

NON-SEDATING BARBITURATE COMPOUNDS AS NEUROPROTECTIVE AGENTS

BACKGROUND OF THE INVENTION

The invention relates to the use of non-sedating barbiturate compounds given in a manner and dosage effective to produce blood levels and brain levels of these drugs and/or their active metabolites sufficient to provide a neuroprotectant effect. In particular, the methods and formulations of the invention permit treatment of cerebral ischemia, head trauma and other acute neurologic injuries, and prevention of resulting neuronal damage.

Ischemia (stroke) is the third leading cause of death in the United States. When blood supply to the brain is reduced below a critical threshold, a cascade of biochemical events leads to irreversible damage to neurons and brain infarction. Research on treatment and prevention of ischemia is extensive but unfortunately it remains at a basic stage and no adequate therapies are yet in practice (10).

Barbiturates in high concentrations have been shown to be neuroprotective in cerebral ischemia in rodents and primates, to reduce the extent of ischemia brain infarction, and to prevent or lessen brain damage (1-4). One theory as to how barbiturates prevent neuronal injury in ischemia is that they inhibit the ischemia-induced uncontrolled release of neurotransmitters, which can attain high, neurotoxic concentrations that cause neuronal death (5).

The literature regarding the neuroprotective effects of anesthetic barbiturates is over two decades old, but the clinical use of barbiturates has been severely limited because of toxicity. The dosages and blood and brain levels necessary to confer neuroprotection are toxic and cause lethargy, stupor, and coma. Even higher doses that might be more effective are lethal (1-4, 6), making barbiturates unsuitable for treatment of ischemia (1). These toxic side effects establish a "functional ceiling" on dosage for barbiturates, and have discouraged further research into the use of anesthetic/sedative barbiturates to protect from ischemia.

Levitt et al., U.S. 4,628,056 describes non-sedating oxypyrimidine derivatives and their use as anticonvulsants, anti-anxiety and muscle relaxant agents. The literature does not suggest the use of such compounds as neuroprotectant agents. Indeed, even in published studies about using sedative barbiturates for neuroprotection there is no reference to non-

sedating barbiturate compounds. It is generally believed that the anticonvulsant and neuroprotective effects of barbiturates are linked to their sedative/hypnotic effects. For example, Lightfoote et al. suggested that the protective effects of pentobarbital are due to the duration of the barbiturate-induced anesthesia (3). This viewpoint has been reinforced by biochemical studies at the cell receptor level that relate all these effects to action at the GABA receptor. Thus, the prior art teaches away from using sedative barbiturates for neuroprotection because of their toxicity, and also teaches away from using non-sedative barbiturates as neuroprotectants because they lack sedating or anesthetic properties.

SUMMARY OF THE INVENTION

In summary, the invention involves non-sedating barbiturates such as for example 1,3-dimethoxymethyl 5,5-diphenyl-barbituric acid (DMMDPB), 1-monomethoxymethyl 5,5-diphenylbarbituric acid (MMMDPB) and diphenyl-barbituric acid (DPB) and their precursors, derivatives and analogs, and their administration over a range of dosages that result in a range of blood levels and brain levels of these drugs and their metabolites making them useful as neuroprotectants. In particular, the invention is directed to the treatment of cerebral ischemia, head trauma and other acute neurologic injuries, using non-sedating barbiturates.

There are many circumstances where individuals at risk of cerebral ischemia are clearly identified in advance, for example: individuals undergoing cardiac surgery or carotid endarterectomy, and individuals with atrial fibrillation, transient ischemic attacks (TIAs), bacterial endocarditis, strokes, or subarachnoid hemorrhage due to a cerebral aneurysm. In such cases, a non-sedating barbiturate is used prophylactically in individuals at risk for ischemic damage. The drugs can also be used after an acute event. These compounds can be given in oral form as a tablet, capsule, liquid or via intravenous or other parental routes.

This invention succeeds where previous efforts to treat cerebral ischemic attack with barbiturates have failed. This invention solves a problem previously thought to be insoluble, that of toxic effects of neuroprotective dosages of barbiturates. The invention avoids the toxicity and sedative effects of barbiturates known in the prior art without loss of efficiency.

This invention satisfies a long-felt need for a non-toxic neuroprotectant, and is contrary to the teachings of the prior art regarding the inability of barbiturates to produce clinically meaningful neuroprotection. According to the invention, it is possible to separate

the anticonvulsant and sedative effects of barbiturates, and neuroprotection correlates much better with the anticonvulsant rather than the sedative effect of barbiturates.

This invention differs from the prior art in the recognition of specific compounds, their modifications and dosages that are effective in neuroprotection but that were not
5 previously recognized.

The present invention is a method for providing neuroprotection to a mammal, preferably a human. The method comprises administering to the mammal a non-sedating barbiturate in a dose effective to provide a neuroprotection effect. Non-sedating barbiturates for use in the invention include one or more selected from the group consisting of 1,3-
10 dimethoxymethyl 5,5-diphenyl-barbituric acid (DMMDPB), 1-monomethoxymethyl 5,5-diphenylbarbituric acid (MMMDPB), and diphenyl barbituric acid (DPB). The precursors, derivatives and analogs of the foregoing compounds, as well as the salts of all the foregoing are also suitable for practicing the invention.

The effective neuroprotective dose of the non-sedative barbiturate preferably exceeds
15 the coma-producing dose of a sedative barbiturate. Depending on the specific need of the mammal, the dose of the non-sedative barbiturate may exceed a dose that would be lethal with a sedative barbiturate. This unexpected and seemingly paradoxical effect of the present method is further reflected in the relative dosage levels that are possible with the methods of this invention.

Also, the neuroprotective dose of the non-sedative barbiturate exceeds the minimum
20 anticonvulsant dosage of the barbiturate. In some embodiments of the present invention the effective dose of the non-sedative barbiturate is in the range of from about 2 times to about 5 times the anticonvulsant dosage. In yet other contexts where the need of the mammal requires, the effective dose of the non-sedative barbiturate is in the range of from about 5
25 times to about 10 times the anticonvulsant dosage of the non-sedative, or even higher so long as the dose is clinically acceptable.

Advantageously, the neuroprotective effect of the present methods can be used to mitigate the effect of cerebral ischemic. The non-sedating barbiturate can be administered orally, intravenously, transdermally, in combination with an adjuvant, or transpulmonarily
30 by means of a particulate or aerosol inhalant. Moreover, within the scope of the invention, the non-sedating barbiturate can be administered preventively, prophylactically or therapeutically, at a clinically acceptable dose. The compound may be administered prophylactically before evident neuronal damage, or therapeutically after onset of neuronal

damage. The neuroprotective effect diminishes, or protects the subject from, neuronal damage caused by head trauma or cerebral ischemia. The compound may be administered in conjunction with cardiac surgery or carotid endarterectomy. The mammalian subject may have or be at risk for atrial fibrillation, a transient ischemic attack (TIA), bacterial
5 endocarditis, a stroke, head trauma, or subarachnoid hemorrhage.

Typically, to achieve neuroprotection the non-sedating barbiturate is administered in a dose sufficient to obtain blood concentrations of at least about 30 µg/ml of barbiturate, preferably at least about 100 µg/ml, more preferably at least about 250 µg/ml, and possibly as high as 200-300 µg/ml, or even higher. In contrast, the reported therapeutic range for
10 phenobarbital is lower, 10-30 µg/ml blood levels (6). Thus, preferred ranges are at or above about 25, 30, 50, 75, 100, 200, 250, or 300 µg/ml.

The invention includes a pharmaceutical composition comprising a non-sedating barbiturate administered in an amount effective to have a neuroprotectant effect. Preferably, the non-sedating barbiturate is administered in oral doses in the range of from about 25 to
15 about 1,500 mg/kg/day body weight. Preferably the dose is greater than about 25 mg/kg/day, or greater than about 100 mg/kg/day, or greater than 250 mg/kg/day. A preferred dose is one that is pharmacologically equivalent to a dose of about 1000 mg/kg/day in the rat. Thus, dosage forms may be sufficient individually or in multiple doses to provide a dose equal to or above about 15, 20, 25, 50, 70, 100, 250, 500, 1000, or 1500
20 mg/kg body weight per day.

In human trials it has been unexpectedly found that DMMDPB, one of the neuroprotectant compounds, is much better absorbed in humans than in rats or dogs. It has further been found that the half life of DMMDPB, as well as the half life of MMMDPB and DPB are greater than the half-lives found in rats or dogs. Specifically, with dosages of 20
25 mg/kg/day, the half-lives of DMMDPB, MMMDPB, and DPB are approximately 20hrs, 20hrs, and 50hrs respectively after a two week exposure in humans. Similarly, the maximum concentration (C_{max}) of the drug in the blood following 7 days of dosing in the range of 20 mg/kg/day are 1.2 µg/ml, 36 µg/ml and 43 µg/ml respectively.

The unexpectedly high absorption and prolonged half-life in humans makes it
30 possible to achieve substantial blood levels with lower than expected oral dosages. Thus, for example, it is possible to obtain total barbiturate blood levels (i.e., DMMDPB + MMMDPB + DPB) > 53 µg/ml with dosages of about 15 mg/kg/day; and total barbiturate levels > 72 µg/ml with dosages in the range of 20 mg/kg/day. Blood levels of non-sedating barbiturates

greater than 100 µg/ml are achieved with dosages between about 40 and about 100 mg/kg/day, and are within the scope of the invention. With parenteral administration of non-sedating barbiturates, similar blood concentrations are obtained with daily dosages of less than 25 mg/kg/day. However, first day loading dosages may still need initial dosages of
5 greater than 25 mg/kg.

The invention provides an article of manufacture comprising a container comprising a pharmaceutical composition and a label with indications for use as a neuroprotectant, the pharmaceutical composition comprising a non-sedating barbiturate compound in an amount effective for neuroprotection upon administration to a subject in need of neuroprotection;
10 and a pharmaceutically acceptable carrier or excipient.

Another embodiment is a method for providing neuroprotection comprising (a) diagnosing a patient's need for cerebral neuroprotection, (b) selecting a non-sedative barbiturate, and (c) providing to the patient a dose of the non-sedative barbiturate sufficient to raise the concentration in the patient's brain to a level effective to provide
15 neuroprotection.

Further objectives and advantages will become apparent from a consideration of the description and examples.

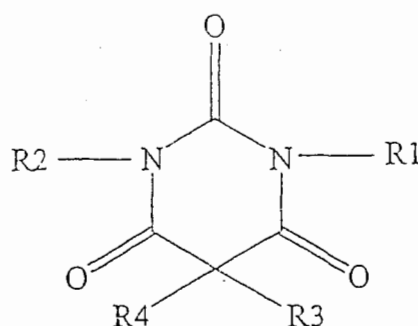
DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

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In describing preferred embodiments of the present invention, specific terminology is employed for the sake of clarity. However, the invention is not intended to be limited to the specific terminology so selected. It is to be understood that each specific element includes all technical equivalents which operate in a similar manner to accomplish a similar purpose.
25 Each reference cited here is incorporated by reference as if each were individually incorporated by reference.

The term "non-sedative barbiturate" encompasses the family of 5,5-diphenyl barbituric acid anticonvulsant compounds described in Levitt et al., U.S. 4,628,056, and metabolic precursors and metabolites, and derivatives and structural analogs (including
30 addition salts thereof) having a non-sedative neuroprotectant activity. Other barbituric acid derivatives that are non-sedating are also within the scope of the invention.

Derivatives, precursors, and analogs of barbituric acid include barbituric acids of the formula:



wherein one or more nitrogen is substituted with lower alkyl, or a lower alkoxy substituted lower alkyl group; or at least one of R^1 and R^2 , together with the nitrogen, forms a carbamate, an amide, or an acetal of the formamide derivative, i.e. R^1 or R^2 is CO_2R , COR or $\text{CH}(\text{OR})_2$. Methylether groups are preferred R^1 and R^2 groups and methoxymethyl is more preferred. Methyl is also a preferred value for R^1 and/or R^2 . Other derivatives of barbituric acids according to the invention are carbamates, amides and acetals where one or both of R^1 and R^2 is CH_2OR^5 , wherein R^5 is lower alkyl, alkylaryl or benzyl; CO_2R^6 , wherein R^6 is lower alkyl or aryl; COR^7 , wherein R^7 is hydrogen, lower alkyl or aryl; or $\text{CH}(\text{OR}^8)_2$, wherein R^8 is a lower alkyl group.

Preferred values for R^3 and R^4 are aryl, phenyl, phenyl substituted with a halogen or lower alkyl group, benzyl, benzyl wherein the aromatic ring is substituted with a halogen or lower alkyl group, lower alkyl or lower alkyl substituted with an aromatic moiety. Aryl represents any carbocyclic ring, such as phenyl, naphthyl and higher analogues, as well as heteroaromatic rings substituted with one or more heteroatoms such as sulfur, oxygen and nitrogen. According to the invention, nonsedating barbituric acid derivatives are those where at least one of R^3 and R^4 is an aromatic ring or an aromatic ring containing moiety e.g. aryl, phenyl, substituted phenyl, benzyl, substituted benzyl or arylalkyl. Preferred substituents on the aromatic rings are methyl, ethyl, and fluorine. Phenyl and substituted phenyl are preferred for R^3 and R^4 . Embodiments where R^3 and R^4 are both phenyl are most preferred.

In preferred compounds, one of R^1 and R^2 is hydrogen, or one or both of R^1 and R^2 is methyl or alkoxyethyl, preferably methoxymethyl. At least one and preferably both of R^3 and R^4 is preferably phenyl or substituted phenyl, tolyl, fluorophenyl, ethylphenyl.

As can be readily understood, salts of the above compounds are also contemplated, including organic salts, such as acid addition and base addition salts.

In order to fall within the scope of this genus, the compound must (1) be a barbituric acid chemical derivative, (2) not be sedating, in the sense that the subject remains awake and alert at useful doses, that is, not anesthetized, and (3) manifest neuroprotective activity in an animal model described herein or in a human at a dose that is not toxic to the relevant animal species, or show activity in an *in vitro* assay now known or later discovered that is accepted as a model for *in vivo* neuroprotection.

These barbituric acid derivatives may be both prodrugs and active ingredients in the subject, thus combining to produce the desired pharmacodynamic effect of neuroprotection. Sustained levels are readily obtained with such compounds.

Thus, certain barbiturate compounds have been developed and have anticonvulsant activity without being sedating even at very high brain concentrations (that would be lethal with other barbiturates). According to the invention, such compounds are used to neuroprotect an animal at risk for or suffering from one or more ischemic episodes such as that modeled by middle cerebral artery occlusion, while these compounds do not cause the toxic effects of other barbiturates when present at concentrations required for prevention of ischemic brain damage.

As described herein, non-sedative barbiturate drugs lessen or prevent ischemic brain damage in a rat model of focal cerebral ischemia produced by middle cerebral artery occlusion. This demonstrates utility in humans.

In a reproducible, predictive model of cerebral ischemia known in the art, selective neuronal damage is produced in the striatum and cerebral cortex by bilateral carotid occlusion accompanied by systemic hypotension. The resulting cerebral ischemia causes a release of excitotoxic neurotransmitters and dopamine in striatum. Pentobarbital inhibited this ischemia-induced release, pointing to one possible mechanism of barbiturate neuroprotection. (5) A neuroprotective dose of pentobarbital was found to be 70 mg/kg. Inhibition of neurotransmitter release by several neuroprotective anesthetic agents (isoflurane, etomidate, propofol) was also known.

The above and similar animal models (see Examples) can be used

- (1) to analyze whether a non-sedative barbiturate with anticonvulsant properties but little or no anesthetic activity can provide neuroprotection in the striatum or hippocampus, and
- (2) to determine if the agent prevents or reduces release of neurotransmitters in response to ischemia. Uncontrolled or unmodulated neurotransmitter release is one of the

postulated mechanism of ischemic damage. For non-sedating barbiturates that inhibit release of neurotransmitters, this approach can serve as a biochemical assay for predicting utility of a compound according to the invention, and the invention encompasses such methods.

5 A neuroprotective effect according to the invention can be demonstrated and characterized by performing a dose-response study and measuring statistically significant differences in neuronal damage at the various doses of the drug. Dose-response curves generated in such studies can be used to compare the relative degree of neuroprotection and sedation of a test compound.

10 Cerebral ischemia is induced in rats by occlusion of the middle cerebral artery ("MCA") (7-9). The occlusion can be performed in an irreversible or reversible manner. In the latter case, after a period of obstruction, blood flow is restored. These animal preparations are thus appropriate to model various types of strokes in humans and to permit determination of a drug's neuroprotective action. Such models permit observation of the
15 prevention of brain damage and the evaluation of the drugs as being useful for humans who are at risk for ischemic stroke for reduction of subsequent brain damage induced by an ischemia event. Because they prevent brain damage in models of irreversible ischemia and reversible ischemia with restoration of blood flow, the compounds of the invention are also useful for treating acute ischemic stroke either alone or in combination with other agents, for
20 example, thrombolytics such as tissue plasminogen activator that reduce the extent of brain infarction when circulation is restored.

The term "treatment" is intended to encompass administration of compounds according to the invention prophylactically to prevent or suppress an undesired condition, and therapeutically to eliminate or reduce the extent or symptoms of the condition.
25 Treatment according to the invention is given to a human or other mammal having a disease or condition creating a need of such treatment. Treatment also includes application of the compound to cells or organs *in vitro*. Treatment may be by systemic or local administration.

The non-sedative barbiturate compositions of the present invention, may be formulated into "pharmaceutical compositions" with appropriate pharmaceutically
30 acceptable carriers, excipients or diluents. If appropriate, pharmaceutical compositions may be formulated into preparations including, but not limited to, solid, semi-solid, liquid, or gaseous forms, such as tablets, capsules, powders, granules, ointments, solutions,

suppositories, injections, inhalants, and aerosols, in the usual ways for their respective route of administration.

Methods known in the art can be used to achieve time-release of the composition or to prevent metabolism, release or absorption of the composition until it has reached its intended target site. A pharmaceutically-acceptable formulation should be employed that does not inactivate the active drug of the present invention.

In pharmaceutical dosage forms, the compositions may be used alone or in appropriate association, as well as in combination with, other pharmaceutically-active compounds.

The pharmaceutical compositions of the present invention can be delivered via various routes and to various sites in an animal body to achieve the desired neuroprotective effect. Local or systemic delivery can be accomplished by injection, infusion, application or instillation of the composition into one or more body cavities, or by inhalation or insufflation of an aerosol. Parenteral administration can be by intramuscular, intravenous, intraperitoneal, subcutaneous intradermal, or topical administration.

The compositions of the present invention can be provided in unit dosage form, wherein each dosage unit, e.g., a teaspoon, tablet, solution, or suppository, contains a predetermined amount of the active drug or prodrug, alone or in appropriate combination with other pharmaceutically-active agents. The term "unit dosage form" refers to physically discrete units suitable as unitary dosages for human and animal subjects, each unit containing a predetermined quantity of the composition of the present invention, alone or in combination with other active agents, calculated in an amount sufficient to produce the desired effect, in association with a pharmaceutically-acceptable diluent, carrier (e.g., liquid carrier such as a saline solution, a buffer solution, or other physiological aqueous solution), or vehicle, where appropriate. The specifications for the novel unit dosage forms of the present invention depend on the particular effect to be achieved and the drug's particular pharmacodynamics in the particular host.

An "effective amount" of the composition is that required to produce the desired pharmacologic effect in a host. This can be monitored using any of a number of end-points known to those skilled in the art. The "effective dose" will depend on the bioavailability of specific dosage forms delivered by one or another route of administration. The neuroprotective dosage and blood level of the present compounds is at least 2-fold and preferably at least about 5 to 10-fold the anticonvulsant dosage of a sedating barbiturate.

Based on rat data, the anticonvulsant ED₅₀ for phenobarbital is about 50-100 mg/kg. A non-sedating barbiturate dose of 1 g/kg given over 7 days protects against cerebral ischemia in the rat. Similar or lower doses are suitable in humans based on the enhanced absorption in humans discussed above.

5 The amount of each active agent employed in the Examples below provides general guidance for the range that can be utilized by the skilled practitioner to optimize the doses and methods of the present invention. Moreover, such dose ranges do not preclude use of a higher or lower dose of a component, as might be warranted in a particular application. For example, the actual dose and schedule may vary depending on whether the compositions are
10 administered in combination with other drugs, or depending on inter-individual differences in pharmacokinetics, drug disposition, and metabolism. Similarly, amounts may vary for *in vitro* applications. It is within the skill in the art to readily adjust the dose in accordance with the necessities of a particular situation without undue experimentation

 Having now generally described the invention, the same will be more readily
15 understood through reference to the following examples which are provided by way of illustration, and are not intended to be limiting of the present invention, unless specified.

EXAMPLE 1

General Design

 The non-sedative barbiturate (NSB) drug is tested in rats exposed to either reversible
20 or irreversible ischemia. Varying doses of drug are administered. The neuroprotective effect is compared to a negative control (placebo) and a positive control, pentobarbital, a known neuroprotective but sedative barbiturate, given at doses known to reduce infarct volume in cerebral ischemia (1-4).

 Animals are sacrificed several days after the onset of the ischemic insult and the
25 brains examined to determine the volume of brain infarction as an outcome measure of the drug's reduction of ischemic brain damage. The animals are examined clinically and graded prior to sacrifice to determine if the drug has conferred any beneficial effect on relevant functions following ischemic "stroke."

 Four experimental models are preferred for testing the neuroprotective effects of the
30 NSB drug. See Ginsberg, M.D., "Animal Models of Global and Focal Cerebral Ischemia," Chapter 34 in Welsh, K.M.A *et al.*, *Primer on Cerebrovascular Diseases*, Academic Press, New York, 1997; and Pulsinelli WA, Brierley JB. A new model of bilateral hemispheric ischemia in the unanesthetized rat, *Stroke* 1979 May - June 10(3): 267-72.

1. Irreversible ischemia produced by middle cerebral artery (MCA) occlusion ;
2. Reversible ischemia produced by MCA occlusion;
3. Transient global ischemia produced by cross-clamping the aorta for a defined interval; and
4. Transient global ischemia produced by cauterizing both vertebral arteries and reversibly clamping the common carotid arteries.

In each experimental model, groups of rats are treated with either:

1. Negative control (placebo) via nasogastric (NG) tube;
2. Positive control: intraperitoneal (IP) dose of 70 mg/kg pentobarbital; or
3. The NSB compound DMMDPB (or a compound being tested for its utility in the present invention) via NG tube at doses between 500 mg/kg and 1500 mg/kg for 7 days prior to experimental infarctions.

The results are compared.

EXAMPLE 2

Irreversible Cerebral Ischemia

Irreversible MCA occlusion was produced by ligating the carotid artery and then inserting a filament into the origin of the MCA with the animal maintained under halothane anesthesia. Blood flow in the MCA was measured by laser doppler and those animals in which a significant drop in blood flow occurred were considered to have experienced cerebral ischemia, and to be at risk for subsequent damage (*i.e.*, a stroke). Indeed, no clinical strokes occurred in animals that did not experience a precipitous drop in MCA blood flow. All the animals showing with a drop in MCA blood flow experienced strokes.

Animals at risk were then followed behaviorally and scored by clinical findings using the Bederson grading scale as either:

- | | |
|---|-----------------------|
| 0 | no evidence of stroke |
| 1 | mild stroke |
| 2 | moderate stroke |
| 3 | severe stroke |

Those animals that survived for three days were sacrificed and their brains examined. Animals to be sacrificed were given chloral hydrate (35 mg/kg IP, and their brains were fixed by intracardiac perfusion with heparinized 0.9% saline followed by 10% buffered formalin. The brains were removed from the cranial vault with care to leave the arachnoid intact with the intracranial vessels underneath. The fixed brains were frozen at 80°C. Coronal sections 20 μ m thick were cut at 400 μ m intervals in a cryostat at -20°C, dried on a hot plate at 60°C, fixed in 90% ethanol for 10 minutes and stained with hematoxylin and eosin (7). Infarcted brain is pale compared to the rest of the brain. The amount of infarcted brain was determined by microscopic inspection of the brain sections and calculation of infarct volumes in mm³.

The results are shown in Tables 1 and 2 below. The numbers vary between groups because not all animals experienced a drop in MCA blood flow with the procedure. All animals were treated with DMMDPB dosages of 1000 mg/kg/day for 7 days.

TABLE 1

Effect of DMMDPB on Death due to Cerebral Ischemia

Treatment Group	Behavior	n	Death within 24 hr	Survival at 24 hr n (%)	Survival at 48 hr n (%)	Survival at 72 hr n (%)
Control (males)	Sedated	12	9 (75%)	3 (25%)	2 (17%)	1 (8%)
Pentobarbital (males)	Sedated	9	6 (67%)	3 (33%)	3 (33%)	3 (33%)
DMMDPB	Not sedated	17	2 (12%)	15 (88%)	10 (59%)	8 (47%)
Males		14	2 (24%)	12 (76%)	7 (50%)	5 (36%)
Females		3	0 (0%)	3 (100%)	3 (100%)	3 (100%)

Other dose ranging studies in rats treated with DMMDPB for 7 days established that female rats have substantially higher blood levels than male rats. Specifically, at a dosage of DMMDPB of 500 mg/kg the total barbiturate level in males was 59 μ g/ml and females 170 μ g/ml. At a dosage of 1000 mg/kg the total barbiturate level in males was 77 μ g/ml and females 227 μ g/ml; and at a dosage of 2000 mg/kg the total barbiturate level in males was 110 μ g/ml and females 328 μ g/ml. Thus females consistently had blood levels 250%-300% that of males at the same dosage. This data shows a type of "dose response effect" or "blood

level response effect" whereby higher blood levels correlate to higher survival in female rats in the results tabulated above.

TABLE 2

Neurologic status of the first 9 animals of Table 1

Treatment group	Rat#	Weight (g)	Neurologic Status (Bederson grading scale 0-3)			Pathology
			Day 1	Day 2	Day 3	
Placebo	1	260	3	X		Died 24 hrs
	2	260	3	X		Died 24 hrs
	3	240	3	X		Died 24 hrs
Pentobarbital	1	260	0	1	1	SAH (autopsy)
	2	250	2	2	2	Brain collected
DMMDPB	1	270	1	1	1	Brain collected
	2	230	3	3	X	Died 48 hrs
	3	240	2-3	3	X	Died 48 hrs
	4	260	2-3	3	3	Brain collected

Pathology (visual and microscopic examination) shows smaller infarct volumes in animals pretreated with pentobarbital and DMMDPB.

Thus, DMMDPB proved to protect the animals against death. Other data indicated that DMMDPB treated animals did not manifest sedation compared to placebo group. In contract, the pentobarbital animals were anesthetized and immobile. The neuroprotective effects at non-sedating doses were comparable to or better than the effects of the sedative pentobarbital but without the side effects-of sedation, particularly at day two.

These neuroprotective effects of DMMDPB are predictive for monomethoxymethyl diphenyl barbituric acid (MMMDPB) and the presumptive pharmacologically active chemical moiety diphenyl barbituric acid (DPB), which are metabolic products of DMMDPB. Indeed, in animal studies over periods ranging from 1-30 days DMMDPB was rapidly metabolized to MMMDPB and eventually to DPB.

Results from clinical studies with humans demonstrated a pattern of blood levels similar to that seen in animals: DPB>MMMDPB>DMMDPB. Again, the same pattern was shown in that blood levels of DMMDPB were minimal, while MMMDPB and DPB concentration was higher. This animal model of neuroprotection is predictive for humans because: (a) the metabolic behavior of this compound in animals is predictive of human metabolism, and (b) the anticonvulsant activity in animals correlates with anticonvulsant activity in humans.

Although several sedative barbituric compounds previously found to be neuroprotective in such animal models provided some benefit in human studies, their use over even relatively short time periods is precluded by their sedative and other neurological and psychological side-effects. These side effects make prophylactic treatment infeasible for patients identified as being at high risk of stroke. According to the present invention, in contrast, the NSBs have minimal side effects in humans. Thus, it is now established that diphenyl barbituric acid and its precursors, analogues and derivatives constitute a class or family of compounds suitable for neuroprotection of humans.

Example 3

Reversible Cerebral Ischemia Model

Rats are pretreated as in Example 1 (above) and a similar procedure is performed except that the filament occluding the MCA is removed after 30 to 60 minutes, restoring blood flow through the MCA. Rats are then followed clinically for three days, graded for their degree of stroke and then sacrificed as in Example. The brains are removed and examined as described above.

The NSB compounds are shown to be neuroprotective under these conditions.

Example 4

Rats are pretreated as in Example 1 (above) and then, during ether anesthesia, the rats' vertebral arteries are electrocauterized through the alar foramina of the first cervical

vertebra. Reversible clamps are then placed loosely around the common carotid arteries. After 24 hours, working with awake rats, the carotid clasps are tightened to produce 4-vessel occlusion. Following 10-30 minutes of 4-vessel occlusion, the clasps are removed and 72 hours later the animals sacrificed by perfusion fixation. Untreated rats routinely
5 demonstrate ischemic neuronal damage after 20 or 30 minutes of 4-vessel occlusion. Multiple areas of the forebrain, including the H1 and paramedian hippocampus, striatum, and posterior neocortex are evaluated. The non-sedating barbiturates are shown to be neuroprotective under these circumstances.

The embodiments illustrated and discussed in this specification are intended only to
10 teach those skilled in the art the best way known to the inventors to make and use the invention. Nothing in this specification should be considered as limiting the scope of the present invention. The above-described embodiments of the invention may be modified or varied, and elements added or omitted, without departing from the invention, as appreciated by those skilled in the art in light of the above teachings. It is therefore to be understood
15 that, within the scope of the claims and their equivalents, the invention may be practiced otherwise than as specifically described.

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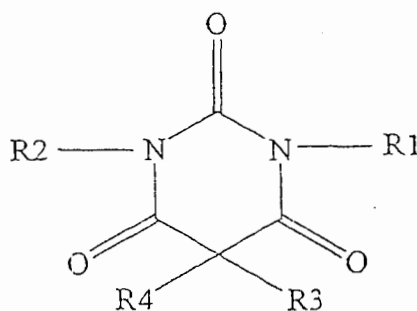
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WHAT IS CLAIMED IS:

1. A method for providing neuroprotection in a mammal at risk for cerebral ischemia, comprising administering to the mammal an effective neuroprotective dose of a non-sedating barbiturate.

2. The method of claim 1, wherein the non-sedating barbiturate is selected from the group consisting of 1,3-dimethoxymethyl 5,5-diphenyl-barbituric acid (DMMDPB), 1-monomethoxymethyl 5,5-diphenylbarbituric acid (MMMDPB), diphenyl barbituric acid (DPB), a precursor thereof, a derivative thereof, an analog, thereof, and salts thereof.

3. A method for providing neuroprotection to a mammal in need of neuroprotection, comprising administering to the mammal a non-sedating barbiturate in a dose sufficient to provide a neuroprotective effect, the non-sedating barbiturate having the structure:



wherein R^1 and R^2 may be the same or different and are independently hydrogen; lower alkyl, optionally substituted by lower alkoxy; CH_2OR^5 , wherein R^5 is lower alkyl, alkylaryl or benzyl; CO_2R^6 , wherein R^6 is lower alkyl or aryl; COR^7 , wherein R^7 is hydrogen, lower alkyl or aryl; or $CH(OR^8)_2$, wherein R^8 is a lower alkyl group; and

wherein R^3 and R^4 may be the same or different and are independently hydrogen; aryl optionally containing one or more heteroatoms selected from the group consisting of N, S and O; phenyl; phenyl substituted with a halogen or lower alkyl group; benzyl; ring substituted benzyl having one or more halogens, lower alkyl groups or both; lower alkyl; or lower alkyl substituted with an aromatic moiety;

provided that at least one of R^1 and R^2 is an aromatic ring containing moiety;
and salts thereof.

4. The method of claim 3, wherein R^1 and R^2 may be the same or different and
are independently phenyl; phenyl substituted with a halogen or lower alkyl group; benzyl;
5 ring substituted benzyl having one or more halogens, lower alkyl groups or both; lower
alkyl; or lower alkyl substituted with an aromatic moiety; provided that at least one of R^1 and
 R^2 is phenyl or substituted phenyl.

5. The method of any of claims 1-4, wherein said dose is higher than about 50
10 mg/kg body weight.

6. The method of claim 1, wherein said dose is higher than about 70 mg/kg body
weight.

7. The method of claim 1, wherein said dose is higher than about 100 mg/kg
15 body weight.

8. The method of claim 1, wherein said dose is higher than about 250 mg/kg
body weight.

9. The method of claim 1, wherein said dose is higher than about 500 mg/kg.

10. The method of claim 1, wherein the compound is administered at a dose of at
least about 1000 mg/kg body weight.

11. The method of any of claims 1-4, wherein the compound is administered in a
dose sufficient to produce blood concentrations of at least about 30 $\mu\text{g/ml}$ of said compound
or an active metabolite thereof.

12. The method of claim 1, wherein the compound is administered in a dose
sufficient to achieve blood levels of at least about 50 $\mu\text{g/ml}$ of barbiturate of said compound
or an active metabolite thereof.

13. The method of claim 1, wherein the compound is administered in doses sufficient to produce blood concentrations of at least about 100 µg/ml of said compound or an active metabolite thereof.

5 14. The method of claim 1, wherein the compound is administered in doses sufficient to produce blood concentrations of at least about 1,000 µg/ml of said compound or an active metabolite thereof.

10 15. The method of any of claims 1-4, wherein the compound is administered orally.

16. The method of any of claims 1-4, wherein the compound is administered intravenously.

15 17. The method of any of claims 1-4, wherein the compound is administered in combination with an adjuvant.

18. The method of any of claims 1-4, wherein the compound is administered prophylactically before evident neuronal damage.

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19. The method of any of claims 1-4, wherein the compound is administered therapeutically after onset of neuronal damage.

20. The method of claim 3, wherein the neuroprotective effect diminishes, or
25 protects the subject from, neuronal damage caused by head trauma or cerebral ischemia.

21. A method according to claim 1, wherein the compound is administered in conjunction with cardiac surgery or carotid endarterectomy.

30 22. A method according to claim 1, wherein the mammal has or is at risk for atrial fibrillation, a transient ischemic attack (TIA), bacterial endocarditis, a stroke, head trauma, or subarachnoid hemorrhage.

23. A method of providing neuroprotection comprising
- (a) identifying a subject in need of cerebral neuroprotection,
 - (b) selecting a non-sedative barbiturate compound capable of neuroprotecting said subject, and
 - (c) administering to said subject an effective neuroprotective dose of said compound, thereby providing said neuroprotection.