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**United States Patent** [19][11] **Patent Number:** 5,378,729

Kohn et al.

[45] **Date of Patent:** Jan. 3, 1995[54] **AMINO ACID DERIVATIVE  
ANTICONVULSANT**[75] **Inventors:** Harold L. Kohn, Houston; Darrell  
Watson, Belton, both of Tex.[73] **Assignee:** Research Corporation Technologies,  
Inc., Tucson, Ariz.[21] **Appl. No.:** 710,610[22] **Filed:** Jun. 4, 1991**Related U.S. Application Data**[63] Continuation-in-part of Ser. No. 354,057, May 19,  
1989, abandoned, and Ser. No. 392,870, Aug. 11, 1989,  
abandoned, which is a continuation of Ser. No. 80,528,  
Jul. 31, 1987, abandoned, which is a continuation-in-  
part of Ser. No. 916,254, Oct. 7, 1986, abandoned,  
which is a continuation-in-part of Ser. No. 702,195,  
Feb. 15, 1985, abandoned, said Ser. No. 354,057, is a  
continuation-in-part of Ser. No. 80,528, Feb. 15, 1985.[51] **Int. Cl.<sup>6</sup>** ..... A61K 31/535; A61K 31/445;  
C07D 211/72; C07D 261/04[52] **U.S. Cl.** ..... 514/231.2; 514/315;  
514/397; 514/406; 514/415; 514/424; 514/461;  
514/468; 514/486; 546/292; 548/371.4;  
548/245; 564/148; 564/152; 564/154[58] **Field of Search** ..... 564/148, 155, 154, 152;  
548/616, 245, 371.4; 514/461, 548, 549;  
546/292[56] **References Cited****U.S. PATENT DOCUMENTS**

2,676,188	4/1954	Bruce et al.	564/155
2,721,197	10/1955	Sheehan	564/155
3,340,147	9/1967	Martin et al.	514/616
3,657,341	4/1972	Thorne et al.	260/558 A
3,707,559	12/1972	Mazur et al.	564/158
4,018,826	4/1977	Gless, Jr. et al.	564/215
4,260,684	4/1981	Schult	564/155
4,303,673	12/1981	Biedermann et al.	564/155
4,513,009	4/1985	Roques et al.	564/155
4,595,700	6/1986	Donald et al.	514/616
4,618,708	10/1986	Roques et al.	564/154
4,873,241	10/1989	Napier et al.	514/237.8

**FOREIGN PATENT DOCUMENTS**

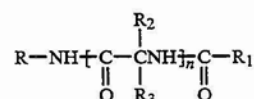
0885303	3/1981	Belgium
0194464	2/1980	European Pat. Off.
0007441	10/1980	European Pat. Off.
0038758	10/1981	European Pat. Off.
0042626	12/1981	European Pat. Off.
0046707	3/1982	European Pat. Off.
0263506	10/1987	European Pat. Off.
0400400	5/1990	European Pat. Off.
1927692	12/1969	Germany
0393355	10/1965	Switzerland
1051220	12/1966	United Kingdom

**OTHER PUBLICATIONS**Chemical Abstracts, vol. 92; No. 7:51712r (Feb. 18,  
1990).Chemical Abstracts, vol. 96; No. 5:35710r (Feb. 1,  
1982).Chemical Abstracts, vol. 101; No. 9; 72124v (Aug. 27,  
1984).Chemical Abstracts, vol. 91; No. 21:175147; (Nov. 19,  
1979).

(List continued on next page.)

*Primary Examiner*—Marianne M. Cintins*Assistant Examiner*—T. Criares*Attorney, Agent, or Firm*—Scully, Scott, Murphy &  
Presser[57] **ABSTRACT**

The present invention relates to compounds exhibiting central nervous system (CNS) activity which are useful in the treatment of epilepsy and other CNS disorders. The compounds of this invention have the following general formula:



and pharmaceutically acceptable salts thereof.

**150 Claims, No Drawings**

## OTHER PUBLICATIONS

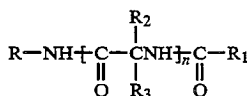
- Kohn, et al. (1988) *Brain Research* 457: 371-375, Marked Sterospecificity in a New Class of Anticonvulsants.  
Chemical Abstracts, vol. 97; 145266d (1982).
- White, et al. (1981) *JACS*, 103:4231-4239, Active-Site-Directed Inhibition of alpha-Chymotrypsin by Deaminatively Produced Carbonium Ions: An Example of Suicide of Enzyme-Activated-Substrate Inhibition.
- Legall, et al. (1988) *Int. J. Protein Res.*, 12:279-291 Synthesis of Functionalized Non-Natural Amino Acid Derivatives via Amidoalkylation Transformations.
- Conley, et al. (1987) *J. Med. Chem.*, 30(3):574-580, Functionalized DL-Amino Acid Derivatives, Potent New Agents for the Treatment of Epilepsy.
- Garcia, et al. (1984) *Tetrahedron Letters*, 25(42) 4841-4844, New Synthetic "Tricks" Triphenylphosphine-Mediated Amide Formation from Carboxylic Acids and Azides.
- Rebek, et al. (1979), *J. Am. Chem. Soc.*, 101(3):737, On the Rate of Site-Site Interactions in Functionalized Polystyrenes.
- Cortes, et al. (1985) *J. Med. Chem.*, 28:601-606, Effect of Structural Modification of the Hydantion Ring on Anticonvulsant Activity.
- Ikeda, et al. (1977) *Tetrahedron*, 33(5): 489-495, Photochemical Synthesis of 1,2,3,4-Tetrahydroisoquinolin-3-ones from N-Chloroacetylbenzylamines.
- Katritzky, et al. (1990) *J. Org. Chem.* 55: 2206-2214, Benzotriazole-Assisted Synthesis of Monoacyl Animals and Their Peptide Derivatives.
- Lipshutz et al. (1983) *J. Am. Chem. Soc.* 105; 7703-7713, Heterocycles as Masked Diamide/Dipeptide Equivalents, Formation and Reactions of Substituted 5-(Acylamino) Oxazoles as Intermediates en Route to the Cyclopeptide Alkaloids.
- Lipshutz et al. (1983) *J. Org. Chem.* 48:3745-3750, An Approach to the Cyclopeptide Alkaloids (Phenylcyclopeptides) via Heterocyclic Diamide/Dipeptide Equivalents, Preparation and N-Alkylation Studies of 2,4(5)-Disubstituted Imidazoles.
- Rogues, (1987) 193rd ACS National Meeting Amer. Chem. Society, Apr. 15-10, 1987, Use of Various Metallopeptidase Inhibitors to Study the Physiological Rate of Endogenous Neuropeptides.
- Kohn, et al. (1990) *J. Med Chem.* 33:919-926, Preparation and Anticonvulsant Activity of a Series of Functionalized  $\beta$ -Aromatic and  $\alpha$  Heteroaromatic Amino Acids.
- Lipshutz et al. "Heterocycles in Synthesis . . . Imidazoles" *Journal of the American Chemical Society*, vol. 106, No. 2, pp. 457-459 CA 102(19): 160030n (1985).

## AMINO ACID DERIVATIVE ANTICONVULSANT

This invention was made with Government support under NS15604 awarded by the National Institutes of Health. The Government has certain rights in the invention.

The present application is a continuation-in-part of copending U.S. patent application Ser. No. 07/354,057 filed on May 9, 1989 and a CIP of U.S. patent application Ser. No. 07/392,870 filed on Aug. 11, 1989 both now abandoned. U.S. patent application Ser. No. 07/354,057 filed on May 19, 1989, now abandoned being a continuation-in-part of U.S. patent application having Ser. No. 07/080,528, filed on Jul. 31, 1987 now abandoned, which is a continuation-in-part of U.S. patent application Ser. No. 06/916,254, filed Oct. 7, 1986, now abandoned which is a continuation-in-part of U.S. patent application Ser. No. 06/702,195, filed Feb. 15, 1985 now abandoned said U.S. patent application Ser. No. 07/392,870 filed Jul. 11, 1989, abandoned being a continuation application of U.S. patent application having Ser. No. 07/080,528, filed on Jul. 31, 1987, now abandoned, which is a continuation-in-part of U.S. patent application Ser. No. 06/916,254, filed Oct. 7, 1986, now abandoned, which is a continuation-in-part of U.S. patent application Ser. No. 06/702,195 filed on Feb. 15, 1985 now abandoned.

The present invention relates to compounds and pharmaceutical compositions having central nervous system (CNS) activity which are useful in the treatment of epilepsy and other CNS disorders. More specifically, the compounds of this invention can be characterized as protected amino acid derivatives having the following general formula:



R is hydrogen, lower alkyl, lower alkenyl, lower alkynyl, aryl, aryl lower alkyl, heterocyclic, heterocyclic lower alkyl, lower alkyl heterocyclic, lower cycloalkyl, lower cycloalkyl lower alkyl, and R is unsubstituted or is substituted with at least one electron withdrawing group, or electron donating group;

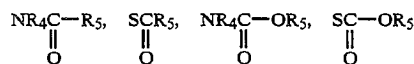
R<sub>1</sub> is hydrogen or lower alkyl, lower alkenyl, lower alkynyl, aryl lower alkyl, aryl, heterocyclic lower alkyl heterocyclic, lower cycloalkyl, lower cycloalkyl lower alkyl, each unsubstituted or substituted with an electron donating group or an electron withdrawing group and

R<sub>2</sub> and R<sub>3</sub> are independently hydrogen, lower alkyl, lower alkenyl, lower alkynyl, aryl lower alkyl, aryl, heterocyclic, heterocyclic lower alkyl, lower alkyl heterocyclic, lower cycloalkyl, lower cycloalkyl lower alkyl, or Z-Y wherein R<sub>2</sub> and R<sub>3</sub> may be unsubstituted or substituted with at least one electron withdrawing group or electron donating group;

Z is O, S, S(O)<sub>n</sub>, NR<sub>4</sub>, PR<sub>4</sub> or a chemical bond;

Y is hydrogen, lower alkyl, aryl, aryl lower alkyl, lower alkenyl, lower alkynyl, halo, heterocyclic, heterocyclic lower alkyl, lower alkyl and Y may be unsubstituted or substituted with an electron donating group or an electron withdrawing group, provided that when Y is halo, Z is a chemical bond, or

ZY taken together is NR<sub>4</sub>NR<sub>5</sub>R<sub>7</sub>, NR<sub>4</sub>OR<sub>5</sub>, ONR<sub>4</sub>R<sub>7</sub>, OPR<sub>4</sub>R<sub>5</sub>, PR<sub>4</sub>OR<sub>5</sub>, SNR<sub>4</sub>R<sub>7</sub>, NR<sub>4</sub>SR<sub>7</sub>, SPR<sub>4</sub>R<sub>5</sub> or PR<sub>4</sub>SR<sub>7</sub>, NR<sub>4</sub>PR<sub>5</sub>R<sub>6</sub> or PR<sub>4</sub>NR<sub>5</sub>R<sub>7</sub>,



R<sub>4</sub>, R<sub>5</sub> and R<sub>6</sub> are independently hydrogen, lower alkyl, aryl, aryl lower alkyl, lower alkenyl, or lower alkynyl, wherein R<sub>4</sub>, R<sub>5</sub> and R<sub>6</sub> may be unsubstituted or substituted with an electron withdrawing group or an electron donating group and

R<sub>7</sub> is R<sub>6</sub> or COOR<sub>8</sub> or COR<sub>8</sub>

R<sub>8</sub> is hydrogen or lower alkyl, or aryl lower alkyl, and the aryl or alkyl group may be unsubstituted or substituted with an electron withdrawing group or an electron donating group and

n is 1-4 and

a is 1-3.

The predominant application of anticonvulsant drugs is the control and prevention of seizures associated with epilepsy or related central nervous system disorders. Epilepsy refers to many types of recurrent seizures produced by paroxysmal excessive neuronal discharges in the brain; the two main generalized seizures are petit mal, which is associated with myoclonic jerks, akinetic seizures, transient loss of consciousness, but without convulsion; and grand mal which manifests in a continuous series of seizures and convulsions with loss of consciousness.

The mainstay of treatment for such disorders has been the long-term and consistent administration of anticonvulsant drugs. Most drugs in use are weak acids that, presumably, exert their action on neurons, glial cells or both of the central nervous system. The majority of these compounds are characterized by the presence of at least one amide unit and one or more benzene rings that are present as a phenyl group or part of a cyclic system.

Much attention has been focused upon the development of anticonvulsant drugs and today many such drugs are well known. For example, the hydantions, such as phenytoin, are useful in the control of generalized seizures and all forms of partial seizures. The oxazolinediones, such as trimethadione and paramethadione, are used in the treatment of nonconvulsive seizures. Phenacemide, a phenylacetylurea, is one of the most well known anticonvulsants employed today, while much attention has recently been dedicated to the investigation of the diazepines and piperazines. For example, U.S. Pat. Nos. 4,002,764 and 4,178,378 to Allgeier, et al. disclose esterified diazepine derivatives useful in the treatment of epilepsy and other nervous disorders. U.S. Pat. No. 3,887,543 to Nakanishi, et al. describes a thieno [2,3-e][1,4]diazepine compound also having anticonvulsant activity and other depressant activity. U.S. Pat. No. 4,209,516 to Heckendorn, et al. relates to triazole derivatives which exhibit anticonvulsant activity and are useful in the treatment of epilepsy and conditions of tension and agitation. U.S. Pat. No. 4,372,974 to Fish, et al. discloses a pharmaceutical formulation containing an aliphatic amino acid compound in which the carboxylic acid and primary amine are separated by three or four units. Administration of these compounds in an acid pH range are useful in the treatment of convulsion disorders and also possess anxiolytic and sedative properties.

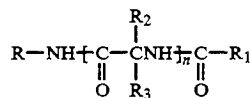
Unfortunately, despite the many available pharmacotherapeutic agents, a significant percentage of the population with epilepsy or related disorders are poorly managed. Moreover, none of the drugs presently available are capable of achieving total seizure control and most have disturbing side-effects. Clearly, current therapy has failed to "seize control" of these debilitating diseases.

It is therefore one object of the present invention to provide novel compounds exhibiting CNS activity, particularly anticonvulsant activity.

Another object of this invention is to provide pharmaceutical compositions useful in the treatment of epilepsy and other CNS disorders.

A further object of this invention is to provide a method of treating epilepsy and related convulsant disorders.

These and other objects are accomplished herein by providing compounds of the following general formula:



wherein R, R<sub>1</sub>, R<sub>2</sub>, R<sub>3</sub>, R<sub>4</sub>, R<sub>5</sub>, R<sub>6</sub>, R<sub>7</sub>, R<sub>8</sub>, n, Z, Y are as defined hereinabove.

The present invention contemplates employing the compounds of Formula I in compositions of pharmaceutically acceptable dosage forms. Where the appropriate substituents are employed, the present invention also includes pharmaceutically acceptable addition salts. Moreover, the administration of an effective amount of the present compounds, in their pharmaceutically acceptable forms or the addition salts thereof, can provide an excellent regime for the treatment of epilepsy, nervous anxiety, psychosis, insomnia and other related central nervous disorders.

The alkyl groups when used alone or in combination with other groups, are lower alkyl containing from 1 to 6 carbon atoms and may be straight chain or branched. These groups include methyl, ethyl, propyl, isopropyl, butyl, isobutyl, tertiary butyl, amyl, hexyl, and the like.

The aryl lower alkyl groups include, for example, benzyl, phenethyl, phenpropyl, phenisopropyl, phenbutyl, and the like, diphenylmethyl, 1,1-diphenylethyl, 1,2-diphenylethyl, and the like.

The term aryl, when used alone or in combination, refers to an aromatic group which contains from 6 up to 18 ring carbon atoms and up to a total of 25 carbon atoms and includes the polynuclear aromatics. These aryl groups may be monocyclic, bicyclic, tricyclic or polycyclic and are fused rings. Polynuclear aromatic compound is meant to encompass bicyclic, tricyclic fused aromatic ring system containing from 10-18 ring carbon atoms and up to a total of 25 carbon atoms. The aryl group includes phenyl, and the polynuclear aromatics e.g., naphthyl, anthracenyl, phenanthrenyl, azulenyl and the like. The aryl group also includes groups like ferrocenyl.

Lower alkenyl is an alkenyl group containing from 2 to 6 carbon atoms and at least one double bond. These groups may be straight chained or branched and may be in the Z or E form. Such groups include vinyl, propenyl, 1-butenyl, isobutenyl, 2-butenyl, 1-pentenyl, (Z)-2-pentenyl, (E)-2-pentenyl, (Z)-4-methyl-2-pentenyl,

(E)-4-methyl-2-pentenyl, pentadienyl, e.g., 1,3 or 2,4-pentadienyl, and the like.

The term alkynyl include alkyne substituents containing 2 to 6 carbon atoms and may be straight chained as well as branched. It includes such groups as ethynyl, propynyl, 1-butylnyl, 2-butylnyl, 1-pentynyl, 2-pentynyl, 3-methyl-1-pentynyl, 3-pentynyl, 1-hexynyl, 2-hexynyl, 3-hexynyl and the like.

The term cycloalkyl when used alone or in combination is a cycloalkyl group containing from 3 to 18 ring carbon atoms and up to a total of 25 carbon atoms. The cycloalkyl groups may be monocyclic, bicyclic, tricyclic, or polycyclic and the rings are fused. The cycloalkyl may be completely saturated or partially saturated. Examples include cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, cyclooctyl, cyclodecyl, cyclohexenyl, cyclopentenyl, cyclooctenyl, cycloheptenyl, decalynyl, hydroindanyl, indanyl, fenchyl, pinenyl, adamantyl, and the like. Cycloalkyl includes the cis or trans forms. Furthermore, the substituents may either be in endo or exo positions in the bridged bicyclic systems.

The term "electron-withdrawing and electron donating" refer to the ability of a substituent to withdraw or donate electrons relative to that of hydrogen if the hydrogen atom occupied the same position in the molecule. These terms are well understood by one skilled in the art and are discussed in *Advanced Organic Chemistry*, by J. March, John Wiley and Sons, New York N.Y., pp. 16-18 (1985) and the discussion therein is incorporated herein by reference. Electron withdrawing groups include halo, including bromo, fluoro, chloro, iodo and the like; nitro, carboxy, lower alkenyl, lower alkynyl, formyl, carboxyamido, aryl, quaternary ammonium, trifluoromethyl, aryl lower alkanoyl, carbalkoxy and the like. Electron donating groups include such groups as hydroxy, lower alkoxy, including methoxy, ethoxy and the like; lower alkyl, such as methyl, ethyl, and the like; amino, lower alkylamino, di(loweralkyl) amino, aryloxy such as phenoxy, mercapto, lower alkylthio, lower alkylmercapto, disulfide (lower alkylthio) and the like. One skilled in the art will appreciate that the aforesaid substituents may have electron donating or electron withdrawing properties under different chemical conditions. Moreover, the present invention contemplates any combination of substituents selected from the above-identified groups.

The term halo includes fluoro, chloro, bromo, iodo and the like.

The term acyl includes lower alkanoyl.

As employed herein, the heterocyclic substituent contains at least one sulfur, nitrogen or oxygen, but also may include one or several of said atoms. The heterocyclic substituents contemplated by the present invention include heteroaromatics and saturated and partially saturated heterocyclic compounds. These heterocyclics may be monocyclic, bicyclic, tricyclic or polycyclic and are fused rings. They may contain up to 18 ring atoms and up to a total of 17 ring carbon atoms and a total of up to 25 carbon atoms. The heterocyclics are also intended to include the so-called benzoheterocycles. Representative heterocyclics include furyl, thienyl, pyrazolyl, pyrrolyl, imidazolyl, indolyl, thiazolyl, oxazolyl, isothiazolyl, isoxazolyl, piperidyl, pyrrolinyl, piperazinyl, quinolyl, triazolyl, tetrazolyl, isoquinolyl, benzofuryl, benzothienyl, morpholinyl, benzoxazolyl, tetrahydrofuryl, pyranyl, indazolyl, purinyl, indolinyl, pyrazolidinyl, imidazolinyl, imadazolidinyl, pyrrolidinyl,

nyl, furazanyl, N-methylindolyl, methylfuryl, pyridaziny, pyrimidinyl, pyrazinyl, pyridyl, epoxy, aziridino, oxetanyl, azetidiny, the N-oxides of the nitrogen containing heterocycles, such as the nitric oxides of pyridyl, pyrazinyl, and pyrimidinyl and the like. The preferred heterocyclic are thienyl, furyl, pyrrolyl, benzofuryl, benzothiényl, indolyl, methylpyrrolyl, morpholinyl, pyridyl, pyrazinyl, imidazolyl, pyrimidinyl, or pyridazinyl. The preferred heterocyclic is a 5 or 6-membered heterocyclic compound. The especially preferred heterocyclic is furyl, pyridyl, pyrazinyl, imidazolyl, pyrimidinyl, or pyridazinyl. The most preferred heterocyclic is furyl and pyridyl.

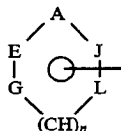
The preferred compounds are those wherein n is 1, but di, tri and tetrapeptides are also contemplated to be within the scope of the claims.

The preferred values of R is aryl lower alkyl, especially benzyl, and the preferred R<sub>1</sub> is H or lower alkyl. The most preferred R<sub>1</sub> group is methyl.

The most preferred electron donating substituent and electron withdrawing substituent are halo, nitro, alkanoxy, formyl, arylalkanoxy, arylloxy, carboxyl, carbalkoxy, carboxamide, cyano, sulfonyl, sulfoxide, heterocyclic, guanidine, quaternary ammonium, lower alkenyl, lower alkynyl, sulfonium salts, hydroxy, lower alkoxy, lower alkyl, amino, lower alkylamino, di(loweralkyl)amino, amino lower alkyl mercapto, mercaptoalkyl, alkylthio; and alkylidithio. The term "sulfide" encompasses mercapto, mercapto alkyl and alkylthio, while the term disulfide encompasses alkylidithio. These preferred substituents may be substituted on any one of R<sub>1</sub>, R<sub>2</sub>, R<sub>3</sub>, R<sub>4</sub>, R<sub>5</sub> or R<sub>6</sub>, R<sub>7</sub> or R<sub>8</sub> as defined herein.

The ZY groups representative of R<sub>2</sub> and R<sub>3</sub> include hydroxy, alkoxy, such as methoxy, ethoxy, aryloxy, such as phenoxy; thioalkoxy, such as thiomethoxy, thioethoxy; thioaryloxy such as thiophenoxy; amino; alkylamino, such as methylamino, ethylamino; arylamino, such as anilino; lower dialkylamino, such as, dimethylamino; trialkyl ammonium salt, hydrazino, alkylhydrazino and arylhydrazino, such as N-methylhydrazino, N-phenylhydrazino, carbalkoxy hydrazino, aralkoxycarbonyl hydrazino, aryloxycarbonyl hydrazino, hydroxylamino, such as N-hydroxylamino (—NH—OH), lower alkoxy amino [(NHOR<sub>18</sub>) wherein R<sub>18</sub> is lower alkyl], N-lower alkylhydroxyl amino [(NR<sub>18</sub>)OH wherein R<sub>18</sub> is lower alkyl], N-lower alkyl-O-lower alkyl hydroxyamino, i.e., [N(R<sub>18</sub>)OR<sub>19</sub> wherein R<sub>18</sub> and R<sub>19</sub> are independently lower alkyl] and o-hydroxylamino (—O—NH<sub>2</sub>); alkylamido such as acetamido, trifluoroacetamido, lower alkoxyamino, (e.g. NH(OCH<sub>3</sub>)); and heterocyclicamino, such as pyrazoylamino.

The heterocyclic groups representative of R<sub>2</sub> and R<sub>3</sub> have the formula



or those corresponding partially or fully saturated form thereof wherein n is 0 or 1

A, Z, L and J are independently CH, or a heteroatom selected from the group consisting of N, O, S, and

G is CH, or a heteroatom selected from the group consisting of N, O and S,

but when n is O, G is CH, or a heterocyclic selected from the group consisting of NH, O and S with the proviso that at most two of A, E, L, J and G are heteroatoms.

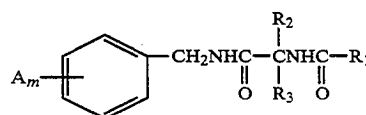
5 If the ring depicted hereinabove contains a nitrogen ring atom, then the N-oxide forms are also contemplated to be within the scope of the invention.

When R<sub>2</sub> or R<sub>3</sub> is a heterocyclic of the above formula, it may be bonded to the main chain by a ring carbon atom. When n is O, R<sub>2</sub> or R<sub>3</sub> may additionally be bonded to the main chain by a nitrogen ring atom.

It is preferred that one of R<sub>2</sub> and R<sub>3</sub> is hydrogen

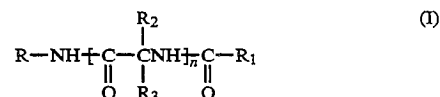
10 In a preferred embodiment, one of R<sub>2</sub> and R<sub>3</sub> is hydrogen and that the other is heterocyclic. It is preferred that one of R<sub>2</sub> and R<sub>3</sub> is a heterocyclic having Formula XI. The preferred heterocyclics include furyl, thienyl, benzothiényl, benzofuryl, oxazolyl, thiazolyl, isoxazolyl, indolyl, pyrazolyl, isoxazolidinyl, benzothiényl, benzofuryl, morpholinyl, indolyl, pyrrolyl, furfuryl, and methylpyrrolyl, pyridyl, pyrazinyl, imidazolyl, pyrimidinyl or pyridazinyl. In another preferred embodiment, one of R<sub>2</sub> and R<sub>3</sub> is alkyl (e.g. methylisopropyl), aryl (e.g., phenyl), 2-thiomethylethyl, lower alkoxy (e.g., ethoxy, methoxy), anilino, propenyl, alkylamino (e.g., ethylamino or methylamino). In another preferred embodiment, one of R<sub>2</sub> and R<sub>3</sub> is hydrogen and the other is heterocyclic lower alkyl, lower alkenyl, amino, lower alkoxy amino, N-lower alkylhydroxyamino, lower alkoxyamino, N-lower alkyl-O-lower alkylhydroxyamino or aralkoxycarbonylhydrazino,

Preferred compounds of the present invention have the following general formula:



40 wherein R<sub>1</sub> is H or lower alkyl, R<sub>2</sub> and R<sub>3</sub> are as defined above and A is hydrogen or an electron donating group or electron-withdrawing group and m is 0-5. It is preferred that A is hydrogen (i.e., m=0). However, values of m equalling 1, 2 or 3 are also preferred.

45 Preferred embodiments include compounds of Formula I



50 wherein R and R<sub>1</sub>, independently, are hydrogen, lower alkyl, lower alkenyl, lower alkynyl, aryl lower alkyl, aryl, heterocyclic, lower alkyl heterocyclic, each unsubstituted or substituted with at least one substituent;

55 R<sub>2</sub> and R<sub>3</sub>, independently, are hydrogen, lower alkyl, lower alkenyl, lower alkynyl, aryl lower alkyl, aryl, heterocyclic, lower alkyl heterocyclic, each unsubstituted or substituted with at least one substituent; halogen or a heteroatom containing oxygen, nitrogen, sulfur or phosphorous substituted with hydrogen, lower alkyl or aryl, said lower alkyl or aryl groups being substituted or unsubstituted; and

n is 1 to 4.

Another preferred embodiment is a compound having Formula I

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Sync your system to PACER to automate legal marketing.