeneration of the dopaminergic neurons (Fig. 10a and 10b). The protective effects of 2-Hydroxy-TTBA are also observed at 7 d following administration of MPTP (Fig. 10c and 10d).

<Example 9>

Inhibition of mitochondrial ROS production in the spinal cord after traumatic spinal cord injury

Sprague-Dawley rats (250 - 300 g, female) received traumatic spinal cord injury by compression of the dorsal spinal cord. T8 segment of spinal cord was
exposed on the dorsal side, compressed with 20 g for 10 min, and then mitochondrial CM-H$_2$X Ros was immediately injected into the spinal cord. 2-Hydroxy-TTBA (50 mg/kg, ip) or a vehicle control (control) was injected with mitochondrial CM-H$_2$X Ros. Levels of mitochondrial ROS in the dorsal horn neurons were analyzed 48 hr later as described above after immunolabeling with anti-NeuN antibody, a neuron-specific marker. Administration of 2-Hydroxy-TTBA significantly reduced mitochondrial ROS production in the spinal cord neurons after the traumatic spinal cord injury (Fig. 11)(mean ± SEM, N=12 rats/condition). *, Significant difference from the control, p<0.05 using independence t test.

As described above, tetrafluorobenzyl derivatives or its pharmaceutically-acceptable salts in the present invention can be used as NMDA receptor antagonists, anti-oxidants, and inhibitors of zinc neurotoxicity.

The concrete diseases applicable with tetrafluorobenzyl derivatives or its pharmaceutically-acceptable salts are described as follows.

Application examples described below are part of examples of this invention. This invention is not limited to application examples.
<Application example 1> Stroke

Interrupted blood supply to brain or stroke induces neuronal death primarily through over-activation of \( \text{Ca}^{2+} \)-permeable glutamate receptor induced by accumulation of glutamate, an excitatory neurotransmitter, in synaptic cleft. It has been well documented that NMDA receptor antagonists decrease the neuronal cell death by ischemic stroke accounting for 80% of stroke [Simon et al., Science, 226:850-852 (1984); Park et al., Ann Neurol., 24:543-551 (1988); Wieloch, Science, 230:681-683 (1985); Kass et al., Exp. Neurol., 103:116-122 (1989); Weiss et al., Brain Res., 380:186-190 (1986)]. It has also been reported that ROS and zinc are main mechanism of neuronal death following stroke. Anti-oxidants or inhibitors of zinc toxicity protect ischemic injury in animal models of stroke [Flamm, E. S. et al., Stroke, 9(5):445-447(1978); Kogure, K. et al., Prog. Brain Res., 63:237-259(1985); Chan, P. H., J. Neurotrauma., 9 suppl2:S417-423(1992)]. Therefore, the compounds in the present invention showing multiple protective effects against excitotoxicity, oxidative stress, and zinc toxicity can be used as therapeutic drugs for stroke.

<Application example 2> Trauma
Excitotoxins are closely related to the degeneration of neuronal cells following traumatic brain injury (TBI) and traumatic spinal cord injury (TSCI). Quinolinic acid, an NMDA receptor agonist present in human body, is increased 50 to 500 times in TBI patients [Sinz et al., J. Cereb. Blood Flow Metab., 18:610-615 (1988)]. It has been reported that NMDA receptor antagonists decrease the neuronal death following TBI and TSCI [Faden et al., J Neurotrauma, 5:33-45 (1988); Okiyama et al., J Neurotrauma, 14:211-222 (1997)]. Anti-oxidants also inhibit tissue damage following TBI or TSCI [Faden & Salzman, Trends Pharmacol. Sci., 13:29-35(1992)]. Therefore, the compounds in the present invention showing multiple protective effects against excitotoxicity and oxidative stress can be used as therapeutic drugs for TBI and TSCI.

<Application example 3> Epilepsy

Administration of kainate, an agonist of AMPA and kainate glutamate receptors, induces seizure and neuronal cell death in several brain areas including the hippocampal formation. NMDA receptor antagonists were shown to inhibit convulsion and seizure in several epileptogenic animal models [Anderson et al., J. Neurophysiol, 57:1-21 (1987); Wong et al., Neurosci
Anti-oxidants inhibit seizure and seizure-induced neuronal death [He et al., *Free Radic. Biol. Med.* 22:917-922(1997); Kabuto et al., *Epilepsia*, 39:237-243(1998)]. Therefore, the compounds in the present invention showing multiple protective effects against excitotoxicity and oxidative stress can be used as therapeutic drugs for epilepsy and seizure-induced neuronal death.

<Application example 4> Amyotrophic lateral sclerosis

Neurobiol. Dis., 7:375-383(2000)]. Therefore, the compounds in the present invention showing multiple protective effects against excitotoxicity and oxidative stress can be used as therapeutic drugs for ALS.

5

<Application example 5> Parkinson's disease (PD)

PD is a neurodegenerative disease showing the disorder of motor function by a selective death of dopaminergic neurons in the substantia nigra. Several antagonists of NMDA receptors protect dopaminergic neurons from the dopaminergic neurotoxin MPTP (1-methyl-4-phenyl-1,2,3,6- tetrahydropyridine) [Lange et al., Naunyn Schmiedebergs Arch.Pharmacol. 348:586-592 (1993); Brouillet and Beal. Neuroreport. 4:387-390 (1993)]. NMDA receptor antagonists also ameliorate levodopa-induced dyskinesia and thus can improve the therapeutic effects of levodopa [Papa and Chase, Ann.Neurol. 39:574-578 (1996); Marin et al., Brain Res. 736:202-205 (1996)]. Oxidative stress as well as excitotoxicity has been proved as a main mechanism of neuronal cell death in PD patients [Schapira et al., biochem. Soc. Trans., 21:367-370(1993)]. Therefore, the compounds in the present invention showing multiple protective effects against excitotoxicity and oxidative stress can be used as therapeutic drug for PD.
<Application example 6> Huntington's disease (HD)

HD is a progressive neurodegenerative disease predominantly affecting small- and medium-sized interneurons in the striata. These pathological features of HD are observed in vivo and in vitro following administration of NMDA receptor agonists, raising the possibility that NMDA receptor-mediated neurotoxicity contributes to selective neuronal death in HD [Koh et al., Science 234:73-76 (1986); Beal et al., Nature 321:168-171 (1986); Beal et al., J. Neurosci. 11:1649-1659 (1991)]. Since evidence is being accumulated that oxidative stress, such as mitochondrial dysfunction and generation of ROS, causes neuronal death observed in PD, it is possible that the drugs inhibiting ROS are used for therapy of HD [Jenner, Pathol. Biol. 44:57-64(1996); Albers & Beal, J. Neural. Transm. Suppl., 59:133-154(2000)]. Therefore, the compounds in the present invention showing multiple protective effects against excitotoxicity and oxidative stress can be used as therapeutic drugs for HD.

<Application example 7> Alzheimer's disease (AD)

The degeneration of glutamatergic neurons in the cerebral cortex and hippocampal formation and of cholinergic neurons in the basal forebrain, extracellular deposit of amyloid plaque, and

<Application example 8> Ocular diseases and cataract

Cataract is a senile disease accompanying lenticular opacity. Oxidative stress has been considered as a primary mediator of cataract. In fact, anti-oxidants have been applied to treat cataract [Varma et al., Curr. Eye Res. 3:35-57(1984); Anderson et al., Adv. Exp. Med. Biol., 366:73-86(1994)].

Therefore, the compounds in the present invention showing multiple protective effects against excitotoxicity and oxidative stress can be used as therapeutic drugs for ocular diseases.

<Application example 9> Drug addiction

The activation of mesolimbic dopaminergic neurons in the ventral tagmental area that project to the nucleus accumbens is essential for the process of drug addiction that requires neuronal excitability [Self D. W. and Nestler E. J., Annu. Rev. Neurosci., 18:463-95 (1995)]. Several lines of evidence supports that NMDA receptor antagonists can be applied to reduce neuronal adaptability to abused drugs and have been suggested as new therapeutic agents for drug addiction [Boening et al., Alcohol Clin. Exp. Res., 25:127S-131S (2001); Vorel et al., Science, 292:1175-8 (2001)].

<Application example 10> Depression

Tricyclic antidepressants and MAO (monoamine...
oxidase) inhibitors, antidepressants, increase neurotransmission of noradrenaline and serotonin. The existing therapeutic drugs induce a lot of side effects through interacting with other nervous system and have no therapeutic effect in 30% of depressed patients [Pacher et al., Curr. Med. Chem., Feb. 8 (2):89-100 (2001)]. Recently, NMDA receptor antagonists were shown to be applicable as new therapeutic drugs for depression [Le D. A. and Lipton S. A., Drugs Aging, 18:717-724 (2001); Petrie et al., Pharmacol. Ther., 87:11-25 (2000)].

<Application example 11> Pain

Neuropathic pain results from the increased neurotransmission following peripheral injury and neural tissues damage in relation with surgery, cancer patients, and trauma etc [Hempenstall K. and Rice A. S., Curr. Opin. Investig. Drugs., Mar;3 (3):441-8 (2002); McDonnell et al., Curr. Oncol. Rep., 2:351-7 (2000)]. Since the activation of NMDA receptor is necessary for the processing of neuropathic pain, it has been reported that NMDA receptor antagonists can be used as therapeutic drugs to treat neuropathic pain [Parson C. G., Eur. J. Pharmacol., 429:71-8 (2001); Hewitt, Clin. J. Pain., 16:S73-9 (2000)].
Multiple sclerosis, Meningitis, Encephalitis or Hydrocephalus

multiple sclerosis, meningitis, encephalitis or hydrocephalus.

The present invention has been described in an illustrative manner, and it is to be understood that the terminology used is intended to be in the nature of description rather than of limitation. Many modifications and variations of the present invention are possible in light of the above teachings. Therefore, it is to be understood that within the scope of the appended claims, the invention may be practiced otherwise than as specifically described.

**Industrial Applicability**

As described hereinbefore, Tetrafluorobenzyl derivatives or pharmaceutically-acceptable salts, and pharmaceutical composition containing the same as the effective component can prevent and treat the neurodegenerative diseases such as amyotrophic lateral sclerosis, Parkinson's diseases, Huntington's disease or Alzheimer's disease, the convulsive neuronal diseases such as epilepsy etc, and brain injury by stroke, trauma, or hydrocephalus, ocular diseases by glaucoma and retinopathy, mental diseases by drug addiction and depression, neuropathic pain, inflammatory diseases by meningitis and encephalitis as
described above.
THE CLAIMS DEFINING THE INVENTION ARE AS FOLLOWS:

1. A tetrafluorobenzyl derivative represented by the following chemical formula 1:

```
[Chemical Formula 1]
```

wherein,

R₁, R₂, and R₃ are hydrogen or halogen,
R₄ is hydroxy, alkyl, alkoxy, halogen, alkoxy substituted with halogen, alkanoyloxy or nitro, and
R₅ is carboxyl acid, ester having C₁-C₄ alkyl, carboxyamide, sulfonic acid, halogen or nitro; or a pharmaceutically acceptable salt thereof.

2. The tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to claim 1, wherein R₁, R₂ and R₃ are halogen.

3. The tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to either claim 1 or claim 2, wherein R₁, R₂ and R₃ are F.
4. The tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to any one of claims 1 to 3, wherein \( R_4 \) is hydroxyl, alkyl, alkoxy or halogen.

5. The tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to any one of claims 1 to 3, wherein \( R_4 \) is hydroxyl or alkoxy.

6. The tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to any one of claims 1 to 3, wherein \( R_4 \) is hydroxy.

7. The tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to any one of claims 1 to 3, wherein \( R_4 \) is carboxyl, carboxamide, sulfonic acid or halogen.

8. The tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to any one of claims 1 to 3, wherein \( R_4 \) is carboxyl.

9. The tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to either claim 5 or claim 6, wherein \( R_4 \) is carboxyl.

10. The tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to claim 1, wherein the tetrafluorobenzyl derivative is
2-hydroxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethyl-benzylamino) -benzoic acid,
2-nitro-5-(2,3,5,6-tetrafluoro-4-trifluoromethyl-benzylamino) -benzoic acid,
2-chloro-5-(2,3,5,6-tetrafluoro-4-trifluoromethyl-benzylamino) -benzoic acid,
2-bromo-5-(2,3,5,6-tetrafluoro-4-trifluoromethyl-benzylamino) -benzoic acid,
2-hydroxy-5-(2,3,5,6-tetrafluoro-4-methyl-benzylamino) -benzoic acid,
2-methyl-5-(2,3,5,6-tetrafluoro-4-trifluoromethyl-benzylamino) -benzoic acid,
2-methoxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethyl-benzylamino) -benzoic acid,
5-(2,3,5,6-tetrafluoro-4-trifluoromethyl-benzylamino) -benzoic acid,
2-trifluoromethoxy benzoic acid,
2-nitro-4-(2,3,5,6-tetrafluoro-4-trifluoromethyl-benzylamino) phenol,
2-chloro-4-(2,3,5,6-tetrafluoro-4-trifluoromethyl-benzylamino) phenol,
2-hydroxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethyl-benzylamino) -benzamide,
2-hydroxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethyl-benzylamino) -benzenesulfonic acid,
methyl 2-hydroxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethyl-benzylamino) -benzoate,
2-ethanoyloxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethyl-benzylamino) -benzoic acid,
2-propanoyloxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethyl-benzylamino) -benzoic acid,
2-cyclohexan carbonyloxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethyl-benzylamino)-benzoic acid.

11. The tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to claim 2, wherein the tetrafluorobenzyl derivative is 2-hydroxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethyl-benzylamino) benzoic acid.

12. The tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to claim 2, wherein the tetrafluorobenzyl derivative is 2-hydroxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethyl-benzylamino)-benzene sulfonic acid.

13. The tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to claim 2, wherein the tetrafluorobenzyl derivative is 2-hydroxy-5-(2,3,5,6-tetrafluoro-4-trifluoromethyl-benzylamino)-benzoamide.

14. The tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to claim 2, wherein the tetrafluorobenzyl derivative is 2-chloro-4-(2,3,5,6-tetrafluoro-4-trifluoromethyl-benzylamino)-phenol.

15. A pharmaceutical composition, comprising a tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to any one of claims 1 to 14 and a pharmaceutically acceptable carrier.
16. A method of treatment or prevention of neurodegenerative diseases in a patient, comprising administering to the patient a tetrafluorobenzyl derivative or its pharmaceutically acceptable salt of any one of claims 1 to 14 as an effective ingredient, wherein the neurodegenerative diseases are selected from the group consisting of amyotrophic lateral sclerosis, Parkinson's disease, Huntington's disease, Alzheimer's disease, and combinations thereof.

17. A method for the treatment or prevention of epilepsy in a patient, comprising administering to the patient a tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to any one of claims 1 to 14 as an effective ingredient.

18. A method for the treatment or prevention of stroke in a patient, comprising administering to the patient a tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to any one of claims 1 to 14 as an effective ingredient.

19. A method for the treatment or prevention of traumatic brain injury, traumatic spinal cord injury or hydrocephalus in a patient, comprising administering to the patient a tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to any one of claims 1 to 14 as an effective ingredient.
20. A method for the treatment or prevention of glaucoma, retinopathy or macular degeneration in a patient, comprising administering to the patient a tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to any one of claims 1 to 14 as an effective ingredient.

21. A method for the treatment or prevention of drug addiction in a patient, comprising administering to the patient a tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to any one of claims 1 to 14 as an effective ingredient.

22. A method for the treatment or prevention of depression in a patient, comprising administering to the patient a tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to any one of claims 1 to 14 as an effective ingredient.

23. A method for the treatment or prevention of pain in a patient, comprising administering to the patient a tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to any one of claims 1 to 14 as an effective ingredient.

24. A method for the treatment or prevention of encephalitis, meningitis, and/or multiple sclerosis in a patient, comprising administering to the patient a tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to any one of claims 1 to 14 as an effective ingredient.
25. An N-methyl-D-aspartate (NMDA) receptor antagonist, comprising a tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to any one of claims 1 to 14 as an effective ingredient.

26. An antioxidant comprising a tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to any one of claims 1 to 14 as an effective ingredient.

27. A zinc toxicity inhibitor comprising a tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to any one of claims 1 to 14 as an effective ingredient.

28. Use of a tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to any one of claims 1 to 14 in the manufacture of a medicament for treatment or prevention of neurodegenerative diseases selected from the group consisting of amyotrophic lateral sclerosis, Parkinson's disease, Huntington's disease, Alzheimer's disease, and combinations thereof.

29. Use of a tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to any one of claims 1 to 14 in the manufacture of a medicament for treatment or prevention of epilepsy.
30. Use of a tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to any one of claims 1 to 14 in the manufacture of a medicament for treatment or prevention of stroke.

31. Use of a tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to any one of claims 1 to 14 in the manufacture of a medicament for treatment or prevention of traumatic brain injury, traumatic spinal cord injury or hydrocephalus.

32. Use of a tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to any one of claims 1 to 14 in the manufacture of a medicament for treatment or prevention of glaucoma, retinopathy or macular degeneration.

33. Use of a tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to any one of claims 1 to 14 in the manufacture of a medicament for treatment or prevention of drug addiction.

34. Use of a tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to any one of claims 1 to 14 in the manufacture of a medicament for treatment or prevention of depression.
35. Use of a tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to any one of claims 1 to 14 in the manufacture of a medicament for treatment or prevention of pain.

36. Use of a tetrafluorobenzyl derivative or its pharmaceutically acceptable salt according to any one of claims 1 to 14 in the manufacture of a medicament for treatment or prevention of encephalitis, meningitis and/or multiple sclerosis.

37. A derivative of claim 1, a composition of claim 15, a method of treatment of any one of claims 16 to 24, or a use of any one of claims 28 to 36, substantially as hereinbefore described.
FIGURES

FIG. 1a

Log [2-Hydroxy-TTBA], μM

[NMDA], 300μM

FIG. 1b

Log [2-Hydroxy-TTS, 2-Hydroxy-TTA, 2-Ethan-TTBA, 2-Propan-TTBA, 2-Cyclohexan-TTBA], μM

[NMDA], 300μM
FIG. 2a

A.

300 μM NMDA

0 μM

10
50
100
500
1 mM

Degree of Blockade (% of Control)

2.5
50
100

2-Hydroxy-TTBA (μM)

FIG. 2b

B.

300 μM NMDA

100 μM 2-Hydroxy-TTBA

300 μM NMDA

250 pA

5 s
FIG. 3a

Alone -1 -0.5 0 0.5 1 1.5 2

Log [2-Hydroxy-TTBA, trolox], μM

[Fe$^{2+}$], 50μM

FIG. 3b

Log [2-Hydroxy-TTB, 2-Hydroxy-TTA,
2-Chloro-TPP, 2-Ethan-TTBA, 2-Propan-TTBA
2-Cyclohexan-TTBA], μM

[Fe$^{2+}$], 50μM
FIG. 4

% LDH release vs. Log [2-Hydroxy-TTBA, Aspirin, Trolox]

FIG. 5

% LDH release vs. Log [2-Hydroxy-TTBA], μM

[Zn^{2+}], 100 μM
FIG. 6a

\[ \text{Log [2-Hydroxy-TTBA,Trolox], \mu M} \]

[DPPH], 100 \mu M

FIG. 6b

\[ \text{Log[2-Hydroxy-TTS, 2-Hydroxy-TTA, 2-Chloro-TTP], \mu M} \]

[DPPH], 100 \mu M
6/12

FIG. 7a

FIG. 7b

The number of degenerating cells

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<th>DH</th>
<th></th>
<th>VH</th>
<th>DH</th>
</tr>
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<td></td>
<td></td>
<td>2-Hydroxy-TTBA</td>
<td></td>
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* Significant difference
**FIG. 8a**

![Graph showing infarct volume change over time.](image)

**FIG. 8b**

![Graph showing infarct volume change with different doses.](image)
FIG. 8c

![Graph showing infarct volume (mm^3) over time with 2-HydroxyTTBA (5 mg/kg)](image)

FIG. 8d

![Graph showing infarct volume (mm^3) with 2-HydroxyTTBA (mg/kg)](image)
FIG. 8e

![Bar graph showing infarct volume (mm³) with control and 2-Hydroxy-TTBA (mg/kg) at different doses.](image-url)
FIG. 9a

10/12

Control 2-Hydroxy-TTBA

Pixel intensity/neuron

FIG. 9b

The number of degenerating neurons in CA1

Control 2-Hydroxy-TTBA
FIG. 10c

FIG. 11

Fluorescence Level

avg. pixel intensity/neuron

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<th>Control</th>
<th>2-Hydroxy-TTBA</th>
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</thead>
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<tr>
<td>25</td>
<td>15</td>
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DHN